Production of native creatine kinase B in insect cells using a baculovirus expression vector

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Abstract

A full-length human creatine kinase B (B-CK) cDNA was used to produce a recombinant baculovirus (AcDZ1-BCK). Sf9 cells infected with this recombinant expressed a homodimeric protein composed of 43 kDa subunits which, under optimal conditions, formed up to 30% of the total soluble cellular protein. Upon analysis by PAGE, zymogram assay and gel filtration chromatography the recombinant protein behaved like authentic dimeric human BB-CK protein. Studies with a newly produced monoclonal antibody (CK-BYK/21E10) directed against an epitope in the N-terminus of the protein confirmed the identity of the product. The recombinant BB-CK protein was purified to over 99% homogeneity from the total protein extract of AcDZ1-CKB infected cells in one single step involving anion exchange column chromatography on MonoQ in FPLC. Dialysed protein had a specific activity of 239 U/mg protein. (Mol Cell Biochem 143: 59-65, 1995)

Key words: creatine kinase B, baculovirus, SF9 cells, monoclonal antibody

Introduction

Different isoforms of creatine kinase (CKs; EC 2.7.3.2) play an important role in the energy metabolism of vertebrate cells by catalyzing the reversible exchange of high-energy phosphate between ATP and creatine phosphate (CrP) [1]. CKs, the homo- or heterodimeric cytosolic BB, BM and MM enzymes [2] or the dimeric and octameric mitochondrial proteins [3], are indeed most prominently expressed in cells and tissues with a high and fluctuating energy demand such as cells in the central nervous system (CNS), cardiac and skeletal muscle, kidney, retina and spermatozon [2 and refs. therein]. Consequently, expression of the different enzymatic subforms is subject to complex tissue and cell-type specific regulation. The B-CK subunit forms an especially interesting study object in this respect. Not only are transcriptional and translational principles involved in its expression, also the modulation of enzymatic activity by protein modification [phosphorylation; 4] and non-covalent protein and membrane associations [5, 6] do play a role. B-CK studies may therefore form a paradigm for the study of other CK-isoforms or enzymes with a similar role in the ATP regulatory networks. The BB-CK isoenzyme has also a direct link to clinical applications. The determination of BB-CK in circulation is of importance in the investigation of neuromuscular (genetic) disorders, the monitoring of different events in the process of cardiac infarction, strokes, CNS and muscular traumas and the development of malignancies [7-11]. Development of reliable standardized tests for B-CK activity in human serum thus far has been hampered by the limited availability of pure enzyme as a control standard for enzyme activity determinations and by the lack of antibodies specific against B-CK.

The baculovirus expression vector system developed by
Smith et al. (1983) has been used for high level expression of eukaryotic proteins, including multimeric proteins and proteins which undergo post-translational modifications [13, 14]. Here we present the use of the baculovirus expression vector system for the production of large quantities of authentic, enzymatically active, human BB-CK dimeric protein. Furthermore, we report on the production of a monoclonal antibody (CK-BYK/21E10) highly specific against the N-terminal portion of B-CK which was used to establish the identity of the in vitro produced protein. The availability of pure subunits and a highly specific monoclonal antibody opens up future possibilities for the study of the pleiotropy of associations and modifications involved in the subcellular compartmentalization of B-CK. Moreover, it forms a valuable asset in the development of standardized immuno-assay kits for clinical chemical monitoring.

Material and methods

Viruses and cells

Sf9 cells, derived from the armyworm Spodoptera frugiperda, were grown at 27°C in flasks using Grace’s Insect medium (Gibco) [14] supplemented with 3.3 g/l yeastolate, 3.3 g/l lactalbumin hydrolysate (Gibco), 5.6 g/l BSA and 10% fetal bovine serum. The wild-type strain E2 of Autographa californica multiple-nucleocapsid nuclear polyhedrosis virus (AcMNPV) was used. The wild-type and recombinant viruses were propagated and titrated (plaque assay, p.f.u. determination) on loosely adherent monolayers of Sf9 cells as described [15].

Construction of the recombinant expression vector

One forward primer (5'-CTTGGATCCATGCCCTTCAC-3') and one reverse primer (5'-TCCGGCTTCAATTCTGGGCAGGCA-3') were designed for PCR amplification of the 1148 bp DNA fragment spanning the complete B-CK open reading frame (start and stop codon underlined). GGATCC sequences (bold print) were included in the primers to bracket the segment between flanking BamHI restriction sites. For PCR amplification, alkaline denatured [16] plasmid pCKB-131 DNA [17] served as template and standard conditions were used for generation and isolation of the DNA product [18]. Next, the 1158 bp BamHI fragment (including primer sequences) was cloned downstream of the polyhedrin promoter in the baculovirus transfer vector pAcDZ1 which contains an E. coli LacZ marker for facilitating recombinant-virus identification [19]. To introduce the B-CK DNA insert into the BamHI cleaved pAcDZ1 vector standard cloning methods were used [18]. The orientation and integrity of the reading frame was checked by supercoiled DNA sequencing [16] using B-CK specific primers.

Generation of recombinant AcMNPV

One μg of purified pAcDZ1-BCK transfer vector DNA was cotransfected with 2 μg pure wt AcMNPV DNA into Sf9 cells (80–85% confluency) using lipofectin transformation (GIBCO-BRL) according to the manufacturer’s protocol [see also Peng et al., 20]. Transfected cells were assayed for the production of recombinant AcMNPV-BCK virus using routine β-galactosidase staining procedures. Recombinant virus was further isolated from culture supernatant through four rounds of plaque purification. A soft agar overlay was added to assist in picking of pure clones. Recombinant plaques were scored as lacking polyhedra and positive for β-galactosidase [21]. One purified plaque was chosen and expanded to a large seeding stock of recombinant virus (stored at 4°C).

Production and purification of B-CK protein from baculovirus infected Sf9 cells

For optimization of production of B-CK protein, Sf9 cells were infected at different multiplicities of infection (moi’s ranging from 1–10). To this end, Sf9 cells were seeded in 25 cm²-flask in 4 ml medium at a density of 1.10⁶ cells/ml and were infected with recombinant virus at the appropriate dilution for 1 h. For harvest, the cells were collected by brief centrifugation at the appropriate time interval after infection (48–72 h), washed in cold phosphate-buffered saline (PBS, 7.3 mM NaH₂PO₄, 55 mM KCl, 47 mM NaCl, 6.8 mM CaCl₂, pH 6.2), resuspended in 8 ml of 0.05 M Tris-HCl (pH = 7.5), 5 mM 2-mercaptoethanol in the presence of 50 μg/ml PMSF and lysed by 3 cycles of freeze-thawing at −80°C. The lysate was centrifuged at 4,000 xg for 10 min and either used directly or stored frozen until use. Recombinant B-CK protein production was verified by SDS-PAGE [22] and standard CK-activity assays. The total protein concentration in the lysate or in purified fractions was measured using the method of Bradford et al. [23].

For preparative purification, cell lysates were adjusted to 10 mM MgCl₂ and subjected to a 1-h treatment with 10 μg/ml DNAsel followed by an additional centrifugation at 12,000 rpm to remove protein aggregates, cell debris and unbroken cell organelles. Approximately 3.5 ml of the lysate (7.6 mg of protein) was then applied on a MonoQ FPLC-column and protein was eluted using a 0–0.8 M NaCl gradient in the above Tris/HCl-buffer. B-CK activity in in-
Individual fractions were determined and individual fractions were resolved by SDS-PAGE. The fractions highest in 43 kDa B-CK protein were pooled, dialysed against phosphate-buffered saline (PBS), and protein concentrations determined.

Production of a peptide immunogen and of a monoclonal antibody against human B-CK

The peptide immunogen BW-17 (Ser-Asn-Ser-His-Asn-Ala-Leu-Lys-Leu-Arg-Phe-Pro-Ala-Glu-Asp-Glu-Phe) corresponding to a unique stretch of amino acids (residues 4—20) from the N-terminal end of the human B-CK was produced by standard solid phase synthesis [24]. The purity of the peptide was assessed by determination of its amino acid composition and by routine HPLC analysis. Next, activated BW-17 peptide was coupled to BSA using MHS as described (25), yielding a ratio of 16 moles of peptide per mole of BSA.

For immunisation of Balb/C mice, 80 μg of the BW-17-BSA conjugate in 300 μl PBS was mixed in with 300 μl Feesn adjuvant and injected intraperitoneally at day one. Preimmune serum was taken prior to immunisation. Booster injections of 80 μg immunogen in incomplete Freunds adjuvant were given at days 14 and 28. Fusion experiments for hybrydoma production were started at around day 33. The standard methodology for fusion of immunized Balb/C spleen cells with SP2/0AG14 myeloma cells was used [26]. Hybridomas were screened by standard ELISA on plates coated with (i) recombinant B-CK/β-galactosidase fusion protein produced in E. coli, (ii) synthetic BW-17 peptide conjugated to casein or thyroglobulin, (iii) Sf-9 lysate from wt or recombinant AcMNPV-BCK or (iv) commercially available M-CK protein (Boehringer) as a negative control. Promising clones were recloned twice by limiting dilution and their specificity was further verified on Western blots as described below. Ultimately, monoclonal (MoAb) CK-BYK/21E10 was chosen for its high specificity and stable growth characteristics. Nude Balb/C mice were used to produce ascites fluid. MoAb CK-BYK/21E10 is an IgG2b.

Typing of purified recombinant B-CK protein: SDS-PAGE analysis

Protein was resolved by SDS-PAGE in 10% (w/v) slab gels and then, for Western blotting, electrophoretically transferred to a nitrocellulose membrane (Schleicher and Schull) (1 h; 250 mA) in a buffer containing 25 mM Tris-HCl (pH = 8.3), 192 mM glycine, and 20% (v/v) methanol. Alternatively, for routine analysis the gels were directly stained with Coomassie Brilliant Blue or silver stain [27] to verify the purity of the B-CK preparation. For immunoblot analyses, nitrocellulose sheets were first blocked with 1% gelatin in Tris-buffered saline/Tween 20 (10 mM Tris-HCl pH 8.0, 150 mM NaCl, 0.05% Tween 20; TBST) followed by incubation with MoAb antibody CK-BYK/21E10 at the appropriate dilution (1:50,000 for ascites MoAb). Filters were washed extensively in TBST and then incubated with a secondary antibody consisting of alkaline phosphatase-conjugated goat anti-mouse IgG (Promega Biotech). The color was developed with nitroblue tetrazolium/5-bromo-4-chloro-3-indolyl phosphate substrate using the Protoblot AP system according to the specifications of the manufacturer (Promega).

Protein sizing by gel filtration

Protein in the dialyzed pure pooled peak fractions of the Mono-Q column (950 μg/ml; 2.5 ml total) was brought on a FPLC-Superdex 200 column (gelfiltration column of Pharmacia/LKB FPLC system) to see whether B-CK was present in its monomeric or dimeric form. Retention in the elution was calibrated with a series of known proteins such as aldolase, bovine serum albumin and ovalbumin. The various fractions of the Superdex 200 elution protein were tested in standard CK-activity assays for the specific activity of the protein (in standard CK Units per mg protein/l). Finally, the immunogenicity of B-CK in the pooled fraction of the MonoQ column was retested by Western blotting.

Determination of CK-activity

Accurate determination of CK-activity was performed using the CK-NAC activated method (CBR-program Boehringer Mannheim; CBR-CK NAc-activated kit no. 475742) adapted for a Cobas Myra automatic analyser (Hoffman La Roche, Basel, Switzerland).

Zymogram typing of CK isoenzymes

Typing of cytosolic CK isoenzymes in cell lysates was done by applying the lysate to cellulose acetate membrane strips (Sartorius) prior to electrophoretic separation at 200 V for 1 h in Tris-barbital buffer pH 8.6 as described by Harm et al. [28]. The positions of CK activity were visualized by activity staining according to Kanemitsu et al. [29].

Results and discussion

Recombinant baculovirus expressing the B-CK protein was isolated as described under ‘Materials and methods’. To determine whether insect Sf9 cells infected with this plaque-
purified recombinant AcMNPV-BCK indeed expressed protein of the expected size (43 kDa), total crude cell lysates were resolved by PAGE. Upon staining a predominant and distinct band of 43 kDa corresponding to the expected molecular weight of creatine kinase B (i.e. 43 kDa; 381 amino acids; [17]) was found (Fig. 1A, lane 2). This band was several fold more intense than a weak 43 kDa band which occurs as an endogenous protein in lysates of mock-infected cells (Fig. 1A, lane 1). The new 43 kDa band appeared first at 24 h after infection, reached an optimal concentration around 48 h after infection and gradually disappeared after 72 h, presumably as a consequence of lysis of the cells. Determination of B-CK activity in cell-lysates revealed that the enzymatic activity of the protein followed exactly the same time course (Fig. 1B). Uninfected SF-9 cells did not show CK activity (not shown).

Optimization of production of B-CK protein revealed that infection with a MOI of 1 gave exactly the same result as a MOI of 10. We could only achieve a maximum p.f.u. of 3.10⁷/ml. Normally, in baculovirus preparations easily 10–20 fold higher titers can be obtained. Though we cannot provide direct evidence it is conceivable that excessive overproduction of B-CK interferes with normal cellular ATP levels needed to sustain transcription and translation processes required for completion of the viral replication cycle. Estimates of the maximal amount of recombinant protein based on the intensity of signals on stained gels suggest that the 43 kDa band of B-CK subunits accounted for approximately 30% of the soluble protein in the lysate under optimal conditions. These results affirm maxima reported for other eukaryotic cytoplasmic proteins expressed from the polyhedrin promoter in this system [30, 31].

To provide independent evidence for the authenticity of the recombinant protein we used a monoclonal antibody, MoAb CK-BYK/21E10, which is specific against the human B-CK protein. The generation of this monoclonal is briefly described in ‘Materials and methods’. Relevant characteristics of the antibody showing its specificity on Western blots of human cell and tissue extracts are given in Fig. 1C. Further description of the antibody and its use in immunohistochemical analysis will be detailed elsewhere (Sistermans et al. Cell and Tissue Res, in press). Immunoblotting (Fig. 1C) demonstrated that the recombinant protein was specifically recognized by the anti-B-CK peptide monoclonal an-
Fig. 2. Purification of B-CK from recombinant virus infected S/9 cells and characterisation of the purified protein. A) Elution profile on the MonoQ-FPLC column. AcMNPV-BCK recombinant virus infected S/9 cell lysate was prepared and applied on a FPLC-MonoQ anion exchange column of the Pharmacia/LKB FPLC system as described under 'Materials and methods'. The OD280 elution profile after application of a NaCl gradient is shown. The vertical 0–100% NaCl scale ranges from 0–0.8 M NaCl. B) SDS/PAGE and immunoblot analysis of the different peak fractions. Panel to the left, proteins stained with silver stain. Panel to the right, proteins immunoreactive with monoclonal CK-BYK/21E10. Lane 1, pooled peak fraction of the MonoQ column (2 μg); lane 2 B-CK recombinant virus infected S/9 lysate (2 μg); lane 3, molecular weight markers as described in the legend of Fig. 1. The weak band (right panel, lane 2) at ± 40 kDa is a proteolytic B-CK cleavage product, which was occasionally observed upon prolonged storage of crude lysates. C) Elution profile of the purified protein on a FPLC-Superdex 200 gel filtration column. Pooled purified protein (2.38 mg) was dialysed against PBS and applied on the Superdex 200 column (Pharmacia/LKB FPLC system). The OD280 elution and activity distribution profiles are shown. B-CK enzymatic activity was determined as described in 'Materials and methods' on an automatic Cobas Myra analyser. Peaks in the elution profiles of reference marker proteins (aldolase (158 kDa) and bovine serum albumin (68 kDa)) are marked with an arrow indicating the native molecular weight in kDa. D) Zymogram assay comparison of recombinant and native BB-CK protein. Migration positions of the MM-, BM- and BB-CK-isoenzymes in the standard marker mix (Boehringer kit) are indicated to the left; also the position of the mitochondrial (ubiquitous) CK isoform which is co-expressed with BB-CK in most cells and tissues is given; lane 1, dialysed pooled pure BB-CK protein (400 ng); lane 2, extract (4 μg protein equivalent) of the human OVCAR-3 cell line which produces authentic human BB-CK protein.
The baculovirus expression vector system is an attractive system that has been reported [33]. Ultimately, these studies may provide us with more insight in B-CK's biological properties and the role of protein-structural flexibility in partitioning and intracellular compartmentalization phenomena. In addition, the baculovirus expression vector system is an attractive system for producing large amounts of pure native B-CK protein which can be used as standard in serological tests for brain and cardiac infarction [8, 9]. Until now, transient production of CK proteins in cDNA transfected COS cells [34] was the method of choice for production of CKs for potential use as clinical standards. Also the newly generated monoclonal antibody CK-BYK/21E10 can be considered a useful tool in these tests.

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