EMAP-II Expression Is Associated with Macrophage Accumulation in Primary Uveal Melanoma

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PURPOSE. Primary uveal melanoma may contain arcs, loops, and networks of periodic acid-Schiff (PAS)-positive patterns, along which numerous macrophages are present. Their recruitment into tumor tissue is mediated by chemotactic cytokines, for which vascular endothelial growth factor (VEGF)-C and endothelial monocyte–activating polypeptide (EMAP)-II are candidates. In this study, the extent of VEGF-C and EMAP-II immunoreaction was related to infiltration of macrophages.

METHODS. Serial sections of 25 primary uveal melanoma lesions were analyzed by immunohistochemistry.

RESULTS. The analysis showed no correlation of VEGF-C immunoreaction and localization of macrophages. However, accumulation of macrophages occurred at sites of EMAP-II expression, especially in areas containing nests of tumor cells, surrounded by arcs, loops, and network patterns. In tumors with a strong EMAP-II immunoreaction, the adhesion molecule intracellular adhesion molecule (ICAM)-1 was strongly expressed on endothelial cells. EMAP-II-positive endothelial cells did not express VEGF receptor-2. However, extensive release of von Willebrand factor was observed. Signs of apoptosis were found neither in tumor cells nor endothelial cells.

CONCLUSIONS. In uveal melanoma, macrophages accumulate at sites of EMAP-II expression. Based on the results, it may be hypothesized that this process of chemotaxis is facilitated by EMAP-II-dependent expression of ICAM-1 on vascular endothelial cells and concomitantly leads to localized vascular damage, as indicated by release of von Willebrand factor. (Invest Ophthalmol Vis Sci. 2003;44:1801–1806) DOI:10.1167/iovs.02-0624

In both primary uveal and cutaneous melanoma, nine different patterns of extracellular matrix deposition have been identified by conventional periodic acid-Schiff (PAS) staining.1–3 These different patterns appear to be of prognostic significance. Especially, the presence of PAS-positive arcs, loops, and network patterns has been associated with poor survival.1–4

In a recent study,5 we suggested that these fibrovascular septa constitute a fluid-conducting meshwork and contain endothelial cells, stromal cells, and macrophages along the arcs, loops, and network patterns. Other reports6,7 also have noted the abundant presence of macrophages in uveal melanoma. Despite this abundance,5–7 their role in uveal melanoma is largely unknown. Macrophages may be involved in angiogenesis (reviewed in Ref. 8), which is supported by a recent study that showed that the number of macrophages was related to microvascular density.9 High numbers of macrophages located in the tumor have also been related to poor prognosis in primary uveal melanoma6,7 and in other tumor types.9–11 Furthermore, their strict colocalization with the arcs, loops, and network patterns5,6 suggests either a role in the development of these patterns or, alternatively, the use of these patterns to invade the tumor lesion. It is unknown how invasion of macrophages in uveal melanoma is mediated. As recently reported, in xenografted melanoma, macrophages were recruited by vascular endothelial growth factor (VEGF)-C,12 which is also expressed in uveal melanoma.13

Besides VEGF-C, endothelial monocyte–activating polypeptide (EMAP)-II may also be involved in the process of macrophage invasion because of its ability to attract monocytes and granulocytes.14,15 EMAP-II mRNA has been detected in normal tissue and in tumors.16–21 It was originally described as a tumor-derived cytokine, isolated from the MethA tumor supernatant.14,15 It is involved in embryonic development,16,22,23 inflammation,21,24 and autoimmune disease.25 The possible effects of EMAP-II on tumor growth are complex.20 On the one hand, release of EMAP-II by hypoxic or apoptotic tumor cells may be involved in macrophage influx and subsequent macrophage-mediated angiogenesis in hypoxic or apoptotic areas.14,15,17 Indeed, hypoxic prostate adenocarcinoma cells release EMAP-II.20 Infiltration of blood-stream monocytes in tumor tissue is mediated by endothelial adhesion proteins ICAM-1 and VCAM-1 and, to a lesser extent, P- and E-selectin (for extensive review, see Ref. 27). If EMAP-II expression mediates recruitment of monocytes and resident macrophages to malignant tumors, this would require concomitant expression of endothelial adhesion proteins. It was already described previously that P- and E-selectin are upregulated on endothelial cells under the influence of recombinant EMAP-II in vitro.14 On the other hand, EMAP-II-mediated influx of macrophages may enhance the immune response against a tumor and thereby counteract tumor growth. Furthermore, EMAP-II may also inhibit tumor growth by targeting the tumor vascular bed through induction of endothelial cell apoptosis,18 by binding to α-adenosine triphosphate (ATP) synthase20 and priming the tumor vasculature for (local) destruction by TNF-α.19,29,30 Evidence of these previously proposed mechanisms is circumstantial, however, and therefore speculative. Most data in recent literature were obtained by studying the effects of applied, recombinant EMAP-II. Thus, the effects of endogenous EMAP-II produced in the tumor interstitium are unclear.

To elucidate the role of endogenous, tumor-derived EMAP-II in primary uveal melanoma, we evaluated both the occurrence and localization of EMAP-II, ICAM-1, and VCAM-1 and determined the presence of macrophages by immunohistochemistry. In addition, levels of apoptosis, endothelial activation (as indicated by expression of VEGF receptor (VEGFR)-2), and signs of endothelial damage (as characterized by the release of von Willebrand Factor [vWF]) were studied. Finally, we investigated VEGF-C expression in relation to the presence and localization of macrophages.
Materials and Methods

Tissues

Tissues were obtained according to guidelines of the Dutch legislation, and our work adhered to the tenets of the Declaration of Helsinki. Frozen specimens from 25 uveal melanomas were obtained from the disease archives of the University Medical Centre Nijmegen where they were stored at -150°C. Presence of disease in all specimens had been determined by a pathologist. All primary melanomas were obtained by surgery, and data of the clinical outcome were obtained for 16 patients with a follow-up time from 1 to 15 years (median 4.5 years). The uveal melanomas varied from 7 to 28 mm in diameter (median, 22 mm) and included 25 choroidal and 2 ciliary melanoma lesions. Uveal melanoma lesions were divided in two parts along the maximal diameter. One part was formalin fixed and the other part was snap frozen. Hematoxylin and cosin (H&E) staining was used on paraffin-embedded sections, and the 25 uveal melanomas were classified as 5 spindle cell type and 20 epithelioid or mixed type. Azan and PAS histochemistry without counterstaining on unbleached paraffin-embedded sections showed that 10 (44%) uveal melanomas contained loops and networks matrix patterns (Fig. 1A).\(^1,2,5,13\) It appeared that in lesions previously classified as arcs, loops, and network patterns,\(^1,2\) the presence of arcs (with and without branching) was minimal. In addition, the arcs, loops, and network patterns were focally present in a tumor. Therefore, it is possible that sampling errors occurred, leading to evaluation of a section containing no network patterns, whereas they were present in other parts of the tumor. Taken together, these considerations may explain the relatively low incidence of arcs, loops, and network patterns in our specimens. In our experience,\(^5,13\) the extent of melanin pigmentation did not significantly hamper the detection of extracellular matrix by PAS and Azan histochemistry. The use of PAS and Azan histochemistry in identifying matrix patterns was recently confirmed by immunofluorescence and electron microscopy.\(^5\)

Antibodies

Antibodies used for immunohistochemistry are listed in Table 1. Anti-CD34 (QBEnd/10), anti-CD68 (Kp1 and PG-M1), and anti-vWF were from Dako (Glostrup, Denmark). Anti-ICAM-1 (PN-E12.1) was raised in our laboratory,\(^7\) and anti-VCA M-I was from Immunotech SA (Mar seilles, France). Anti-VEGFR-2 (Clone KDR-2) was from Sigma (St. Louis, MO). Goat polyclonal antibodies to VEGF-C were purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Rabbit polyclonal antibodies EMAP-II (SA 2846) were obtained from Matthias Clauss (Max Planck Institute, Bad Nauheim, Germany). The rabbit was boosted once against a purified recombinant peptide containing 20 amino acids located at the amino terminal site of the mature human EMAP-II form. Control experiments demonstrated the specificity of the antibodies to both mouse and human EMAP-II and its precursor p43, as previously shown.\(^17,54\)

Immunohistochemistry

Four-micrometer cryosections were air dried and fixed in acetone at room temperature for 10 minutes. Only strongly pigmented sections were bleached by incubating the sections in 3.0% (vol/vol) hydrogen peroxide and 1.0% (wt/vol) disodium hydrogen phosphate for 18 hours at room temperature. Cryosections were incubated with 3% hydrogen peroxide for 30 minutes and subsequently with 20% normal goat (EMAP-II and vWF staining) or horse serum (other staining) for 10 minutes. Successive sections of each specimen were incubated with QBEnd/10 (diluted 1:100), anti-EMAP-II (diluted 1:80) polyclonal antibodies, Kp1 (diluted 1:100), PG-M1 (diluted 1:50), PN-E12.1 (undiluted supernatant), anti-VCAM-1 (diluted 1:40), anti-vWF (diluted 1:300), or anti-VEGF-C (diluted 1:20) for 60 minutes at room temperature. Anti-VEGFR-2 (diluted 1:400) was incubated overnight at 4°C. After the first and all following incubation steps, sections were rinsed with ample phosphate-buffered saline (PBS). Then secondary, 1:200 diluted biotinylated affinity-purified anti-rabbit IgG (for the EMAP-II and vWF stain-

![Figure 1](https://example.com)
TABLE 1. Primary Antibodies Used: Antigens and Type of Cells Stained

<table>
<thead>
<tr>
<th>Antigen</th>
<th>Antibody</th>
<th>Type of Cell</th>
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<tbody>
<tr>
<td>CD34</td>
<td>QBEnd/10</td>
<td>Endothelial</td>
</tr>
<tr>
<td>EMAPII</td>
<td>Anti-EMAPII*</td>
<td>Tumor, endothelial</td>
</tr>
<tr>
<td>CD68</td>
<td>Kp1</td>
<td>Macrophages</td>
</tr>
<tr>
<td>CD68</td>
<td>PG-M1</td>
<td>Macrophages</td>
</tr>
<tr>
<td>VEGF-C</td>
<td>Anti-VEGF-C*</td>
<td>Macrophages</td>
</tr>
<tr>
<td>ICAM-1</td>
<td>PN-E12.1</td>
<td>Endothelial</td>
</tr>
<tr>
<td>VCAM</td>
<td>anti-VCAM-1</td>
<td>Endothelial</td>
</tr>
<tr>
<td>vWF</td>
<td>anti-vWF*</td>
<td>Endothelial</td>
</tr>
<tr>
<td>VEGFR-2</td>
<td>anti-VEGFR-2</td>
<td>Endothelial</td>
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* Polyclonal antibody.

RESULTS

Vascular Staining

In all 25 uveal melanomas, the vasculature was identified by anti-CD34 mAb staining on frozen sections (Table 1) and morphologic properties, as described previously. Often, the endothelial layer was discontinuous, compatible with endothelial damage (Fig. 1B). Although blood vessel density varied among different lesions, no evident increase of blood vessel density in distinct areas within a tumor lesion was observed. Azan and PAS histochemistry (Fig. 1A) and CD34-immunohistochemical staining of serial paraffin-embedded sections indicated that the blood vasculature was closely associated with loops and network patterns, as described previously.

Colocalization of EMAPII Expression and Macrophages

EMAP-II expression was detected in 23 (92%) of 25 uveal melanomas as a cytoplasmic staining of tumor cells (Figs. 1C, 1D). In 15 tumors, EMAPII expression was localized in circumscript areas. Especially, tumor cells directly surrounding blood vessels were strongly positive for EMAPII. In eight melanomas, all tumor cells were intensely positive for EMAPII (referred to as ubiquitous immunoreaction). By comparison to parallel endothelial CD34 staining in serial sections, EMAPII was also present on blood vascular endothelium in three tumors in the areas with local expression of EMAPII, whereas in six tumors with ubiquitous EMAPII staining of tumor cells, endothelial cells were also ubiquitously positive. Especially in areas of loops and network patterns, all tumor cells showed strong EMAPII immunoreaction (Fig. 1D). Strong endothelial EMAPII immunoreaction was also observed in areas with necrosis (not shown). Finally, background staining patterns that could interfere with EMAPII expression analysis were not observed when using other polyclonal antibodies (e.g., anti-vWF polyclonal antibodies).

In all 25 melanomas, macrophages could be detected by CD68 staining and by morphologic characteristics (compare serial sections in Figs. 1E, 1F). When using Kp1 antibody, tumor cells were also weakly positive for CD68 in eight melanoma lesions (Fig. 1F), whereas incubation with the PG-M1 mAb did not result in CD68 staining of melanoma cells (Fig. 1H). These data confirm earlier results. Both macrophages and melanophages were strongly associated with the extracellular loops and network patterns, which contain laminin (Figs. 1G, 1H).

In the 15 tumors with local differences in EMAPII staining intensity, macrophages were consistently more abundant in areas with higher EMAPII levels, especially around blood vessels (compare serial sections in Figs. 1C, 1E). Counting of the number of macrophages in four of these tumors demonstrated a significant difference (P < 0.0001) between those in tumor areas with negative to low EMAPII immunoreaction (72 ± 49 macrophages/mm²; mean ± SD) and those in areas with strong EMAPII immunoreaction (457 ± 129 macrophages/mm²). Macrophages were detected around both EMAPII-negative (n = 12) and -positive (n = 3) blood vessel endothelium. In eight melanomas with ubiquitous EMAPII immunoreaction, macrophages were equally distributed over the tumor tissue. In these tumors, macrophages were present around EMAPII-negative and -positive (n = 7) vessels as well. In areas containing loops and network extracellular matrix patterns, nests of tumor cells were intensively positive for EMAPII immunostaining. In addition, EMAPII immunoreaction was lower in areas containing no loops and network patterns compared with areas with these patterns within one lesion.

ICAM-1 and VCAM-1 Immunostaining

Coexpression of ICAM-1 and EMAPII was found on tumor cells in all 23 EMAPII-positive cases (Figs. 2A, 2B). Conversely, in areas with no or low tumoral EMAPII immunoreaction, ICAM-1 was not detected on tumor cells. In only 3 of 15 tumors with local EMAPII expression was ICAM-1 expression observed on EMAPII-positive endothelium, whereas this was the case in 6 of 8 tumors (Figs. 2A, 2B) with generalized immunoreaction. ICAM-1 was often expressed on a subset of endothelial cells...
conptosis was absent. In the other two tumors, apoptosis was showed staining, whereas in the remaining tumor cells, apopto-
one of the tumors were apoptotic endothelial cells found. In
Apoptosis was detectable in only 3 of 15 uveal melanomas. In
Detection of Apoptosis
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We analyzed 16 EMAP-II VEGFR-2 and vWF Immunostaining
(Figs. 2C, 2D). In this group of tumors, EMAP-II
sion of EMAP-II positivity, VEGFR-2, and absence of vWF-re-
VEGFR-2 and vWF Immunostaining
We analyzed 16 EMAP-II-positive uveal melanomas for the
presence of VEGFR-2 and vWF as markers of endothelial activ-
atation and damage, respectively. In 14 melanomas, comparison
to parallel EMAP-II staining demonstrated that EMAP-II-positive
endothelial cells showed no expression of VEGFR-2 (Figs. 3A,
3B). However, strong staining of vWF in endothelial cells and
underlying tissue indicated a release of vWF by this endothelium
(Figs. 3A, 3C). Conversely, EMAP-II-negative endothelial
cells expressed VEGFR-2 in five melanomas (Figs. 3D, 3E)
whereas no evident damage was detected in these cells by vWF
staining (not shown). In only two tumors, a weak colocaliza-
tion of EMAP-II positivity, VEGFR-2, and absence of vWF-re-
lease was observed. Patterns similar to those of EMAP-II stain-
ing were not observed when using other polyclonal antibodies
(e.g., anti-vWF polyclonal antibodies).

Detection of Apoptosis
Apoptosis was detectable in only 3 of 15 uveal melanomas. In
none of the tumors were apoptotic endothelial cells found. In
one tumor, an evidently necrotic area positive for EMAP-II,
showed staining, whereas in the remaining tumor cells, apop-
tosis was absent. In the other two tumors, apoptosis was
confined to a few tumor cells dispersed over the entire tumor
area (not shown).

Absence of Colocalization of VEGF-C Expression
and Macrophages
Finally, we analyzed 12 melanomas for expression of VEGF-C.
In six melanomas, VEGF-C was expressed in restricted areas.
No evident colocalization of accumulation or absence of mac-
rophages and VEGF-C expression was observed in any lesion
(data not shown).

Correlation with Clinical Outcome
In only one patient, liver metastasis was found 2 years after
eutelation. This tumor contained PAS-positive loops and net-
work patterns and ubiquitous EMAP-II immunoreaction of tu-
mor and endothelial cells and macrophages was equally dis-
tributed over the lesion. In addition, ICAM-1 was expressed by
the tumor and a subset of endothelial cells. No VCAM-1 ex-
pression or apoptosis was observed. EMAP-II-positive endo-
thelial cells lacked expression of VEGFR-2 but stained strongly for
vWF. VEGF-C expression was not evaluated.

DISCUSSION
Recently, we described that the arcs, loops, and network
eextracellular matrix patterns may represent a fluid-conducting
meshwork in xenografted and primary uveal melanoma.5 Our
findings indicated that these structures closely parallel the
previously described vascular channels.39 Although it has been
demonstrated that melanoma cells may express nonmelano-
cytic markers like endothelial-cell-associated CD34 and mac-
rophage-associated CD68,39,40 we could not confirm the pres-
ence of blood-conducting channels lined by tumor cells.
Instead, septa consisting of extracellular matrix components
that could be visualized by PAS and Azan histocytology sur-
rrounded tumor cell nests. Remarkably, many macrophages
(i.e., CD68-positive cells) were found associated with this
meshwork in melanoma xenografts. We now have studied the
presence and localization of macrophages in uveal melanoma,
focusing on the mechanisms underlying infiltration by this cell

![Image](http://iovs.arvojournals.org/pdfaccess.ashx?url=/data/journals/iovs/933226/ on 05/08/2017)
type. Our results also show that in uveal melanoma, macrophages are abundantly present and colocalize with the extracellular matrix patterns containing laminin.5,41

In uveal melanoma containing loops and network extracellular matrix PAS-positive patterns, the monocyte chemotactic protein EMAP-II was abundantly present in the cytoplasm of tumor cells and was accompanied by local macrophage accumulation. Additional analysis of uveal melanomas lacking these patterns showed that different levels of local EMAP-II staining within one tumor were present and that macrophages were preferentially located in areas with highest EMAP-II immunoreaction. These data support the hypothesis that tumor cells recruit local resident tissue macrophages and peripheral monocytes by release of EMAP-II. This hypothesis was corroborated by several previous studies on the expression of EMAP-II in tumor cells and the ability of EMAP-II to attract macrophages.14,15 Indeed, infiltration of the tumor may be facilitated by the loops and network patterns by their serving as a gateway. However, secondary changes including extensive retinal detachments, glioma, ruberosis, and intraocular hemorrhage may contribute to the influx of macrophages as well, because most of the lesions analyzed were large. Because VEGF-C expression did not colocalize with absence or accumulation of macrophages, our data suggest that VEGF-C does not play a substantial role in macrophage invasion in uveal melanoma.

EMAP-II is normally retained intracellularly and can be released and partially processed by several triggers, including apoptosis and hypoxia.17,20 Because apoptotic cells were hardly present in the tumor lesions, this does not seem to be a decisive factor in EMAP-II expression in uveal melanoma. Hypoxic tumor cells may express hypoxia inducible factor (HIF)-1α and release vascular endothelial growth factor (VEGF)-A. In this respect, it is noteworthy that we observed no release of VEGF-A in the primary uveal melanoma lesions.13 However, to the best of our knowledge, expression of HIF-1 in uveal melanoma has not been reported thus far.

Purified recombinant mature EMAP-II protein activates endothelial cells and enhances the expression of adhesion molecules P- and E-selectin.14 These adhesion molecules mediate invasion of different types of leukocytes into the underlying tissue (for review, see Ref. 27). Macrophage infiltration is primarily mediated by ICAM-1 and VCAM-1 adhesion molecules, however. Remarkably, one of our first observations was that EMAP-II and ICAM-1 are coexpressed on tumor cells. This indicates that either both these molecules are upregulated by the same stimulus, or that EMAP-II expression induces ICAM-1 expression by an autocrine mechanism. Moreover, ICAM-1 was predominantly expressed on the vascular endothelium in those tumors that showed intense ubiquitous staining for EMAP-II. In tumors with focal EMAP-II immunoreaction, the endothelium was often negative for both EMAP-II and ICAM-1, suggesting that a certain threshold level of EMAP-II binding to endothelial cells is necessary for induction of ICAM-1. Functional experiments in cell cultures are necessary to confirm these findings. Because only four tumors expressed vascular cell adhesion molecule (VCAM) on their vessels, it is unlikely that this adhesion molecule is induced by EMAP-II expression and contributes to infiltration of monocytes.27

To elucidate further the role of endogenously produced EMAP-II in primary uveal melanoma, we evaluated additional parameters that have been associated with EMAP-II expression in the literature—that is, the induction of apoptosis and of angiogenesis. As stated earlier, EMAP-II is involved in macrophage influx in several types of carcinomas. This has been related to tumor progression5,10 and may be explained partly by macrophage-induced neovascularization13 (for review see Ref. 44). We hypothesized that in primary uveal melanoma, hypoxic tumor cells secrete EMAP-II, contributing to recruit-


