The human oncoprotein PRAME unveiled: interaction networks and genomic targets

Adalberto Costessi
To the inspiring and dedicated teachers
I had the opportunity to meet
and to the memory of Gabriella Greblo.

“A teacher affects eternity;
he can never tell where his influence stops.”

Henry Adams
The human oncoprotein
PRAME unveiled
interaction networks and genomic targets

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AIM

The human protein PRAME (preferentially expressed antigen of melanoma) is frequently overexpressed in tumours. Importantly, high expression levels are associated to poor clinical outcome in melanoma, neuroblastoma, and breast cancer. However, its function and molecular mechanisms in normal and cancer cells are not yet well understood and the cellular pathways affected by PRAME are still elusive. Although PRAME has been indicated as a promising potential target for cancer therapy and anti-tumour immunization protocols, efforts to develop strategies to interfere with the oncogenic activity of PRAME are hindered by a lack of information about its function.

The aim of this thesis was to investigate the molecular function of human PRAME using cell line model systems and a powerful combination of proteomic and genomic techniques. We developed innovative biochemical approaches to dissect the protein-protein interaction network of PRAME, and we setup chromatin immunoprecipitation assays coupled to next-generation sequencing technologies to reveal binding profile of PRAME to the human genome in cancer cells.
OUTLINE

Chapter 1 provides a general introduction and an overview of the knowledge available on PRAME and the PRAME-like gene family. This chapter introduces also the proteomic and genomic techniques developed and applied in this thesis.

Chapter 2 describes the setup and application of epitope-tagging affinity purification and mass spectrometry analysis to the identification of the protein interactors of PRAME.

In chapter 3 we extend the results of chapter 2 and characterize PRAME as a component of Cullin2-based ubiquitin ligases. Furthermore, we present the first genome-wide ChIP-seq profiling of PRAME that associates this oncoprotein to the NFY signaling pathway.

Chapter 4 presents the results of improved protein-complex purification protocols and analysis of the PRAME interactome to a deeper level. We characterize direct interactions between PRAME and the human EKC/KEOPS complex and we show that EKC subunits associate with PRAME binding sites on chromatin.

Chapter 5 reports the development of rabbit antibodies against PRAME and their validation for use in several assays. We further discuss the experimental strategies that can be undertaken to generate genome-wide ChIP-seq profiles of a protein of interest, including epitope-tagging approaches. Finally, we investigated the reported function of PRAME as dominant repressor of retinoic acid signaling using proliferation assays and PRAME knockdown systems.

Chapter 6 summarizes the results of the previous chapters and discusses the implications of this thesis for further research.

In the Appendix, the work performed for this thesis is presented in a chronological context, as a reference for similar projects in the future.
LIST OF ABBREVIATIONS AND GENE SYMBOLS

Ab, antibody
ChIP, chromatin immunoprecipitation
CRL, Cullin-RING ligases
CUL2, cullin 2
EKC, Endopeptidase-like Kinase Chromatin-associated
ELOB/TCEB2, elongin B/transcription elongation factor B (SIII), polypeptide 2
ELOC/TCEB1, elongin C/transcription elongation factor B (SIII), polypeptide 1
IP, immunoprecipitation
KEOPS, kinase, putative endopeptidase and other proteins of small size
LAGE3, L antigen family, member 3
MS, mass spectrometry
NFY, nuclear transcription factor Y
OSGEP, O-sialoglycoprotein endopeptidase
PRAME, preferentially expressed antigen in melanoma
RBX1, ring-box 1
RING, really interesting new gene
TP53RK, TP53 regulating kinase
TPRKB, TP53RK binding protein
WB, western blot
CHAPTER 1

GENERAL INTRODUCTION
Cancer is a general term that refers to more than hundred different diseases that are all characterised by uncontrolled cell proliferation. Collectively, cancers constitute the second leading cause of human mortality worldwide (WHO, 2012). Cancers are mostly classified based on the tissue of origin and histological analysis of tissue sections or cells, based on which diagnosis and prognosis are made. Some cancers are also tested for specific molecular markers of known prognostic value, i.e. ERalpha and HER2 for breast cancer. The advent of array-based techniques, and more recently high-throughout sequencing, to quantitatively measure the levels of messenger RNA transcripts has substantially increased the power to sub-classify cancers based on gene expression profiles (McDermott et al, 2011). Importantly, specific gene expression signatures have been characterised that are predictive of prognosis and can therefore guide therapeutic interventions, like the Mammaprint 70-gene profile for breast cancer (McDermott et al, 2011; van ’t Veer et al, 2002).

The human oncoprotein PRAME

The human gene PRAME (preferentially expressed antigen of melanoma) is frequently overexpressed in tumours and constitutes a molecular marker of poor clinical outcome for several cancers like melanoma, neuroblastoma and breast cancer (Kilpinen et al, 2008; Oberthuer et al, 2004; Epping & Bernards, 2006; Doolan et al, 2008; Haqq et al, 2005). The discovery of the PRAME gene was reported in 1998 by the Coulie lab (Ikeda et al, 1997). The authors identified PRAME as the gene encoding an antigenic peptide recognised by tumour-specific cytolytic T-lymphocytes isolated from a melanoma patient. The gene encodes a putative protein of 509 amino acids with unknown function. In healthy adult tissues, PRAME is expressed at high levels only in the testis, while low levels were detected in the adrenals, ovary, endometrium, and very low levels in skin, brain, breast, and kidney (Ikeda et al, 1997). However, PRAME is frequently overexpressed in a wide range of solid and haematological tumours including melanoma, head and neck carcinoma, ovarian cancer, breast carcinoma, lung carcinoma, renal carcinoma, neuroblastoma, and various acute and chronic haematological malignancies (Ikeda et al, 1997; Kilpinen et al, 2008).

Based on its peculiar expression pattern, PRAME belongs to the category of cancer-testis antigens, similarly to the MAGE, BAGE and GAGE gene families. The expression in healthy tissues other than testis, however,
differentiates PRAME from other cancer-testis antigens, whose expression is restricted to normal testis. Since the testis does not bear HLA molecules, it is invisible to the immune system and high expression of PRAME (and other CTA proteins) in this tissue does not trigger an immune response. It has been suggested that the low expression of PRAME in other normal tissues is insufficient for recognition by cytolytic T-lymphocytes, and only upon malignant overexpression would the immune system be able to mount an anti-PRAME immune response (Ikeda et al, 1997; van Baren et al, 1998).

**PRAME expression in cancer**

A large number of studies investigated the transcript levels of *PRAME* in cohorts of cancer patients. While normal skin and benign nevi were found negative, high PRAME expression was found to correlate with the stage of melanoma lesions (Haqq et al, 2005), and a similar trend was reported for neuroblastoma: 93% of primary lesions and 100% of advanced cases overexpressed PRAME (Oberthuer et al, 2004). On the contrary, PRAME levels were lower in neuroblastoma cases characterised by a good prognosis and a high frequency of spontaneous regression (Fischer et al, 2006).

Importantly, significant associations were found between higher PRAME expression levels and poor clinical outcome (i.e. shorter overall and disease-free survival) in neuroblastoma (Oberthuer et al, 2004), as well as serous ovarian adenocarcinoma patients (Partheen et al, 2008). High PRAME mRNA levels were also found to be an independent prognostic factor of poor clinical outcome in breast cancer by two independent studies (Doolan et al, 2008; Epping et al, 2008).

*PRAME* transcripts were detected in about 30-40% of acute and chronic leukaemias, and hypomethylation of the *PRAME* locus seems to have a role in this process (Roman-Gomez et al, 2007; Schenk et al, 2007). Studies on patients affected by BCR-ABL-positive chronic myeloid leukaemia (Ph+ CML) indicated that *PRAME* was activated in the advanced stages and blast crisis (Radich et al, 2006; van Baren et al, 1998). The fact that *PRAME* was not expressed in the early stages, when the BCR-ABL translocation is already present, prompted the authors to suggest that the BCR-ABL translocation is not sufficient to activate *PRAME*, and the later activation could be due to a secondary correlation with the progression of the disease. However, two studies reported that *PRAME* expression could be induced by transfection of BCR-ABL in cell line models (De Carvalho et al, 2011; Watari et al, 2000).
Therefore, it cannot be excluded that additional (still BCR-ABL-mediated) signals are necessary for full activation of PRAME in CML blasts.

Although a growing body of evidence indicates that PRAME contributes markedly to the oncogenic phenotype of several cancers, some studies have shown that PRAME expression can be associated also with leukaemia subtypes characterised by a favourable prognosis. In particular, high PRAME levels were reported in patients with the genomic translocations t(8;21) and t(15;17), which result in the expression of the AML1-ETO and PML-RARA fusion proteins, respectively (van Baren et al., 1998). A larger recent study on patients carrying the PML-RARA translocation reported that high PRAME levels at diagnosis were associated with a better clinical outcome (Santamaría et al., 2008).

Finally, several studies indicated that PRAME expression levels are a useful marker to monitor the presence of cancer cells in the body (also called “minimal residual disease”) in patients during and after chemotherapy (Steinbach et al., 2002; Santamaria et al., 2008).

Molecular studies of PRAME

There are indications that epigenetic processes are involved in the regulation of PRAME expression. DNA hypomethylation of two different regions of the PRAME locus was reported to correlate with expression in cell lines as well as in the progression of CML. Notably, hypomethylation induced by 5-aza-dC treatment resulted in PRAME upregulation in several cell lines, suggesting that all the factors required for PRAME expression were present in the cells, and that DNA methylation mediates the silencing of this locus (Ikeda et al., 1997; Roman-Gomez et al., 2007; Schenk et al., 2007).

Five different transcript variants of PRAME have been described (Schenk et al., 2007). These variants show two different transcription start sites (positions +1 and +151), but they all encode the same amino acid sequence. Regulation of expression of the different variants has not yet been reported, and their functional relevance is unclear (Schenk et al., 2007). Recent analysis of small non-coding RNAs revealed that PRAME can be targeted by the microRNA miR-211, and downregulation of miR-211 correlates with upregulation of PRAME in melanoma cell lines (Sakurai et al., 2011).

Although numerous studies reported expression patterns of PRAME in human malignancies, the molecular and cellular functions of PRAME are not yet clear. In a functional genetic screen, ectopic expression of PRAME
was found to repress the growth inhibitory effects of histone deacetylase inhibitors (HDACi) (Epping et al, 2007). HDACi induce proliferation arrest and apoptosis in cancer cells but not in normal healthy cells; therefore, they are promising compounds for use in the treatment of cancer (Kim & Bae, 2011). Ectopic expression of human PRAME in mouse embryonic fibroblasts (MEFs) prevented growth arrest and apoptosis upon treatment with the HDACi PXD101, while PRAME overexpression did not affect proliferation in standard culturing conditions (Epping et al, 2007). Luciferase reporter assays indicated that PRAME could repress retinoic acid signaling induced by PXD101, while PRAME silencing in A375 melanoma cells sensitized the cells to PXD101 treatment in vitro as well as in a xenograft model (Epping et al, 2007).

The authors reported in a separate publication that PRAME interacts with the retinoic acid receptor alpha (RARA) and suggested that PRAME mediates repression of retinoic acid signaling via recruitment of the transcriptional repressor EZH2 to RARA target genes (Epping et al, 2005). In particular, the authors reported in vitro protein-protein interactions between PRAME and the ligand binding domain of RARA fused to GST, while a mammalian two-hybrid system detected an interaction between the ligand binding domain of RARA and the C-terminus of PRAME (aa. 416-509). Co-immunoprecipitation experiments further suggested that this interaction requires the C-terminal LRELL amino acid motif in PRAME and the presence of the ligand all-trans retinoic acid (atRA). Luciferase reporter assays indicated that expression of PRAME represses retinoic acid signaling and that this effect is mediated by protein-protein interactions with the transcriptional repressor EZH2. A ChIP assay suggested that PRAME associates with the RARB2 promoter, a known target of retinoic acid signaling. Finally, PRAME silencing in A375 melanoma cells was reported to restore retinoic acid signaling and inhibit retinoic acid induced growth arrest both in proliferation assays in vitro and in a xenograft model in vivo (Epping et al, 2005). Contrary to the model put forward by the Bernards lab, Wadelin et al. could not reproduce the interactions between PRAME and the ligand binding domain of RARA in GST pulldown and yeast two-hybrid assays (Wadelin et al, 2010). Furthermore, studies on primary AML and breast cancer samples indicated that PRAME expression did not associate with a down-regulation of retinoic acid signaling (Epping et al, 2008; Steinbach et al, 2007).

Notably, it was reported that overexpression of SOX9 in melanoma cell lines lead to increased sensitivity to retinoic acid and at the same time
downregulation of PRAME (Passeron et al, 2009). However, the authors did not provide a causative link between these two findings. Therefore, it is still unclear whether the downregulation of PRAME plays any role in the increased sensitivity to retinoic acid caused by SOX9 overexpression, or whether it is an unlinked event.

In conclusion, a putative role for PRAME in the retinoic acid signaling pathway remains unclear, and additional studies are required to shed light on these mechanisms.

**PRAME in leukaemia**

The molecular functions of PRAME in leukaemia cells are also controversial, since conflicting reports suggested that PRAME might induce caspase-independent cell death (Tajeddine et al, 2005), or rather promote cell survival repressing pro-apoptotic genes (Tanaka et al, 2011).

Tajeddine et al. reported that PRAME causes decreased proliferation and caspase-independent cell death in CHO-K1, HeLa and KG1 cell lines. Using a microarray-based approach, transfection of PRAME in leukaemia KG1 cells was found to downregulate expression of the *HSP27*, *CDKN1A/p21* and *S100A4* genes, which are inhibitors of apoptosis. However, these associations were not reported in a microarray gene expression analysis of paediatric AML samples with high versus low PRAME expression (Goellner et al, 2006). Moreover, an independent study on paediatric AML samples by real-time PCR found that high levels of PRAME were not associated with down-regulation of *HSP27*, *S1000A*, *CDKN1A/p21* or *RARB* (Steinbach et al, 2007).

Tajeddine et al. also reported that downregulation of PRAME by siRNA increased proliferation and tumorigenicity of K562 leukaemia cells in a xenograft nude mice model. On the contrary, Tanaka et al. showed that siRNA transfection in the same cell line suppressed proliferation in vitro and promoted apoptosis. Another study in agreement with Tajeddine et al. found that silencing of PRAME promoted cell proliferation of K562 cells; however, the same effect was obtained upon overexpression of PRAME in HL60 cells (Oehler et al, 2009). Moreover, PRAME silencing on K562 cells expressing increased numbers of RARA receptors did not show any difference in proliferation in the same assays (Oehler et al, 2009). It is possible that the functional relevance of PRAME differs between different cell types and cancers, and further analyses are required in order to understand the mechanisms and pathways in which PRAME is involved.
Evolution of the PRAME gene family

*PRAME* is the founding member of a large family of genes with a peculiar evolutionary history. In human, the *PRAME* gene family is comprised of the single *PRAME* gene on chromosome 22, and a cluster of 22 *PRAME*-like genes and 10 pseudogenes on chromosome 1 (Birtle *et al*, 2005).

Molecular evolution analyses established that the *PRAME*-like gene cluster originated from a translocation dating shortly before the divergence between rodents and primates (~85-95 million years ago). In fact, synthentic regions of rodent and primate genomes contain *PRAME*-like homologues, while dog and chicken genomes do not (Birtle *et al*, 2005). After the first translocation event, the mouse and primate loci underwent an independent and very rapid expansion by “birth-and-death” processes, i.e. gene duplication and pseudogene creation. This process explains the finding that mouse *Prame*-like genes are monophyletic, similarly to the human *PRAME*-like genes. In other words, while ~80% of mouse genes have a single human orthologue, there is no pair of mouse and human *PRAME* genes with a simple 1:1 orthology. This evidently hinders the development of mouse model systems for the study of human *PRAME* genes.

Furthermore, independent duplication events have been detected in all branches of the primate lineage (Rhesus Macaque Genome Sequencing and Analysis Consortium *et al*, 2007). Extensive duplications occurred early in the primate evolution as well as in recent chimpanzee evolution, while the macaque branch seems to have been less dynamic (Figure 1). The chimpanzee and hominin branches separated around 6-7 million years ago, and two independent large segmental duplications occurred in the terminal human branch in the past 3 million years. This implies that a number of human *PRAME*-like genes do not have a 1:1 orthology even with chimpanzee *PRAME* genes.

Besides the extraordinarily rapid and independent expansion of *PRAME* genes in the rodent and the primate lineages, there is also evidence for copy number variation in the *PRAME*-like gene cluster among human populations (Birtle *et al*, 2005). This indicates that there have been additional recent duplications that are not yet fixed in the human population, nor captured in the human genome reference sequence.

Evidence has also been found for positive selection of amino acid residues in *PRAME* (Birtle *et al*, 2005; Rhesus Macaque Genome Sequencing and Analysis Consortium *et al*, 2007). Homology modelling suggested that the
positively-selected residues identified are likely to cluster on an accessible surface of PRAME proteins, underscoring a potential role in binding interactions. Moreover, since positive selection is a common feature of genes involved in immunity and reproduction, it was suggested that PRAME genes might have roles in these processes (Birtle et al, 2005; Wadelin et al, 2010). Given its peculiar characteristics, it was suggested that the PRAME gene family has played a key role in species evolution and that the differences between human and chimpanzee PRAME genes may have contributed to the functional divergence along the hominin lineage (Birtle et al, 2005).

While the evolution of the PRAME-like gene cluster has been studied in detail, the origin of the PRAME gene on chr22 is unclear. Inspection of the orthologous genomic regions in the mouse, rat and chimp genomes, did not identify PRAME orthologues, and gaps in the genome builds have not been reported that could accommodate a missing PRAME gene (Birtle et al, 2005).
PRAME-like genes and early development

Despite the extraordinary evolutionary history of PRAME genes, their biological and molecular function is still poorly understood. While no reports have been published yet on human PRAME-like genes, a number of studies in the mouse identified roles for this gene family in oogenesis (Dadé et al., 2003), spermatogenesis (Wang et al., 2001) and stemness (Bortvin et al., 2003; Cinelli et al., 2008).

A number of mouse PRAME-like genes were first identified as genes with specific expression in the oocyte, and therefore they were first called “oogenesin” (Dadé et al., 2003; Minami et al., 2003). Analysis of their predicted amino acid sequence suggested that the encoded proteins belong to the leucine-rich repeat (LRR) superfamily and fold in a non-globular horse-shoe structure (Dadé et al., 2003).

Four PRAME-like genes were found to play roles in pluripotency and development of mouse embryos cloned by nuclear transfer (Bortvin et al., 2003). In a study aimed at identifying genes involved in establishing or maintaining developmental pluripotency, the authors performed an in silico analysis to identify genes with an expression pattern similar to that of Oct4 (Bortvin et al., 2003). OCT4 is a transcription factor expressed in pluripotent cells of the early embryo and adult germ cells that is known to play an essential role in the control of developmental pluripotency. Ten genes were found that satisfied the search criteria, among which Pramel4, Pramel5, Pramel6, Pramel7. Analysis of expression of these genes established that they are active in pluripotent embryonic tissues (blastocyst, morula and 4-cell embryo), while they are not expressed in differentiated somatic tissues. Importantly, analysis of their activity in mouse embryos cloned from somatic or embryonic stem (ES) cells revealed that successful reactivation of the full set of Oct4-genes correlated with successful development of the cloned embryos to term. The authors concluded that although the biological functions of these genes have yet to be elucidated, they serve as informative markers for pluripotent cell types and cloned embryos (Bortvin et al., 2003).

Another study found that the MBD3 component of the nucleosome remodelling and histone deacetylation (NuRD) complex mediates silencing of Pramel6 and Pramel7 expression in mouse embryonic stem cells (Kaji et al., 2006). These genes were found to be expressed at very low levels in wild type ES cells, while knockout of Mbd3 resulted in elevated transcript levels as well as increased H3K9 and H4 acetylation levels at their genomic loci,
but not at the silent Pramel5 promoter. This correlated with LIF-independent maintenance of pluripotency in culture and a defect in lineage commitment of Mbd3-/- ES cells. Although a causative role for Pramel6 and Pramel7 in these processes was not addressed, expression analysis suggested that Pramel6 and Pramel7 contribute in deciding the fate of ES cells (Kaji et al, 2006).

An independent study confirmed that Pramel6 and Pramel7 are prevalently expressed in embryonic pluripotent cells, and further determined that whereas Pramel6 is typically expressed in all cells of the morula and blastocyst, Pramel7 is expressed only in the inner part of the morula and in the inner cell mass of the blastocyst, similarly to Nanog, a factor known to be involved in pluripotency (Cinelli et al, 2008). Notably, ectopic overexpression of the STAT3 transcription factor resulted in overexpression of Pramel6 and Pramel7 and allowed the derivation of pluripotent embryonic stem cell lines from mouse strains that are otherwise “non permissive” to this procedure. Remarkably, overexpression of Pramel7 in ES cells was sufficient to sustain maintenance of pluripotency when cells were cultured in the absence of LIF, indicating an important role for this protein in maintaining ES cells in a pluripotent state (Cinelli et al, 2008).

**Ubiquitin pathways**

Modification of proteins by the addition of ubiquitin (a highly conserved 76-amino-acid polypeptide) or ubiquitin-like proteins has emerged as a major mechanism by which virtually all events in cells can be regulated, including cell cycle progression, gene transcription, DNA repair, protein turn-over, receptor trafficking, and apoptosis (Deshaies & Joazeiro, 2009; Budhidarmo et al, 2012).

Ubiquitination (also known as ubiquitylation) is best-known for targeting proteins for degradation by the 26S proteasome, but other functions include modulation of protein interactions and alteration of subcellular distribution.

Members of ubiquitin pathways play also important roles in cancer and a number of them have been identified as either oncogenes (i.e. MDM2) or tumour suppressor genes (i.e. BRCA1 and VHL), and the first pharmacological inhibitors of the proteasome have entered the clinic as anti-cancer drugs, and significant efforts are undertaken in this field of drug discovery (Joazeiro et al, 2006).

Attachment of ubiquitin to target proteins is performed by a cascade of three enzymes (E1-E2-E3) (Deshaies & Joazeiro, 2009; Metzger et al, 2012
and references therein). The first step is the activation of ubiquitin by one predominant ubiquitin-activating enzyme E1. Activated ubiquitin is then transferred from E1 to one of the 30-40 mammalian ubiquitin-conjugating enzymes (UBCs or E2s), and eventually to the final substrates that are bound specifically to one of the several hundreds different ubiquitin ligases (E3s). Substrates can be modified by a single ubiquitin on one or more sites (monoubiquitination and multi-monoubiquitination, respectively) or can be tagged with chains of ubiquitin (polyubiquitination). Chain formation can occur through the seven internal lysine residues on ubiquitin (K6, K11, K27, K29, K33, K48, and K63) or its N-terminus (M1). Ubiquitin chains are recognized by specialised ubiquitin-binding domains (UBDs) that form transient, noncovalent interactions either with ubiquitin moieties or with the linkage region in their chains and determine the function of the modification (Ikeda et al., 2010). K48-linked polyubiquitin chains were originally described as the signal that targets substrates for proteasomal degradation, while atypical K11, K63, and M1-linked chains were associated to DNA repair, cell cycle progression and inflammation. However, recent reports have challenged the notion that specific chain topologies regulate specific cellular processes, and it still unclear how a specific type of linkage is generated and how this is translated to the fate of the modified protein (Ikeda et al., 2010).

The ubiquitination cascades result in the formation of a covalent bond between the C-terminus of ubiquitin and, most frequently, the epsilon-amino group of a lysine side chain in the substrate. The ubiquitination of the N-terminus of proteins or other amino acids like threonine, serine, or cysteine has also been described. E3-mediated attachment of ubiquitin to substrates is highly regulated in response to cellular cues and one substrate can be also ubiquitinated by different ubiquitin ligases while, on the other hand, one ligase can target several proteins. Moreover, E3s themselves can undergo both self-ubiquitination and be ubiquitinated by heterologous ligases. Examples are the RING finger 1B component of the polycomb repressive complex 1 (PRC1), which self-ubiquitinates to increase its activity, while MDM2 self-ubiquitination targets itself and the p53 proto-oncogene for proteasome degradation (Weissman et al., 2011).

Since E3s dictate the specificity toward a variety of substrates and at the same time interact with an E2, they are the most structurally complex and diverse enzymes in the pathway (Zimmerman et al., 2010). E3 ligases can be classified into two main structural classes that bear either a HECT (ho-
mologous to E6-AP carboxy terminus) domain or a RING (really interesting new gene) domain, although a few have a U-box, which is structurally and functionally similar to the RING domain (Budhidarmo et al, 2012). HECT-containing E3 ligases, of which there are ~20 in humans, form an intermediate conjugate with the ubiquitin received from a charged E2 before transferring it to the target. In contrast, the more prevalent (~500) RING-containing E3 ligases bring together an ubiquitin-charged E2 and a substrate protein and facilitate the transfer of ubiquitin directly from the charged E2 to the substrate (Deshaies & Joazeiro, 2009).

Cullin-RING E3 ubiquitin ligases

Among the E3 RING ligases, the structurally-related multisubunit cullin-RING ubiquitin ligases (CRLs) are particularly significant (Petroski & Deshaies, 2005; Sarikas et al, 2011). This is a diverse group of E3 complexes characterized by containing both a cullin family protein and a RING finger protein (RBX1 or RBX2). The human genome encodes for six cullin proteins, CUL1, CUL2, CUL3, CUL4A, CUL4B, and CUL5, and two additional related proteins, PARC and CUL7, which are considered atypical cullins.

Extensive genome-wide analysis of the cullin family has suggested that three ancestral cullin genes, termed ‘Culα’, ‘Culβ’ and ‘Culγ’, appeared in early eukaryotic evolution, from which the cullin genes evolved after the split of the unikonts (which include animals and fungi) and the bikonts (which include plants) (Marín, 2009). The human CUL1, CUL2, CUL5, CUL7 and PARC genes were derived from one common ancestral gene (Culα), and the available data indicate that the emergence of multiple Culα genes in animals has been accompanied by a diversification of their protein partners and complex subunits. The CUL3 and CUL4A/4B genes evolved from two distinct ancestors, the Culβ and Culγ gene, respectively (Marin, 2009).

CRLs are the most abundant E3s as it is predicted that about 400-500 different E3 are encoded in the human genome. Their diversity is due to the fact that many alternative complexes can be generated in a combinatorial way: multiple related substrate receptor proteins may substitute for each other to form similar but functionally distinct complexes.

The roles of the cullin protein in this type of complex are quite well understood. Structural data indicate that cullins have an overall elongated crescent-shaped structure consisting of a stalk-like N-terminal domain (NTD), and a globular C-terminal domain (CTD) (Zimmerman et al, 2010) (Figure
2). The NTD binds adaptor proteins and substrate receptor subunits, which confers substrate specificity, while the CTD binds the RING-finger containing protein RBX1 or RBX2, which recruits the ubiquitin-conjugating enzyme E2. In this context, cullins act as backbones that facilitate ubiquitination by correctly positioning the RING finger protein and the substrate (Marín, 2009; Zimmerman et al., 2010; Petroski & Deshaies, 2005) (Figure 2).

Cullin-RING ligases can be classified in a number of main CRL classes based on the different adaptors, substrate receptors, and RING-finger proteins used (Marín, 2009; Petroski & Deshaies, 2005):

1. Cullin1/RBX1/SKP1/F-box-receptor CRLs (historically known as SCF complexes) that contain SKP1 or related proteins as adaptors and proteins with F-box motifs as substrate receptors;

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**Structural model of cullin-RING (CRL) ubiquitin ligases.**

The cullin-based E3 ubiquitin ligases position a substrate in close proximity to the RING-bound E2 ubiquitin-conjugating enzyme (not shown), which catalyzes the transfer of ubiquitin to the substrate.

(A) Schematic model of the CRL modular architecture (adapted from Sarikas et al., 2011). (B-E) Structural models of CRLs with the scaffold cullin proteins, RING-box proteins, adaptor proteins and substrate receptors (from Zimmerman et al., 2010).
2. Cullin2/RBX1/ElonginBC/VHL-box-receptor and Cullin5/RBX2/ElonginBC/SOCS-box-receptor CRLs (BC-box CRLs), which contain two Elongin proteins as adaptors and BC-box containing proteins as substrate receptors (Petroski & Deshaies, 2005; Okumura et al, 2012; Mahrour et al, 2008);
3. Cullin3/RBX1/BTB-receptor CRLs, characterized by lacking additional adaptors and containing proteins with BTB (broad complex/tramtrack/bric-a-brac) domains as substrate receptors, directly bound to the cullins;
4. Cullin4/RBX1/DDB1/WD40-proteins CRLs, which contain DDB1 as adaptor and proteins with WD40 domains as substrate receptors, also called DCAFs (DDB1–CUL4A-associated factors).

Despite sharing the identical adaptor proteins Elongin B and Elongin C, CUL2 and CUL5 direct the assembly of E3 ligases with distinct substrate receptor proteins: CUL2 with von Hippel-Lindau (VHL) or related BC-box proteins, and CUL5 with suppressors of cytokine signaling (SOCS)-box proteins. Sequence comparison of the VHL and SOCS-box proteins revealed that they each include a degenerate, ~12 amino acid sequence motif with consensus sequence [S,T,P]Lxxx[C,S,A]xxxΦ (Mahrour et al, 2008; Kamura et al, 2004). This motif, which is referred to as the BC-box, is required for binding to the Elongin BC complex. Downstream of the BC-box, a Cul2- or Cul5-box is present, which mediates the specific interaction to the corresponding cullin scaffold (Mahrour et al, 2008; Kamura et al, 2004). The most studied substrate of EloBC-cullin ligases is the von Hippel-Lindau (VHL) protein, the product of the VHL tumour suppressor gene. More than half of sporadic clear cell renal carcinomas contain inactivating mutations of VHL, and the VHL complex has ubiquitin ligase activity and targets the hypoxia-inducible factor-α (HIF-α) family of transcription factors (HIF-1–3α) for proteasomal degradation. At normal oxygen levels, proline residues within the oxygen-dependent degradation domain of HIF-α are hydroxylated and recognized by pVHL for polyubiquitination and proteasome-mediated degradation (Okumura et al, 2012).

The ligase activity of CRLs was found to be modulated by the covalent conjugation of an ubiquitin-like molecule, Nedd8 (Deshaies & Joazeiro, 2009). This modification, termed neddylation, activates the E3 ligase activity by promoting substrate polyubiquitination, likely via the induction of extensive conformational changes. Neddylation is a reversible modification that can
be enzymatically removed by the COP9 complex, and cullins are thought to undergo dynamic regulation by neddylation-deneddylation cycles. Another level of regulation of CRLs activity is performed by the CAND1 protein (Cullin-Associated and Neddylation-Dissociated-1), which inhibits the E3 ligase activity of CRLs by binding to both the C-terminus as well as the N-terminus of all cullins in their un-neddylated forms (Deshaies & Joazeiro, 2009).

**The EKC/KEOPS complex**

Two independent laboratories have identified an ancient and highly conserved multiprotein complex named KEOPS (Downey *et al*, 2006) or EKC (Kisseleva-Romanova *et al*, 2006), starting from genetic clues in yeast. This complex has orthologues from Archaea to Eukarya and so far it has been implicated in telomeres maintenance, transcriptional regulation, and t^6^A modification of tRNAs (Kisseleva-Romanova *et al*, 2006; Downey *et al*, 2006; Daugeron *et al*, 2011; Yacoubi *et al*, 2011; Srinivasan *et al*, 2011). Yeast EKC comprises four subunits which are also conserved in the human genome (human orthologues are indicated in brackets): Pcc1p (LAGE3, also known as ES03), the ATPase Kae1p (OSGEP), the kinase Bud32p (TP53RK, also known as PRPK), and Cgi121p (TPRKB). In addition, yeast EKC also includes Gon7p (also known as Pcc2p), which appears to be fungi-specific (Kisseleva-Romanova *et al*, 2006).

The laboratory of Domenico Libri isolated Pcc1, a protein of small size with no predicted functional domains in a search for suppressors of a splicing defect and demonstrated that its function is not related to splicing. Their genetic and biochemical experiments led to the description of the five-subunit EKC complex. Furthermore, the authors showed that Pcc1p is required for efficient transcription of several yeast genes induced by pheromones or galactose, and that deletion of Pcc1 inhibited the recruitment of the SAGA and Mediator co-activators (Kisseleva-Romanova *et al*, 2006).

A parallel genome-wide screen in yeast identified the cg121 null mutant as a suppressor of the telomere-capping defect of cdc13-1 cells (Downey *et al*, 2006). Deletion of CGI121 or other EKC subunits resulted in short telomeres in an otherwise wild-type context. A detailed analysis of three-dimensional crystal structures of archaeal EKC subunits revealed that the complex has a linear topology Pcc1-Kae1-Bud32-Cgi121, and Pcc1 is an obligate dimer that can support dimerization of the full complex in a V shape (Figure 3). Dimerization of the Pcc1p and Kae1p subunits was found to be necessary for
cell viability (Mao et al., 2008), and an intact interaction surface between Kae1p and Bud32p was required for both the transcriptional and telomere maintenance functions (Hecker et al., 2008).

Intriguingly, the OSGEP subunit is also present in bacteria (YgjD) and eukaryotic genomes express an OSGEP paralogue (Qri7/OSGEPL1) that localizes to mitochondria (Reinders et al., 2006). Comparative genomic studies identified OSGEP as one of the very few genes present in all genomes sequenced so far (Galperin, 2008), suggesting an extremely conserved function. Very recently, several groups have reported a crucial role for the YgjD/Kae1/OSGEP protein family in the biosynthesis of N6-threonylcarbamoyl adenosine (t^6A) (Daugeron et al., 2011; Yacoubi et al., 2011; Srinivasan et al., 2011): a universal modification at position 37 of tRNAs decoding ANN codons, which is required for accurate translation of messenger RNAs (Yarian et al., 2002).

Human LAGE3 belongs to the NY-ESO gene family together with the closely related LAGE1 and LAGE2 (Alpen et al., 2002), and all three genes are clustered in the same region on chromosome X. While LAGE3 is ubiquitously expressed, LAGE1 and LAGE2 are cancer-testis antigens with high expression in healthy testis and upregulation in a number of cancer tissues, similar to PRAME. Notably, expression of the N-terminal domain of LAGE3 fused to the C-terminus of Pcc1p could partially rescue the growth defects of a Pcc1 N-terminal mutant yeast strain, while full length LAGE3 could not substitute for Pcc1 (Kisseleva-Romanova et al., 2006).

Human TP53RK was first cloned by cDNA subtraction as an interleukin-2 upregulated gene in cytotoxic T-cells, and it was shown to phosphorylate the human oncoprotein p53 at Serine 15 (Abe et al., 2001). Recombinant TP53RK

**Figure 3** Composite model of the archaeal EKC/KEOPS complex in dimeric form (from Mao et al., 2008).
expressed in bacteria was found catalytically inactive, but incubation with COS-7 cell lysates was sufficient to activate the enzymatic activity, suggesting the need for activators and potentially protein interactors. The kinase Akt/ PKB was found to play a role in the activation of TP53RK through phosphorylation of the Ser250 residue (Facchin et al, 2007). Underscoring a high level of functional homology, TP53RK could partially rescue the growth phenotype of Bud32 deletion in yeast, and Bud32 could interact with and phosphorylate human p53 in vitro (Facchin et al, 2003), although yeast does not possess a bone fide p53 homologue.

Human TPRKB was first isolated by a two-hybrid screen using TP53RK as a bait, and was suggested to inhibit the binding of TP53RK to p53 (Miyoshi et al, 2003).

Although the structures of several EKC subunits have been characterised and phenotypes caused by mutations or depletion of EKC have been described in yeast, the precise molecular functions and mechanisms of EKC in Archaea as well as Eukarya are still unclear.

**Purification and identification of protein complexes**

Recent advances in genome sequencing technologies have resulted in hundreds of genomes being completely sequenced, and this number is rapidly growing (http://www.ensemblgenomes.org/). In parallel, computational algorithms have been developed that can efficiently predict and annotate genes, thereby generating comprehensive lists of predicted proteins for many species (the “proteomes”). However, such lists are not sufficient to understand cellular processes, which are dependent on the coordinated action of multiprotein complexes and dynamic signaling networks of interacting proteins. Therefore, experimental approaches have been developed to dissect protein-protein interaction networks (the “interactomes”) in order to fully understand the cellular machinery (Mering et al, 2002; Gingras et al, 2007).

A powerful technique to identify protein-protein interactors is the purification of entire protein complexes by epitope-tagging and affinity-based methods coupled to mass spectrometry identification. This approach has been successfully applied at a high-throughput scale for the dissection of the yeast and human interactomes (Ewing et al, 2007; Gavin et al, 2002), and also to the study of specific proteins of interests (Le Guezennec et al, 2006).

Epitope tags are short amino acid sequences that can be recognised by
specific immunoprecipitation-grade antibodies (i.e. FLAG, HA, MYC) or other affinity reagents (i.e. calmodulin-binding columns, streptavidin beads). The first step entails cloning the coding sequence of the target protein in frame with sequences coding for one or multiple tags in a suitable expression vector, often retroviral. Subsequently, viral particles are produced to transduce the target cells and polyclonal or monoclonal cultures with stable expression of the tagged protein are selected. The tagged protein of interest is purified from protein extracts together with its protein partners, which are subsequently identified by mass spectrometry (Figure 4). The list of interactors identified can be further validated by epitope-tagging of these proteins in a so-called “complex walking” approach. Importantly, proteins can take part to multiple complexes at the same time, but this cannot be inferred from a simple list of interactors. However, several lists of interactors generated by complex walking can be integrated to generate an interaction network. In this way, distinct complexes can be dissected and shared or specific subunits among these complexes can be identified.

**Figure 4**

**Discovery of protein interactors by complex purification and mass spectrometry.**

The main steps of biochemical protein-complex purifications and subsequent identification of the purified proteins by mass spectrometry are illustrated. First, a protein extract is made from a suitable cell system and antibodies (or other affinity reagents) are used to enrich for the protein complex of interest (in colours) and wash away unrelated proteins (grey circles). The proteins purified are separated by electrophoresis and subjected to enzymatic digestion with trypsin. The peptides obtained are identified by mass spectrometry, and data analysis generates a list of proteins specifically present in the original sample.
Importantly, this approach allows to deduce the function of unknown or predicted proteins, since proteins that interact with one another or are part of the same complex are often involved in the same cellular processes. If an uncharacterised target protein interacts specifically with a protein of known function, such function can be generally extended to the target protein of interest and follow-up experiments can be designed based on this data.

**Chromatin immunoprecipitation and high-throughput sequencing**

Chromatin immunoprecipitation (ChIP) is a powerful technique to study protein-DNA interactions in the context of chromatin *in vivo*. Genome-wide profiling of protein-DNA interactions (ChIP-seq) together with quantitative measurements of transcriptomes (mRNA-seq) are increasingly used to decipher the regulatory networks that underlie biological processes in health and disease (Park, 2009).

In a ChIP assay, proteins are covalently crosslinked to their binding sites on chromatin by incubation of the cells or tissues with formaldehyde (Figure 5). Crosslinked chromatin is then extracted from the cell nuclei and fragmented by sonication or enzymatic digestion into pieces of a few hundred base pairs. The fragmented chromatin is used as input in an immunoprecipitation reaction using antibodies against a target protein, i.e. a transcription factor, histone, or polymerase. In this way, the DNA associated to the protein of interest is enriched and purified. The presence of genomic DNA regions of interest (binding sites) can be determined by real-time quantitative PCR assays using specific primers. However, this approach is inherently limited to a priori knowledge or hypotheses on putative binding sites for the protein studied.

The introduction of microarrays (ChIP-chip) and more recently next-generation sequencing approaches (ChIP-seq) has enabled the discovery of unknown binding sites and the rapid and efficient mapping of protein-DNA interactions at the genomic scale in the same experiment. For ChIP-seq experiments on the Illumina sequencing platform, the DNA obtained from a ChIP experiment is subject to an adaptor ligation step, which adds specific sequences to both ends of the DNA molecules. After a PCR amplification step, the “sequencing library” is obtained. The library is hybridized on a glass surface via sequences present in the adaptors, and each molecule is subsequently amplified by isothermal bridge amplification, generating clusters of about 1000 identical molecules. The clustered flowcell is then
installed on the sequencer (Genome Analyzer or HiSeq) and subject to a sequence-by-synthesis protocol, generating millions of sequence reads in one experiment within a few days.

The ChIP-seq technique is characterized by a number of key features: first of all, ChIP-seq has single-nucleotide resolution, which allows the precise determination of a binding site and a more accurate discovery of relevant sequence motifs. Very low amounts of ChIP DNA (10-50 ng) and a limited number of PCR amplification cycles are required. Although the PCR can introduce some GC-bias, the background levels of a ChIP-seq experiment are significantly lower than ChIP-chip, and the dynamic range is theoretically unlimited. Taken together, these factors result in a very high sensitivity for this technique, which allows the identification of binding sites with moder-
ate enrichment that would not be detected with an array approach. This is not only relevant for the study of low-affinity binding sites, but also for the profiling of proteins that do not bind DNA directly but rather via protein-protein interactions and of proteins that establish only weak interactions with chromatin. The genomic coverage in ChIP-Seq is only limited by alignability of the sequence reads to the reference genome and not by probe design and cross-hybridization issues. If needed, the specificity of the alignments can be further increased by generating longer or paired-end reads. Moreover, ChIP-seq data can be reanalysed at a later stage if a new genome build becomes available. Finally, the steady drop in the costs of Next-Generation sequencing together with recent protocols allowing barcoding and multiplexing of samples have made ChIP-seq experiments more cost-effective, especially for higher eukaryotes and large genomes.

A critical aspect of ChIP-seq experiments is that a suitable ChIP-grade antibody against the protein of interest is needed. Nowadays, thousands of antibodies are available from many different commercial suppliers, but only a limited number of them works in native immunoprecipitation assays, and even fewer work in ChIP. Therefore, antibodies must be thoroughly tested before use in order to assess their specificity (cross-reaction with other proteins containing a similar epitope is a common problem) and sensitivity (signal vs. background in ChIP experiment). Therefore, proper negative controls are crucial in order to draw valid conclusions. Several different types of negative controls are being used in the literature, but not all of them have the same power. Some researchers compare the ChIP-qPCR recoveries of the antibody of interest with an unrelated antibody on the same genomic locus. While this type of negative control might establish a background level related to the primers used in qPCR, it does not address the specificity (i.e. cross-reactivity) of the antibody of interest or the background levels related to the antibody itself. Primers designed on genomic loci that are not bound by the protein of interest can better indicate background levels related to the antibody. Other useful controls to test the specificity of the antibody are experiments performed on cells that lack the protein of interest, for instance upon knockout or knockdown. Similarly, ChIP assays can be performed on epitope-tagged proteins in a cell line expressing the tagged protein while the wild type cell line is used as negative control. Alternatively, serum or antibodies harvested before the immunization protocol against the protein of interest (preimmune serum) can establish if the signal detected in ChIP is an aspecific signal relat-
ed to the animal used for the immunization, or rather a specific signal caused by antibodies raised in the immunization. A positive result can arguably represent a signal related to the protein of interest, but cross reactivity with proteins containing the same epitope cannot be formally excluded.

Next-generation sequencing approaches have been successfully applied also to genome-wide expression profiling (mRNA-seq) (Pepke et al, 2009). Also in this case, the larger dynamic range, higher sensitivity and the digital nature of the mRNA-seq data have constituted distinct advantages compared to array-based approaches. The increasing use and applications of these sequencing technologies and the large-scale data collection should form the basis for truly integrative systems biology analyses. Developments of the computation aspects will allow researchers to combine and correlate hundreds or thousands of protein binding profiles, transcriptomes, and protein-protein interaction datasets, with the final goal to decipher molecular regulatory pathways and eventually address important biological questions.
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IDENTIFICATION of protein interactors of the human oncoprotein PRAME by epitope-tag affinity purifications

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ABSTRACT

Preferentially expressed antigen of melanoma (PRAME) is frequently overexpressed in tumors and high PRAME levels correlate with poor clinical outcome of several cancers. However, the molecular mechanisms and pathways in which PRAME acts are still unclear. In order to investigate the molecular function(s) of PRAME, we set out to identify its protein interactors. To this end, we developed a biochemical protein-complex purification approach using epitope-tagged versions of PRAME and immunoprecipitation assays coupled to mass spectrometry analysis. In this chapter, we report the setup and optimization of these techniques, which led us to identify specific interactions between PRAME and Cullin2-based E3 ubiquitin ligases.
INTRODUCTION

The human protein PRAME (Preferentially expressed antigen in melanoma) is frequently overexpressed in cancer, and high expression levels often correlate with poor clinical outcome (Ikeda et al., 1997; Kilpinen et al., 2008; Oberthuer et al., 2004; Haqq et al., 2005). PRAME is the founding member of a large family of PRAME-like genes, characterized by a recent evolutionary history and a high degree of conservation (Birtle et al., 2005; Consortium et al., 2007).

Computational analysis of the human genome identified 22 PRAME-like genes and 10 pseudogenes (Birtle et al., 2005). Analysis of the primary amino acid sequence of PRAME revealed that it is a leucine-rich repeat protein (Epping et al., 2005; Birtle et al., 2005) and it was suggested that PRAME could fold into a horse-shoe structure (Birtle et al., 2005; Wadelin et al., 2010). However, defined functional motifs could not be identified and the molecular function(s) of PRAME and PRAME-like proteins are still unclear.

An experimental approach to gain insight into the function of an unknown protein is to identify its protein-protein interactors, since proteins that interact with one another or are part of the same complex are generally involved in the same cellular processes (Phizicky et al., 2003). If the interactors identified have a known function, this function can often be extended to the unknown protein and specific follow-up experiments can be designed. If the interactors identified are also uncharacterized proteins, inspection of their amino acid sequences might reveal functional domains or motifs that can eventually give clues to putative functions. A powerful approach to determine protein interactors and entire protein complexes involves the stable expression of epitope-tagged versions of the protein of interest in a suitable cell system using retroviral delivery. Subsequently, the protein and its interacting partners are biochemically purified using antibodies or affinity-based reagents which specifically and efficiently bind to the epitope tags, and the proteins are identified by mass spectrometry and database searches (Aebi-sold & Mann, 2003; Rigaut et al., 1999).

In this chapter, we describe the application of epitope-tagging and mass spectrometry analysis to the unbiased identification of the protein interactors of PRAME. We successfully setup a single-step protein complex purification protocol and discovered that the human oncoprotein PRAME establishes interactions with Cullin2-based E3 ubiquitin ligases.
RESULTS AND DISCUSSION

The ProtA-TEV-3xMyc epitope tag in FM6 cells

Since PRAME is overexpressed in the vast majority of human melanomas, we set out to identify its protein interactors applying a tap-tag (tandem affinity purification) approach in the human melanoma cell line FM6 (Rigaut et al., 1999). A modified pZOME retroviral system was used to generate FM6-pZ3XN-PRAME cell lines stably expressing PRAME fused to a combination of N-terminal tags: 2xProtA-2xTEV-3xMYC (ProtA-TEV-3xMYC-PRAME; 72 kDa) (Fig. 1A). A similar tag with a single MYC epitope was successfully used for the characterization of proteins complexes involved in transcriptional regulation (Le Guezennec et al., 2006).

The purification strategy includes a first incubation of protein extracts with crosslinked IgG beads for which the protein A domains have a high affinity (Fig. 1B). After incubation, several washes are performed to discard unbound proteins, and recombinant TEV (Tobacco Etch Virus) protease is added to cleave the tags at its recognition site. This proteolytic cleavage releases the bait protein and its interacting partners in solution. The last step is a MYC immunoprecipitation to further purify the complex.

Western blot of protein extracts from FM6-pZ3XN-PRAME cells showed a prominent band at the expected size, and a lower band with a significantly lower intensity (Fig. 1C). These extracts were used to optimize the conditions for TEV cleavage. The amounts of TEV required and the incubation temperature were tested in preliminary experiments (not shown). Western blot analysis of whole cell extracts after incubation with TEV showed that cleavage was most efficient after overnight incubation at 4°C (Fig. 1D, lanes 7 and 8).

Next, a small-scale purification experiment was performed to test the efficiency of IgG binding and TEV cleavage. Figure 1E shows that after incubation of protein extract with IgG beads, about 50% of ProtA-TEV-3xMYC-PRAME was bound to IgG beads (lane 4), while about 50% was still present in the supernatant (lane 2). When TEV protease was added to the beads and incubated overnight, however, almost no cleavage was detected in the eluates (lanes 6 and 9) and the intact protein was still bound to the beads. Independent experiments confirmed that TEV could efficiently cleave its recognition site in ProtA-TEV-3xMYC-PRAME from soluble protein extracts, but not when the tagged protein was bound to IgG beads.

A possible explanation for the inefficient cleavage is that the 27 kDa TEV protease might not be able to efficiently access its target sites when ProtA-TEV-3xMYC-PRAME is immobilized on IgG beads. In conclusion, although this epitope tag had
**TAP approach in FM6-p3XN-PRAME cells.**

(A) Schematic representation of ProtA-TEV-3xMYC-PRAME.

(B) Overview of the tandem affinity purification (TAP) procedure. The target protein is light blue, its protein interactors orange. The first purification is based on binding of the proteinA domains of the tagged protein to IgG beads. Most unbound proteins (in grey) are washed off and the target protein complex is released by TEV protease cleavage. The complex is further purified by a second purification using beads with Myc antibodies and elution with molar excess of MYC peptide (red square).

(C) Protein levels of PRAME in cell lines. Western blot analysis of endogenously and epitope-tagged PRAME. Left panel: FM6-p3XN-PRAME stable cell lines and HEK293T cells transiently transfected with pcDNA5-StrepII-2xTEV-MYC3xHA-PRAME (TAG-PRAME, 69 kDa) are analyzed with MYC antibody. Right panel: whole cell extracts from the cell lines indicated are probed with anti-PRAME antibody (P4-1:1000 dilution). Equal amounts of extracts were loaded for all samples, and this was checked by Ponceau S staining of the membranes.

(D) TEV cleavage on protein extracts. Whole cell extracts from FM6-p3XN-PRAME cells were subjected to TEV cleavage at 4°C for 4 hours or overnight. For each reaction, 18 μl of whole cell extracts were mixed with 59 μl IP150/0.1 buffer and 1 μl of TEV protease (preparation batch 14-2), or 2 μl of the same protease preparation diluted 1:1 in glycerol (TEV glyc), or 1 μl buffer (no TEV control). The effect of protease inhibitors was tested by incubating extracts with 1 μl TEV and 1x protease inhibitors cocktail. After incubation, 14 μl laemli buffer 4x was added and the sample boiled to stop the reaction; 25 μl of all reactions were analyzed by western blot with MYC antibody.

(E) TEV cleavage of epitope tagged PRAME bound on IgG beads is not efficient. IgG beads were incubated for 1h with PBS (TEV) or PBS with 1% BSA (No TEV and TEV+BSA). Beads were subsequently washed with IP150/0,1 buffer without protein inhibitors. 15 μl of beads were mixed with 100 μl IP150/0.1 and 50 μl of FM6-p3XN-PRAME whole cell extract, and rocked for 2h at 4 degrees. The supernatant was collected, and the beads washed with 500 μl of IP150/0.1. Elution was performed by adding 100 μl of IP150/0.1 and 2 μl of TEV protease with rocking at 4 degrees overnight. The supernatant (TEV eluate) was collected, and beads washed and boiled in 100 μl laemli buffer. 20% fractions of all samples were analyzed by western blot with MYC antibody.
**Figure 2**

(A) Map of the LZRS-ZeoR retroviral plasmid. The multi cloning site (MCS) is indicated in the square. The sequence coding for TAG-PRAME was cloned as a BamH1-XhoI fragment.

(B) Nucleotide and amino acid sequences of the StreptI-2x1EV-MYC-3xHA tag.

(C) Western blot analysis of single cell clones of HeLaS3-TAG-PRAME and K562-TAG-PRAME cells with HA.Y11 antibody against the HA TAG. Arrows indicate the two main bands for TAG-PRAME; asterisks indicate background bands. Clone numbers are indicated; WT, wild type cells (negative control); poly, polyclonal culture; ctrl, HEK293T cells transiently transfected with pcDNA5-TAG-PRAME. Equal amounts of whole cell extracts were loaded and checked by Ponceau S staining of the membrane (not shown).

(D) Gene expression analysis of PRAME by real-time RT-PCR. Left panel, cDNA prepared from the cell lines indicated was analyzed with two different primers for PRAME and the results normalized against GAPDH. Right panel, independent experiment with normalization against a pool of reference genes (GAPDH, ACTB, GUSB). HeLaS3 cells growing in suspension or adhesion (plate) were tested.
been successfully applied to the purification of protein complexes (Le Guezennec et al., 2006), it could not be applied to the analysis of PRAME protein interactors.

**The StrepII-2xTEV-MYC-3xHA epitope tag in K562 cells**

We next tested a different N-terminal combinatorial tag: StrepII-2xTEV-MYC-3xHA (further referred to as TAG) that could allow the use of several combinations of two-step purification strategies (Le Guezennec et al., 2006). Stable cell lines expressing StrepII-2xTEV-MYC-3xHA-PRAME (TAG-PRAME, 69 kDa) were generated using the retroviral LZRS-ZeoR vector in the human leukaemia K562 cells, which express PRAME at high levels, and the cervical cancer HeLaS3 cells, which express PRAME at very low to undetectable levels (Fig. 2A and B).

Single cell colonies were selected and tested for expression: K562-TAG-PRAME clone 12 and HeLaS3-TAG-PRAME clone 3 were chosen for further experiments (Fig. 2C). Gene expression analysis by real-time RT-PCR indicated that TAG-PRAME was expressed at the same levels as endogenous PRAME in K562-TAG-PRAME clone 12 cells, and about 3-fold higher in HeLaS3-TAG-PRAME clone 3 cells (Fig. 2D).

Preliminary results indicated that large amounts of protein extracts would be needed for purifications of sufficient quantities of PRAME and associated proteins. We therefore set up protocols for growth and harvest of suspension cells from multiple 5-liter spinner flasks in parallel. For this purpose, we optimized a nuclear/cytoplasmic protein extraction protocol for both K562 and HeLaS3 cells and we assessed the quality of all extractions by western blot analyses for the nuclear protein HDAC2 and the cytoplasmic marker α-tubulin. Representative experiments are presented in Figure 3A. TAG-PRAME was detected in both the cytoplasmic and nuclear fractions, similarly to the endogenous PRAME protein (not shown).

Notably, cytoplasmic extracts yielded on average between 3 and 4-fold more proteins than nuclear extracts (in mass). Therefore, for quality control of the protein extractions, equivalent fractions in volume - and not the same amount of micrograms - of nuclear and cytoplasmic extracts were analyzed by western blot. In this way, it is possible to directly compare the two fractions and assess the potential contamination of the cytoplasmic extract by nuclear proteins and vice versa.

**Two-step TAG-PRAME purifications**

At the time of our study, it was generally assumed that a single immunoprecipitation (IP) step does not sufficiently enrich for the bait, because too many aspecific interactors are still present in the eluates. This would result in a mixture
Tandem affinity purification approach from K562-TAG-PRAME cells.

(A) Western blot analysis of nuclear and cytoplasmic protein extracts from K562 cell lines. (B-D) Western blot analyses of pilot experiments for TAG-PRAME complex purification with a two-step IP approach; 3% of each sample were analyzed, unless indicated otherwise. All membranes were probed with mouse HA.Y11 antibody.

Figure 3

of proteins with a too high complexity to allow the successful identification of bait and interacting proteins by mass spectrometry. Therefore, most studies used a tandem affinity purification protocol in order to increase the purity of the protein complex of interest (Rigaut et al., 1999). In line with these studies, we performed several small-scale pilot experiments with tandem IPs, each one with two rounds of peptide elution. The two eluates from the first IP were pooled and used as input for a second IP with a different antibody. We tested three antibodies in different combinations: mouse MYC produced in house from hybridoma cultures, rabbit HA.
from Abcam (ab9110), and the PRAME affinity-purified antibody P4. Representative experiments are shown in Figure 3B-D: MYC-HA, HA-MYC, and P4-MYC purifications.

In order to assess the efficiency of the protocol, equivalent fractions for all steps were analysed by western blot: input (nuclear extract), supernatant after incubation of the extract with beads-Ab, first and second peptide eluates, and beads after elution. Comparison of the protein levels in input and supernatant gives information on the efficiency of the binding of the tagged protein to the beads, while comparison of eluates and beads gives information on the efficiency of elution.

Lanes 1 to 5 (Fig. 3B-D) show that TAG-PRAME could be efficiently bound and peptide-eluted with any of the three antibodies used in the first IP. Notably, later experiments revealed that peptide elution from the HA-ab9110 antibodies was only possible when using a specific lot of the antibody, and later lots did not allow peptide elution any more.

Although the first IP steps were efficient for all three antibodies, the final yields after the second purifications were considered too low to proceed with a large-scale purification (compare final eluates in lanes 7 with input levels in lanes 1). With this low efficiency, too large amounts of protein extracts and antibodies would be required to obtain enough purified proteins for detection by mass spectrometry.

**Single-step TAG-PRAME purification and mass spectrometry**

Based on the previous results, we set out to experimentally test whether a single IP purification approach could be applied to the identification of PRAME protein partners. In order to reduce the background levels, we used relatively stringent buffers to wash the antibody-beads after incubation with the protein extracts and before peptide elution. Figure 4A shows that increasing the salt concentration of the wash buffer from 150 to 300 mM reduced very significantly the number of background interactions detected by SDS-PAGE and silver stain of peptide eluates from a MYC IP on either K562-TAG-PRAME or K562 wild type extracts. A salt concentration of 500 mM NaCl did not improve the results further, and we therefore chose the 300 mM concentration for follow up experiments. Notably, we did not identify clear bands specific for the TAG-PRAME eluates, consistent with the notion that TAG-PRAME is present at very low abundance in such eluates as compared to background proteins.

Next, we performed a large-scale immunoprecipitation experiment with crosslinked MYC beads and 10 ml of nuclear extracts from K562-TAG-PRAME cells,
Single-step immunoprecipitations from K562 cells.

(A) Background levels in single-step IP protocol are decreased by 300 mM NaCl buffer. MYC IPs were performed with 300 μl nuclear extracts from the cell lines indicated and 20 μl MYC-beads. After incubation, the beads were washed with buffers of different salt concentration, as indicated. Peptide eluates were resolved by SDS-PAGE and protein bands visualized by silver staining.

(B) Western blot analysis of large scale MYC IP from K562-TAG-PRAME nuclear extracts. The fraction of samples analyzed is indicated. The mouse HA.Y11 antibody was used.

(C) Silver staining of eluates from large scale MYC IP. Both first and second eluates from large scale IPs on the nuclear extracts indicated were resolved on SDS-PAGE and proteins visualized by silver staining. Molecular weights (kDa) of the marker are indicated on the left.

**Figure 4**
corresponding to about 6 litres of culture and $6 \times 10^9$ cells. Immunoprecipitation from 4 ml of nuclear extracts of parental K562 cells (about 2.5 litre culture and $2.5 \times 10^9$ cells) was performed as negative control. Western blotting of an aliquot of the peptide eluates confirmed the efficiency of the IP protocol (Fig. 4B). We resolved the eluates on SDS-PAGE and visualized the proteins by silver staining (Fig. 4C). Several bands were identified, the most intense being present in both the TAG-PRAME and the negative control samples, representing background interactions with beads or antibodies. However, a number of fainter bands were specifically present in the TAG-PRAME eluates and not in the negative control. The entire gel lanes were excised in multiple fractions and processed for mass spectrometry analysis.

The peptides identified were used in a MASCOT database search and the proteins found in the TAG-PRAME and WT eluates were compared to generate a list of proteins specific for the TAG-PRAME sample (Table 1). PRAME was successfully identified with 7 different peptides, establishing that a single IP/MS approach can be successfully applied to the purification of PRAME protein complexes.

Remarkably, peptides were found in the TAG-PRAME eluates that matched all known subunits of Cullin2 E3 ubiquitin ligases (Petroski & Deshaies, 2005): Cullin2 (CUL2), Transcription elongation factor B polypeptide 1 (TCEB1, also called Elongin C), Transcription elongation factor B polypeptide 2 (TCEB2, also called Elongin B), and RING-box protein 1 (RBX1), suggesting a functional link between PRAME and this complex. These interactions were confirmed by western blot of independent immunoprecipitation experiments and were further characterized in other chapters of this thesis.

Taken together, our results show that a specific epitope tag may not always be useful for the purification of a protein of interest. The ProtA-2xTEV-MYC tag had been previously used for the purification of protein complexes (Le Guezennec et al, 2006), but in the case of PRAME the TEV cleavage step proved too inefficient. On the contrary, we successfully used the N-terminal StrepII-2xTEV-MYC-3xHA tag and identified PRAME interactors using a single-step purification approach with crosslinked MYC beads and peptide elution. We identified specific interactions between PRAME and all the known subunits of Cullin2-based E3 ubiquitin ligases, suggesting a functional link between ubiquitination pathways and the oncogenic properties of PRAME.
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**Table 1**

**PRAME putative protein interactors from the first large scale MYC IP.** The table lists all proteins identified with at least 2 peptides and a selection of proteins identified with only 1 peptide.
MATERIALS AND METHODS

Cell culture and plasmids
HEK293T and FM6 cells were cultured in DMEM, K562 and HeLaS3 in RPMI medium (Gibco, Invitrogen) at 37°C in 5% CO2. Both media were supplemented with 5% Glutamax, 10% fetal bovine serum, 100 U/ml of penicillin, 100 µg/ml of streptomycin (Gibco, Invitrogen). Stable cell lines were generated by retroviral transduction, as described (Costessi et al, 2011). Generation of plasmids pcDNA5-TAG-PRAME and LZRS(Zeo)-TAG-PRAME, and preparation of nuclear/cytoplasmic extracts are described elsewhere (Costessi et al, 2011).

Antibodies and western blot
Mouse monoclonal MYC (9E11) were produced in-house from hybridoma cultures; the PRAME antibody P4 was generated by peptide immunization of rabbits and affinity-purification. The following commercial antibodies were used: rabbit HA (Abcam ab9110), mouse monoclonal HA.Y11 (Covance MMS-101P; 1:3000 dilution for WB), alpha-tubulin (Santa Cruz B-7, sc-5286; 1:2000 dilution for WB), HDAC2 (Biomol SA-402; 1:3000–1:5000). Proteins were separated on conventional 10% SDS-PAGE gels and western blots were visualized by ECL (GE Healthcare) or Odyssey (LiCor). All antibodies were diluted in PBS/T with 5% skimmed milk for western blot analyses.

Tandem affinity purifications of StrepII-2xTEV-MYC-3xHA-PRAME
Pilot purifications with two tandem IP steps were performed on nuclear extracts of K562-TAG-PRAME cl.12 cells. For the first IP reaction, 300 µl nuclear extract were mixed with 700 µl IPP150/0.5 buffer (300 mM NaCl, 10 mM Tris-HCl pH 8, 0.5 mM EDTA, 0.5% NP-40) supplemented with protease and phosphatase inhibitors (1 mM PMSF, 1 mM Na3VO4, 1 mM NaF, 1x Protease Inhibitor Cocktail, Roche), and either 50 µl of PRAME P4 antibody, 9 µl HA antibody (Abcam ab9110) or 100 µl of crosslinked MYC-beads. After either 2 hours or overnight incubation, 100 µl of ProtG beads (Sigma) were added to P4 and HA IPs and incubated for 2 hours. Subsequently, the reactions were spun down and supernatant (containing unbound proteins) collected. The beads were washed 3 times with IPP150/0.5, and subjected to 2 round of peptide elution with 500 µl elution buffer (IPP150/0.5 with inhibitors and 100 µg/ml peptide) for 30 minutes in a thermoshaker at 28°C and 1000 rpm. After elution, the beads (containing bound proteins that were not eluted) were boiled in 250 µl laemli buffer. The two eluates from the first IP were pooled and used as input to the second IP, using half the amount of antibody and beads: either 50 µl MYC-beads, 4.5 µl HA antibody, or 25 µl P4 antibody. Incubation and elutions were performed as in the first IP step with 100 µl of elution buffer.

Large scale single-step MYC immunoprecipitation
Large scale IP was performed by mixing 10 ml nuclear extracts from K562-TAG-PRAME cells (from about 6 liter cultures) with 50 µl crosslinked MYC-beads, and 4 ml nuclear extracts from K562 wild type cells with 15 µl MYC-beads. After overnight incubation at 4°C in rotation, the beads were spun down and washed 3 times with 5 ml IPP300/0.5, and once with IPP150/0.5. Two consecutive peptide elutions were performed with either 100 µl (TAG-PRAME) or 30 µl (wild type) elution buffer, as described above.

Silver staining
The peptide eluates were resolved by SDS-PAGE and the proteins were fixed by incubating the gel in fixation solution (40:10:50 v/v/v ethanol: acetic acid: water) for 20-30 minutes at room temperature and gentle rocking. The gel was washed twice with washing solution (30% v/v ethanol in water) on a shaking platform for 20 minutes, rinsed twice with water, and incubated with water for 20 minutes on a shaking platform.

The gel was incubated in sensitizing solution (0.02% sodium thiosulfate) for 1-2 minutes, rinsed 3 times with water, and incubated for 30 minutes in chilled 0.1% AgNO3 solution at 4°C. The gel was quickly rinsed twice with water and incubated in developing solution (0.05% formaldehyde in 2% sodium carbonate). When a sufficient degree of staining was obtained the reaction was stopped with 5% acetic acid. The gel was washed several times in 1% acetic acid and stored in water.
Preparation of samples for mass spectrometry

The lanes from the SDS-PAGE gel corresponding to the first eluates were excised with a clean scalpel, chopped into small pieces and transferred into 1,5 ml tubes. The gel particles were equilibrated for 5 minutes with 100 µl ABC (50 mM ammonium bicarbonate) and centrifuged to discard the liquid. Then, 100 µl acetonitrile was added and incubated for 5 minutes to shrink the gel particles. The ABC and acetonitrile steps were repeated twice, or until the particles turned white. The gel pieces were swelled in 100 µl reduction buffer (10mM dithiothreitol in 50mM NH₄HCO₃) and incubated for 20 minutes at 56°C to reduce the protein. All liquid was discarded and the gel pieces were shrunk in 100 µl acetonitrile. The acetonitrile was replaced by 100 µl alkylation buffer (55 mM iodoacetamide in 50 mM NH₄HCO₃) and incubated for 20 minutes at room temperature in the dark. The gel particles were washed with 100 µl ABC (50mM Ammonium Bi Carbonate) for 5 minutes and ABC was replaced by 100µl acetonitrile; these steps were repeated twice. Gel pieces were rehydrated at room temperature in digestion buffer (50 mM NH₄HCO₃) containing 12,5 ng/µl trypsin (Promega). After 30 minutes, 10-50 µl ABC was added to cover the gel pieces and samples were incubated overnight at 37°C. Peptides were extracted from the gel matrix by adding 100 µl of 2% TFA (trifluoroacetic acid) for 10 minutes at room temperature; gel particles were spun down and the supernatant recovered. The peptide extraction was repeated a second time. Then, 100 µl acetonitrile was added for 5-10 minutes at room temperature to shrink the gel pieces, 100 µl 2% TFA was added for 5-10 minutes and supernatant was recovered after centrifugation. All samples were concentrated in a vacuum centrifuge (Savant) to a maximum of 200 µl, which was transferred to a stagetip. The peptides were eluted in 2-4 µl buffer B (80% acetonitrile, 1% TFA) and diluted 10 times in buffer A (3% acetonitrile, 0,5% acetic acid, 1% TFA). Mass spectrometry analysis was performed as previously described (Olsen et al, 2004).

ACKNOWLEDGEMENTS

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REFERENCES


CHAPTER 3

THE TUMOR ANTIGEN PRAME

is a subunit of a Cul2 ubiquitin ligase and associates with active NFY promoters


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ABSTRACT

The human tumor antigen PRAME (Preferentially expressed antigen of melanoma) is frequently overexpressed in tumors. High PRAME levels correlate with poor clinical outcome of several cancers, but the mechanisms by which PRAME could be involved in tumorigenesis remain largely elusive. We applied protein-complex purification strategies and identified PRAME as a substrate recognition subunit of a Cullin2-based E3 ubiquitin ligase. PRAME can be recruited to DNA in vitro, and genome-wide chromatin immunoprecipitation experiments revealed that PRAME is specifically enriched at transcriptionally active promoters that are also bound by NF-Y and at enhancers. Our results are consistent with a role for the PRAME ubiquitin ligase complex in NF-Y-mediated transcriptional regulation.
INTRODUCTION

*PRAME* is the founding member of a large family of PRAME genes (Birtle et al., 2005). PRAME genes have been implicated in oogenesis (Dadé et al., 2003), spermatogenesis (Wang et al., 2001) and stemness (Bortvin et al., 2003; Cinelli et al., 2008), but their function is poorly understood. Human PRAME was initially identified as the protein triggering an autologous anti-tumor immune response in a melanoma patient (Ikeda et al., 1997). Since then, many reports have shown that PRAME is not expressed in most normal adult tissues, while it is overexpressed in a wide variety of solid and hematological malignancies (Kilpinen et al., 2008). Although not prognostic for all cancers (Steinbach et al., 2002; Tajeddine et al., 2005; van Baren et al., 1998), high PRAME expression has been found to correlate with the stage of melanoma lesions (Haqq et al., 2005) and with shorter overall survival of neuroblastoma (Oberthuer et al., 2004) and serous ovarian adenocarcinoma patients (Partheen et al., 2008). High PRAME mRNA levels were also shown to be an independent prognostic factor of poor clinical outcome in breast cancer (Doolan et al., 2007; Epping et al., 2008). In addition, PRAME is significantly activated in BCR-ABL-positive chronic myeloid leukemia (Ph+ CML) patients at the onset of advanced phase and blast crisis (Radich et al., 2006), and hypomethylation of the PRAME locus seems to play a role in this process (Roman-Gomez et al., 2007; Schenk et al., 2007). A growing body of evidence indicates that PRAME contributes markedly to the oncogenic phenotype of several cancers. PRAME has been recently implicated in retinoic acid signaling and polycomb-mediated transcriptional repression in melanoma cells (Epping et al., 2005); however, these mechanisms do not seem to play a role in breast cancer (Epping et al., 2008) and acute myeloid leukemia (Steinbach et al., 2007). While evidence that expression of PRAME is restricted mainly to tumor cells has designated it as a promising therapeutic target, efforts to develop strategies to interfere with the oncogenic activity of PRAME are hindered by a lack of information about its function(s) (Paydas, 2008).

In this study, we combined biochemical and genomics approaches to identify PRAME as a subunit of Cullin2-based E3 ubiquitin ligases. In addition, we show that PRAME is recruited in cells to epigenetically and transcriptionally active promoter regions bound by NFY, a transcription factor that is essential for early embryonic development (Bhattacharya et al., 2003) and has been implicated in the maintenance of the high proliferative capacity of ES cells (Grskovic et al., 2007) as well as in the inhibition of differentiation (Gurtner et al., 2003).
RESULTS

PRAME associates with Cullin2-based E3 ubiquitin ligases

To investigate the molecular function of human PRAME, we used a proteomic approach to identify PRAME-interacting proteins. To this end, we stably expressed epitope-tagged versions of PRAME suitable for protein purifications in two cellular backgrounds: human K562 leukemia cells, which endogenously express detectable levels of PRAME, and human embryonic kidney HEK293 cells, which lack detectable endogenous PRAME (Fig. 1A). When analyzed by size exclusion chromatography, both endogenous and epitope-tagged PRAME from K562 cells showed a broad elution profile, with a substantial fraction of the protein eluting earlier than expected of free PRAME, which has a predicted molecular mass of 58 kDa (Fig. 1B). This observation suggests that PRAME may be part of one or more protein complexes.

We purified PRAME-associated proteins from K562-TAG-PRAME or HEK293-HA-PRAME cells by anti-epitope immunoaffinity chromatography and analyzed them by mass spectrometry. As shown in Tables 1 and 2, eluates from K562-TAG-PRAME and HEK293-HA-PRAME included a collection of proteins not found in control samples. Among the most highly represented were Cullin family member 2 (Cul2), Elongins B (TCEB2) and C (TCEB1), and RING finger protein Rbx1, all of which are subunits of the large family of multisubunit, Cullin-based, ubiquitin ligases. In these ubiquitin ligases, the Elongin BC dimer acts as an adaptor that links an Elongin BC-box containing substrate recognition subunit, or BC-box protein, to heterodimeric modules composed of either Cullin 2 and Rbx1, or Cullin 5 and Rbx2 (Kamura et al., 2004; Kamura et al., 1998; Lonergan et al., 1998). The RING finger domains of the related Rbx1 and Rbx2 proteins in turn recruit and activate an E2 ubiquitin conjugating enzyme for ubiquitination of substrates (Kamura et al., 1999; Petroski & Deshaies, 2005).

The interaction of TAG-PRAME with ubiquitin ligase subunits was confirmed by western blotting of TAG-eluates. As shown in Figure 1C, these fractions contained PRAME, Elongins B and C, and two forms of Cul2 that most likely correspond to unmodified and a more slowly migrating Neddylated-Cul2, a hallmark of active ubiquitin ligases. Supporting the physiological relevance of these interactions, endogenous PRAME, Cul2, Elongin B, and Elongin C were all detected in anti-PRAME immunoprecipitates from nuclear extracts of parental K562 cells (Fig. 1D). Notably, while PRAME was clearly
**PRAME assembles with Cullin2 ubiquitin ligases.**

(A) Ectopic expression of tagged versions of PRAME suitable for purifications. Whole-cell extracts of K562 cells or HA-immunoprecipitates from 293 cells were analyzed by western blot with anti-PRAME antibodies. K562 cells expressing StrepII-MYC-3xHA-PRAME (TAG-PRAME) at near-endogenous levels were selected, and 293 Flp-In cells stably expressing HA-PRAME were generated using the Invitrogen Flp-In system.

(B) Superose 6 gel filtration analysis of nuclear extracts from K562-TAG-PRAME and K562 wild type cells. Western blot was performed with anti-HA or affinity-purified anti-PRAME antibodies. Both TAG-PRAME and endogenous PRAME partially elute in high-molecular weight fractions. The numbers represent eluates fractions. A less prominent and faster migrating band for TAG-PRAME was detected, that most likely corresponds to a C-terminal truncation.

(C) TAG-PRAME co-immunoprecipitates Cullin2-EloBC complex. Nuclear extracts from K562 wild type and K562-TAG-PRAME cells were subjected to anti-TAG immunoprecipitation and western blotting as indicated; 0.8% of input and 33% of IP were loaded. A more slowly migrating form of Cullin2 was enriched after IP; this band is consistent with being post-translationally neddylated Cullin2. Asterisks indicate antibody heavy chains.

(D) Endogenous PRAME interacts with Cullin2-EloBC complex. Nuclear extracts from K562 wild type cells were subjected to immunoprecipitation with anti-PRAME or pre-immune sera and western blotting as indicated; 1% of input, 1% of supernatant and 20% of IP were loaded. PRAME was detected using a light-chain specific secondary antibody. PRAME was readily depleted from the extract after incubation with the specific antibodies (compare lanes 1, 2 and 4). Asterisks indicate protein A that disassociated from the beads after elution; the open arrow indicates PRAME, s/n, supernatant, IP, immunoprecipitates.
depleted from the extract after incubation with the specific antibodies (Fig. 1D, compare lanes 1, 2, and 4), Cul2, Elongin B, and Elongin C were not, consistent with their function as components of the many Cullin-based ubiquitin ligases present in cells (Kamura et al, 1998; Mahrour et al, 2008).

**PRAME is a BC-box protein**

Although we identified all of the other subunits of Cul2-based ubiquitin ligases as PRAME-interacting proteins, we did not detect any of the previously identified BC-box proteins in our fractions. BC-box proteins are characterized by a sequence motif of about 10 amino acids, called the BC-box, that binds to Elongin C. In addition, a downstream Cullin-box specifies binding to either Cul2 or Cul5 (Kamura et al, 2004; Mahrour et al, 2008). Multiple se-
**Figure 2**

| A | PRAME   | NP_996839 | 23 | PRLQLDAGSRLKDE---4---AALRELPRVFF   | 57 |
|   | VHLhs   | NP_000542 | 155 | YVRKMIQLQVRSVL---4---YRLDFVREYEDIE | 189 |
|   | FEM1B   | NP_056137 | 594 | KMDKLCFAARGAND---1---NYQDQTPRTEEFG | 625 |
|   | ZYG11BL | NP_006327 | 6  | PEELMACTDFCLRNL---15---HPDIFLPSECDRW | 51  |
|   | APBBF2  | NP_006371 | 10 | PEELNTYRISAQVDNY---4---RGIRSEILNQFDY | 44  |
|   | KLHDC2  | NP_055130 | 361 | PKSLVRLSLEAVICFK---4---NSWNCLPKHLLHSVN | 395 |
|   | KLHDC3  | NP_476502 | 335 | SPSLKTLCKLAVIQYN---1---LDQSCLPHDIRWELN | 365 |

**BC-box**

| B | FLAG-PRAME WT | - + - |
|   | FLAG-PRAME mBC | - - + |
|   | His-Cul2 | + + + |
|   | EloBC | + + + |
|   | FLAG-PRAME | + + + |
|   | Cul2 | |
|   | EloC | |

**Cul2-box**

| C | TAG-PRAME: | INPUT | TAG IP |
|   |  | WT | mBC | WT | mBC |
|   | FLAG IP | Cu2 | EloB | EloC |

**D**

| D | ATP | - + + + + |
|   | E1+E2 | + - + + + |
|   | Flag-PRAME+EloBC | + + - + + |
|   | His-CUL2+Myc-RBX1 | + + + - + |

**E**

**PRAME is a BC-box substrate receptor for Cul2-EloBC ligases.**

(A) Multiple sequence alignment of known Elongin BC-associating proteins with PRAME. Sequences were aligned using the blosum62mt2 alignment matrix with the AlignX program of the Vector NTI software package. Color coding is based on the AlignX default similarity table. Identical amino acids are highlighted in yellow, very similar ones in blue, and similar amino acids are shown in green. The canonical BC-box and Cul2-box sequences are indicated below the alignment. (B) PRAME has a functional BC-box. SF21 cells were co-infected with the baculoviruses as indicated. Total cell lysates and anti-FLAG immunoprecipitates were analyzed by western blotting. IP, immunoprecipitation; mBC, BC-box mutant. (C) BC-box defective PRAME lacks interaction with endogenous Cul2-EloBC ligases. Immunoblot analysis of 12CAS HA immunoprecipitates from nuclear extracts of K562 cells stably expressing TAG-PRAME wild type or TAG-PRAME BC-box mutant (PRAME mBC). Asterisks indicate antibody heavy chains. (D) The Cul2-PRAME ligase has in vitro ubiquitination activity. Epitope-tagged subassemblies of the PRAME Cul2-based ubiquitin ligase complex were purified from insect cells and used in an in vitro ubiquitin assay (as described in Methods). The results were analyzed by western blot using rabbit anti-Flag and mouse anti-ubiquitin antibodies. (E) Model summarizing Cul2-EloBC-PRAME interactions.
sequence alignments of PRAME with several known BC-box proteins identified putative BC- and Cul2-boxes near the PRAME N-terminus (Fig. 2A), raising the possibility that PRAME might function as a substrate recognition subunit of a Cul2-based ubiquitin ligase.

To assess the functionality of the predicted motifs, we introduced point mutations in key residues of the putative BC-box (L26P/A30F) or the putative Cul2-box (L48A/P49A). We observed that the BC-box mutation greatly reduced the binding of PRAME to Elongin C and Cul2 in baculovirus-infected insect cells (Fig. 2B). We next generated K562 and HEK293 cells stably expressing epitope-tagged PRAME carrying the BC-box or Cul2-box mutations and performed immunoprecipitations from nuclear extracts. In these cells, both mutations disrupted the interaction of PRAME with endogenous Cullin2 complex components (Fig. 2C and Table 2), demonstrating the functionality of the motifs.

In order to test the biochemical activity of the Cullin2-PRAME complex, we performed in vitro ubiquitination assays using purified recombinant proteins. As shown in Fig. 2D, we observed significant synthesis of poly-ubiquitin chains only when all the subunits were combined in the presence of ATP, confirming that the Cullin2-PRAME complex is a catalytically active ubiquitin ligase.

Taken together, these results indicate that PRAME’s BC- and Cul2-boxes are essential for the assembly of a PRAME-containing Elongin BC ubiquitin ligase, and they identify PRAME as a BC-box substrate recognition subunit of a Cul2-based ubiquitin ligase (Fig. 2E).

**PRAME associates with chromatin**

Cell fractionation and cesium chloride gradient purification of cross-linked chromatin from K562 cells showed that PRAME associates with chromatin (data not shown). In addition, we observe that PRAME can be recruited to DNA in vitro. Consistent with the absence of recognizable DNA binding domains in PRAME and other PRAME complex subunits, purified PRAME complexes did not bind directly to DNA immobilized on agarose beads. When incubated with bead-bound DNA in the presence of nuclear extract, however, PRAME complexes were retained on the DNA beads, suggesting that activity(s) in the extract are needed to bridge PRAME’s interaction with DNA (Suppl. Fig. 1A).

To investigate the chromatin-binding profile of PRAME, we performed ChIP assays followed by deep sequencing (ChIP-seq) using anti-PRAME
Genome-wide profiling of PRAME association with chromatin.

(A) Overview of PRAME binding sites. UCSC genome browser views of ChIP-seq profiles generated with preimmune (negative control) and PRAME antibodies in K562 cells. Specific PRAME peaks are detected at the promoters of the DNM2, EIF2AK3, TOP2A, and MYST2 genes.

(B) Genomic distribution of PRAME binding sites (left) compared to the distribution of genomic DNA (right). The majority of sites (43%) are located within ±1kb from a transcription start site. (C) Promoter binding of PRAME occurs at the TSS. Frequency plot of the distances of PRAME binding sites from TSS shows that 83% of PRAME promoter sites are located between -400 bp and +100 bp, with a peak at -87bp.

(D) Motif analysis identifies a CCAAT motif under PRAME promoter peaks. Top, logo of the JASPAR motif matching the most enriched sequence identified by GenomeMotifs. Bottom, the positional preference plot represents the frequency of the locations of the CCAAT motifs with respect to the summits of PRAME promoter summits, and indicate that the CCAAT motif is enriched at the center of PRAME binding regions.
antibodies or pre-immune serum as control for the specificity of the signals obtained. Overlapping sequence tags were joined and displayed as tag-density profiles in the UCSC genome browser. Strikingly, inspection of the profiles revealed that the promoter regions of many protein-coding genes had specific enrichment for PRAME, as exemplified by the screenshots in Figure 3A. Validation experiments by ChIP-qPCR confirmed the high quality of the sequencing data (Suppl. Fig. 1B). Furthermore, ChIP-seq with stably expressed, epitope-tagged PRAME resulted in very similar if not identical patterns of PRAME enrichment (data not shown). To substantiate our finding, we used the MACS algorithm (Zhang et al., 2008) at a false discovery rate of $10^{-6}$ and identified a total of 14132 PRAME binding sites. Location analysis showed that 43% of the PRAME sites are closer than 1 kb upstream or downstream of a transcription start site (TSS) (Fig. 3B). Importantly, promoters were 9-fold enriched as compared to the background distribution of genomic DNA.

Recently, Heintzman et al. identified enhancer elements in K562 cells based on ChIP-on-chip histone modification signatures and binding of the enhancer-associated acetyltransferase p300 (Heintzman et al., 2009). Most enhancers were cell type specific, and they were enriched in genomic domains containing genes specifically expressed in K562 cells. When we compared the binding profile of PRAME to the predicted enhancers we found a significant overlap between promoter-distal PRAME binding sites and enhancer regions: 44% of the distal PRAME sites overlapped predicted enhancers, whereas only 4.1% of random genomic sites did (11-fold enrichment). Moreover, similarly to the annotated enhancers, the distal PRAME sites were not randomly distributed in the genome, but clustered in the vicinity of PRAME-bound promoters (Suppl. Fig. 1C), suggesting possible promoter-enhancer interactions.

**PRAME associates with NF-Y promoters**

We next analyzed all PRAME binding sites falling within 1 kb of promoters. When we plotted their distance to the closest TSS, we found that 83% of promoter-proximal sites bound by PRAME are concentrated in a narrow window between -400 bp and +100 bp from TSS, peaking around -87 bp relative to the TSS.

Promoters are characterized by the presence of transcription factor binding sites that critically contribute to the regulation of gene expression. Since neither PRAME, nor other PRAME complex subunits contain recognizable DNA-binding domains, we reasoned that other transcription factor(s) might mediate the recruitment of PRAME to these regions. We therefore performed
de novo motif discovery with GimmeMotifs (van Heeringen & Veenstra, 2010) using the DNA sequences underlying the promoter regions bound by PRAME. This analysis identified a CCAAT sequence as the top-scoring motif, which stood out with highly significant p-values ($p<10^{-20}$) and the highest enrichment and MNCP scores as compared to a pool of random promoters (Fig. 3D and Suppl. Fig. 2). The CCAAT sequence can be bound by the Nuclear Transcription Factor Y (NFY), a heterotrimeric transcription factor ubiquitously expressed in cycling cells (Maity & de Crombrugghe, 1998). The NFYA subunit harbors the DNA-binding domain that recognizes the CCAAT sequence, while NFYB and NFYC are characterized by histone-like folds (Maity & de Crombrugghe, 1998).

Similar de novo motif analysis performed on the distal PRAME binding sites identified two significant motifs: a GATA/Evi1 motif, and a Fos motif. ChIP-seq experiments with antibodies against the NFYA and NFYB subunits revealed that PRAME sites at promoters are indeed bound by NFY in vivo. Screenshots of the sequencing data show clear enrichment of NFY at the representative PRAME sites DN2 and EIF2AK3, as well as PRAME enrichment at the classical NFY promoters TOP2A and CDC25C (Fig. 4A). Consecutive ChIP experiments (ChIP-re-ChIP) using crosslinked NFYA antibodies detected PRAME and NFY concomitantly on the same regions, indicating that these proteins are simultaneously localized at the same promoter regions (Fig. 4B).

Using the MACS algorithm we identified a total of 28844 regions that were bound by NFYA and/or NFYB. The distance distributions of both NFYA and NFYB binding sites from TSSs showed a distinct peak around -78/-84 bp very similar to PRAME (Fig. 4C). We next counted the number of sequence tags within NFY-binding sites and performed a regression curve analysis (Suppl. Fig. 3). The binding of NFYA and NFYB were very closely correlated at the vast majority of the sites, as expected for stoichiometric binding and in agreement with in vitro experiments showing that all subunits are required for efficient DNA binding (Sinha et al, 1995).

Globally, 71% of all PRAME binding sites overlapped NFY regions. Importantly, virtually all PRAME-bound promoters were also bound by NFY, corroborating and extending the relevance of the CCAAT motif (Fig. 4D). In contrast, a sizeable number of NFY promoters did not contain PRAME peaks (Fig. 4D). We performed functional annotation of the associated genes and found that NFY-only genes as well as NFY genes with PRAME enrichment are associated with metabolic and cancer-related pathways. Furthermore, NFY-only genes are specifically enriched for immunological processes and intercellular junc-
**PRAME associates with NFY-bound promoters genome-wide.**

(A) Overview of NFY binding to PRAME loci. UCSC genome browser views show clear binding of PRAME, NFYA, and NFYB at the promoters of the *DNM2, EIF2AK3, TOP2A*, and *MYST2* genes. 
(B) Sequential ChIP (ChIP-re-ChIP) experiments show co-occupancy of PRAME and NFY. The first ChIP was performed with covalently crosslinked NFYA-beads, followed by either anti-PRAME serum (NFYA-PRAME) or preimmune serum as negative control (NFYA-preimmune). 
(C) Promoter binding of NFY and PRAME occurs at the TSS. Frequency plot as in Fig. 3C shows very similar distributions of distances from TSS for NFY and PRAME. 
(D) PRAME associates with a subset of NFY-promoters genome-wide. Venn diagram of the overlap between NFY-bound (green) and PRAME-bound (blue) promoters shows that PRAME promoters overlap almost completely with NFY promoters. On the contrary, a minority of NFY promoters are not bound by PRAME. An example is the *TSPAN15* gene (right).
tions, while NFY genes with PRAME binding are significantly associated with numerous transcription-related pathways and cell cycle regulation (Supplementary Table 1).

**PRAME is specifically recruited to transcriptionally active NFY promoters**

Post-translational modifications of the histone proteins, like lysine methylations and acetylations, are implicated in the regulation of gene transcription. Therefore, we used histone marks to assess the transcriptional status of PRAME loci. For this purpose we processed ChIP-seq datasets generated by the ENCODE project for K562 cells (see methods), and we quantitated the signals at all NFY promoters. We found that the active histone marks H3K9ac and H3K4me3 showed a strong correlation with the binding of PRAME (Fig. 5A). In contrast, NFY-only promoters had near-background levels of these marks.

To investigate the transcriptional status of the associated genes, we quantitated the occupancy in gene bodies of RNA Polymerase II and of the histone marks H3K36me3, which has been linked to transcriptional elongation (Krogan *et al.*, 2003), and H3K27me3, which has been linked to repression (Barski *et al.*, 2007). As shown in Figure 5B, NFY genes without PRAME binding displayed low levels of PolII and H3K36me3, and high levels of the repressive H3K27me3 mark. In contrast, PRAME-bound genes had significantly higher occupancy of both PolII and H3K36me3 (Fig. 5B and D, and Suppl. Fig. 4), suggesting that PRAME is associated specifically with transcriptionally active genes. To substantiate these findings, we analyzed gene expression levels by RNA-seq in our K562 cells. In agreement with the ChIP-seq data, we detected few, if any, transcripts from NFY-only genes, while PRAME-bound NFY genes were expressed at high levels (Fig. 5C).

We next tested if components of the Cullin2 complex are present together with PRAME on chromatin. In the absence of ChIP-grade antibodies, we established stable K562 cell lines expressing Elongin C tagged with a double TY1 epitope (TTE-EloC), for which a ChIP-grade antibody was validated in our lab. ChIP-qPCR experiments showed specific binding of EloC at all the PRAME promoters tested (Fig. 5E). As a control, we performed similar experiments with TTE-PRAME.

Taken together, our data supports a model in which a Cul2/EloBC/PRAME complex is specifically bound to the subset of epigenetically and transcriptionally active NFY promoters and nearby enhancer regions.
**Figure 5**

**A**

<table>
<thead>
<tr>
<th>Tag density (Tags/1kb)</th>
<th>H3K9ac</th>
<th>H3K4me3</th>
</tr>
</thead>
<tbody>
<tr>
<td>No PRAME</td>
<td></td>
<td></td>
</tr>
<tr>
<td>PRAME</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Tag density (Tag density) on H3K9ac and H3K4me3 modifications at NFY promoters.

**B**

![Box plots of tag density for H3K36me3, RNA Pol II, and H3K27me3](image)

**C**

![Gene expression (NE) plots](image)

**E**

![Gene expression (NE) plots for different conditions](image)

**PRAME discriminates between active and inactive NFY loci.**

(A) PRAME binding positively correlates with H3K9ac and H3K4me3 modifications at NFY promoters. Left, heat map representing median tag densities (tags/kb) of H3K9ac and H3K4me3 at promoter-associated NFY binding sites without (“No PRAME”) or with (“PRAME”) PRAME binding. Higher levels of both histone marks are found if PRAME is present. Right, UCSC genome browser views of two representative genes: TSPAN15 and PHB.

(B) PRAME binding of NFY promoters correlates with high H3K36me3 and RNA Pol II, and with low H3K27me3 in the gene body. Box plots show the tag density (tags/kb) over the bodies of the genes associated with NFY promoters without or with PRAME binding.

(C) PRAME binding correlates with high levels of gene expression. Box plot of Normalized Expression values computed from RNA-seq experiments show that PRAME-bound NFY genes are expressed at higher levels than those without PRAME.

(D) Genome browser view of the genome-wide data combined. Two neighboring tail-to-tail NFY genes are shown: SPRYDA is bound by PRAME and is active, while GLS2 is not bound by PRAME and is in a repressed state.

(E) Model summarizing the genome-wide data.
DISCUSSION

Our proteomic data reveals that the human oncoprotein PRAME is a BC-box substrate recognition subunit of a Cul2-based E3 ubiquitin ligase, suggesting a functional link between ubiquitination and the oncogenic properties of PRAME. Notably, the Von Hippel-Lindau tumor suppressor protein is one of the best-characterized BC-box proteins, and its tumor suppressor activity is due in part to its role as the substrate recognition subunit of a Cul2-based ubiquitin ligase (Kaelin, 2007). Intriguingly, all the annotated PRAME-like proteins display a complete conservation of the BC-box and Cul2-box at their N-terminus (Suppl. Fig. 5), suggesting that all members of the large PRAME protein family might function as substrate recognition subunits of Cul2-based ubiquitin ligases, possibly displaying different degrees of specificity and/or redundancy in substrate selection.

To date, ubiquitination has been demonstrated to play important roles in gene regulation, affecting several different steps in the transcriptional process (Collins & Tansey, 2006; Conaway et al, 2002). Many activators are subject to ubiquitin-mediated degradation at a rate that parallels their transactivation abilities (Salghetti et al, 2001). Histones, transcription factors, and RNA polymerase II (RNAPII) are targets of ubiquitination, and ubiquitination...
events are required for the initiation-competent RNAPII complex to exchange factors and become competent for cotranscriptional mRNA processing (Muratani et al., 2005). A recent study indicated that the stability of NFYA can be modulated by ubiquitination and degradation via the proteasome, and that degradation-resistant NFYA mutants increased cell proliferation (Manni et al., 2008). Interestingly, some of the lysine residues in NFYA that are target of ubiquitination could also be acetylated by p300, suggesting a regulatory competition between these post-translational modifications (Manni et al., 2008).

Our genome-wide analyses reveal that PRAME is strongly enriched at NFY promoters, and binding of PRAME predicts a permissive epigenetic landscape and transcriptional activity at these regions. Moreover, we found that PRAME is also significantly enriched at adjacent enhancer regions, suggesting that it could participate in promoter-enhancer interactions. Recently, Ceribelli et al. reported a ChIP-on-chip analysis of NFYB in HeLa cells with 2.2% coverage of the human genome. Similar to our K562 data, the majority of the NFY-bound promoters identified in HeLa cells were transcriptionally active and carried H3K9-14ac and/or H3K4me3 marks, while about 27% of NFY-bound promoters were devoid of these active marks and mostly silent. Further experiments implicated NFY not only in the expression of active genes, but also in the repression of silent ones, revealing an unexpected dual nature of the NFY complex (Ceribelli et al., 2008). The basis of this dual functionality, however, is still unclear, and genome-wide data suggest that the activity and inactivity of NFY promoters is determined by additional factors and mechanisms that are yet to be identified.

PRAME is the first factor reported to discriminate between active and inactive NFY loci. Our genome-wide analyses, summarized in Figure 5F, reveal that PRAME is specifically recruited to NFY loci associated with the promoters of genes that are highly transcribed and that have high levels of H3K9ac and H3K4me3 at their promoters and high H3K36me3 but low H3K27me3 levels in their gene bodies. This data suggest a role for PRAME in transcriptional regulation, and possibly in the switch between activity and inactivity of NFY loci. It is tempting to speculate that PRAME might mediate ubiquitination, and possibly degradation, of one or more of the factors that are involved in the regulation of NFY promoters. Despite evidence that PRAME and NFYA can co-occupy the same genetic loci (Fig. 4B), we have not been able to detect stable and direct interactions between PRAME and NFY subunits by coimmunoprecipitation following their expression in transfected mammalian cells or
in baculovirus-infected insect cells (data not shown). We have also observed that PRAME can be recruited in vitro to a DNA fragment containing the NFY-dependent HSPA5 promoter by a mechanism that depends on activity(s) present in a nuclear extract. It seems likely that PRAME-NFY interactions, if they exist, occur only on DNA and depend on additional, yet to be identified factors, since (i) purified NFY is not sufficient to recruit PRAME to HSPA5 promoter DNA, and (ii) supplementing nuclear extract with exogenously added NFY does not further enhance PRAME recruitment (Suppl. Figure 1A).

Interactions have been reported between NFY and the histone acetyltransferases GCN5 and PCAF (Currie, 1998), and single-gene studies showed that NFY can activate at least some of its target genes in cooperation with the acetyltransferase p300, which is able to acetylate histone tails as well as NFY subunits (Caretti et al., 2003; Gurtner et al., 2008; Li et al., 1998; Salsi et al., 2003). While these acetyltransferases constitute potential targets of PRAME, other unknown factors could also take part to these mechanisms. Notably, p300 is not only recruited to promoters, but it is also a hallmark of enhancer elements. Heintzman and coworkers used p300 ChIP-on-chip binding profiles to identify an enhancer-specific chromatin signature, which was successfully used to predict enhancer elements in several cell lines, including K562 (Heintzman et al., 2007). Significantly, we found that about half of the promoter-distal PRAME sites are localized at predicted enhancers, suggesting that PRAME might be implicated in promoter-enhancer crosstalk.

In an effort to gain insight into PRAME’s potential role in transcription regulation, we knocked down PRAME with retroviral shRNA vectors and measured the mRNA levels of several target genes. Despite a 70% decrease of PRAME transcripts, we did not observe reproducible changes in the transcript levels of the target genes tested (data not shown). This is perhaps not surprising given that PRAME is a member of a large family of closely related proteins that could have overlapping or redundant functions in cultured cells. Further, the fact that the PRAME gene family underwent an extraordinary expansion specifically in the rodents and primates (Birtle et al., 2005; Consortium et al., 2007) suggests that they are more likely to fine-tune transcriptional regulatory processes rather than to be essential for transcription. Regrettably, the lack of a one-to-one orthology between the human and mouse PRAME genes (Birtle et al., 2005) has thus far precluded the development of suitable knock-out mouse models for studying PRAME function.
Notably, it is becoming clear that many proteins associate with the promoter regions of genes, but their depletion does not seem to cause significant or widespread transcriptional defects. For example, the yeast Set1 histone methyl transferase is widely associated with active promoters, but deletion of the Set1 gene causes expression changes of 1.5-fold or more in only about twenty genes (Miller et al., 2001). Similarly, the Jmjd3 histone demethylase was recently shown to associate with the promoters of LPS-induced genes, but the expression of most of them was unaffected by knock-out of Jmjd3 (De Santa et al., 2009). These observations are consistent with the model that some factors are not strictly required for the bulk of transcription, but instead have ancillary or regulatory functions that become apparent only in very specific conditions or subpopulations of cells.

While numerous reports established a role for PRAME in human cancers, several studies indicated that PRAME-like genes have roles in the early stages of spermatogenesis (Wang et al., 2001) and oogenesis (Dadé et al., 2003), as well as in embryonic development and embryonic stem cells (Bortvin et al., 2003). Recently, overexpression of Pramel7 in E14 embryonic stem cells was reported to maintain a pluripotent state in the absence of the anti-differentiation factor LIF (Cinelli et al., 2008). These findings are consistent with the notion that stem cells and neoplastic tissues share many properties, and several oncogenic pathways were recently found to also regulate self-renewal mechanisms in stem cells (Reya et al., 2001). Interestingly, recent studies suggest important roles for NFY in stem cell biology. Knock-out mice demonstrated that NFYA is essential for early mouse development (Bhattacharya et al., 2003), and NFYA was found to promote the self-renewal of hematopoietic stem cells (Zhu et al., 2005). Moreover, a systematic analysis of cis-regulatory elements identified the CCAAT box and NFY as crucial regulators of both mouse and human stem cells-specific genes, and NFY was involved in the maintenance of the proliferative capacity of ES cells (Grskovic et al., 2007).

Taken together, our data unambiguously shows that the PRAME oncoprotein is part of a Cullin2-based ubiquitin ligase complex and that PRAME localizes to active NFY promoters, paving the road for understanding the functions of PRAME in normal cells and in human malignancies. Future experiments will address the role of the PRAME ubiquitin ligase at TSSs, its role in transcription of NFY-regulated genes, and its contribution to the oncogenic properties of PRAME.
MATERIALS AND METHODS

Cell culture and stable cell lines
HEK293 cells were cultured in DMEM and K562 in RPMI medium (Gibco, Invitrogen) at 37°C in 5% CO2. Both media were supplemented with 5% Glutamax, 10% fetal bovine serum, 100 U/ml of penicillin, 100 µg/ml of streptomycin (Gibco, Invitrogen). Stable HEK293-HA-PRAME cells were generated using pcDNA5/FRT/HA-PRAME and the Flp-in system (Invitrogen). K562 stable cell lines were generated with retroviral constructs according to the following 5-days protocol. On the first day, HEK293T cells at 50-60% confluence in a 6-cm dish were transfected using Lipofectamine2000 (Invitrogen) and 1,8 µg of MD-MLV, 300 ng VSV-G, 3.6 µg retroviral construct, 100 ng pGFP. After 24 h (day 2), the virus-containing supernatant was harvested, centrifuged for 10 min at 3000 rpm and used for a first round of transduction of 10^6 K562 cells in the presence of 8 µg/ml of Polybrene (Sigma) for 24 h. A second round of transduction was performed on day 3, and on day 4 the target cells were transferred to fresh RPMI and selection was started on day 5 with either 6-10 µg/ml of puromycin (Sigma) or 1-2 mg/ml Zeocin (Invitrogen).

Antibodies and western blot
PRAME antisera and affinity-purified antibodies were generated by immunizing rabbits with the previously described peptide FPEPEAAQPMTKKRKVDG (Epping et al., 2005) (V27 serum). Mouse monoclonal HA-12CA5 and Myc-9E11 were produced in-house from hybridoma cultures. Commercial antibodies used were: mouse monoclonal FLAG-M2 (Sigma), c-Myc monoclonal (Roche Applied Science), rabbit HA (Bethyl Laboratories, or Abcam ab9110), mouse monoclonal HA.11 (Covance MMS-101P), rabbit Cul2 (Zymed Laboratories 51-1890, mouse ElonginC (BD Transduction Laboratories 610760), goat ElonginB (Santa Cruz P-16, sc23407), alpha-tubulin (Santa Cruz B-7, sc-5286), HDAC2 (Biomol SA-402). Proteins were separated by conventional SDS-PAGE or NuPAGE 4-12% Bis-Tris gels (Invitrogen) run with MES buffer. Western blots were visualized by ECL (GE Healthcare) or Odyssey (LiCor).

Plasmids and cloning
Full-length human PRAME was PCR-amplified and cloned into NcoI and Xhol sites of pcDNA5/FRT/T0/StrpII-2xTEV-Myc-3xHA (Le Guezennec et al, 2006) to generate pcDNAs-TAG-PRAME. A BamHI-Xhol fragment encoding TAG-PRAME was further subcloned into LZRS(Zeo) to generate the retroviral expression vector LZRS(Zeo)-TAG-PRAME. Full-length human PRAME (ATCC) was subcloned with N-terminal HA epitope tags into pcDNA5/FRT (Invitrogen). Mutagenesis to generate PRAME mutants was performed using the QuikChange II XL site-directed mutagenesis kit (Stratagene) according to the manufacturer’s instructions. To create the pTTE retroviral vector for ChiP, the ProtA-2xTEV-Myc cassette of the pZXN plasmid (Le Guezennec et al, 2006) was removed by digestion with BamHI and SalI. The resulting backbone was ligated to overlapping synthetic oligos with BamHI and Xhol overhangs, encoding two consecutive TY1 epitopes, a single ERalpha epitope and an extended MCS. PRAME was cloned using the EoRI site to generate pTTE-PRAME plasmid; EoLC was cloned using BglII and Xhol to generate pTTE-EoLC plasmid. All constructs were checked by sequencing.

Gel filtration analysis
Size exclusion chromatography was performed with a Superose 6 column on a SMART system (Amersham Pharmacia Biotech) with buffer containing 300 mM NaCl, 10% glycerol, 20 mM TRIS, 0,2 mM EDTA, 1 mM DTT and 1 mM PMSE. 50 µl nuclear extract were separated in 100 µl fractions and analyzed by SDS-PAGE and western blot.

Purification of protein complexes from K562 cells
K562 cells were grown in spinner flasks and harvested at a confluence of about 10^6 cells/ml. For each protein extraction, about 10^6 cells were centrifuged for 15 minutes at 1600 rpm and washed with PBS. Cell pellets were resuspended in 2,5 volumes of hypotonic buffer T1 (10 mM Hepes pH 7,9, 10 mM KCl, 0,1 mM MgCl₂, 0,1 mM EDTA pH 8, 1 mM DTT, 10% glycerol) supplemented with phosphatase and protease inhibitors (1 mM NaVO₃, 1 mM NaF, 1 mM PMSE, Protease inhibitor cocktail (Roche)). After 15 minutes of incubation on ice, the cell membranes were disrupted.
with 20 strokes of Pestle A, or until 80-90% of the cells were stained by trypan blue. After centrifugation for 20 minutes at 3000 rpm at 4°C, the supernatant was harvested and regarded as the cytoplasmic fraction. The pellet containing the nuclei was resuspended in 1 volume of 12 buffer (20 mM Hepes pH 7.9, 420 mM NaCl, 1.5 mM MgCl₂, 0.1 mM EDTA pH 8, 1 mM DTT, 10% glycerol) supplemented with inhibitors, and incubated in rotation for 1 hour at 4°C to extract the nuclear proteins. The suspension was cleared by ultracentrifugation for 30 minutes at 100000 g at 4°C (Sorvall RC M150 GX). The supernatant was the nuclear fraction. The quality of the cytoplasmic/nuclear extractions was determined by western blot of alpha tubulin (cytoplasmic marker) and HDAC2 (nuclear marker).

About 5–10 ml (60–120 mg) of nuclear extracts were incubated overnight at 4°C with anti-Myc (9E11) antibodies covalently crosslinked to protA-sepharose CL-4B (GE Healthcare) (see below). The following day, beads were washed three times with 10 ml buffer IPP300/0.5 (300 mM NaCl, 10 mM Tris-HCl pH 8, 0.5 mM EDTA and 0.5% NP-40), and the protein complexes were eluted with 20 µg/ml of Myc peptide in a thermoshaker for 30 minutes at 28°C.

Crosslinking of antibody to beads

ProtA-sepharose CL-4B (GE Healthcare) or protG-sepharose (Sigma) beads were incubated with antibody in PBS at room temperature for 1-2 h, then washed once with 10 volumes of sodium borate solution (0.2 M, pH 9). Chemical cross-linking of the antibodies to the beads was performed resuspending the beads in a sodium borate solution containing 5.2 mg/ml DMP (dimethyl-pimelimidate, Sigma) and incubating for 30 minutes at room temperature with gentle rotation. The beads were washed once in 10 volumes ethanolamine (0.2 M, pH 8), incubated with ethanolamine for 2 h at room temperature, washed twice with glycine (0.1 M, pH 3) to remove uncrosslinked antibodies, and eventually with PBS to neutralize the pH.

Mass Spectrometry and emPAI calculation for K562 complexes

The eluates of large scale immunoprecipitations from K562 cells were concentrated by speedvac and run shortly (~1 cm) on SDS-PAGE gels to remove detergents and the excess of peptide. The gel lane was then fixed, cut in small pieces, reduced and alkylated. Proteins were digested overnight with trypsin (Promega) and eluted from the gel with trifluoroacetic acid. Peptides were sequenced using a nano-high-pressure liquid chromatography Agilent 1100 nanoflow system connected online to a 7-Tesla linear quadrupole ion trap-Fourier transform (FT) mass spectrometer (Thermo Electron, Bremen, Germany) essentially as described (Olsen et al, 2004). Proteins were identified using the Mascot search algorithm (Matrix Science) with IPI human database v3.37, which contains 69 164 unique entries. First ranked peptides (Mascot peptide scores ≥20) were parsed from Mascot database search html files with MSQuant (http://www.msquant.sourceforge.net) to generate unique first ranked peptide lists that were further filtered for absolute calibrated mass error <5 and peptide delta score >5. To remove redundancy at the protein level and to uniquely assign peptides to one IPI Db entry, the peptides were remapped to the IPI human Db using the program Protein Coverage Summarizer (http://nccr.pnl.gov/software/) as described (Lasonder et al, 2008). To determine the protein abundance, we computed the exponentially modified Protein Abundance Index (emPAI) values for our data (Ishihama et al, 2005; Lasonder et al, 2008); emPAI values for all proteins were calculated as 10−PAI (PAI = observed peptides/observable peptides). The number of observable peptides was calculated by in silico tryptic digestion of the IPI human Db v3.37 using the program Protein Digestion Simulator (http://nccr.pnl.gov/software/).

Purification of protein complexes from 293 cells and MudPIT analysis

To prepare protein samples for mass spectrometry, 293-HA-PRAME and control wild type cells were grown to 70–80% confluence in 10–20 roller bottles then washed, lysed and subjected to anti-HA agarose immunoaffinity chromatography as described (Mahrour et al, 2008). TCA-precipitated proteins were urea-denatured, reduced, alkylated and digested with endoproteinase Lys-C (Roche) followed by modified trypsin (Roche) as described (Florens & Washburn, 2006; Washburn et al, 2001). Peptide mixtures were loaded onto 100 µm fused silica microcapillary columns packed with 5-µm C₁₈ reverse phase (Aqua, Phenomenex), Strong cation
exchange particles (Partisphere SCX, Whatman), and reverse phase. Loaded microcapillary columns were placed in-line with a Quaternary Agilent 1100 series HPLC pump and a LTQ ion trap mass spectrometer equipped with a nano-LC electrospray ionization source (ThermoFinnigan). Fully automated 10-step MudPIT runs were carried out on the electrosprayed peptides, as described (Florens & Washburn, 2006). Tandem mass (MS/MS) spectra were interpreted using SEQUEST against a database of 61,430 sequences, consisting of 37,742 human proteins (downloaded from NCBI on 2008-03-04), 177 usual contaminants (such as human keratins, IgGs, and proteolytic enzymes), and to estimate false discovery rates, 30,716 randomized amino acid sequences derived from each non-redundant protein entry. Peptide/spectrum matches were sorted and selected using DTASelect with the following criteria set: spectra/peptide matches were only retained if they had a DeltCn of at least 0.08, and minimum XCorr of 1.8 for singly-, 2.0 for doubly-, and 3.0 for triply-charged spectra. In addition, peptides had to be fully-tryptic and at least 7 amino acids long. Combining all runs, proteins had to be detected by at least 2 such peptides, or 1 peptide with 2 independent spectra. Under these criteria the final false discovery rate at the peptide level is 0.151 ± 0.009. Peptide hits from multiple runs were compared using CONTRAST. To estimate relative protein levels, we took advantage of evidence that Normalized Spectral Abundance Factors (NSAFs) vary with the relative abundance of proteins in a sample. NSAFs were calculated for each detected protein, using the formula:

\[
(\text{NSAF})_k = \frac{\text{SpC}_k}{\text{Length}_k} \sum_{i=1}^{N} \left( \frac{\text{SpC}_i}{\text{Length}_i} \right)
\]

where \(\text{SpC}\) = spectral count, \(L\) = protein length in amino acids, and \(i = \) all proteins detected in the MudPIT runs, as described (Florens et al., 2006; Paolelli et al., 2006).

Expression of recombinant proteins in SF21 insect cells

cDNAs encoding wild type or mutant PRAME, Cul2, Elongins B and C were subcloned into pBacPAK8 and recombinant baculoviruses were generated. SF21 cells were cultured at 27°C in SF-900 medium supplemented with 10% fetal bovine serum. SF21 cells were co-infected with the recombinant baculoviruses indicated in the figures. 50 hours after infection, cells were collected, lysed, and processed as described (Mahrour et al., 2008).

In vitro ubiquitination assays

The PRAME-Elongin BC complex and the His-T7-Xpress-Cullin2/Myc-RBX1 complex were obtained by co-infecting SF21 cells with baculoviruses encoding for each individual protein and purifying through Flag tag or nickel-agarose chromatography, respectively. Different combinations of these complexes were then incubated for 1 h at 37°C with ~50 nM Uba1, ~0.2 µM UbcH5a, and 0.25 mg/ml ubiquitin, in 20 µl reactions containing 50 mM Tris-HCl, pH 7.6, 50 mM NaCl, 5 mM MgCl₂, 0.5 mM EDTA (pH 7.9), 5% glycerol, 2 mM ATP, 0.5 mM dithiothreitol, 0.3 units/ml pyrophosphatase, 60 mM creatine phosphate, and 0.3 mg/ml creatine phosphokinase. N-terminally (His)₆-tagged mouse E1 was expressed in SF21 insect cells and purified by nickel-agarose chromatography as described. Human UbcH5a with an N-terminal 6-histidine tag and a C-terminal Flag tag were expressed in E. coli strain BL21(DE3) and purified by nickel agarose.

ChIP and qPCR

Chromatin harvests, ChIPs, and qPCR analyses were performed as previously described (Denissov et al., 2007) with the following minor adjustments. Cells were crosslinked for 15 min in 1% formaldehyde, they were resuspended in ChIP incubation buffer at a concentration of 16 millions/ml and sonicated using a Bioruptor sonicator (Diagenode) for 15 min with settings: high power, 30 s ON, 30 s OFF. For every ChIP reaction, 200 µl of sonicated chromatin were used. The immunoprecipitated chromatin was eluted with 200 µl of elution buffer at room temperature for 20 minutes and DNA was purified with Qubit fluorometer and Quant-iT dsDNA HS Assay (Invitrogen). The following antibodies were used for ChIP: preimmune serum (preV27, "no antibody" control) and PRAME antiserum (V27); NFYA (Santa Cruz H-209, sc-10779); NFYB (pAb-007-100) and BB2 (against TY1 epitope) were from Diagenode.
For sequential ChIP experiments (ChIP-re-ChIP), in order to avoid antibody carry-over, the first ChIP was performed with antibody covalently crosslinked to ProteinA/G-Sepharose beads (Santa Cruz) with DMS as described above. Immunoprecipitated chromatin was then eluted in a smaller volume and further processed as a normal ChIP with PRAME immune serum or the preimmune serum as negative control. Real-time qPCR was performed using the SYBR Green mix (Bio-Rad) with the MyIQ thermocycler (Bio-Rad). Specific primers for qPCR (Biolegio) were designed as previously described (Denissov et al, 2007) and are listed in Supplementary Table 2. Primers amplifying exon 12 of albumin were used as negative control.

RNA-seq library preparation and data analysis

Total RNA was extracted from K562 cells with the RNeasy kit and on-column DNase treatment (Qiagen). 100 µg of total RNA were subjected to two rounds of poly(A) selection using the Oligotex mRNA Mini Kit (Qiagen) and the concentration of poly(A)+ RNA was measured with a Qubit fluorometer (Invitrogen). 200 ng of purified poly(A)+ RNA were diluted in 160 µl RNase-free water and fragmented by addition of 40 µl 5x fragmentation buffer (200 mM Tris acetate pH 8.2, 500 mM potassium acetate and 150 mM magnesium acetate) and incubation at 94°C for 120 seconds. After purification with RNeasy Minelute Kit (Qiagen), the quality of fragmented RNA was analyzed on an Experion (Bio-Rad). Double stranded cDNA was synthesized from the fragmented RNA with Superscript III (Invitrogen) using random hexamers, and then used for Illumina sample prepping and sequencing (see below).

A total of 7,246,489 RNA-seq reads were uniquely mapped to hg18 and used for bioinformatical analysis. Normalized gene expression values (NE) for RefSeq genes were computed with Genomatix Region Miner tools.

Illumina high-throughput sequencing

A starting amount of 10-20 ng of ChIP DNA or double stranded cDNA was used for sample prepping according to standard protocols. Briefly, end repair was performed with T4 DNA ligase, Klenow DNA polymerase and T4 PNK; a 3’ protruding adenosine was added using Klenow DNA polymerase, and T-overhang adaptors were ligated. DNA molecules with a size of 270-300 bp were selected from gel, purified and amplified with 14 cycles of PCR. Quality controls were performed by qPCR of DNA before and after sample prepping, and by analysis of the PCR products with an Experion (Bio-Rad). Cluster generation and sequencing-by-synthesis (35bp) were performed using the Illumina Genome Analyzer II according to standard protocols (Illumina). The image files were processed to extract DNA sequence data, and sequences were mapped to the human hg18 reference genome using the eland program allowing at most one base mismatch. Sequence reads were filtered based on quality controls and sequences that could not be mapped uniquely on the reference genome were discarded. For visualization purposes, ChIP-seq reads were directionally extended to the length of the fragments used for sequencing and the number of overlapping sequence reads was determined for each base pair in the genome, averaged over a 10 bp window, and visualized in the University of California Santa Cruz genome browser (http://genome.ucsc.edu). ChIP-seq and RNA-seq data are published in the GEO database (GSE26439), and in the published history “PRAME” in Galaxy (http://main.g2.bx.psu.edu/).

ChIP-seq analysis and peak detection

Bioinformatical analysis was performed using a combination of in-house developed scripts, a Genomatix Genome Analyzer server (http://www.genomatix.de), and Galaxy tools (http://main.g2.bx.psu.edu/).

ChIP-seq experiments generated the following number of uniquely-mapped reads: 5.5 millions for PRAME, 5.6 millions for preimmune (control), 8.9 millions for NFYA, and 8.8 millions for NFYB. In order to compensate for the small difference in sequencing depth between the PRAME and preimmune datasets, the number of tags of preimmune ChIP-seq was equalized to the number of tags of PRAME ChIP-seq by uniform removal of tags as described (Nielsen et al, 2008). PRAME binding sites (peaks) were subsequently identified with the MACS peak calling algorithm (Zhang et al, 2008) using the two-sample method with a p-value (e) of 10^-6, which compared the PRAME profile to the preimmune (control) profile. NFYA and NFYB binding sites were determined with MACS using the standard...
method with extended lambda (10 kb, 50 kb, 100 kb) and p-value (e) of $10^{-6}$. MACS identified a total of 141 32 PRAME peaks, 25 151 NFYA peaks, and 163 350 NFYB peaks. NFYA and NFYB peaks were combined in a total of 28 844 NFY binding sites that were tested for tag density. The number of sequence reads were counted in 500 bp regions centered on the summits of the 28844 NFY binding sites (with the summit being the site of highest enrichment as determined by MACS), and the data were visualized in dot plots using R.

**Location analysis and motif discovery**

To analyze the genomic distribution of the binding sites identified, we first determined the fraction of peak summits localized within a 2 kb window centered around the TSS of primary transcripts (TSS ± 1 kb) using the Genomatix Genome Inspector. The genomic distribution of the binding sites located outside promoter regions ("distal sites") relative to genes was analyzed with PinkThing (http://pinkthing.cmbi.ru.nl/). The background distribution of genomic DNA was determined applying the same approach to pools of random genomic sites (n > 33 000).

The distances of promoter-associated binding sites from TSSs were computed with the Genomatix Genome Inspector tool, and plots were generated with R.

The localization of PRAME at enhancer elements was determined by overlaps between PRAME peaks summits and enhancer regions recently predicted for K562 cells (Heintzman et al, 2009). The genomic coordinates of the published enhancer regions were converted from hg17 to hg18 with the Galaxy Lift-over function, resulting in 24 545 enhancers.

**De novo** motif discovery was performed using the GimmeMotifs pipeline (van Heeringen & Veenstra, 2010), which incorporates a number of widely-used motif analysis tools, including MDmodule, MEME, Weeder, MotifSampler, BioProspector. Based on the distribution of PRAME binding sites, we performed motif searches on the DNA sequences underlying PRAME summits located within ~400 bp and +100 bp of a primary TSS (n=4981). A CCAAT motif scored the highest values of enrichment and MNCP scores, as compared to pools of background sequences with the same genomic distribution.

**ENCODe datasets**

We retrieved and analyzed publicly-available ChIP-seq datasets of H3K9ac, H3K4me3, H3K36me3, H3K27me3, and RNA-PolII generated from K562 cells by the “ENCODe Chromatin Group at Broad Institute and Massachusetts General Hospital” (http://hgdownload.cse.ucsc.edu/goldenPath/hg18/encodeDCC/wgEncodeBroadChipSeq/).

Starting from 27 903 RefSeq genes with a RNA-seq expression value, we identified 7 214 genes longer than 1 kb with a NFY peak summit in the proximal promoter region. Next, we quantified the signals of PRAME in a 500 bp window under the NFY binding site, H3K9ac and H3K4me3 in a 3 kb window, H3K36me3 and H3K27 in the gene body, and RNA-PolII in the gene body excluding the first 500 bp.

**Immobilized template assay method**

A 5'-biotinylated DNA fragment of ~670 bp containing nucleotides -280 to +150 (relative to the annotated TSS) of the human HSPA5 promoter downstream of ~240 of DNA derived from the pGEM-T Easy Vector (Promega) was immobilized on streptavidin-coupled magnetic beads (Dynabeads M-280 Streptavidin, Invitrogen) following the manufacturer’s instructions. This DNA fragment contains 3 ER stress elements (ERSE), each of which includes binding sites for NFY and ATF6. Reactions containing different combinations of Hela nuclear extract, Flag-PRAME complex (~50ng), and NFY (~100ng) were incubated with ~200ng of immobilized DNA for 30 minutes at 30ºC in buffer containing 50mM NaCl, 40mM Heps-NaOH pH 7.9, 15% Glycerol, 2mM MgCl$_2$, 0.15mM EDTA, 1mM DTT, 0.02% NP-40, and 0.1mg/ml BSA. The magnetic beads were washed three times in the same buffer without BSA. Bound proteins were eluted by boiling the beads in 1X SDS loading buffer and analyzed by western blotting. NFY was purified by anti-Flag immunoaffinity chromatography from Sf21 cells coinfectd with baculoviruses encoding Flag- NFYA, Myc-NFYB, and HA-NFYC. Flag-PRAME and PRAME-associated proteins (including Elongins B and C, Cul2, and Rbx1) were purified using anti-Flag agarose from nuclear extracts from 293-FRT cells stably expressing Flag-PRAME. NE, HeLa nuclear extract.
ACKNOWLEDGEMENTS

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REFERENCES


SUPPLEMENTARY INFORMATION

Supplementary Figure 1

(A) PRAME binds a DNA fragment containing the NFY-dependent HSP5 promoter by a mechanism that depends on activity(s) present in HeLa nuclear extract. A DNA fragment containing the HSP5 promoter was immobilized on Streptavidin-coupled magnetic beads and incubated with various combinations of Flag-PRAME complex, HeLa nuclear extract and recombinant NFY expressed in and purified from insect cells coinfecte
**Supplementary Figure 2**

**Motif found**

<table>
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<th>Positional preference plot</th>
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**De novo motif discovery at PRAME binding sites.**

Significant sequence motifs identified at PRAME binding sites at TSS. "Motif found" is the motif identified in the de novo discovery; "Jaspar match" is the annotated motif present in the JASPAR database that is most similar to the discovered motif (a p-value for the similarity test is shown); "significance and enrichment" report statistics of the enrichment of the discovered motif in the input sequences versus two different sets of backgrounds; the "positional preference plot" shows the frequency of the location of the discovered motif relative to the summit of PRAME binding sites.
**NFY ChIP-seq data analysis.**

(A) Correlation between NFYA and NFYB binding detected by ChIP-seq. NFYA and NFYB tag densities were determined in 500 bp regions centered around the summits of NFYA and/or NFYB binding sites (n=28844). The data is displayed after sorting on NFYA (left) or NFYB (right) tag densities. The plots show that NFYA and NFYB signals are very well correlated for the majority of the binding sites. A subpopulation of binding sites shows high signals for NFYA, but low for NFYB (~5-10% of the sites, depending on the threshold applied), which might reflect dynamic in vivo binding of NFY subunits in the context of chromatin. PRAME binding sites didn't show a bias towards or against these regions.

(B) Genomic distribution of NFY binding sites.

(C) NFYA, NFYB, and PRAME binding occur at the same site, and not generally within promoters. Each NFYA promoter summit is centred on the zero, and the distances to NFYB (green) or PRAME (blue) summits plotted as frequency density.
UCSC genome browser views of all genome-wide datasets combined.
Overviews of the active NFY loci TOP2A and STAT2, and of the inactive NFY loci TSPAN15 and SLC4A8.
Complete conservation of the BC-box and Cul2-box in the PRAME-like protein family.

All human PRAME-like proteins share a conserved BC-box and Cul2-box at their N-terminus. The amino acid sequence of human PRAME was aligned with ClustalX to the previously published multiple alignment of the human PRAME-like proteins encoded on human chromosome 1 [Birtle et al. 2005]. The alignment was edited with CILC Sequence Viewer to indicate dissimilar residues in grey, and to identical in yellow the identical residues within the BC- and Cul2-boxes.
### Supplementary Table 2  
**ChIP-qPCR primers**

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CHAPTER 4

THE HUMAN EKC/KEOPS COMPLEX

Is Recruited to Cullin2 Ubiquitin Ligases by the Human Tumour Antigen PRAME

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ABSTRACT

The human tumour antigen PRAME (preferentially expressed antigen in melanoma) is frequently overexpressed during oncogenesis, and high PRAME levels are associated with poor clinical outcome in a variety of cancers. However, the molecular pathways in which PRAME is implicated are not well understood. We recently characterized PRAME as a BC-box subunit of a Cullin2-based E3 ubiquitin ligase. In this study, we mined the PRAME interactome to a deeper level and identified specific interactions with OSGEP and LAGE3, which are human orthologues of the ancient EKC/KEOPS complex. By characterizing biochemically the human EKC complex and its interactions with PRAME, we show that PRAME recruits a Cul2 ubiquitin ligase to EKC. Moreover, EKC subunits associate with PRAME target sites on chromatin. Our data reveal a novel link between the oncoprotein PRAME and the conserved EKC complex and support a role for both complexes in the same pathways.
INTRODUCTION

The human oncoprotein PRAME (preferentially expressed antigen in melanoma) was first identified and cloned as the antigen responsible for an anti-tumour immune response in a melanoma patient (Ikeda et al, 1997). Follow-up experiments revealed that PRAME is expressed at low levels in few normal adult tissues like adrenals, ovaries, and endometrium, and at high levels only in the testis (Ikeda et al, 1997; Kilpinen et al, 2008). However, overexpression of PRAME is frequently found in a wide variety of human cancers, including acute and chronic haematological tumours, synovial sarcoma, lung, breast, and renal carcinoma (Allander et al, 2002; Ikeda et al, 1997). Importantly, high PRAME levels were found to correlate with advanced stages of disease in melanoma (Haqq et al, 2005), neuroblastoma (Oberthuer et al, 2004), serous ovarian adenocarcinoma (Partheen et al, 2008), and chronic myeloid leukaemia (Radich et al, 2006), and to constitute an independent prognostic factor of poor clinical outcome in breast cancer (Doolan et al, 2008; Epping et al, 2008). In contrast, high levels of PRAME were found to correlate with good prognosis in leukaemia cases carrying the t(15;17) PML-RAR translocation (acute promyelocytic leukaemia) (Santamaria et al, 2008).

Although these findings suggested a role for PRAME in human malignancies, the detailed molecular mechanisms and pathways involved are not yet clear. PRAME was reported to repress retinoic acid signaling in melanoma cell lines (Epping et al, 2005), but this was not confirmed for breast cancer or leukaemia cases (Steinbach et al, 2007; Epping et al, 2008). Conflicting reports on leukaemia cells suggested that PRAME might induce caspase-independent cell death (Tajeddine et al, 2005), or repress apoptosis-related genes to promote cell survival (Tanaka et al, 2011).

Recently, through biochemical characterization of PRAME-containing protein complexes, we established that this oncoprotein is a component of Cullin2-based E3 ubiquitin ligases and belongs to the family of BC-box proteins, associating PRAME to a clear biochemical activity and pathway (Costessi et al, 2011). PRAME establishes direct interactions with other ligase subunits through conserved N-terminal motifs: a BC-box (aa. 25-34) mediates interactions with the ElonginB-ElonginC heterodimer, and a downstream Cul2-box (aa. 48-56) mediates interactions with the Cullin2 scaffold protein. Genome-wide chromatin immunoprecipitation experiments further revealed that Cul2-PRAME ubiquitin ligases specifically associate with active promot-
ers regulated by the transcription factor NFY and with proximal enhancers (Costessi et al., 2011).

Two independent laboratories have identified an ancient and highly conserved multiprotein complex named KEOPS (Downey et al., 2006) or EKC (Kisseleva-Romanova et al., 2006), which has orthologues from Archaea to Eukarya and has been implicated in telomeres maintenance, transcriptional regulation, and t^6A modification of tRNAs. Yeast EKC comprises four subunits which are also conserved in the human genome (human orthologues are indicated in brackets): Pcc1p (LAGE3, also known as ESO3), the ATPase Kae1p (OSGEP), the kinase Bud32p (TP53RK, also known as PRPK), and Cgi121p (TPRKB). In addition, yeast EKC also includes Gon7p (also known as Pcc2p), which appears to be fungi-specific (Kisseleva-Romanova et al., 2006).

Intriguingly, the OSGEP subunit is also present in bacteria (YgjD) and eukaryotic genomes express an OSGEP paralogue (Qri7/OSGEPL1) that localizes to mitochondria (Reinders et al., 2006). Comparative genomic studies identified OSGEP as one of the very few genes present in all genomes sequenced so far (Galperin, 2008), suggesting an extremely conserved function. Very recently, several research groups have reported a crucial role for the YgjD/Kae1/OSGEP protein family in the biosynthesis of N6-threonylcarbamoyl adenosine (t^6A) (Daugeron et al., 2011; Yacoubi et al., 2011; Srinivasan et al., 2011): a universal modification at position 37 of tRNAs decoding ANN codons, which is required for accurate translation of messenger RNAs (Yarian et al., 2002).

Human LAGE3 belongs to the NY-ESO gene family together with the closely related LAGE1 and LAGE2 (Alpen et al., 2002), and all three genes are clustered in the same region on chromosome X. While LAGE3 is ubiquitously expressed, LAGE1 and LAGE2 are cancer-testis antigens with high expression in healthy testis and upregulation in a number of cancer tissues, similarly to PRAME.

The aim of the present study was to mine the protein-protein interactome of the PRAME oncoprotein to a deeper level by protein complex purification and mass spectrometry. Our experiments revealed that PRAME specifically interacts with the EKC complex in human cells and that it recruits Cul2-based E3 ubiquitin ligases to EKC. We further show that EKC subunits co-localize with PRAME at the promoter regions of transcriptionally active human genes, supporting a common functional role for these complexes on chromatin. Overall, our findings provide novel and important insights in the molecular pathways in which the oncoprotein PRAME is active.
RESULTS

PRAME interacts with the EKC/KEOPS complex

We have recently applied biochemical purification strategies and characterized PRAME as a component of Cullin2 E3 ubiquitin ligases (Costessi et al, 2011). Here, we improved the immunoaffinity purification protocols and mined the PRAME interactome to a deeper level (see materials and methods). We isolated epitope-tagged PRAME (StrepII-Myc-3xHA-PRAME, referred to as TAG-PRAME) from K562 cells, which express high levels of endogenous PRAME, and from HeLaS3 cells, which express endogenous PRAME at very low to undetectable levels. As expected, mass spectrometry analysis of HA immunoprecipitates detected Cullin2 ligase components with the highest emPAI: Cullin2, Elongin B, Elongin C, and Rbx1 (Table 1). Notably, western blot analysis showed that Cullin2 and Elongin C were expressed to similar levels in nuclear extracts from the two cell lines (not shown).

Furthermore, a number of additional proteins were specifically detected in eluates of K562-TAG-PRAME and HeLaS3-TAG-PRAME cells, but not in the control parental cell lines. These included OSGEP (O-sialoglycoprotein endopeptidase) and LAGE3 (L antigen family member 3), which are the human orthologues of the Kae1 and Pcc1 subunits of the recently described EKC complex (Table 1). OSGEP and LAGE3 were present with a similar abundance in TAG-PRAME eluates from both K562 and HeLaS3 cell lines, consistent with similar expression levels. However, in these eluates we did not identify peptides matching TP53RK or TPRKB, which are the predicted human orthologues of the EKC subunits Bud32 and Cgi121, respectively.

Other potential interactors identified in PRAME eluates were BAG2 and UBL4A (Table 1). BAG2 (BAG family molecular chaperone regulator 2) is a nucleotide exchange factor for the chaperone protein Hsp70 that contributes to the processing of misfolded proteins (Xu et al, 2008). Similarly, UBL4A (Ubiquitin-like protein 4A) is part of a chaperone protein complex that promotes correct membrane targeting of tail-anchored proteins (Wang et al, 2011; Mariappan et al, 2010). Peptides for both BAG2 and UBL4A were identified in all anti-HA eluates from K562-TAG-PRAME cells, but were not detected in the anti-MYC eluates. Additional experiments are therefore needed to establish whether these proteins are real interactors of PRAME or rather false positives. For the purpose of this study, we further concentrated on EKC subunits.
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**Table 1**

PRAME-interacting proteins identified by Mass Spectrometry of purified complexes from K562 and HeLaS3 cells. PRAME-associating proteins were purified from nuclear extracts of K562-TAG-PRAME by anti-MYC or anti-HA immunoprecipitation and from HeLaS3-TAG-PRAME cells by anti-HA immunoprecipitation and subjected to mass spectrometry. Immunoprecipitations from wild type K562 and HeLaS3 cells were used as a control for immunospecificity of the purifications. Components of Cullin2 ligases and EKC complex are indicated in bold.
The specificity of the interactions between PRAME and EKC subunits was confirmed by transient transfections in HEK293T cells and coimmunoprecipitation experiments of TAG-PRAME with OSGEP and LAGE3 fused to a TTE (TY1-TY1-ER) epitope tag (Fig. 1A). We also used a baculovirus expression system to co-express FLAG-PRAME with different combinations of the four human EKC subunits in SF21 insect cells. Consistent with the transient transfection experiments, we observed that PRAME could bind to OSGEP and LAGE3 independent of TP53RK and TPRKB. Interestingly, although both OSGEP and LAGE3

**Figure 1**

**A**

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**B**

**PRAME interacts with OSGEP and LAGE3.**

(A) Coimmunoprecipitation assays verify the interaction of PRAME with the EKC subunits OSGEP and LAGE3. Constructs expressing the indicated proteins were transiently transfected in 293T cells. Anti-TAG immunoprecipitations were performed with rabbit HA antibody (Abcam). Anti-FLAG immunoprecipitations were performed, and total cell lysates and eluates were analyzed by western blotting. Panels showing anti-HA immunoblots are from the same exposure of the same blot, as are the panels showing anti-cMyc immunoblots. Asterisks indicate light chains of the antibodies used in the immunoprecipitation.

(B-C) PRAME directly interacts with EKC complex through OSGEP and LAGE3. SF21 cells were co-infected with baculoviruses expressing the indicated proteins. Anti-FLAG immunoprecipitations were performed, and total cell lysates and eluates were analyzed by western blotting. Panels showing anti-HA immunoblots are from the same exposure of the same blot, as are the panels showing anti-cMyc immunoblots. Asterisks indicate light chains of the antibodies used in the immunoprecipitation.

**C**

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The specificity of the interactions between PRAME and EKC subunits was confirmed by transient transfections in HEK293T cells and coimmunoprecipitation experiments of TAG-PRAME with OSGEP and LAGE3 fused to a TTE (TY1-TY1-ER) epitope tag (Fig. 1A). We also used a baculovirus expression system to co-express FLAG-PRAME with different combinations of the four human EKC subunits in SF21 insect cells. Consistent with the transient transfection experiments, we observed that PRAME could bind to OSGEP and LAGE3 independent of TP53RK and TPRKB. Interestingly, although both OSGEP and LAGE3

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**B**

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**Table 2**

Human EKC-interacting proteins identified by Mass Spectrometry of purified complexes from nuclear extracts of K562 and HeLaS3 cells. Proteins interacting with each of human EKC subunits are indicated in bold.
could bind PRAME independent of the other (Fig. 1B, lanes 3 and 6), LAGE3 appears to enhance the PRAME-OSGEP interaction, since substantially more OSGEP bound to PRAME when LAGE3 was also expressed (Fig 1B, compare lanes 1, 4, and 5 to lane 6, and Fig. 1C, compare lane 4 to lanes 5 and 6). Finally, we observed small amounts of TP53RK and TPRKB copurifying with FLAG-PRAME dependent on both OSGEP and LAGE3 (Fig. 1B, lanes 1, 3, and 6).

Taken together, our results identified stable interactions between PRAME and human orthologues of the EKC/KEOPS complex.

**EKC purifications from K562 cells**

We next studied the expression patterns of EKC, Cullin2 ligase subunits and PRAME in normal and cancer cells by querying the GeneSapiens gene expression database (Kilpinen et al, 2008). This database contains mRNA expression data of most human genes across almost 10000 array experiments from 175 healthy and pathological tissues. Consistent with the literature, we found that PRAME is expressed only in very few normal adult tissues while it is upregulated in several cancers (Suppl. Fig. S1). On the contrary, LAGE3, OSGEP, TP53RK, TPRKB, and CUL2 seem to be expressed in most if not all tissues (Suppl. Figures S1-S4).

To explore further the protein-protein interactions between PRAME and EKC, we generated K562 cell lines stably expressing epitope-tagged versions of each of the four human predicted human EKC subunits and performed protein complex purifications. As shown in Table 2, each EKC subunit co-purified with the other three subunits of the complex, indicating that a complete EKC complex is present in human cells.

Furthermore, TAG-OSGEP and TAG-LAGE3 (but not TAG-TP53RK or TAG-TPRKB) copurified with significant amounts of endogenous PRAME, consistent with TAG-PRAME purifications. Remarkably, TAG-OSGEP and TAG-LAGE3 associated also with Cullin2 ligase components (Table 2). The few peptides for Cul2-PRAME components identified in TAG-TP53RK and TAG-TPRKB eluates were not supported by western blot of the same samples (not shown).

Interestingly, peptides matching the putative protein C14orf142 were specifically identified in eluates from OSGEP, LAGE3 and TP53RK (Table 2). Similarly to LAGE3 and TPRKB, C14orf142 is predicted to be a small protein without clear structural domains. It is tempting to speculate that this protein might constitute a novel EKC interactor in human cells.

Although we did not identify peptides matching Cul2 by mass spectrometry of TAG-OSGEP eluates, we did detect small amounts of Cul2 by western
blotting of the same samples (Fig. 2, lane 6). Intriguingly, the level of Cul2 detected in TAG-OSGEP eluates was substantially lower than in TAG-PRAME eluates (Fig. 2, lanes 4 and 6), while Elongins B and C were present in similar amounts in all purifications. These findings might indicate that OSGEP establishes less stable interactions with Cullin2 as compared to PRAME. Consistent with the mass spectrometry results, TAG-PRAME did not copurify with TPRKB (Fig. 2, lane 4), while a clear band was detected in TAG-OSGEP eluates.

Our proteomic experiments show for the first time that a complete EKC complex is present in human cells, and that the OSGEP and LAGE3 subunits are present in a complex together with PRAME and Cullin2 ligase components.

**Topology of the human EKC complex**

In order to study the topology of the interactions within the human EKC complex, we expressed pairs of EKC subunits in the baculovirus system and performed immunoprecipitations (Fig. 3). As shown in figure 3, we observed reciprocal coimmunoprecipitation of HA-LAGE3 and c-Myc-OSGEP and of HA-TPRKB and c-Myc TP53RK, respectively. In addition, c-Myc TP53RK copurified with HA-OSGEP upon anti-c-Myc immunopurification, although this interaction was not detected in the reciprocal experiment, raising the possibility that the c-Myc epitope was occluded and/or that binding of the antibody to the epitope disrupts the interaction. Taken together, these experiments established that human EKC has the linear configuration LAGE3-OSGEP-TP53RK-TPRKB, consistent with the three-dimensional structures of archael EKC orthologues that were recently reported (Mao et al., 2008).
Interestingly, Mao et al. reported that archael Pcc1, the orthologue of human LAGE3, forms highly interdigitated homodimers and hypothesized that dimerization of Pcc1 might support dimerization of the full EKC complex. Moreover, the crystal structures indicated that one moiety of Pcc1 could bind only one moiety of Kae1. We tested the dimerization properties of human EKC using combinations of different epitope tags in transient transfection experiments. As shown in figure 4A, FLAG-OSGEP coimmunoprecipitated TTE-OSGEP when LAGE3 was cotransfected (lane 4), establishing that multiple OSGEP moieties are present in the same biochemical complex. Importantly, the concomitant expression of PRAME did not affect these dimerization properties (Fig. 4A, compare lanes 4 and 5). Taking into consideration the structural information available for archaeal EKC, our data suggest the formation of a tetrameric LAGE3-OSGEP assembly (Fig. 4A).

In the same experiments, we noticed that OSGEP levels were significantly lower in the absence of LAGE3 (Fig. 4A, compare lanes 2 and 4), suggesting that the stability of OSGEP could be critically dependent on interactions with other proteins. To address this hypothesis, we transfected FLAG-OSGEP together with each of the other EKC subunits or PRAME. Fig. 4B shows that LAGE3, TP53RK and PRAME could all enhance the stability of OSGEP. TPRKB did not alter OSGEP levels, consistent with the absence of a direct interaction between these two proteins. Interestingly, LAGE3 exerted the highest positive
OSGEP protein levels are modulated by protein-protein interactions.

(A) Multiple OSGEP moieties are present in the same complex. Constructs expressing the indicated proteins were transiently transfected in 293T cells. Total cell lysates and FLAG eluates were analyzed by western blotting. A plasmid expressing GFP was cotransfected to control for the transfection efficiency. (B) OSGEP levels are affected by protein-protein interactions. 293T cells were transfected with constructs expressing FLAG-OSGEP and the proteins indicated. The bar graph shows FLAG-OSGEP levels quantified by immunoblot with the Odyssey system (values are the average of two independent experiments). A plasmid expressing GFP was cotransfected to control for the transfection efficiency. (C) and (D) LAGE3-OSGEP interface mutants decrease OSGEP protein levels. Transient transfections in 293T cells with the mutant constructs indicated and immunoblot by Odyssey of total cell lysates. Graphs report intensities of FLAG-OSGEP quantified with Odyssey (A.U., arbitrary units).
effect, which correlates with the possibility to form tetrameric LAGE3-OSGEP complexes, while TP53RK can bind only one moiety of OSGEP. The finding that PRAME had a relatively modest effect on OSGEP levels might reflect a less stable interaction between OSGEP and PRAME, than between OSGEP and TP53RK.

We investigated further the effect of LAGE3 on the protein levels of OSGEP in transient transfection experiments with mutant constructs. Based on the crystallographic structures of archael EKC (Mao et al., 2008), we introduced point mutations in aminoacid residues predicted to be involved in the protein-protein interactions between OSGEP and LAGE3: R94A in OSGEP, and L129R or T133R in LAGE3. Additionally, we generated OSGEP mutant P35A/P36A (referred to as PPAA) and LAGE3 mutants K95A and K109A, which are not predicted to contribute to the OSGEP-LAGE3 interaction. As shown in figure 4C, the OSGEP mutant R94A showed lower steady-state levels, which could not be enhanced by co-expression of wild-type LAGE3, while OSGEP PPAA behaved like the wild type. Similarly, levels of wild-type OSGEP were not enhanced upon co-expression of either T133R or L129R LAGE3 mutants, while the LAGE3 lysine mutants behaved the same as wild type LAGE3.

Taken together, our data suggests (i) that the human EKC complex is organized in the same way as its archaeal counterpart, (ii) that dimerization of LAGE3 can mediate formation of higher order structures containing at least two molecules of OSGEP, (iii) that formation of these higher order structures is independent of PRAME, and (iv) that the protein levels of OSGEP can be significantly modulated by protein-protein interactions.

**PRAME bridges Cullin2 ligases to the EKC complex**

We next studied the role of the Cullin2 complex in the association of PRAME with EKC. Transient transfection experiments showed that the interaction of PRAME with OSGEP and LAGE3 does not require an intact BC-box, since a BC-box defective PRAME (PRAME M2) coimmunoprecipitated OSGEP and LAGE3 as efficiently as wild type PRAME (Fig. 5A).

Next, we purified TAG-LAGE3 protein complexes from HeLaS3 cells, which express very low to undetectable levels of endogenous PRAME. As expected, all EKC subunits copurified with TAG-LAGE3, while no peptides were identified for PRAME (Table 2). Importantly, components of Cullin2 ligases did not co-purify with LAGE3 in these cells. The abundance of EKC subunits was similar in the eluates from K562-TAG-LAGE3 and HeLaS3-TAG-LAGE3 cells, consistent with similar protein levels in the two cell lines.
These results, together with the finding that TAG-OSGEP and TAG-LAGE3 associate with endogenous PRAME and Cullin2 ligase components in K562 cells, suggested that PRAME could be involved in recruiting a Cullin2 ligase to the EKC complex.

To test this hypothesis, we knocked down endogenous PRAME with stable transfection of retroviral constructs in K562-TAG-OSGEP and K562-TAG-LAGE3 cell lines and performed immunoprecipitations. Figure 5B clearly shows that downregulation of PRAME significantly reduced the interaction between each EKC subunit and Cul2, EloB and EloC, establishing that PRAME is required for these interactions. As expected, the interaction of TAG-OSGEP and TAG-LAGE3 with TPRKB was not affected by PRAME knockdown. Notably, we found that the level of Cul2 detected in TAG-OSGEP eluates was substantially lower than in TAG-LAGE3 eluates (Fig. 5B, lanes 1 and 4), despite similar amounts of EloB and EloC. These results are similar to the previous comparison between TAG-OSGEP and TAG-PRAME eluates (Fig. 2, lanes 4 and 6).
Taken together, our data reveal that PRAME can bridge Cullin2 ligases to the EKC complex (Fig. 5C).

**Cul2-PRAME ligases and EKC subunits co-localize to active promoters**

The yeast orthologues of OSGEP (Kae1p) and LAGE3 (Pcc1p) have been reported to associate with the promoters and transcribed regions of several genes in a transcription-dependent manner (Kisseleva-Romanova et al., 2006). Genome-wide chromatin immunoprecipitation experiments revealed that PRAME and Cullin2 complex components are specifically enriched at transcriptionally active promoters that are also bound by the transcription factor NFY, and at nearby enhancers (Costessi et al., 2011).

In the absence of validated ChIP-grade antibodies against EKC subunits, we generated K562 cell lines stably expressing the proteins of interest fused to the TTE epitope tag (TY1-TY1-ER), which can be efficiently and specifically used in ChIP-qPCR experiments (Costessi et al., 2011). To determine whether human OSGEP and LAGE3 are present on chromatin, we performed ChIP assays with the BB2 antibody (which recognizes the TY1 tag) and tested a panel of PRAME-bound promoters. The results in figure 6 clearly show that all PRAME-bound promoters tested are efficiently recovered in both K562-TTE-OSGEP and K562-TTE-LAGE3 cells, but not in parental K562 cells or control K562-TTE cells.

Our genomic experiments extend the biochemical interactions to the chromatin environment, establishing that PRAME and EKC subunits associate with the same promoter regions.

**Figure 6**

<table>
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<th>OSGEP and LAGE3 are present at PRAME-bound promoters on chromatin.</th>
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<td>ChIP-qPCR experiments using BB2 antibody were performed on K562 cells lines stably expressing TTE-OSGEP or TTE-LAGE3. As control for specificity, ChIPs were performed on the parental cells (wt) and on cells transduced with the vector expressing the tag only (K562-TTE) using a panel of primer sets for PRAME binding sites. Values are expressed as mean recoveries ± standard deviation from three independent experiments.</td>
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We recently reported the first purifications of PRAME-containing protein complexes by epitope-tagged immunoprecipitations and mass spectrometry, which revealed that PRAME is a BC-box substrate receptor component of Cullin2-based E3 ubiquitin ligases (Costessi et al., 2011). In the present study, we identified novel interactions of Cul2-PRAME ligases with the human EKC/KEOPS complex, we discovered that PRAME recruits ligase components to EKC, and we showed that these proteins associate with the same promoter regions, strongly suggesting a functional link between these complexes in the same pathway.

EKC is an ancient and extremely conserved complex with orthologues in Archaea and Eukarya. It has been implicated in telomere maintenance, transcriptional regulation, and t6A modification of tRNAs in yeast (Kisseleva-Romanova et al., 2006; Downey et al., 2006; Daugeron et al., 2011; Yacoubi et al., 2011; Srinivasan et al., 2011). Our interaction data support a linear architecture LAGE3-OSGEP-TP53RK-TPRKB for the human EKC complex and dimerization around the LAGE3 subunit, with two OSGEP moieties per complex. Our results are fully consistent with recently reported crystal structures of archaeal EKC proteins (Mao et al., 2008).

Consistent with the notion that all subunits contribute to the activity of the complex, yeast EKC was reported to constitute a stable entity. Downey et al. reported that deletion of either Bud32p, Cgi121p, or Gon7p resulted in shorter telomeres. An independent study found that Pcc1p was required for efficient transcription of several yeast genes induced by pheromones or galactose, and that deletion of Pcc1 inhibited the recruitment of the SAGA and Mediator co-activators (Kisseleva-Romanova et al., 2006). Dimerization of the Pcc1p and Kae1p subunits was found to be necessary for cell viability (Mao et al., 2008), and an intact interaction surface between Kae1p and Bud32p was required for both the transcriptional and telomere maintenance functions (Hecker et al., 2008). Most recently, the yeast EKC complex was shown to be essential for a universal chemical modification of tRNAs, called threonyl carbamoyl adenosine (t6A) (Srinivasan et al., 2011; Yacoubi et al., 2011; Daugeron et al., 2011). This modification occurs at position 37 of tRNAs decoding ANN codons, including the first ATG of all protein-coding mRNAs, and is required for efficient translation (Yarian et al., 2002). The t6A modification activity was dependent on Pcc1p, the ATPase activity of Kae1p, and the kinase activity of...
Bud32p, but not on Cgi121p (Srinivasan et al, 2011).

In our experiments, all human EKC orthologues co-purified together and with similar abundance both in K562 and HeLaS3 cells, demonstrating for the first time that a complete and stable EKC complex is present in human cells. However, we consistently found that PRAME complexes contained primarily OSGEP and LAGE3, and very low or undetectable levels of the other two subunits TP53RK and TPRKB. This might reflect a PRAME-mediated disruption of an otherwise stable EKC complex, or a preferential binding of PRAME to an already existing (and possibly dynamic) EKC submodule that was not previously detected in yeast. It is tempting to speculate that PRAME could mediate ubiquitination of EKC components via recruitment of the Cul2 E3 ligase complex. In an attempt to test this hypothesis, we performed ubiquitination assays upon transient transfections in HEK293 cells in presence or absence of the proteasome inhibitor MG132, and in vitro assays with purified proteins. As expected for BC-box proteins, we detected polyubiquitin chains associated with PRAME itself. However, the high background signals in these assays did not allow us to detect a putative ubiquitination of EKC subunits. Hence, our assays could not rule out that EKC might be ubiquitination targets of Cul2-PRAME, and more efforts are needed to test this hypothesis experimentally.

Interestingly, dynamic regulation of EKC components has been reported: the kinase activity of Bud32p was repressed by association with Kae1p (Hecker et al, 2008), but promoted by binding to Cgi121p (Mao et al, 2008). Moreover, we found that the protein levels of OSGEP can be significantly modulated by association with other proteins (Fig. 4), with LAGE3 exerting the most pronounced positive effect, followed by TP53RK and PRAME. Interestingly, the effect of LAGE3 correlates with the formation of tetrameric complexes in which an LAGE3 dimer binds two OSGEP moieties (Fig. 4A), which are likely not in direct contact with each other (Mao et al, 2008).

The results of our transient transfection experiments and those obtained using the baculovirus system suggested (i) that PRAME is able to interact independently with LAGE3 and OSGEP and (ii) that LAGE3 enhances the association of OSGEP with PRAME. Considering the dimerization properties of the LAGE3-OSGEP submodule, we therefore considered the possibility that multiple PRAME moieties might be present in the same protein complex. In this case, both epitope-tagged PRAME and endogenous PRAME should be present in the same protein complex in K562-TAG-PRAME cells, since the expression
level of the two proteins were similar. Contrary to this hypothesis, however, TAG-PRAME did not co-immunoprecipitate endogenous PRAME in our experiments (see Fig. 2 lane 4). Although we cannot formally exclude the possibility that TAG-PRAME might compete with the endogenous protein, our data are most consistent with the idea that there is only one PRAME molecule per complex. Notably, our data suggest that the interaction between PRAME and LAGE3-OSGEP is very stable, since it was efficiently detected in all assays and despite the long incubations times in our protein-complex purification protocols. However, we cannot exclude that high levels of PRAME might be required for the interactions to take place in cells. Taking into account the published crystal structures of archaeal EKC, we propose the models depicted in Figure 5C for the architecture of the human EKC complex and the relationship between EKC and Cul2-PRAME ligases. Further experiments are needed to confirm the stoichiometry of PRAME-EKC complexes, and to characterize the interaction surfaces involved.

Besides the functional experiments in yeast, no information is available yet about EKC functions in other organisms. At present, only studies focusing on single subunits of the human complex have been reported. LAGE3 appears to be ubiquitously expressed, while the two closely related LAGE1 and LAGE2 are cancer-testis antigens with high expression confined to healthy testis and cancer tissues (Alpen et al., 2002). Interestingly, expression of the N-terminal domain of LAGE3 fused to the C-terminus of Pcc1p could partially rescue the growth defects of a Pcc1 N-terminal mutant yeast strain, while full length LAGE3 could not substitute for Pcc1 (Kisseleva-Romanova et al., 2006). The OSGEP gene was found expressed in all somatic tissues tested and it lies in a head-to-head orientation with the tightly regulated APEX gene, which encodes a multifunctional DNA repair enzyme (Ikeda et al., 2002). Analyses of their bidirectional promoter identified a CCAAT box crucial for OSGEP transcription. Furthermore, OSGEP could partially rescue the growth defects of the yeast kae1Δ mutant (Kisseleva-Romanova et al., 2006), underlining degrees of functional conservation. Human TP53RK was first cloned by cDNA subtraction as an interleukin-2 upregulated gene in cytotoxic T-cells, transcripts were detected in a number of normal tissues and cancer cell lines, and it was shown to phosphorylate the human oncoprotein p53 at Serine 15 (Abe et al., 2001). Recombinant TP53RK expressed in bacteria was found catalytically inactive, but incubation with COS-7 cell lysates was sufficient to activate the enzymatic activity, suggesting the need for activators and po-
tentially protein interactors. The kinase Akt/PKB was found to play a role in the activation of TP53RK through phosphorylation of the Ser250 residue (Facchin et al., 2007). Underscoring a high level of functional homology, TP53RK could partially rescue the growth phenotype of Bud32 deletion in yeast, and Bud32 could interact with and phosphorylate human p53 in vitro (Facchin et al., 2003), although yeast does not possess a bona fide p53 homologue. Human TPRKB was first isolated by a two-hybrid screen using TP53RK as a bait, and was suggested to inhibit the binding of TP53RK to p53 (Miyoshi et al., 2003). Expression of TP53RK was detected in the high-PRAME expressing cell line K562 as well as in the low-PRAME expressing cell line KG1 (Miyoshi et al., 2003). Gene expression data obtained from the GeneSapiens database further suggests that EKC subunits are expressed in most if not all human tissues (Suppl. Figures S2-S4).

Considering the extreme conservation of EKC components during evolution, it can be expected that at least some of the functions and pathways already identified for this complex in yeast could be valid for human cells as well. However, it cannot be excluded that human EKC acquired additional, yet to be identified functions during evolution. The absence of efficient knock-down or knock-out models have so far hindered advanced functional analysis of EKC in higher eukaryotes. Future studies will have to address these issues, and in particular to understand the connections between apparently different functions ascribed to this complex, like phosphorylation of the p53 oncogene, chemical modification of tRNAs, and physical association with promoter regions.

Kae1p and Pcc1p have been found to associate with active chromatin and to regulate expression of some of the yeast promoters to which they are recruited (Kisseleva-Romanova et al., 2006). A pcc1-4 mutation prevented activation of GAL1 by impairing recruitment of the SAGA histone acetyltransferase and mediator without affecting recruitment of the activator Gal4p. In contrast, while Pcc1p was recruited to the constitutively active PMA1 and the heat-shock inducible HSP104 genes in a transcription-dependent manner, transcription of these genes was not affected in the pcc1-4 mutant.

Our ChIP assays revealed that LAGE3 and OSGEP are also present on chromatin and, in particular, that they colocalize with PRAME at many of its target sites, consistent with our protein-protein interaction data. Since neither PRAME nor EKC proteins possess known DNA-binding domains, it is
not clear whether they can bind DNA directly. Genome-wide ChIP-seq experiments have recently revealed that PRAME associates with transcriptionally active NFY-regulated promoters and nearby enhancers (Costessi et al., 2011). Therefore, the transcription factor NFY might play a direct or indirect role in the targeting of a Cul2-PRAME-EKC complex to chromatin. It is also not clear whether there is a hierarchy in the association of PRAME and EKC proteins with chromatin. In order to address this point, we established stable cell lines with downregulation of endogenous PRAME using the published retroviral pRetroSuper system in either K562 wild type or K562-TTE-LAGE3 cells and we performed ChIP experiments. Unfortunately, despite a 70% downregulation of PRAME mRNA levels, our preliminary results did not support a solid conclusion on the hierarchy of binding to chromatin for PRAME and EKC. More experiments, and likely in a different cell and knockdown system, are required to address this point.

Importantly, EKC proteins have an ancient evolutionary origin and seem to be ubiquitously expressed in adult tissues, while PRAME appeared more recently in evolution, and its expression seems to be tightly confined to particular developmental stages and tissues and to be particularly deregulated in oncogenesis. These characteristics are consistent with a model in which EKC can associate with chromatin independently of PRAME. Only when PRAME is expressed would a Cul2-PRAME ubiquitin ligase be recruited to chromatin by virtue of specific protein-protein interactions between LAGE3-OSGEP and PRAME.

In an attempt to study the putative function of EKC subunits on target gene transcription, we established K562 cells with stable downregulation of OSGEP or LAGE3 using pRetroSuper vectors and performed genome-wide mRNA-seq analyses. Despite a 50-80% knockdown efficiency for OSGEP and LAGE3, we did not detect genes with significant expression changes when compared to a GFP knockdown control. Our inconclusive data could of course be the result of inefficient downregulation of the target genes. Although the K562 cell line was successfully used to dissect the protein-protein interaction network, we cannot exclude that this cell line might not be suitable for the study of the finer putative chromatin functions and signaling of EKC and PRAME.

The development of ChIP-grade antibodies against EKC components will be a crucial step to address important questions as to whether PRAME target
sites are a subset of EKC sites (or vice versa), and to address the functional relationships between EKC, Cul2-PRAME ligases and NFY on transcriptional regulation of target genes, and potentially other pathways like the recently described modification of tRNAs.

Taken together, our data reveal a novel link between the human oncoprotein PRAME and the ancient EKC complex. Our data clearly show that PRAME can recruit Cullin2 ubiquitin ligases to EKC, and that these complexes associate with the same genomic regions, supporting a functional link in the same pathways. Our results add a new twist to both PRAME and EKC biology and provide an important contribution to understanding the molecular mechanisms in which these proteins are active, both in healthy and cancer cells.
MATERIALS AND METHODS

Cell culture and stable cell lines
HEK293T cells were cultured in DMEM and K562 and HeLaS3 in RPMI medium (Gibco, Invitrogen) at 37°C in 5% CO2; cell lines were obtained from ATCC. Both media were supplemented with 5% Glutamax, 10% fetal bovine serum, 100 U/ml of penicillin, 100 µg/ml of streptomycin (Gibco, Invitrogen). K562 and HeLaS3 stable cell lines were generated with retroviral constructs as described (Costessi et al, 2011). Transient transfections were performed with Lipofectamine2000 (Invitrogen).

Antibodies and western blot
Mouse monoclonal HA-12CA5 and Myc-9E11 were produced in-house from hybridoma cultures, PRAME antibodies were previously described (Costessi et al, 2011). Commercial antibodies used were: mouse monoclonal FLAG-M2 (Sigma), c-Myc monoclonal (Roche Applied Science), rabbit HA (Bethyl Laboratories), mouse monoclonal HA.11 (Covance MMS-101P), rabbit Cul2 (Zymed Laboratories 51-1890), mouse ElonginC (BD Transduction Laboratories 610760), goat ElonginB (Santa Cruz P-16, sc-23407), mouse TPRKB (Abcam ab68245), mouse BB2 against TY1 tag (Diagenode), GFP (Santa Cruz). Proteins were separated by conventional SDS-PAGE or NuPAGE 4-12% Bis-Tris gels (Invitrogen) run with MES buffer. Western blots were visualized by ECL (GE Healthcare) or Odyssey (LiCor) for quantification.

Plasmids and cloning
A modified version of the retroviral vector LZRS(Zeo) with an improved MCS was made by ligation of a synthetic oligo to generate LZRSn(Zeo). The sequence coding for the StreplII-TEV-MYC-3xHA tag was excised with BamHI and NotI from pcDNA5-TAG-PRAME and subcloned in LZRSn(Zeo) to generate LZRSn-TAG, where the EcoRI site is in the same frame as in the pTTE retroviral vector (Costessi et al, 2011).

The TTE cassette was subcloned from pTTE (Costessi et al, 2011) to pcDNA3.1(+) using BamHI and Xhol sites to generate pcDNA-TTE for transient transfections.

Full-length human OSGEP was amplified by PCR from cDNA prepared from K562 cells to generate a NotI-EcoRI-OSGEP fragment, which was cloned into pFLAG-CMV2 (pFLAG-OSGEP). OSGEP was subsequently subcloned as an EcoRI-Sall fragment into the EcoRI and Xhol sites of pTTE and pcDNA-TTE, to generate pTTE-OSGEP and pcDNA-TTE-OSGEP, respectively. OSGEP was cloned as a NotI-Sall fragment downstream the TAG in LZRSn(Zeo), generating LZRSn-TAG-OSGEP.

LAGE3 CDS was obtained from Invitrogen, PCR amplified to introduce EcoRI and Xhol sites, and subcloned in the same sites of pTTE, pcDNA-TTE, and LZRSn-TAG-LAGE3.

Full-length TP53RK/PRPK and TPRKB were amplified by PCR from cDNA of K562 cells and subcloned as EcoRI and Xhol fragments to generate all constructs used.

Mutagenesis to generate OSGEP and LAGE3 mutants was performed using the QuikChange II XL site-directed mutagenesis kit (Stratagene) according to the manufacturer’s instructions.

All constructs were checked by sequencing.

Purification of protein complexes and mass spectrometry analysis
Protein complex purifications and mass spectrometry were performed from nuclear extracts (unless otherwise indicated) essentially as described (Costessi et al, 2011). For HeLaS3 cells the protein extraction protocol was performed with a tighter douncer (pestle B). The quality of cytoplasmic and nuclear extractions were tested by immunoblotting of equivalent volume fractions for alpha tubulin (cytoplasmic marker) and HDAC2 (nuclear marker).

For TAG-PRAME purifications with MYC antibody, about 10 ml of protein extracts (~100 mg) were incubated with 50µl (MYC1 experiment) or 600µl (MYC2 experiment) of crosslinked MYC beads and peptide elution was performed. For protein complex purifications with HA antibody, about 5-6 ml of protein extracts were incubated with 600µl HA-12CA5 crosslinked beads and acidic glycine (pH 2.9) was used for elution.

The data analysis steps of peptide-to-protein remapping and emPAI calculation were automated with a PERL script to generate output tables of unique protein IDs with peptide frequencies and emPAI values for each samples (see Supporting Information Protocol S1 and Analysis Script.
S1). The protein list was manually filtered to discard proteins identified in the control samples (false positives).

**Expression of recombinant proteins in Sf21 insect cells**

CDNAs encoding wild type or mutant PRAME, OSGEP, LAGE3, TP53RK and TPRKB were subcloned into pBacPAK 8 and recombinant baculoviruses were generated. SF21 cells were cultured at 27°C in SF-900 medium supplemented with 10% fetal bovine serum. SF21 cells were co-infected with the recombinant baculoviruses indicated in the figures. 50 hours after infection, cells were collected, lysed, and processed as described (Mahrour et al, 2008).

**ChIP and qPCR**

Chromatin harvests, ChIPs and qPCR analyses were performed as previously described (Costessi et al, 2011). The BB2 antibody recognizing the TY1 epitope (Diagenode) was used to ChIP TTE-tagged proteins from K562 stable cell lines.
ACKNOWLEDGMENTS

We are grateful to Edwin Lasonder for his advice on mass spectrometry data analysis and to Blaise Alako for automating the data analysis with a PERL script. We thank Gert Vriend and Hanka Venselaar (CMBI, Nijmegen) for their advices on protein structures, and Anita Kaan for technical assistance with DNA cloning. We also thank our colleagues for helpful discussions and suggestions.
REFERENCES


receptor. *Immunity* 6: 199–208


SUPPLEMENTARY INFORMATION

Protocol S1.
Detailed protocol for comparative analyses of mass spectrometry data.
Available online at www.plosone.com

Analysis script S1.
PERL script mapPep2Prot_v0.3.pl for analysis of mass spectrometry data.
Available online at www.plosone.com
Detailed expression profiles of MAG, TBP, CUL2, and PRAME from the GenesApariens database.

Normalized relative gene expression levels in healthy (green) and cancer (red) tissues are plotted as boxplots. MAG and TBP are shown as references: MAG is a known neuronal marker gene and shows an expression profile that is highly specific for the central nervous system. On the contrary, TBP is a ubiquitously expressed gene. Similarly to TBP, the expression profile of CUL2 indicates ubiquitous expression in both healthy and cancer tissues. On the contrary, PRAME is expressed mainly in the healthy testis, and is upregulated in a number of cancers.
Detailed expression profiles of EKC subunits from the GeneSapiens database. Normalized relative gene expression levels in healthy (green) and cancer (red) tissues are plotted as boxplots. The expression profiles of LAGE3, OSGEP, TPRKB, and TP53RK indicate expression in all or most healthy and cancer tissues.
Figure S3. Gene expression correlation plots between *PRAME* and subunits of Cul2 ligases or the EKC complex.

Co-expression plots were generated from the GeneSapiens database for *PRAME* and Cul2 or each of the EKC subunits. Correlations plots are shown for healthy tissues (A), and cancer tissues (B). The plots indicate that the expression of the genes tested do not correlate linearly. In particular, a large number of samples are characterized by low or no *PRAME* expression, while Cul2 and EKC subunits are expressed at higher levels.
Supplementary figure 4

Figure S4. Gene expression correlation plots between subunits of Cul2 ligases and subunits of the EKC complex.

Co-expression plots were generated from the GeneSapiens database for CUL2 and TCEB1, and couples of EKC subunits. Correlations plots are shown for healthy tissues (A), and cancer tissues (B).
CHAPTER 5

DEVELOPMENT OF REAGENTS AND TOOLS for the study of the human oncoprotein PRAME

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Expression of the human oncoprotein PRAME has been investigated in several tumours and numerous cohorts of cancer patients. However, studies of PRAME function at the molecular and cellular level are scarce and are also hindered by the lack of validated antibodies. In this chapter, we report the development and validation of PRAME antibodies raised against five different peptides. We successfully raised sera and affinity-purified antibodies for use in western blot, immunoprecipitation and ChIP-seq with high sensitivity and specificity. We established that PRAME is a chromatin-associated protein and we generated the first genome-wide ChIP-seq binding profiles of epitope-tagged as well as endogenous PRAME in K562 cells. Finally, we investigated the reported interplay between PRAME and retinoic acid signaling in the A375 melanoma cell line using two independent silencing constructs and complementation with a knockdown-resistant PRAME. In contrast to published reports, our data suggest that the growth phenotype identified in PRAME knockdown cells treated with retinoic acid is likely caused by “off-target” knockdown of a gene unrelated to PRAME by one of the silencing constructs used.
INTRODUCTION

The human *PRAME* gene was first identified in 1998 as the gene coding for the antigen responsible for the anti-tumour immune response in a melanoma patient (Ikeda et al., 1997). Since then, numerous studies have investigated the transcript expression levels of *PRAME* in patients affected by different types of cancers, establishing that this gene is often overexpressed in a wide range of malignancies. Investigation of the molecular and cellular functions of PRAME, however, has been hindered by the lack of validated reagents for use in experimental research. For instance, several assays rely on antibodies with high specificity for the protein of interest, and it takes significant efforts to develop and test antibodies when these are not available from commercial suppliers. Therefore, we invested on the development of a toolbox of reagents, including antibodies, expression plasmids, and suitable epitope tags.

In this chapter, we report the development of a number of rabbit polyclonal antibodies against human PRAME and their validation for western blot, immunoprecipitation, and ChIP assays. We further discuss strategies that can be undertaken to generate genome-wide ChIP-seq binding profiles of proteins of interest, and we present an epitope-tagged approach based on a double TY1 epitope that we successfully applied for ChIP-seq profiling of PRAME in K562 cells. A recent study reported that downregulation of PRAME sensitized A375 melanoma cells to retinoic acid-induced growth arrest, suggesting a repressive function for PRAME on retinoic acid signaling (Epping et al., 2005). Here, we report data generated with two independent knockdown constructs, which do not support the published model and rather suggest that the phenotype observed is likely caused by an “off-target” effect of the knockdown vector used in the literature.
RESULTS AND DISCUSSION

Development of anti-PRAME peptide antibodies

At the beginning of our study, antibodies for human PRAME were not commercially available. Therefore, we performed an immunization protocol in rabbits to raise anti-PRAME peptide antibodies (see Table 1 and Figure 1). Peptides 2, 3, 4, and 5 were designed in regions of PRAME that showed low sequence homology with other PRAME-like proteins, in order to reduce the chance of cross-reactivity; peptide 7 was previously described (Ikeda et al., 1997). Each peptide was administered to two rabbits in parallel, and crude sera were collected every three weeks for a maximum of ten harvests (S1 to S10).

Sera were first tested in dot blot assays to assess if the sera contained antibodies directed against the peptides administered. The results showed that all rabbits mounted an efficient immune response against the corresponding peptide (data not shown).

Next, we tested the sera in western blot assays using protein extracts from 293T cells transiently transfected with pcDNA5-TAG-PRAME (StrepII-TEV-MYC-3xHA-PRAME, 73 kDa) and control untransfected cells. Importantly, 293T cells do not express significant levels of endogenous PRAME. Figure 2 shows the results obtained with sera from bleed S5 (panel A) and bleed S8 (panel B).
Validation of crude sera in western blot assays.

(A-B) Western blot experiments with PRAME crude sera using extracts from HEK293T cells (indicated as C, control) and from HEK293T transiently transfected with pcDNA5-TAG-PRAME (indicated as PR). Results are shown for sera obtained with bleed 5 (A) and bleed 8 (B). About 20 micrograms of whole cell protein extracts were loaded for the control cell line, and about 40 micrograms for TAG-PRAME transfected cells. Sera were used at a 1:1000 dilution in 5% skimmed milk and PBS/T, and secondary antibody conjugated to HRP. The two main bands recognized for TAG-PRAME are indicated with arrows. Sera recognizing TAG-PRAME are underlined.

(C) Peptide competition assay with serum V19. Whole cell extracts from the cell lines indicated were probed with 1:1000 dilutions of preimmune serum from rabbit V19 (Preimm V19), V19/S5 serum, or V19/S5 serum (7 μl) after a 2 hours incubation at 37°C with 50 μg peptide 2 and 100 μl PBS.

(D) Peptide competition with serum V28, as in (C). Asterisk indicates unrelated bands.

B). Sera V19, V20, V22, V27 and V28 successfully detected TAG-PRAME, while the other sera did not work efficiently in these assays.

The V19 and V20 sera recognized also an additional band of about 50-55 kDa. Notably, this band was not detected by antibodies raised against other peptides, nor by the pre-immune serum from rabbit V19 (Fig. 2C), suggesting that reactivity against this band was mediated by antibodies raised by the immunization protocol against peptide 2. Consistent with this hypothesis, a
peptide competition experiment revealed that the band was not recognized when the V19 serum was pre-incubated with peptide 2 (Fig. 2C). Therefore, we conclude that the V19 and V20 sera cross-react with a 50-55 kDa protein unrelated to PRAME, but containing an epitope similar or identical to peptide 2.

Remarkably, V28 (but not V27) sera recognized two additional bands with an apparent molecular weight of 40-45 kDa (Fig. 2A-B). These bands were specifically recognized in 293T cells transiently transfected with pcDNA5-TAG-PRAME or K562 cells transduced with LZRS-TAG-PRAME, but not in the control untransfected cells (Fig. 2D). The preimmune serum from rabbit V28 detected the same bands (Fig. 2D), indicating that the antibodies against these proteins were already present in the serum of rabbit V28 before the immunization protocol, and were not raised by the immunization with peptide 7. This was confirmed by a peptide competition experiment in which V28 serum was pre-incubated with peptide 7 before western blot detection: as expected, detection of PRAME was inhibited, but the two lower “aspecific” bands were still efficiently detected (Fig. 2D).

### Table 1. Peptides used to generate PRAME antibodies and corresponding sera.

<table>
<thead>
<tr>
<th>Peptide ID</th>
<th>Peptide sequence</th>
<th>Peptide position</th>
<th>Rabbits ID</th>
<th>Affinity-purified Ab</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peptide 2</td>
<td>NRASLYSFPEPEA</td>
<td>139-151</td>
<td>V19, V20</td>
<td>P1 (V19/S5)</td>
</tr>
<tr>
<td>Peptide 3</td>
<td>FIPVEVLVDLFL</td>
<td>172-183</td>
<td>V21, V22</td>
<td>P2 (V22/S5)</td>
</tr>
<tr>
<td>Peptide 4</td>
<td>LLHSIHASSYS</td>
<td>265-277</td>
<td>V23, V24</td>
<td>-</td>
</tr>
<tr>
<td>Peptide 5</td>
<td>RASATLQDVFEDE</td>
<td>375-387</td>
<td>V25, V26</td>
<td>-</td>
</tr>
<tr>
<td>Peptide 7</td>
<td>FPEPEAAQPMTKRRKVDG</td>
<td>146-163</td>
<td>V27, V28</td>
<td>P3 (V27/S6) P4 (V28/S5)</td>
</tr>
</tbody>
</table>

**Purification of anti-PRAME antibodies**

The best performing sera V19/S5, V22/S5, V27/S6 and V28/S5 were further subjected to two different purification protocols. The first protocol consisted of tandem precipitations with caprilic acid and ammonium sulphate to deplete abundant serum proteins and enrich in immunoglobulins. Caprilic acid in mildly acidic conditions precipitates most serum proteins, with the exception of IgG molecules. The soluble antibodies are then precipitated with ammonium sulphate and resuspended in PBS. These IgG-enriched fractions were referred to as “clean sera”: V19cl, V20cl, V22cl, V27cl, V28cl.

The second protocol was a peptide affinity purification that specifically
selects the antibodies that have affinity for the peptide used in the rabbit immunization. The peptide is first crosslinked to an insoluble agarose matrix to generate an affinity column, through which the serum is flowed. Antibodies that recognize the peptide bind to the insoluble matrix, while unrelated antibodies and serum proteins flow-through. Eventually, the immobilized antibodies are eluted from the column with acidic glycine, which destabilizes the antibody-peptide interactions. The antibodies purified with this approach were labelled “purified antibodies”: P1 to P4 (Table 1).

The crude sera, the flow-through fractions from the affinity column and the purified antibodies (P1-P4) were tested in ELISA to assess the efficiency of the protocol (data not shown). The results confirmed that P1 (derived from V19/S5 serum) and P2 (from V22/S5) were enriched in specific antibodies, while the flow-through fractions had little reactivity against the peptides, as expected for an efficient affinity purification. The purified antibodies P3 (from V27/S6) and P4 (from V28/S5) showed also a good reactivity against the peptides; however, also the flow-through fractions displayed reactivity against peptide 7, indicating that a fraction of PRAME antibodies did not bind efficiently the affinity column and were still present in the flow-through. Cleaned sera and affinity purified antibodies were tested in western blot assays in the same conditions used previously for the crude sera (Figure 3).

Taken together, our experiments established that antibodies raised against peptide 2 (V19, V19cl, V20, V20cl and P1) efficiently recognized TAG-PRAME but also an unrelated protein with an apparent molecular weight similar to PRAME (Figures 2 and 3). Therefore, these antibodies are not suitable for detection of the endogenous PRAME protein, but can be used for detection of epitope-tagged PRAME.

Similarly, sera V22 and V22cl reproducibly detected TAG-PRAME and an additional band of about 40-45 kDa. This band was less prominent in western blots with the purified antibody P2, where an aspecific band running just above TAG-PRAME was identified (Figure 3B).

Sera V23, V24, V25, V26 performed poorly in all experiments and did not react against TAG-PRAME in western blots (Figure 2).

All antibodies raised against peptide 7 (V27, V27cl, V28, V28cl, P3 and P4) displayed a very strong reactivity against PRAME (Figures 2 and 3). A band of 55 kDa was detected with lower intensity as compared to TAG-PRAME in transfected 293T cells but not in untransfected cells: this bands most likely represents untagged PRAME expressed from the transiently
Validation of cleaned sera and affinity-purified antibodies in western blot assays.

(A) Western blot with cleaned sera on whole cell extracts from HEK293T cells (C) and HEK293T cells transiently transfected with pcDNA5-TAG-PRAME (PR).
(B) Western blot with affinity-purified antibodies P1 and P2 on whole cell protein extracts from the cell lines indicated.

(C) Western blot with affinity-purified antibodies P3 and P4 on whole cell protein extracts from the cell lines indicated. Epitope-tagged and endogenous PRAME are indicated.
(D) Western blot on nuclear protein extracts separated on a 4-12% Novex gradient gel. An unrelated protein detected by P4 antibody is indicated with an asterisk.
(E) Nuclear and cytoplasmic extracts from K562 cells show that the unrelated band detected by the P4 antibody is mainly present in the nuclear fraction.
(F-G) Whole cell protein extracts were probed with V27/S6 serum or Abcam PRAME antibody in parallel. The asterisk indicates an unrelated protein detected by Abcam antibody.

**FIGURE 3**
transfected plasmid, since the starting ATG codon of PRAME was still present in the pcDNA5-TAG-PRAME construct.

The P3 and P4 purified antibodies performed very well with whole cell extracts of several cell lines: a single band at the expected molecular weight of PRAME was identified in K562 and FM6 melanoma cells, which express endogenous gene, while no band was detected in the PRAME-negative 293T cells (Figure 3C). Subsequent experiments, however, revealed that the P4 antibody recognized also an unrelated protein with the same apparent molecular weight of TAG-PRAME in nuclear extracts of K562 cells. This unrelated protein was less evident in whole cell extracts (Fig. 3C), cytoplasmic extracts (Fig. 3E), and chromatin fraction (Fig. 7B).

During the course of our study, a commercial antibody became available against human PRAME (ab32185, Abcam). Western blot experiments revealed that this antibody could recognize epitope-tagged PRAME (Figure 3F-G). However, this antibody displayed a significant cross-reactivity with a protein of similar molecular weight as PRAME in all cell lines tested. Therefore, antibody ab32185 (Abcam) could not be used for detection of the endogenous PRAME protein, although this antibody has been used in recent publications (De Carvalho et al., 2011; Partheen et al., 2009; Quintarelli et al., 2008; Partheyen et al., 2008; Passeron et al., 2009).

**Validation of P4 antibody specificity with PRAME knockdown cells.**

(A) The target sites of knockdown constructs pRS-P1 and pRS-P2 are indicated on the coding sequence of human PRAME.

(B) Western blot on the cell lines indicated. HeLaS3-TAG-PRAME cl3 and K562-TAG-PRAME cl12 lines were transduced with the pRS knockdown constructs indicated.
PRAME knockdown

The specificity of the affinity purified P4 antibody was further tested upon knockdown of PRAME. For this purpose, stable cell lines were made using two independent retroviral pRetroSuper constructs targeting PRAME: pRS-PRAME1 and pRS-PRAME2 (further indicated as pRS-P1 and pRS-P2; Figure 4). Double-stable cell lines were made by transduction of zeocin-resistant K562-TAG-PRAME cl.12 and HeLaS3-TAG-PRAME cl.3 with pRS-P1 or pRS-P2 retroviral particles and selection of polyclonal cell populations with puromycin. Western blot analysis with HA antibody established that both knockdown constructs efficiently silenced TAG-PRAME (Figure 4). As expected, the 55 kDa band recognized by P4 antibody was also downregulated, demonstrating that this band corresponds to the endogenous PRAME protein. The same experiment showed that PRAME is not detectable in HeLaS3 wild type cells in these conditions, which is consistent with cDNA gene expression analyses (Chapter 2, Figure 2D).

PRAME antibodies in immunoprecipitation assays

Finally, PRAME antibodies were tested in immunoprecipitation assays using nuclear extracts of K562-TAG-PRAME cells (Figure 5). Immunoprecipitation without antibodies, or with rabbit HA antibodies (Abcam) were performed as negative and positive controls, respectively. Supernatant and IP fractions were analyzed by western blotting with mouse HA.Y11 antibody. The results

**Figure 5**

Validation of PRAME antibodies in immunoprecipitation assays.

Immunoprecipitation experiments on nuclear extracts from K562-TAG-PRAME cl.12 cells. The following antibodies were used: HA Abcam (1 μl), P2 (8 μl), P4 (5 μl), V19cl (5 μl), V27cl (5 μl). Equivalent amounts (18%) of nuclear extract (input), supernatant after IP, and IP fractions (beads boiled after IP) were separated by SDS-PAGE and probed with HA.Y11 antibody and secondary antibody specific for mouse k chains. Next, the same membranes were probed with P4 antibody using a secondary antibody specific for rabbit k chains.
show that P4, V19cl and V27cl antibodies successfully immunoprecipitated TAG-PRAME, while P2 purified antibody did not work in this assay. Moreover, also the endogenous PRAME protein was efficiently immunoprecipitated by V27cl, since it was depleted from the supernatant (Figure 5, lane 13). Notably, the immunoprecipitated endogenous PRAME protein is difficult to visualize in the IP fractions, since it runs at the same height as the heavy chains of the antibodies that are highly abundant in such fractions. Taken together, our results established that PRAME antibodies raised against peptide 2 and 7 performed efficiently also in immunoprecipitation assays.

**PRAME association to chromatin**

PRAME was recently reported to associate with the retinoic acid receptor.
alpha (RARA) and the polycomb protein EZH2, and to mediate repression of transcription of RARA target genes (Epping et al, 2005). However, PRAME binding to chromatin was reported only for the RARB promoter in the retinoic-acid responsive A375 melanoma cell line (Epping et al, 2005). Notably, the K562 cell line was reported to be unresponsive to retinoic acid treatment because of low endogenous expression of retinoic acid receptors (Robertson et al, 1991). Evidence for the association of PRAME with chromatin can be obtained using ChIP and ChIP-seq, though a major hurdle is the absence of validated ChIP-grade antibodies against PRAME.

A general approach to obtain a genome-wide binding profile for a factor of interest is illustrated by the decision tree of Figure 6 (Brinkman & Stunnenberg, 2009). First of all, an appropriate cell system needs to be selected, which involves testing expression of the factor of interest and establishing its association with chromatin. If antibodies and (putative) target sites are available, these can be tested in ChIP-qPCR assays. If a signal is obtained by ChIP-qPCR, this can be further improved by optimizing the antibody concentration, the buffers used, crosslinking and sonication conditions. Subsequently, the specificity of the signals can be validated by using antibodies raised against different epitopes of the protein, knock-out or knockdown systems, or epitope-tagged versions of the protein of interest. When these steps are completed, it is possible to proceed with a genome-wide profiling experiment.

If no signal is obtained in an exploratory ChIP-qPCR assay, either the antibody used is not ChIP-grade, or the presumed genomic sites investigated are not bound by the protein of interest. The ChIP-grade properties of the antibody can be tested performing an immunoprecipitation assay on crosslinked chromatin followed by western blot, or preferably on crosslinked chromatin purified by caesium chloride density centrifugation, which separates free soluble proteins and nucleic acids from the sample, and enriches in chromatin-associated proteins. If the target protein can be IP’ed, the antibody is likely ChIP-grade and a “blind” ChIP-seq experiment can be performed. If putative binding sites are discovered, these can be used in ChIP-qPCR experiments to optimize the assay conditions before performing a new profiling experiment.

If the antibody does not IP the protein of interest, the antibody is not ChIP-grade and alternative approaches can be undertaken, i.e. raising new (ChIP-grade) antibodies, or applying epitope tagging in stable cells lines with retroviral expression systems or BAC transgenics.
In our study, we first investigated the association of PRAME to chromatin in the K562 cell line. We prepared formaldehyde-crosslinked chromatin from K562 wild type and K562-TAG-PRAME cells and fractionated it on a CsCl2 density gradient by ultracentrifugation. This technique separates the free soluble proteins from the chromatin-associated proteins based on their different densities. Five different fractions were collected as indicated in Figure 7A and analyzed by agarose gel electrophoresis and western blot. Agarose gel electrophoresis confirmed that chromatin was enriched in fractions 3 and 4, as indicated by the intense staining present in the wells, which represents crosslinked chromatin that cannot enter the gel matrix (Fig. 7A). As expected, western blotting showed that the core histone H3 was also enriched in these fractions (Fig. 7B). TAG-PRAME and endogenous PRAME were present in the free-protein fraction (fraction 1), but they were also detected in the chromatin fractions, consistent with their association with chromatin in vivo.
(Fig. 10B). Consistent with previous western blot results, a protein unrelated to PRAME with a higher apparent molecular size was recognized by the P4 antibody in the nuclear extracts. Importantly, this protein was not detected in the input or fractionated chromatin fractions, suggesting that the use of V27/P4 antibodies for ChIP assays should not result in cross-reactivity with proteins other than PRAME (Costessi et al., 2011).

In the absence of validated binding sites for PRAME in the K562 cell line, we setup an epitope-tagging approach to identify PRAME binding sites in an unbiased way. For this purpose, we established K562 stable cell lines expressing PRAME fused to a composite N-terminal TTE tag, including two TY1 epitopes and one ERalpha epitope, against which efficient ChIP-grade mono-
clonal antibodies are available: BB2 and F3A6, respectively (Fig. 8A). Expression of TTE-PRAME was tested by western blot in the polyclonal cell line K562-TTE-PRAME and the control cell line K562-TTE carrying the empty vector (Fig. 8B). Next, we performed an exploratory ChIP-seq experiment using the BB2 antibody (against the TY1 tag) and chromatin from K562-TTE control and K562-TTE-PRAME cells. Sequencing was performed on a Solexa GAIIx with a single-read protocol for 36 cycles and generated 4,059,488 reads for the TTE control and 3,615,534 reads for TTE-PRAME after alignment to the human hg18 reference genome. Reads numbers were normalized to the lowest number and average profiles were visualized in the UCSC genome browser. Manual inspection of these profiles revealed a number of loci with specific enrichment of sequence tags for TTE-PRAME (Figure 9). ChIP-qPCR assays were performed with primer sets targeting these putative binding sites, and the assay conditions were optimized to improve the signal-over-background ratio. The results confirmed that the sites tested are bona fide binding sites for TTE-PRAME (see chapter 3).
We then performed ChIP-qPCR experiments using our PRAME antibodies and chromatin from K562 wild type cells. As shown in chapter 3 (Supplementary Figure 1), the V27cl serum gave good recoveries, indicating that it is ChIP-grade and establishing that also the endogenous PRAME protein binds to the genomic loci tested.

Based on these results, we performed a ChIP-seq profiling on K562 wild type cells using the PRAME antibody V27, or the preimmune serum from the same rabbit as negative control. We obtained 3 101 777 sequence reads for PRAME and 2 815 020 sequence reads for the preimmune serum. Screen shots in Figure 9 show genomic regions bound by both TTE-PRAME as well as endogenous PRAME.

Next, we used the MACS algorithm (Zhang *et al.*, 2008) to identify binding sites for TTE-PRAME and PRAME using their corresponding control tracks as negative controls. We iterated the peak calling with different stringency settings: the “e-value” was set to 5, 6, or 7, where 5 corresponds to relaxed settings (that can result in a higher false-positives rate), while an e-value of 7 corresponds to more stringent settings. Table 2 shows that significantly more peaks were called for the endogenous PRAME than for TTE-PRAME.

<table>
<thead>
<tr>
<th>e-value</th>
<th>TTE-PRAME peaks</th>
<th>PRAME peaks</th>
<th>Overlap TTE-PRAME with PRAME peaks</th>
</tr>
</thead>
<tbody>
<tr>
<td>e-5 (m16)</td>
<td>7517</td>
<td>24611</td>
<td>41% (3084)</td>
</tr>
<tr>
<td>e-6 (m16)</td>
<td>2467</td>
<td>14153</td>
<td>56% (1380)</td>
</tr>
<tr>
<td>e-7 (m16)</td>
<td>1142</td>
<td>9228</td>
<td>63% (579)</td>
</tr>
</tbody>
</table>

This is consistent with the lower recoveries obtained for TTE-PRAME in ChIP-qPCR experiments and a higher signal-to-noise ratio for the PRAME serum than the BB2 antibody. We next calculated the overlap between binding sites identified for TTE-PRAME and endogenous PRAME. Notably, the percentage of overlapping peaks was dependent on the stringency applied to the peak calling: the overlap was 41% for e5 peaks, 56% for e6 peaks, and 63% for e7 peaks. This suggests that low e-values (low stringency) result in larger pools of putative binding sites, likely containing a higher percentage of false positives, while higher e-values (high stringency) result in the identification of smaller numbers of high-confidence peaks, although binding sites might be missed.
Take together, our data demonstrate that the oncoprotein PRAME is a chromatin-associated factor, we validated an epitope-tagging approach as well as a PRAME antibody for ChIP-seq analyses and generated the first genome-wide profile of PRAME association to chromatin.

**PRAME and retinoic acid signaling**

Recent work indicated that overexpression of PRAME prevents the growth arrest and apoptosis induced by histone deacetylase inhibitors (HDACIs) (Epping et al, 2007). The authors proposed a model in which HDACI-induced growth arrest requires an active retinoic acid signaling pathway, which PRAME would repress via direct protein-protein interactions with the retinoic acid receptor alpha (RARalpha), recruitment of the EZH2 polycomb protein and eventual transcriptional repression of RARalpha target genes (Epping et al, 2005).

In support of their model, the authors reported that knockdown of PRAME in A375 melanoma cells caused downregulation of the retinoic acid receptor alpha (RARa) at the protein level (Epping et al, 2005, Figure 6F),
which was interpreted as a sign of derepression of retinoic acid signaling. Remarkably, the same experiment presented in a follow-up publication showed no difference in RARalpha levels upon PRAME knockdown (Epping et al., 2007, Figure 4E). Although the authors did not comment on this discrepancy, this discrepancy raises doubts about the reproducibility of the reported RARalpha-PRAME interplay.

Further, the authors reported that knockdown of PRAME sensitized A375 cells to the growth inhibiting effects of retinoic acid treatment (Epping et al., 2005). Importantly, complementation of the cells with a knockdown-resistant version of PRAME was shown to re-establish a normal growth curve, indicating that expression of PRAME confers resistance to retinoic acid-induced growth arrest.

We have recently characterized PRAME as a component of cullin2-based E3 ubiquitin ligases (see chapter 3 and Costessi et al., 2011). In order to confirm whether PRAME can act as a repressor of the retinoic acid signaling as reported, we performed growth assays upon downregulation of PRAME in
the presence or absence of retinoic acid.

First, we generated A375 stable knockdown cell lines by retroviral transduction using two different silencing constructs, pRS-P1 and pRS-P2 and compared them to wild type untransduced cells. Construct pRS-P1 was the one used in the publication of Epping et al. (Epping et al, 2005), while pRS-P2 is an independent construct targeting a region of the PRAME transcript partially overlapping with the region targeted by pRS-P1 (Figure 4). Cells were treated with a final concentration of 1 µM all-trans retinoic acid (atRA) or the equal volume of vehicle (ethanol), and knockdown cells were kept under puromycin selection during the course of the experiment. Cumulative cell numbers reported in Figure 10A-B reflect a growth inhibition on A375-pRS-P1 cells upon retinoic acid treatment. On the contrary, A375-pRS-P2 cells were only slightly affected as compared to the wild type control cells, even though PRAME was downregulated to similar levels by both silencing constructs (Figure 10C).

Cell proliferation assays upon PRAME knockdown and control vectors.
(A) Growth curve of A375 stable cell lines upon treatment with 1µM atRA or vehicle (ethanol).
(b) Cumulative cell numbers at the end of the growth assay (day 21).
(c) Gene expression analysis of PRAME in A375 cell lines used for the growth assay. Expression was measured in triplicate and normalized to a pool of 3 control genes (AU, arbitrary units).
Next, we generated wild type and BC-box mutant versions of PRAME that are resistant to knockdown (KDR) in order to test if ectopic re-expression of PRAME could rescue the growth phenotype, as previously reported (Epping et al, 2005, Supplementary Figure 3). For this purpose, we used the LZRS retroviral system to express either wild type or the M2 BC-box mutant TAG-PRAME in A375-pRS-P1, and the empty LZRS vector was used as negative control. Unexpectedly, a growth assay on these cell lines revealed that complementation of A375-pRS-P1 cells with wild type PRAME did not rescue their growth phenotype, in disagreement with the published data (Figure 11A-B). Expression of TAG-PRAME_KDR was confirmed by cDNA expression analysis (Figure 11C).

Given this discrepancy, we setup an independent knockdown experiment in which two control lines were also included using the empty pRS vector and the pRS-GFP vector expressing a silencing hairpin against GFP. New PRAME knockdown cell lines were generated at the same time with pRS-P1 and pRS-P2 vectors. Growth curves of these lines are reported in Figure 12A-B.

First, cells carrying the empty vector grew better than the cells expressing the GFP control hairpin. This result is consistent with reports that overexpression of silencing hairpins can overload the silencing machinery and therefore lead to growth defects and cell mortality in vitro and in vivo, independent from the downregulation of specific target genes (Grimm et al, 2006). This result further underscores the critical importance of proper controls for knockdown experiments.

Second, no significant difference was found between A375-pRS-GFP control and A375-pRS-P2 knockdown cells, while A375-pRS-P1 cells were clearly affected: almost no A375-pRS-P1 cells survived after three weeks of treatment with retinoic acid, consistent with the first experiment. Untreated A375-pRS-P1 cells displayed a slower growth rate than other cell lines and this might be caused by an increased sensitivity to the low levels of retinoic acid present in the medium used.

In order to investigate the lack of phenotype in A375-pRS-P2 lines, we measured PRAME expression levels during the course of the experiment (day 17). The results displayed in Figure 12C established that both pRS-P1 and pRS-P2 vectors lead to a significant and similar downregulation of PRAME. Therefore, the lack of phenotype for A375-pRS-P2 cells could not be explained by inefficient knockdown of PRAME by this vector as compared to
pRS-P1; atRA-treated A375-pRS-P1 cells could not be tested because too few cells were present in the plates.

Taken together, independent experiments with polyclonal cell populations consistently showed that vector pRS-P1 caused a severe growth phenotype in A375 melanoma cells exposed to 1 µM retinoic acid. However, the pRS-P2 vector did not affect cell growth compared to the pRS-GFP control, even though it silenced PRAME expression with equal efficiency as pRS-P1 (~80%).

Moreover, complementation of A375-pRS-P1 cells with knockdown-resistant TAG-PRAME did not alleviate the growth phenotype. Formally, we cannot exclude that overexpression of TAG-PRAME_KDR and the epitope tag used might have prevented an efficient rescue of the growth phenotype. However, this epitope tag has been successfully used for the purification of native PRAME-containing protein complexes, showing that this tag does not alter the protein structure of PRAME or its ability to establish protein-protein interactions.

In contrast to the report of Epping et al., our data suggest that the growth phenotype induced by the pRS-P1 silencing vector is highly unlikely caused by down-regulation of PRAME, but rather by down-regulation of an unrelated gene which is specifically targeted by construct pRS-P1, but not by pRS-P2.
CONCLUSIONS

In this chapter, we described the development of antibodies against the human oncoprotein PRAME, we tested the reported interplay between PRAME and the retinoic acid signaling using knockdown systems, we established the in vivo chromatin association of PRAME and reported the first genome-wide ChIP profiling of PRAME.

We immunized a total of ten rabbits against five different peptides designed in regions of human PRAME with low sequence similarity to other annotated or predicted PRAME-like proteins (Figure 1 and Table 1). Sera and affinity-purified antibodies were extensively tested in dot blot, western blot, immunoprecipitation and ChIP assays and a number of them were characterized by high sensitivity and specificity (Figures 2-5). Taken together, our data indicate that antibodies raised against peptide 2 (V19, V20, P1) and peptide 3 (V22, P2) can be efficiently used to detect epitope-tagged versions of PRAME by western blot, but are not suited for endogenous PRAME because of cross-reactivity with other proteins of similar size. On the contrary, antibodies raised against peptide 7 (V27, V28, P3, P4) are suitable for detection of both epitope-tagged as well as endogenous PRAME. Furthermore, V19, V27, P4 (Figure 5) and V28 (not shown) antibodies performed successfully in immunoprecipitations assays. Notably, our antibodies performed better than a commercially available PRAME antibody, which had been used in a number of recent publications. Our results underscore the importance of extensive testing of both specificity and sensitivity of antibodies.

We further discussed a general experimental strategy to generate ChIP-seq binding profiles of a factor of interest (Figure 6), which we applied to our study of the PRAME oncoprotein. We established that PRAME is present in the crosslinked-chromatin fraction in caesium chloride gradient experiments (Figure 7) and we successfully applied an epitope-tagging approach with a double TY1 epitope to discover genomic binding sites for TTE-PRAME with a “blind” ChIP-seq profiling (Figure 8 and 12). Moreover, we validated the ChIP-profiling properties of one of our PRAME antibodies, and we generated the first profiles for the endogenous PRAME protein in K562 cells (Figure 9).

We investigated the reported function of PRAME as dominant repressor of retinoic acid signaling. In particular, we aimed to investigate if the interaction of PRAME with Cullin2-based ubiquitin ligases plays in role in the
reported repression. We performed cell proliferation assays in the presence of all trans retinoic acid (atRA) on A375 melanoma cells, similarly to recent studies from the Bernards lab (Epping et al., 2005). While published work included only one silencing vector, we used two different knockdown plasmids in our experiments (pRS-P1 and pRS-P2). Notably, the regions of the PRAME transcript which are targeted by the two silencing plasmids showed a significant overlap (Figure 4) and both constructs resulted in ~80% downregulation of PRAME transcript (Figures 4B, 6-8).

Unexpectedly, several independent experiments reproduced the reported growth phenotype only in A375-pRS-P1 cells, but not in A375-pRS-P2 cells (Figure 10 and 12). Moreover, complementation with a knockdown-resistant PRAME did not rescue the growth phenotype of A375-pRS-P1 cells in our assays (Figure 11), although a rescue was previously reported (Epping et al., 2005). Since wild type PRAME did not rescue the growth phenotype, we could not investigate the potential role of a BC-box mutant PRAME in this assay. Taken together, our data suggest that the growth phenotype induced by the pRS-P1 silencing vector might not be caused by down-regulation of PRAME, but rather by down-regulation of an unrelated transcript which is specifically targeted by construct pRS-P1, but not by pRS-P2. A genome-wide mRNA-seq profiling experiment on these cell lines could potentially identify this gene and shed light on the reported growth phenotype.
MATERIALS AND METHODS

Antigen preparation and rabbit immunization

PRAME peptides with a leading ATA group were synthesized at the Dept. Organic Chemistry (Radboud University) and checked by mass spectrometry. The peptides were first conjugated to KLH (Inject Mariculture Keyhole Limpet Hemocyanin, Pierce, product #77600) according to standard protocols.

A total of ten rabbits (two rabbits for each peptide) were subjected to a first immunization by lymphnode injection of 200 µg of KLH-peptide and Freund adjuvant. Every three weeks, an immunization booster was performed by subcutaneous injection of 400 µg of KLH-peptide and Freund adjuvant. Blood was harvested two weeks every each immunization booster. Rabbit V21 died prematurely during the experiment before the fifth serum harvest. The rabbits were subjected to 8 cycles of booster immunization and blood collection, except rabbits V19, V22, V27, and V28 which were subjected to a total of 10 cycles.

Antibody purification with caprilic acid/ammonium sulphate precipitation

4 ml of rabbit sera were mixed with 2 volumes (8 ml) of 60 mM sodium acetate buffer pH 4.0 (for 250 ml of buffer: 1,23 g/l sodium acetate trihydrate, 3,06 g/l acetic acid). The pH was adjusted to 4,8, if necessary, and 300 µl caprilic acid were added drop-wise. The solution was incubated for 30 minutes on a rotating wheel at room temperature, and centrifuged for 20 minutes at 5000 g, 10 °C. This step precipitates abundant serum proteins, while IgG molecules remain soluble. The top lipid layer and pellet were discarded, the supernatant carefully decanted (~11-12 ml) and 150-200 µl Tris 2M (pH 11) was added to neutralize the acidic pH. A half volume (5,5-6 ml) of saturate ammonium sulphate (pH 7,0-7,2) (132 g in 250 ml water) was added slowly to the supernatant, and the solution was incubated for 4 h in rotation at 4 °C. This pre-precipitation step removes large protein aggregates and any proteins that may precipitate with low concentrations of ammonium sulphate. After 30 minutes centrifugation at 3000 g at 4 °C, the supernatant was carefully transferred to a new tube, and enough ammonium sulphate (5,5-6 ml) was added slowly to obtain a 50% ammonium sulphate saturation. This step precipitates IgG molecules. The solution was incubated in rotation overnight at 4 °C, and centrifuged for 30 minutes, 3000 g, 4 °C. The pellet containing IgG was resuspended with 0,8-1 ml PBS and dialyzed versus three changes of PBS overnight. On average, the final antibody solution doubles its volume after dialysis. Dialysis tubing was prepared by boiling twice for 5 min in 5 mM EDTA, 200 mM sodium bicarbonate, and eventually rinsed in sterile water.

Peptide affinity purification of antibodies

An affinity column was prepared by covalent binding of the peptides used in the immunization protocol to the agarose matrix SulfoLink (#2040, Pierce). Crude sera were then passed through the columns, so that peptide-binding IgG bind to the matrix. After several washes, the IgG were eluted with 100 mM glycine pH 2,7. Next, a gel filtration step (G25, Hightrap, Acta, GE Healthcare) was performed to remove glycine from the eluates and exchange the buffer for PBS (NaCl 0,14 M, KCl 0,0027 M, KH₂PO₄ 0,001 M, Na₂HPO₄ 0,01 M) pH 7,4 containing NaN₃ 0,05% w/v and ProClin300 0,05% w/v. The crude sera, flow-through from the affinity column (unbound antibodies) and the purified antibodies were tested by ELISA (see Supplementary Materials).

Western blot

Proteins were separated by conventional SDS-PAGE or NuPAGE 4-12% BisTris gels (Invitrogen) run with MES buffer. After blotting, protein transfer was checked by staining of the membranes with Ponceau S. The PRAME sera were diluted 1:1000 in standard PBS/T with 5% skimmed milk, and incubated for 1 hour at room temperature. Western blots were visualized by ECL (GE Healthcare).

Cell culture, stable cell lines and growth assays

Stable cell lines were generated as described (Costessi et al, 2011). The pRetroSuper plasmid carries a Puromycin resistance gene, while LZRS carries zeocin resistance. For cell proliferation assays, between
300,000 and 900,000 A375 cells were seeded in 9 cm tissue culture dishes. Cells were passages every 2-3 days, when at least one plate reached ~70% confluence and cell densities measured (Beckman Coulter). Cells were detached with 1 ml 0.5% Trypsin-EDTA (GIBCO), incubated 5 min at 37°C and 3 ml medium were added (10% FCS, 1% PenStrep cocktail, DMEM 1640 with GlutaMAX-I, GIBCO). All plates were diluted with the same factor, so that the fastest growing cells would be confluent after 2-3 days. Either 1 µM aTRA or the same volume of ethanol 96% (vehicle) was added to the medium when cells were split. Stable cell lines carrying pRS vectors were kept on puromycin selection (2 µg puromycin/ml medium).

**Chromatin isolation CsCl\(_2\) gradient**

Chromatin was isolated from 150 million K562 and K562-TAG-PRAME cl.12 cells essentially as described (Costessi et al., 2011) with the following modifications: cells were crosslinked for 30 min in 40 ml volume and sonicated in 2 ml buffer D without SDS (1mM EDTA, 0.5 mM EGTA, 20 mM HEPES and protease inhibitors) for 15 min, because SDS is not compatible with caesium chloride gradient sedimentation. Next, 2 ml of chromatin were mixed with 1.25 g CsCl\(_2\) (final concentration 568 mg CsCl\(_2\)/ml), 70 µl sarcosyl 15% and 1-2 µl EtBr, and centrifuged at 260,000 g for 48 h or overweekend. Samples from different layers were collected and dialysed o/n against buffer D with 5% glycerol.
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REFERENCES


CHAPTER 6

DISCUSSION
In this thesis, we explored the molecular function(s) of the human oncoprotein PRAME using an innovative combination of biochemical protein-complex purification techniques and next-generation sequencing. We dissected the protein-protein interaction network of PRAME applying epitope-tagging and retroviral transduction in cell lines (chapter 2). Mass spectrometry analysis provided the first unbiased list of PRAME interactors, including all subunits of Cullin2-based ubiquitin ligases (chapter 2). These interactions were further characterised in chapter 3, where we identified PRAME as a BC-box protein and substrate receptor component of Cullin2-based ubiquitin ligases. This data associated for the first time a defined biochemical activity to the PRAME oncoprotein. Furthermore, we used specific antibodies and ChIP-sequencing techniques to generate the first genome-wide profiling of PRAME interactions with chromatin. We discovered that PRAME is specifically recruited to transcriptionally and epigenetically active promoter regions bound by NFY and nearby enhancers, implicating PRAME in the NFY signaling pathway. Optimisation of the protein affinity purification protocols and a complex walking approach allowed us to identify and characterise the human EKC/KEOPS complex and its interactions with Cul2-PRAME ligases (chapter 4). In chapter 5, we described the development and validation of PRAME antibodies that have been a crucial tool for our study, and we discussed in detail the strategy we undertook to generate the genome-wide ChIP-seq profiles presented in chapter 3. Finally, we reported the results of cell proliferation assays that do not support the previously reported function of PRAME as dominant repressor of retinoic acid signaling (chapter 5).

The results presented in this thesis are discussed in more detail in this chapter together with an outlook on the future perspectives of this field of research.

**Epitope-tagging technology and dissection of protein complexes**

During the past two decades, epitope tagging has become an indispensable technology in many fields of research, including biochemistry, cellular and molecular biology, and biotechnology. Affinity fusion tags have been used as research tools for detection, purification, and immobilisation of recombinant proteins expressed in bacteria, yeast, insect cells, plants, and mammalian cells (Nilsson *et al*, 1997). We applied this technology to purify
PRAME-containing protein complexes and identify its protein interactors by mass spectrometry analysis. In chapter 2, we described our approach based on the LZRS retroviral expression system and the StrepII-TEV-MYC-3xHA composite epitope tag that we used throughout our study. We setup a single-step immunoprecipitation protocol to purify PRAME-containing protein complexes using crosslinked MYC antibodies and peptide elution (chapters 2 and 3). The yield of the protein-complex purifications were further increased by the use of crosslinked HA antibodies and acidic glycine elution (chapter 4). Notably, this improvement of the protocol allowed us to significantly reduce the amount of cells and protein extracts needed for one purification experiment, and we could scale down from about 10 liters of cell culture (~10^10 cells) to about 3 liters (~3x10^9 cells) for each purification experiment.

Taken together, our mass spectrometry data combined with biochemical experiments characterized PRAME as a BC-box protein and component of Cullin2-based E3 ubiquitin ligases. The interactions between PRAME and the Cullin complex are mediated by conserved amino acid motifs (the so-called “BC-box” and “Cul2-box”) at the N-terminus of PRAME. In vitro experiments with purified recombinant proteins further established that a reconstituted Cullin2–PRAME complex is a catalytically active ubiquitin ligase. Moreover, we discovered that PRAME bridges Cul2 ligases to EKC via direct interactions between PRAME and the OSGEP and LAGE3 subunits, supporting a functional link between these complexes (chapter 4).

Cullin-based ubiquitin ligases

Cullin complexes (also referred to as cullin-RING ligases, CRL) are multi-subunit enzymes and constitute the largest known class of ubiquitin ligases (Petroski & Deshaies, 2005; Lipkowitz & Weissman, 2011; Okumura et al, 2012). Owing to the different cullin proteins expressed in human cells and the great diversity of their substrate-receptor subunits, hundreds of distinct cullin-based E3 ubiquitin ligases are thought to exist, which establishes these enzymes as key mediators of post-translational protein regulation (Petroski & Deshaies, 2005). CRLs have been implicated in a wide range of cellular processes, including cell cycle regulation, DNA replication, gene transcription, and development. Moreover, substrate receptor subunits of these ligases are targeted by viral and bacterial pathogens to subvert normal cellular processes. Although most target proteins that bind to CRL substrate receptors are ubiquitinated as a prelude to degradation, proteolysis is not the only...
outcome, and ubiquitination can also lead to post-translational regulation without alteration of protein levels (Petroski & Deshaies, 2005).

All CRLs share a similar modular architecture with a central cullin protein of elongated shape. The cullin N-terminus binds the substrate-selector module, while the C-terminus interacts with a RING box protein (Petroski & Deshaies, 2005). The N-terminal domain of Cullin2 interacts with the heterodimer ElonginB-ElonginC to bind a BC-box protein, which in turn recruits the ubiquitination substrate to the ligase (Okumura et al, 2012). The Von Hippel-Lindau tumour suppressor (VHL) is one of the most well-studied BC-box proteins and mutations in its coding sequence lead to renal carcinomas because of deregulation of the hypoxia-inducible factor-1α (HIF1α) transcription factor (Okumura et al, 2012). In standard normoxic conditions HIF1α is hydroxylated, which creates a binding site for VHL and subsequent ubiquitination by a Cul2-VHL ubiquitin ligase and degradation by the proteasome.

In hypoxic conditions, however, HIF1α is not hydroxylated and cannot be recognised by the Cul2-VHL ligase, resulting in increased protein levels and upregulation of genes mediating adaptation to low oxygen tension, including factors stimulating the formation of new blood vessels. Loss-of-function mutations of VHL result in constitutive higher levels of HIF1α and aberrant activation of its target genes, contributing to angiogenesis and tumour development (Okumura et al, 2012).

Identification of the substrate(s) of Cul2-PRAME ubiquitin ligases constitutes a crucial step in elucidating the detailed molecular mechanisms and pathways of this so-far elusive oncoprotein and should be the priority of follow-up studies. Our mass spectrometry data indicated that the EKC complex is the most abundant and stable interactor of Cul2-PRAME ligases in several cell lines and EKC subunits are therefore potential substrates of PRAME-mediated ubiquitination. As expected for BC-box proteins, PRAME was itself ubiquitinlated in our experiments, but we could not detect a PRAME-mediated ubiquitination of EKC components. However, we cannot exclude that our negative results were caused by technical limitations of the assays used. Therefore, we believe that more and probably different assays should be developed to assess whether EKC subunits can be substrates of PRAME-mediated ubiquitination.

Importantly, our present knowledge of CRLs indicates that interactions
between ligases and substrates are often dependent on post-translation modifications of the substrates like proline hydroxylation, phosphorylation, or glycosylation (Verma et al., 1997; Yoshida et al., 2002; Ivan et al., 2001). It is therefore likely that PRAME substrates are not (or at very low level) amongst the stable interactors identified in our mass spectrometry data, which were generated from cell lines cultured in standard conditions. Specific signaling events might be necessary to trigger the interaction between PRAME and its substrate(s). Moreover, it appears likely that substrates of PRAME are expressed in a tissue-specific manner, and this would potentially account for the reported tissue-specific effects of PRAME upregulation. The apparent discrepancies of PRAME being associated to good as well as to bad prognosis cancers might be explained by differential expression of PRAME targets in different tumour types.

Intriguingly, we found a complete conservation of the BC-box and Cul2-box at the N-terminus of all the annotated PRAME-like proteins. This suggests that all members of the large PRAME protein family are likely to function as substrate recognition subunits of cullin-based ubiquitin ligases, possibly displaying different degrees of specificity and/or redundancy in substrate selection. It is tempting to speculate that the reported roles of PRAME-like genes in early development and stemness might be mediated by functional interactions with CRLs and ubiquitination of proteins involved in these pathways.

**PRAME and the retinoic acid signaling**

A function of PRAME as dominant repressor of the retinoic acid signaling was reported by the Bernards lab (Epping et al., 2005). Using a genetic screen, the authors found that ectopic expression of the retinoic acid receptor alpha (RARA) or PRAME prevented the growth arrest and apoptosis induced by histone deacetylase inhibitors (HDACIs) (Epping et al., 2007). They provided evidence supporting a model in which HDACI-induced growth arrest requires an active retinoic acid signaling pathway, which PRAME would repress via direct protein-protein interactions with the retinoic acid receptor alpha and the EZH2 polycomb protein, resulting in transcriptional repression of retinoic acid target genes like RARB and CDKN1A (also known as p21) (Epping et al., 2005; 2007). However, the suggested mechanisms do not seem to play a general role, since higher PRAME levels were found to correlate with
lower RARA levels in breast cancer patients (Epping et al, 2008). Moreover, the retinoic acid target genes RARB and CDKN1A were not repressed in acute myeloid leukemia patients with high PRAME levels (Steinbach et al, 2007).

Epping et al. reported that the C-terminus of PRAME (aa. 416-509) interacts directly with the ligand-binding domain (LBD) of RARalpha in GST pull downs and a mammalian two-hybrid system (Epping et al, 2005). The authors further showed that the C-terminal amino acid motif LRELL in PRAME was required for the interaction with the LBD of RARa; reciprocal mutations of the LBD of RARa were not included in the study. Using the full-length PRAME protein, however, Wadelin et al could not detect any interaction with the LBD of RARa using similar GST pull down or yeast two-hybrid assays (Wadelin et al, 2010). In agreement with the findings of Wadelin et al, we did not succeed in reproducing the interaction between RARa and PRAME in coimmunoprecipitations assays upon transient transfections in 293T cells, or upon immunoprecipitation of TAG-PRAME from stable cell lines in the presence or absence of ligand.

Epping et al further described that knockdown of PRAME in A375 human melanoma cells resulted in a higher sensitivity to the administration of retinoic acid (Epping et al, 2005) and HDACI (Epping et al, 2007). Notably, knockdown of PRAME in these cells was shown to cause downregulation of the retinoic acid receptor alpha (RARA) at the protein level (Epping et al, 2005, Figure 6F) and this was interpreted as a derepression of the retinoic acid signaling pathway. However, the same experiment presented in a follow-up publication showed no difference in RARA levels upon PRAME knockdown (Epping et al, 2007, Figure 4E), casting doubts about the reported effects. As described in chapter 5, we successfully reproduced the retinoic acid phenotype upon PRAME knockdown in A375 cells. However, this effect was limited to one of the two knockdown constructs used, despite a similar down-regulation of PRAME expression, and we could not rescue the phenotype by ectopic expression of knockdown-resistant version of PRAME. Our data are consistent with an “off-target” effect of the pRS-P1 knockdown construct. In other words, the phenotype observed is likely caused by down-regulation of a transcript unrelated to PRAME, which is specifically targeted by the published construct pRS-P1, but not by pRS-P2. A genome-wide mRNA-seq profiling experiment on these cell lines could potentially identify this gene and shed light on the cause of the growth phenotype observed.
Our protein-complex purifications clearly established that Cullin2 ligases and EKC subunits are the most abundant interactors of PRAME in a number of human cell lines kept in standard culture conditions, while we did not detect RARa or EZH2. However, we cannot exclude that PRAME might establish interactions with RARa and EZH2 upon activation of particular signaling pathways. Notably, Epping et al performed their genetic screen in p53-deficient mouse embryonic fibroblasts stably expressing an oncogenic RasV12 gene (RasV12 MEFs) (Epping et al, 2007). It is therefore possible that the activation of Ras pathways plays an unforeseen role in PRAME pathways.

In order to disentangle the conflicting result, purifications of PRAME-containing protein complexes from RasV12 MEFs should be performed to identify the most abundant PRAME interactors in this cell system in the presence and absence of retinoic acid. Moreover, the effect of PRAME BC-box mutations on the sensitivity to HDACI should also be tested in the same cells.

The enigmatic EKC/KEOPS complex

Our biochemical experiments demonstrate that the EKC complex is conserved and expressed also in human cells. Several crystal structures of yeast and archaeal EKC subunits were reported and supported a model in which the complex dimerises around the Pcc1/LAGE3 subunit assuming a V shape. Our experimental data together with preliminary structural modelling of the human subunits are consistent with a similar dimeric model for the human EKC. Furthermore, we detected specific interactions between the conserved EKC orthologues and an uncharacterised protein of small size, C14orf142, suggesting that this might be a novel EKC subunit or interactor.

The EKC or KEOPS complex has been first identified in yeast by two independent laboratories that described a role for the complex in either transcription regulation (Kisseleva-Romanova et al, 2006) or telomere maintenance (Downey et al, 2006). More recently, the complex has been involved in the biosynthesis of a chemical modification of tRNAs that is critical for efficient protein synthesis (Daugeron et al, 2011; Yacoubi et al, 2011; Srinivasan et al, 2011).

Human EKC comprises four evolutionarily conserved subunits (yeast and human names are given): Pcc1p/LAGE3, Kae1p/OSGEP, Bud32p/TP53RK, Cgi121p/TPRKB. Additionally, the yeast complex comprises a fifth subunit called Gon7p or Pcc2p, which seems to be fungi-specific. While PCC1/LAGE3, BUD32/TP53RK and CGI121/TPRKB are present in archaea and eukarya, but
not in bacteria, orthologues of KAE1/OSGEP are present in all three domains of life (although often annotated with different names). In fact, a study from 2004 found that KAE1/OSGEP was one of the about 60 genes present in all genomes sequenced until then, with the only exceptions of the reduced genomes of the endosymbiotic bacteria Carsonella ruddii and Sulcia muelleri (Galperin & Koonin, 2004; Galperin, 2008). Moreover, KAE1/OSGEP is represented in bacteria by two different homologues that were shown to establish protein-protein interactions: YeaZ (specific to bacteria) and YgjD (orthologue of Kae1/OSGEP) (Handford et al, 2009).

Phylogenetic analysis of 767 Kae1 homologues grouped these proteins in four distinct clusters (Hecker et al, 2007):

1. archaeal Kae1;
2. eukaryotic orthologues of yeast Kae1p and human OSGEP;
3. bacterial orthologues of E. coli YgjD and eukaryotic orthologues of Qri7p/OSGEPL1;
4. bacterial orthologues of E. coli YeaZ.

Interestingly, the eukaryotic orthologues of Kae1p/OSGEP and the archaeal Kae1 proteins appeared significantly more similar to one another than to the two bacteria-specific subfamilies (Hecker et al, 2007). Yeast Qri7p and human OSGEPL1 are nuclear-encoded paralogues that localize to mitochondria, itself a bacterial symbiont (Oberto et al, 2009). The clustering of Qri7p/OSGEPL1 within the bacterial orthologues of YgjD supports the bacterial origin of these genes. Unlike the YgjD/KAE1/OSGEP gene family, PRAME genes seem to have appeared significantly later in evolution and to have undergone several rounds of gene duplication events in the rodent and hominin lineages (Birtle et al, 2005; Rhesus Macaque Genome Sequencing and Analysis Consortium et al, 2007). The peculiar evolutionary history of these gene families suggests that PRAME might function as a modulator of EKC.

The function of YgjD/KAE1/OSGEP proteins has been uncertain for many years and is still unclear, and functions have been ascribed to these proteins that later appeared to be incorrect. First, the Gcp protein was cloned from Pasteurella haemolytica and suggested to be an O-sialoglycoprotein endopeptidase, hence the names OSGEP and KAE1 (kinase-associated endopeptidase) (Abdullah et al, 1991; Abdullah et al, 1992). However, a peptidase activity was never confirmed by independent studies (Hecker et al, 2007). Second, a study of the archaeal orthologue of Kae1 from Pyrococcus abyssi detected apurinic-endonuclease activity in vitro (Hecker et al, 2007).
Although Srinivasan et al detected this enzymatic activity for yeast Kae1p expressed and purified from E. coli, they reported that the endonuclease activity was caused by a low level contamination from the highly active E. coli enzyme Fpg that copurified with Kae1p (Srinivasan et al, 2011).

Two independent laboratories identified the yeast EKC/KEOPS complex starting from genetic clues. The laboratory of Domenico Libri isolated Pcc1, a protein of small size with no predicted functional domains in a search for suppressors of a splicing defect and demonstrated that its function is not related to splicing (Kisseleva-Romanova et al, 2006). Their genetic and biochemical experiments led to the description of the five-subunit EKC complex, the integrity of which was required for normal induction of pheromone- and galactose-responsive genes.

A parallel genome-wide screen in yeast identified the cgi121 null mutant as a suppressor of the telomere-capping defect of cdc13-1 cells (Downey et al, 2006). Deletion of CGI121 or other EKC/KEOPS subunits resulted in short telomeres in an otherwise wild-type context.

Since kae1, pcc1, and pcc2/gon7 temperature-sensitive strains stop dividing at the non-permissive temperature already after a few divisions, Kisseleva et al. suggested that telomere maintenance is unlikely to be the primary function of the complex. The authors did not exclude that the telomere phenotype might result from an altered transcriptional regulation, or that the complex might be involved in some general chromatin remodelling that would eventually affect both gene loci and their transcription as well as the telomeres (Kisseleva-Romanova et al, 2006).

Most recently, the group of Crécy-Lagarde used in silico comparative genomics approaches to identify a critical function for Kae1 in the universal t^6A biosynthesis pathway (Yacoubi et al, 2011). This pathway mediates the chemical modification of position 37 of all tRNAs that pair with ANN codons (thus, including the first ATG of protein-coding genes), with the only exception of prokaryotic fMet tRNA. The modification consists of a carbonyl group and a threonine attached to the amino group of adenine at nucleotide 37 immediately 3’ of the anticodon, and is necessary to prevent mispairing between the first base of the codon and the third base of the anticodon (Yacoubi et al, 2011).

Work from three independent laboratories established that the t^6A modification activity was dependent on Pcc1p, the ATPase activity of Kae1p,
and the kinase activity of Bud32p, but not on Cgi121p (Srinivasan et al., 2011; Yacoubi et al., 2011; Daugeron et al., 2011). Moreover, Qri7p was found to mediate modification of mitochondrial tRNAs (Yacoubi et al., 2011).

Although the most recent data show that the EKC/KEOPS complex is required for t^6A synthesis, this might not be its only role and its function in telomere maintenance might be unrelated to t^6A modification of tRNAs. More specifically, it was hypothesised that the predicted chaperone activity of this complex might contribute to several processes, t^6A formation being just one of them (Srinivasan et al., 2011).

**Outlook on the interplay between PRAME, EKC and NFY**

Studying the relationships between PRAME, Cul2 ligases and EKC, we discovered that PRAME recruited Cullin2-based ligases to EKC via direct interactions with OSGEP and LAGE3. In fact, we consistently found that PRAME established stable interactions with OSGEP and LAGE3, but not with TP53RK or TPRKB. These results might represent a PRAME-mediated disruption of an otherwise complete EKC complex, or association of PRAME to an already existing EKC subassembly. Characterisation of the amino acid residues involved in these interactions should be prioritised in follow-up studies in order to shed light on the precise stoichiometry of the complexes. In particular, it is not known whether PRAME influences the conformation of OSGEP and LAGE3 and whether PRAME might cause steric hindrance and thus inhibit the formation of a complete dimeric EKC. Ideally, resolution of crystal structures containing PRAME, OSGEP and LAGE3 could provide ultimate clues on the three-dimensional relationships of these proteins.

No data is available yet on EKC function(s) in human cells and the absence of efficient knockdown or knock-out models have severely hindered functional analyses of EKC in higher eukaryotes so far. Considering the extreme conservation of EKC subunits, however, it is reasonable to expect that at least some of the functions and pathways identified for this complex in yeast could be conserved in human cells. However, the complex seems to have acquired also additional functions during evolution, like phosphorylation of the proto-oncogene p53 by TP53RK. Future studies will have to address the connections between apparently different functions ascribed to this complex, like phosphorylation of the p53 oncogene, chemical modification of tRNAs, roles in telomere maintenance and physical association with promoter regions.
Kae1p and Pcc1p were found to associate with active chromatin and to regulate expression of some of the yeast promoters to which they are recruited (Kisseleva-Romanova et al., 2006). In a yeast strain deficient for Pcc1p the activator Gal4p was still found on chromatin, but recruitment of Gal1p, the SAGA histone acetyltransferase and Mediator were inhibited, suggesting a role for EKC in gene transcription. We found that epitope-tagged EKC subunits in human cells can specifically associate with chromatin at active promoter regions bound by PRAME and NFY. Unfortunately, stable knockdown strategies did not detect a clear function of EKC or PRAME on transcriptional regulation of those loci. As we elaborated in chapter 4, likely different cell systems and assays are needed to investigate the functional role(s) of PRAME and EKC on chromatin. As soon as an assay will become available to evidence EKC functions in human cells, the role of PRAME on such functions should be tested. For example, EKC function could be measured in a PRAME-negative cell line before and after ectopic expression of PRAME.

Our genome-wide ChIP-seq analyses revealed that PRAME is strongly enriched at promoters bound by the transcription factor NFY, and binding of PRAME predicts a permissive epigenetic landscape and transcriptional activity at these regions. Moreover, we found that PRAME is also significantly enriched at adjacent enhancer regions, suggesting that it could participate in promoter-enhancer interactions. Although our genome-wide ChIP-seq profiles are consistent with a role for PRAME in active transcription, more efforts are required to provide a definitive answer to the burning question if PRAME can function as a transcriptional modulator.

Since neither PRAME nor EKC proteins possess known DNA-binding domains, it is not clear whether they can bind DNA directly and if the transcription factor NFY might play a direct or indirect role in the targeting of a Cul2-PRAME-EKC complex to chromatin. However, we did not find NFY subunits in our mass spectrometry interaction data of PRAME or EKC. Future studies should investigate the molecular determinants of the association of PRAME and EKC to chromatin and the role of NFY, and whether there is a hierarchy in their binding.

Analyses of gene expression levels in human samples indicate that most normal adult tissues express NFY, EKC and Cul2, but not PRAME; therefore, it is expected that all (or a vast majority) of PRAME-expressing cells also express NFY, EKC and Cul2. Taking into consideration these expression patterns and the ancient nature of EKC as compared to the later appearance of
PRAME, we speculate that EKC could fulfill housekeeping (PRAME-independent) functions also in human cells and that it could interact with chromatin in a PRAME-independent manner. PRAME expression would then lead to the recruitment of a Cul2-based ubiquitin ligase to EKC genomic loci by virtue of direct protein-protein interactions, resulting in ubiquitination of proteins present at these sites, potentially affecting EKC pathways and/or NFY signaling. This hypothesis could be tested by comparing the chromatin-binding profiles of EKC subunits in PRAME-negative and PRAME-expressing cells, and development of ChIP-grade antibodies against EKC proteins will be critical tools for such studies.

Intriguingly, several studies suggested important roles for NFY in stem cell biology (Grskovic et al., 2007) and mouse PRAME-like genes were found to be involved in embryonic development and embryonic stem cells (Bortvin et al., 2003). It should therefore be studied if PRAME-like proteins can also play roles in EKC and NFY pathways in a redundant manner with PRAME, or if they rather display specificities of their own. We foresee that the dissection of the protein-protein interaction networks of PRAME-like proteins will definitely provide stimulating and exciting new leads for the field.

Overall, the experimental work outlined in this thesis establishes a biochemical and molecular function for the PRAME gene family and provides a new twist to both PRAME and EKC biology. Our results provide an important contribution to understanding the molecular mechanisms in which these proteins are active, both in healthy and tumour cells.
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Scientific research projects usually span several years during complementary research lines and technical approaches are combined in order to dissect and understand the molecular mechanisms, proteins or genes of interest. A typical scientific project can be described as a collection of experimental tasks performed at different points in time and often in parallel, whose results are eventually bundled together in a scientific report (i.e. an article in scientific journals, a presentation at a conference, a PhD thesis). Scientific reports are generally structured in a logical and/or chronological order: an initial finding is extended with follow-up experiments, and then a final conclusion or model is presented. Notably, articles in peer-reviewed journals focus on the results obtained, while the time and resources required to obtain them are traditionally not discussed. The efforts required to raise an antibody, to setup a novel technique or negative results are rather discussed at meetings and conferences, or at informal gatherings among researchers. On the other hand, the timeline and the financial and human resources needed for a research project have become standard requirements of increasing importance when submitting project proposals to agencies responsible for research funds.

The aim of this appendix is to provide an overview of our PRAME research project in a chronological context and describe how the different lines of research developed over time, and often in parallel. The main activities and milestones are presented in schemes called Gantt chart, a project management tool that represents the start, duration and end of each activity.

Overall, the research presented in this thesis can be divided in four main lines or tasks (Figure 1):

i. the development of PRAME antibodies,
ii. the identification and characterisation of the protein interactors of PRAME,
iii. the identification and characterisation of PRAME interactions with the EKC complex,
iv. the investigation of PRAME association to chromatin.

As depicted in Figure 2, the project started in the fall of 2005 with a focus on the first two tasks: PRAME antibody development and protein complex purification. The peptides needed to raise the antibodies were designed and synthesised, the immunisation scheme started in March 2006 and lasted about eight months until the last serum was harvested. The antibody charac-
Ab development

Peptide design and immunisation

Test Ab

PRAME protein complex

DNA vectors + cell lines

cations & MS

ChIP

CsCl exp, stable cell lines

ChIP-seq exps. and data analysis

EKC protein complex

DNA vectors + cell lines

cations and MS

Publications

CSHL transcription meeting

EMBL transcription meeting

Paper#1 - NCB

Paper#1 - NSMB

Paper#1 - NSMB #2

Paper#1 @ EMBOJ

Paper#2 - PloSOne

Appendix Figure 1

Figure 1
terisation and testing started as soon as the first sera became available (May 2006) and continued through the second half of 2006, when conditions were setup for use of sera and affinity-purified antibodies in western blot and immunoprecipitation assays (chapter 5).

In parallel, we attempted PRAME protein-complex purifications using FM6 stable cell lines that were already available in the lab. As described in chapter 2, this strategy was not successful because of a too low efficiency and we therefore moved to a different epitope tag (StrepII-Myc-3xHA).
combined with the use of suspension cell lines. In the first half of 2006 we generated the retroviral expression plasmids required and we set up the stable transfection protocol. In the second half of 2006 we generated single-cell clones stably expressing epitope-tagged PRAME and established that PRAME is part of high molecular weight complexes by size-exclusion chromatography (K562 Sup6 in Fig. 2). In parallel, we set up the nuclear/cytoplasmic protein extraction protocol and the immunoprecipitation strategy for protein complex purifications (TAP-TAG). These resulted in the first mass spectrometry data of PRAME interactors in July 2007 using a single MYC IP strategy (chapter 3). TAG-PRAME purification experiments were repeated several times later in the project, also in combination with optimisation of single-step HA IP strategy (chapter 4). In total, almost 2 years of experimental work were required to identify the first protein-protein interactors of PRAME. Based on these results, we started in September 2007 a close collaboration with the Conaway lab (Stowers Institute, USA).

Early in 2007 we started to study the association of PRAME to chromatin (Figure 3). As reported in chapter 5, we performed first caesium chloride gradient experiments and then set up the TTE epitope tag approach for ChIP assays. During the course of 2007, we developed the retroviral pTTE vector, cell lines stably expressing TTE-PRAME and towards the end of the year we performed the first genome-wide ChIP-sequencing experiments for TTE-PRAME (ChIP-seq #1). Analysis of the data identified the first putative binding sites which have been instrumental to setup and optimise ChIP assays for the endogenous PRAME protein in wild type K562 cells (ChIP-seq #2). Bioinformatical analysis revealed a link to the NFY transcription factor (chapter 3), whose binding sites were mapped in ChIP-seq experiments in early 2010 (ChIP-seq #3).

In the second half of the project we also characterised the discovered interactions of PRAME with Cullin2-based E3 ubiquitin ligases and with the human EKC complex (Figure 3). These experiments started with the generation of expression plasmids for all four human EKC subunits (Q3 2007), establishment of stable cell lines, ChIP assays and protein complex purifications and mass spectrometry (Figure 3). Between the end of 2008 and the beginning of 2010, mass spectrometry data for all four subunits were collected and analyzed, while biochemical experiments were performed in parallel.
Our results were first presented at the CSHL Eukaryotic transcription meeting in August 2009 and at the EMBL transcription meeting in August 2010 (Figure 1). After submissions of our work to Nature Cell Biology (NSB) and Nature Structural and Molecular Biology (NSMB), we published chapter 3 of this thesis in EMBO Journal (Q2 of 2010), and chapter 4 in PlosOne (Q3 of 2012).
SUMMARY

The genome present in the nucleus of each human somatic cell is made of about 2 meters of DNA organized in 23 chromosome pairs with a total of 6 billion basepairs. About 5% of the human genome contains protein-coding genes that can be activated to produce proteins under strict regulation. The proteins present in a cell (also called the “proteome”) are not lonely islands, but they rather interact with each other in dynamic networks and multiprotein complexes. Importantly, deregulation of gene expression programs and protein interaction networks can lead to disease and cancer.

The focus of this PhD thesis is to unravel the molecular function of the human oncoprotein PRAME (preferentially expressed antigen of melanoma) using cell line model systems and a powerful combination of proteomic and genomic techniques. At the beginning of this study, PRAME was known to be frequently overexpressed in tumours and high expression levels were found to correlate with poor clinical outcome in melanoma, neuroblastoma, and breast cancer patients. Moreover, evidence that expression of PRAME is restricted mainly to tumour cells has designated it as a promising therapeutic target. However, efforts to develop strategies to interfere with the oncogenic activity of PRAME are hindered by a lack of information about its molecular function(s). Notably, analysis of the human genome revealed that a large family of more than twenty PRAME-like genes exists, whose functions are also unclear.

In order to identify the molecular functions of PRAME, we developed innovative high-throughput biochemical approaches to dissect the protein-protein interaction network of PRAME (chapter 2). By applying epitope-tagging in human cell lines and mass spectrometry analysis of purified protein complexes, we discovered that PRAME is a stable component of Cullin2-based E3 ubiquitin ligases (chapter 3). Ubiquitin ligases mediate the addition of the small protein ubiquitin to target proteins, altering their function and often leading to their degradation. Our results established a first strong functional link between PRAME and a defined biochemical pathway. Using the same epitope-tagging strategy we discovered interactions between Cullin2-PRAME ligases and subunits of the conserved EKC/KEOPS complex that we characterised in chapter 4. In particular, knock-down experiments revealed that PRAME functions as a bridge between Cullin2 ligases and the EKC complex.
In parallel to the proteomic techniques, we set up chromatin immuno-precipitation assays coupled to next-generation sequencing technologies (ChIP-seq). For this purpose, we raised specific PRAME antibodies in rabbits to investigate the binding profile of PRAME to the human genome (chapter 3 and 5). The data analysis showed that PRAME (chapter 2) as well as EKC sub-units (chapter 4) are specifically enriched at transcriptionally and epigenetically active promoter regions that are also bound by the transcription factor NF-Y, and at enhancer elements. These results suggest that PRAME might play a role in the NF-Y signaling pathway.

Overall, the experimental work outlined in this thesis establishes a biochemical and molecular function for the PRAME gene family and provides a new twist to both PRAME and EKC biology. Our results provide an important contribution to understand the molecular mechanisms in which these proteins are active, both in healthy and tumour cells, and thus lay a foundation for novel therapeutic strategies against cancer.
SAMENVATTING

Het genoom in de kern van elke menselijke somatische cel bestaat uit ongeveer 2 meters DNA en is ingedeeld in 23 chromosoom paren die in totaal 6 miljard basenparen bevatten. Ongeveer 5% van het menselijke genoom bevat coderende genen die eiwitten kunnen produceren, wat gebeurt onder strikte regulatie. De eiwitten die aanwezig zijn in een cel (ook wel “proteoom” genoemd) zijn niet geïsoleerde eilanden, maar zijn betrokken bij dynamische netwerken en maken deel uit van multi-eiwit complexen. Ontregeling van genexpressie en eiwit interacties kan leiden tot verschillende ziektes, waaronder kanker.

Het onderwerp van dit proefschrift betreft een uitgebreide studie naar de moleculaire functie van het kanker-gerelateerde eiwit PRAME (preferentially expressed antigen of melanoma) met behulp van cellijn modelsystemen en een krachtige combinatie van proteomische en genomische technieken. Aan het begin van deze studie was bekend dat PRAME vaak actief aanwezig is in kankercellen en dat een hoge expressie van PRAME geassocieerd is met een slechte klinische prognose voor melanoom-, neuroblastoom- en borstkankerpatiënten.

PRAME vervult de rol van een veelbelovend therapeutisch doelwit aangezien expressie van dit eiwit voornamelijk beperkt is tot kankercellen. Echter, het ontwikkelen van strategieën om de activiteit van PRAME te beïnvloeden wordt gehinderd door een gebrek aan informatie over de moleculaire functie(s) van PRAME. Verder heeft de analyse van het menselijk genoom een grote familie van meer dan twintig PRAME-achtige genen aan het licht gebracht, waarvan de functies nog onduidelijk zijn.

Om de moleculaire functies van PRAME te onderzoeken, hebben wij innovatieve high-throughput biochemische technieken ontwikkeld om het eiwit-eiwit interactienetwerk van PRAME te ontrafelen (hoofdstuk 2). Door toepassing van epitop-tagging in menselijke celllijnen en massaspectrometrie van gezuiverde eiwitcomplexen, hebben wij ontdekt dat PRAME een stabiele component is van Cullin2 E3 ubiquitine ligases (hoofdstuk 3). Ubiquitine ligases koppelen het kleine eiwit ubiquitine aan doeleiwitten en veranderen
zodoende hun functie of markeren ze om afgebroken te worden. Onze resultaten hebben voor het eerst een sterke relatie gelegd tussen PRAME en een specifieke biochemische activiteit. Met dezelfde epitope-liging strategie hebben wij laten zien dat Cullin2-PRAME ligases een interactie aangaan met het EKC/KEOPS complex (hoofdstuk 4). In het bijzonder hebben knock-down experimenten aangetoond dat PRAME fungeert als een brug tussen Cullin2 ligases en het EKC complex.

Naast de proteomische technieken hebben wij een techniek ontwikkeld waarbij chromatin-immunoprecipitaties gevolgd worden door next-generation sequencing (ChIP-seq). Wij hebben specifieke PRAME antilichamen in konijnen opgewekt om het bindingsprofiel van PRAME op het menselijke genoom te onderzoeken (hoofdstuk 3 en 5). De data analyse liet zien dat PRAME (hoofdstuk 2) en het EKC complex (hoofdstuk 4) specifiek verrijkt zijn op transcriptionele en epigenetisch actieve promotors van genen die ook gebonden zijn door de transcriptiefactor NF-Y, als ook op enhancer elementen. Deze resultaten suggereren dat PRAME een rol zou kunnen spelen bij de NF-Y signalering-route.

Het experimentele werk beschreven in dit proefschrift heeft een nieuwe biochemische en moleculaire functie voor de PRAME gen-familie vastgesteld en geeft een vernieuwd inzicht aan zowel PRAME en EKC biologie. Onze resultaten leveren een belangrijke bijdrage aan het begrijpen van de moleculaire mechanismen waarin deze eiwitten actief zijn, zowel in gezonde als in kancercellen. Hiermee wordt een basis gelegd voor het ontwikkelen van nieuwe therapeutische strategieën tegen kanker.
RIASSUNTO

Il genoma presente nel nucleo di ogni cellula somatica umana è costituito da circa 2 metri di DNA organizzato in 23 coppie di cromosomi che contengono un totale di 6 miliardi di paia di basi. Circa il 5% del genoma umano costituisce geni codificanti proteine che possono essere attivati sotto stretta regolazione. Le proteine presenti in una cellula (nella loro insieme chiamate anche “proteoma”) non sono isole separate ed indipendenti, ma piuttosto interagiscono tra di loro in reti dinamiche e fanno parte di complessi multiproteici. In particolare, la disregolazione dell’espressione genica e delle interazione tra proteine può portare a malattie e cancro.

L’obiettivo di questa tesi di dottorato è di rivelare la funzione molecolare della oncoproteina umana PRAME (preferentially expressed antigen melanoma, antigene preferenzialmente espresso in melanoma) utilizzando linee cellulari come sistema modello ed una potente combinazione di tecniche di proteomica e genomica. All’inizio di questo studio, PRAME era noto per essere frequentemente attivo in tumori e livelli elevati di espressione erano risultati correlati con prognosi negative in pazienti di melanoma, neuroblastoma e cancro al seno. Inoltre, il fatto che PRAME è espresso principalmente in cellule tumorali e non in cellule sane ha indicato PRAME come un obiettivo terapeutico promettente. Tuttavia, gli sforzi per sviluppare strategie terapeutiche per interferire con l’attività oncogenica di PRAME sono ostacolati dalla mancanza di informazione sulle sue funzioni molecolari. Inoltre, analisi del genoma umano ha rivelato l’esistenza di una grande famiglia di più di venti geni simili a PRAME (PRAME-like genes), le cui funzioni non sono ancora chiare.

Al fine di identificare le funzioni di PRAME a livello molecolare, abbiamo sviluppato approcci biochimici innovativi e high-throughput per identificare la rete di interazioni proteina-proteina di PRAME (capitolo 2). Applicando un approccio di “epitope-tagging” in linee cellulari umane e analisi di spettrometria di massa dei complessi proteici purificati, abbiamo scoperto che PRAME è un componente stabile di E3 ubiquitin ligasi basati sulla proteina Cullin2 (capitolo 3). Le ubiquitin ligasi mediano l’aggiunta della piccola proteina ubiquitina a proteine bersaglio, alterando la loro funzione o spesso causandone la degradazione. I nostri risultati hanno quindi stabilito per la prima volta un
forte relazione funzionale tra PRAME ed un’attività biochimica ben definita. Utilizzando le stesse tecniche abbiamo scoperto interazioni tra Cullin2-PRAME ligasi e subunità del complesso EKC/KEOPS, che abbiamo caratterizzato nel capitolo 4. In particolare, esperimenti di knock-down hanno rivelato che PRAME funziona come un ponte tra le Cullin2 ubiquitin ligasi ed il complesso EKC.

In parallelo alle tecniche di proteomica, abbiamo sviluppato saggi di immunoprecipitazione della cromatina seguiti da sequenziamento high-throughput di nuova generazione (ChIP-seq). A questo scopo, abbiamo generato anticorpi specifici contro PRAME in conigli allo scopo di indagare il profilo di legame di PRAME al genoma umano (capitolo 3 e 5). L’analisi dei dati ha mostrato chiaramente che PRAME (capitolo 2) e subunità di EKC (capitolo 4) legano specificamente elementi enhancer e promotori di geni attivi che sono anche legati dal fattore di trascrizione NF-Y. Questi risultati suggeriscono che PRAME potrebbe giocare un ruolo nella via di segnale di NF-Y.

Nel complesso, il lavoro sperimentale descritto in questa tesi stabilisce una funzione biochimica e molecolare per PRAME e la vasta famiglia di geni PRAME-simili e fornisce una nuova svolta nella biologia sia di PRAME che del complesso EKC. I nostri risultati forniscono un importante contributo alla comprensione dei meccanismi molecolari in cui queste proteine sono attive, sia in cellule sane che tumorali, e offrono così possibilità di sviluppare nuove strategie terapeutiche.
CURRICULUM VITAE

Adalberto Costessi was born on April 23rd in Trieste, Italy. He developed a keen interest for scientific subjects and philosophy at the secondary school Liceo Scientifico Galileo Galilei in Trieste, where he graduated cum laude.

In 1998 he enrolled at the Medical Biotechnology curriculum at the University of Trieste in order to study the molecular mechanisms of biological life and disease. Besides the regular studies, he fulfilled the role of vice-president (2002) and president (2003) of the CNSB, the Italian association of biotechnology students. In these roles, Adalberto promoted a national project to evaluate the quality of the university education in biotechnology, coordinated several projects of public communication of biotechnology, and actively participated in the discussions about GMO and patents regulations. In those years, a peculiar interest for space-related research and weightlessness brought Adalberto in close contact with the European Space Agency (ESA). In 2000 his project proposal on the study of fruit flies in weightlessness was selected for participation to the 3rd Student Parabolic Flight Campaign, awarding him the opportunity to perform the experiment during the weightlessness conditions of a parabolic flight.

In 2002 Adalberto joined the Molecular Biology lab of Gianluca Tell at the University of Trieste where he conducted experimental research on the molecular responses of human osteoblasts to extracellular nucleotides and mechanical stress, leading to two peer-reviewed publications and a graduation cum laude in 2003.

In 2002 Adalberto’s research proposal “BOP - Bone proteomics” to study the molecular responses of human osteoblasts in microgravity was awarded the first prize at the International “Success 2002 Student contest” by the European Space Agency. This brought him to The Netherlands in 2004, at the ESA research centre in Noordwijk. There he contributed to the development of an innovative cell-culture system and trained the astronauts that carried out the BOP experiment during a space mission to the International Space Station in April 2005.

In October of the same year, Adalberto decided to start a PhD research project with the feet on the ground under the supervision of prof. Henk Stunnenberg at the Molecular Biology Department of the Radboud University Nijmegen, The Netherlands. During five years of experimental research on the human oncoprotein PRAME he applied and developed biochemical
techniques to unravel the protein-protein interaction network of PRAME as well as high-throughput sequencing approaches to identify the binding sites of PRAME on the human genome.

Since January 2011, Adalberto works as Product Specialist Genome Analysis at BaseClear (Leiden, The Netherlands), an independent accredited laboratory specialized in DNA analysis services. He leads the Next-Generation Sequencing laboratory and is responsible for the application of the newest related technologies.
Embedding siRNA sequences targeting Apolipoprotein B100 in shRNA and miRNA scaffolds results in differential processing and in vivo efficacy.

The human EKC/KEOPS complex is recruited to Cullin2 ubiquitin ligases by the human tumour antigen PRAME.

The tumour antigen PRAME is a component of Cullin2 ubiquitin ligases and is recruited to active NFY promoters.

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ACKNOWLEDGEMENTS

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And then a big hug to Alex and Michela, of course! The friendship with Alex started in Gianluca’s lab, when you were a PhD student and my direct supervisor (the good old times!) And a couple of years later you and Michela decided to find a job in The Netherlands and follow me to the north. Are you guys ever gonna stop blaming me for the Dutch weather? Ok, ok, one more grappa and we all forget the rain...

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lab, he could get that one, and so I eventually started working on PRAME.
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Adalberto
The human protein PRAME (preferentially expressed antigen of melanoma) is frequently overexpressed in tumours. Importantly, high expression levels are associated to poor clinical outcome in melanoma, neuroblastoma, and breast cancer. However, its function and molecular mechanisms in normal and cancer cells are not yet well understood and the cellular pathways affected by PRAME are still elusive. Although PRAME has been indicated as a promising potential target for cancer therapy and anti-tumour immunization protocols, efforts to develop strategies to interfere with the oncogenic activity of PRAME are hindered by a lack of information about its function.

The aim of this thesis was to investigate the molecular function of human PRAME using cell line model systems and a powerful combination of proteomic and genomic techniques. We developed innovative biochemical approaches to dissect the protein-protein interaction network of PRAME, and we setup chromatin immunoprecipitation assays coupled to next-generation sequencing technologies to reveal binding profile of PRAME to the human genome in cancer cells.