The Inhibition of Uveal Melanoma Cell Metastasis by the Insulin-like Growth Factor-I Receptor Blocker Picropodophyllin

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Abstract

Uveal melanoma is the most common primary intraocular malignant tumor in adults. It has an elevated mortality rate due to a high incidence of metastases, which occur preferentially in the liver (1, 2). The treatment regimen today is enucleation and metastasis of uveal melanoma cells. Targeting this receptor may therefore comprise a strategy to treat ongoing disease (today incurable) as well as a strategy to prevent development of metastases in patients with primary disease.

In uveal melanoma, metastasis is the crucial event determining the outcome of patients. Search for agents decreasing the metastatic potential of the primary tumor is therefore of particular interest. Metastasis is a process involving many components, including tumor cell adhesion, migration, extracellular matrix (ECM) proteolysis, and invasion. The tumor cells undergo intravasation, disperse via the vascular and the lymphatic systems, and finally extravasate to invade the secondary sites. In all these steps, proteolytic enzyme systems are involved, including the matrix metalloproteinase (MMP) system and the plasminogen activation system. The migration of a malignant cell through the ECM and the basement membrane requires proteolytic activities (3). Baker et al. (4) have shown that uveal melanomas express both MMP-2 and MMP-9, tissue inhibitor of metalloproteases 2, urokinase plasminogen activator, plasminogen activator inhibitor 1 and 2, as well as different integrins.

The preferential dissemination to the liver raises the possibility that hepatic environmental factors are important for the growth and progression of uveal melanoma. Such conditions may involve growth factors produced in the liver [e.g., insulin-like growth factor 1 (IGF-I) serving as a ligand of the IGF-IR receptor (IGF-IR)]. Ligand binding to the IGF-IR activates the intrinsic tyrosine kinase of the β-subunit leading to phosphorylation of IGF-IR, which in turn activates key signal transduction molecules involved in cell proliferation and growth.
apoptosis protection, such as the mitogen-activated protein kinases and the phosphoinositol-3-kinase (5–8).

Efforts to target IGF-I system in melanoma in the past have been made (9–12). Recently, we detected expression of IGF-IR in clinical samples of uveal melanoma and showed that cultured uveal melanoma cells responded to inhibition of IGF-IR (using, e.g., neutralizing antibodies) by cell death (13). These data pointed to the possibility of using IGF-IR as target for uveal melanoma. Recently, we showed that the cyclodiginic picropodophyllin (PPP) acts as a specific inhibitor of the IGF-IR tyrosine phosphorylation without inhibiting the highly homologous insulin receptor (14–18). PPP was proved to also inhibit phosphorylation of IGF-IR in tumor tissues of xenografted mice and phosphorylation of downstream signaling molecules [extracellular signal-regulated kinase (ERK) and Akt] was decreased (14). Subsequently, the tumor cell xenografts underwent complete regression.

In this study, we aimed to investigate the effects of PPP on uveal melanoma cell cultures and xenografts with special focus on mechanisms important for tumor cell invasion.

**Materials and Methods**

**Reagents.** PPP was synthesized as described (14) and was dissolved in DMSO (0.5 mmol/L) before addition to cell cultures. The phosphotyrosine (PY99) and polyclonal antibodies to the β-subunit of IGF-IR (H-60) were from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA).

**Cell cultures.** Three cell lines obtained from human primary uveal melanomas (OCM-1, OCM-3, and 92-1) were previously described (13). R+ cells (overexpressing the human IGF-IR) were from Dr. Renato Baserga (Thomas Jefferson University, Philadelphia, PA). R+ cell line was cultured in the presence of G-418 (Promega, Madison, WI).

**Immunoprecipitation and Western blotting.** Cells were cultured to subconfluence in 6-cm plates. After indicated treatments, cells were analyzed for IGF-IR, ERK1/2, and Akt phosphorylation as described elsewhere (19, 20).

**Cell survival assay.** Cell viability determinations were done using the Cell Proliferation Kit II (Roche, Inc., Indianapolis, IN), which is based on colorimetric change of the yellow tetrazolium salt 2,3-bis[2-methoxy-4-nitro-5-sulfophenyl]-2H-tetrazolium-5-carboxanilide inner salt in orange formazan dye by the respiratory chain of viable cells (21).

**Integrin expression profile.** Expression profile of collagen-specific integrins (α1, α2, α5, αv, β1, β3, and αvβ3) was assessed by fluorescence-activated cell sorting using antibodies in the Chemicon Investigator Kit (Chemicon Europe, Hampshire, United Kingdom).

**ECM adhesion assays.** Adhesion of uveal melanoma cells to ECM proteins (collagen type I, collagen type IV, fibronectin, laminin, and vitronectin) was assessed using a CytoMatrix screening kit (Chemicon International, Temecula, CA) following the protocol of the manufacturer. Cells were nonenzymatically disaggregated from culture flasks using cell dissociation solution (Sigma-Aldrich, Stockholm, Sweden) and resuspended at a concentration of 5 × 10^5/mL in RPMI 1640 with 0.1% bovine serum albumin. Cell suspension (100 μL) was added to each well and the plate was incubated at 37°C for 1 hour. After washings and staining (0.2% crystal violet in 10% ethanol for 5 minutes), the levels of adhesion were determined by measuring the absorbance at 540 nm on a microplate reader. Triplicate wells were assessed for each treatment; experiments were repeated thrice and the mean value was calculated. In all cases, adhesion to wells coated with bovine serum albumin acted as negative controls and levels of adhesion to ECM substrates were calculated relatively to the controls.

**Gelatin zymography.** The gelatinolytic activity of MMP-2 was analyzed by zymography as previously described (22). Unheated aliquots of conditioned media (50× concentrated) were electrophoresed on a 7.5% SDS polyacrylamide gel containing 1 mg/mL gelatin (Bio-Rad, Stockholm, Sweden) before and after 2-hour treatment with the MMP-2 activator p-aminophenylmercuric acetate (APMA; 1 mmol/L). Recombinant MMP-2 (Chemicon International) was used as control. Molecular weight standards were run simultaneously. The gels were stained with Coomassie blue and destained with acetic acid-methanol until the desired color intensity was obtained. Gelatinolytic enzymes were detected as transparent bands on the background of Coomassie blue-stained gelatin.

**Assessment of cell migration and invasion in vitro.** To assay the effect of PPP on migration and invasion of uveal melanoma cells, BD BioCoat Matrigel chambers were used following the instructions of the manufacturer (BD Biosciences, San Diego, CA). In brief, cells (5 × 10^3/mL) treated with PPP or solvent for 1 hour were washed and put into chambers containing an 8-μm pore size PET membrane without (for assay of migration) or with a thin layer of Matrigel serving as a reconstituted basement membrane (for assay of basement membrane invasion) or with basement membrane covered by confluent human endothelial human umbilical vascular endothelial cells (for assay of endothelial invasion) for overnight incubation (23). Medium containing 10% serum or IGF-I (as a chemoattractant) is added to the well outside of the chambers.

**In vivo experiments.** Ten-week-old pathogen-free severe combined immunodeficient mice were used and housed within plastic isolators in a sterile facility. OCM-1 cells were injected s.c. at 10^7 cells per mice in a 0.2-mL volume of sterile saline solution. Experimental treatments with PPP were done as we previously described (14). All of the experiments were done according to the ethical guidelines for laboratory animal use and approved by the institutional ethical committee.

**Experimental reproducibility.** All experiments were repeated at least thrice with similar results.

**Results**

**Effects of PPP on IGF-IR phosphorylation and survival of uveal melanoma cells.** We analyzed phosphorylation of IGF-IR in cultured uveal melanoma cells stimulated with IGF-I as well as the dose-response effects of PPP on IGF-IR phosphorylation in these systems. OCM-1 and 92-1 cells had first been serum depleted overnight and then treated with solvent (negative control) or PPP for 60 minutes at various concentrations and finally stimulated with IGF-I for 5 minutes, which is the stimulation time for maximal IGF-IR phosphorylation (15). Both cell lines responded with a clear IGF-IR activation (Fig. 1A). Consistent with results obtained from several other cell types (14, 15, 18), PPP dose-dependently reduced IGF-I-induced receptor phosphorylation (Fig. 1A). In contrast, the IGF-IR expression was not affected by this short treatment with PPP (Fig. 1A). As assessed by densitometry, 500 nmol/L PPP reduced the level of IGF-IR phosphorylation by >50% (data not shown). The PPP treatment also led to decreased phosphorylation of the downstream molecules Akt and ERK1/2 (Fig. 1B).

Next, we investigated the effects of PPP on survival of cultured uveal melanoma cells. The cells were treated with different concentrations of PPP for 48 hours. Cell survival was assessed by 2,3-bis[2-methoxy-4-nitro-5-sulfophenyl]-2H-tetrazolium-5-carboxanilide inner salt. Figure 1C shows that the IGF-IR inhibitor efficiently decreased viability of the three studied cell lines (OCM-1, OCM-3 and 92-1). A PPP concentration of 500 nmol/L reduced cell survival by 60% to 90%. Consistent with the effects on IGF-IR phosphorylation, the IC50 values are between 50 and 500 nmol/L (Fig. 1C).

We also analyzed the effects of IGF-I on viability of uveal...
melanoma cells. OCM-1 cells were serum depleted with and without IGF-I over a 48-hour period. The presence of IGF-I drastically increased viability of the serum-depleted cells (Fig. 1D). However, if PPP (500 nmol/L) was present, this effect of IGF-I was completely abrogated (Fig. 1D).

Effects of PPP on invasion mechanisms in vitro. Invasion involves several crucial mechanisms. One early step is cell adhesion (through integrins at the cell surface) to ECM components. It was recently shown that α1, α2, and α3 integrins showed higher expression in uveal melanoma cells with invasive phenotype compared with noninvasive cells, and inhibition of these α integrins decreased binding to collagen IV, fibronectin, and laminin (24). This is interesting because these three ECM proteins have been reported to surround the tumor cells in primary and metastatic uveal melanomas (25). We first investigated the expression profiles of collagen-specific integrins in OCM-1 and OCM-3. Interestingly, both cell lines expressed fully detectable to high amounts of α1, α2, and α3 integrins (Fig. 2A). Because IGF-IR has previously been shown to control integrin-mediated cell adhesion to ECM proteins (26), we next analyzed the effects of PPP on uveal melanoma cell adhesion to collagen type IV, fibronectin, and laminin in comparison with collagen type I and vitronectin. The cells were treated for 1 hour with three different concentrations of PPP (0, 50, and 500 nmol/L). In a separate experiment, it was shown that 1-hour treatment with PPP does not decrease cell survival (data not shown). PPP induced a strong dose-dependent inhibition of adhesion to fibronectin, laminin,

![Image](https://www.aacrjournals.org/clinicanmuneandcancerreseaarch/figs/1445/2006/12(4)1385-1401_f1.large.jpg)

**Fig. 1.** Effects of PPP on phosphorylation of IGF-IR and growth of uveal melanoma cells. A, cultured OCM-1 and 92-1 cells were serum depleted overnight and then treated with PPP at the indicated concentrations (0–2,500 nmol/L) for 1 hour. IGF-I was then added for a 5-minute incubation, whereupon cells were harvested for immunoprecipitation of IGF-IR. The immunoprecipitates were fractionated by gel electrophoresis and then immunoblotted with a phosphotyrosine antibody or an antibody to the ε-subunit of the IGF-IR (loading control). B, OCM-1 cells were serum depleted overnight and then treated with PPP at the indicated concentrations (0–2,500 nmol/L) for 1 hour. IGF-I was then added for a 5-minute incubation. The cell lysates were analyzed for pAkt/Akt and pERK1/2/ERK1/2. C, cultured OCM-1, OCM-3, and 92-1 cells were cultured in complete medium containing 10% serum in 96-well plates. When the cells had reached subconfluency, they were treated with PPP at the indicated concentrations (50–2,500 nmol/L) for 48 hours. Cells were assayed for viability using the 2,3-bis[2-methoxy-4-nitro-5-sulfophenyl]-2H-tetrazolium-5-carboxanilide inner salt assay. Points, mean of triplicates; bars, SD. D, OCM-1 cells were serum depleted with or without IGF-I (100 ng/mL). The effects of PPP (500 nmol/L) on unstimulated and IGF-I stimulated cells were also investigated. After incubation for 48 hours, cell viability was assayed by 2,3-bis[2-methoxy-4-nitro-5-sulfophenyl]-2H-tetrazolium-5-carboxanilide inner salt. Columns, mean of triplicates; bars, SD.
and collagen IV (Fig. 2B). Vitronectin and collagen I, on the other hand, were not significantly affected.

MMP-2 is generally required for invasive and metastatic capability of malignant cells and has been suggested to be controlled by the IGF-I signaling (22). Therefore, we investigated the effect of IGF-I stimulation on gelatinolytic activity of MMP-2 secreted from uveal melanoma cells and whether PPP could interfere with this event. OCM-1 and OCM-3 cells were incubated in serum-free medium with and without IGF-I and PPP at different concentrations for 6 hours. It was confirmed

Fig. 2. Effect of PPP on uveal melanoma cell adhesion. A. OCM-1 and OCM-3 cells were analyzed for expression profile of collagen-specific integrins using the CHEMICON kit. B. OCM-1 and OCM-3 were cultured in 96-well plates precoated with fibronectin (FN), vitronectin (VN), laminin (LM), collagen type I (CG1), or collagen type IV (CG4). The cells were treated for 1 hour with two different concentrations of PPP (50 and 500 nmol/L) or were left untreated. *, P < 0.05; **, P < 0.01; ***, P < 0.005; ****, P < 0.001. The effects on cell adhesion of these ECM proteins were determined as described in Materials and Methods. Columns, mean; bars, SD. C. OCM-1 and OCM-3 cells were serum depleted overnight and then treated with IGF-I (10 ng/mL) and different concentrations (0, 50, 500, and 2,500 nmol/L) of PPP for 6 hours. The negative control (C-) was treated with IGF-I and PPP. The conditioned media were collected and treated with the MMP-2 activator APMA (1 mmol/L; bottom) or remained untreated (top). As negative control (C-), cells were incubated with serum-free medium without IGF-I. Activated and inactivated recombinant MMP-2 were used as positive controls. The gelatinolytic activity of MMP-2 in the conditioned media was detected as transparent bands on the background of Coomassie blue–stained gelatin.
that 6-hour treatment with PPP does not significantly decrease cell survival (data not shown). The conditioned media were collected and treated with APMA (an MMP-2 activator) or untreated before electrophoresis. Recombinant MMP-2, activated or not by APMA, was used as control. Accordingly, gelatin zymography revealed two bands corresponding to the native (72-kDa) and activated (62-kDa) forms of MMP-2 (Fig. 2C). No MMP-2 expression was seen in cells that were not stimulated with IGF-I (C-). In the presence of IGF-I but in the absence of APMA, both the native and active MMP-2 appeared (Fig. 2C, top). PPP decreased in a dose-dependent manner the amount of active MMP-2 (Fig. 2C, top). In APMA-treated samples, only the active MMP-2 was seen after IGF-I stimulation and was dose-dependently decreased by PPP (Fig. 2C, bottom). Taken together, these data suggest that both the IGF-I-induced activation and expression of MMP-2 are repressed by PPP.

Next, we investigated whether PPP could affect migration of uveal melanoma cells in vitro using matrix chambers containing a pored membrane using 10% serum as a chemoattractant (27, 28). Cells were incubated only with solvent or with PPP (50 or 500 nmol/L) for 1 hour and then added to the chambers overnight. It was confirmed that 1-hour treatment with PPP did not lead to reduced cell viability (data not shown). Figure 3A (left) shows that 100% of both OCM-1 and OCM-3 control cells had migrated through the membrane. However, after treatment with 50 and 500 nmol/L PPP, only 55% to 65% and 15% to 20% of the uveal melanoma cells, respectively, could pass the membrane (Fig. 3A). These responses are also illustrated by the microphotographs in Fig. 3A (right). The results imply that PPP efficiently reduces cell migration.

By adding a reconstituted basement membrane on the pored membrane, we repeated the experiment to assay the effect of
PPP on basement membrane invasion. Of the control cells, 55% to 60% could pass the basement membrane after an overnight incubation, but if the cells had been treated with PPP for 1 hour only, 20% and 5% of the cells could invade through the basement membrane depending on the PPP concentration (50 and 500 nmol/L, respectively; Fig. 3B). Finally, the reconstituted basement membrane was covered by confluent endothelial cells to assay the invasive ability of uveal melanoma cells on this barrier. Using this in vitro model, the capability of tumor cells to invade through vessel endothelium can be evaluated. As shown in Fig. 3C, the addition of endothelial barrier decreased the invasive capability of the untreated uveal melanoma cells only marginally compared with the basement membrane barrier (cf. Fig. 3B), indicating that the tumor cells can easily pass through the endothelial cell layer. The inhibitory effect of PPP on the endothelial invasion was of the same magnitude as that on the basement membrane invasion. As a control experiment to examine quality of the endothelial layer, we investigated the ability of nontumorigenic cells to cross the endothelial cell/basement membrane combination. When serum was used as a chemoattractant, these cells (R+) exhibited 100% migration and 80% of them could pass the basement membrane.
membrane barrier but failed to invade through the endothelial cell/basement membrane combination (Fig. 3D). However, when IGF-I was used as a chemoattractant, ~45% of the cells migrated and <20% invaded the basement membrane but no cells could invade the endothelial layer (Fig. 3D). We also investigated the effects of PPP on migration and basement membrane invasion of cells exposed to IGF-I as a chemo-attractant. The experiment was done on OCM-1 cells. As shown, IGF-I caused 100% migration and ~60% invasion through the basement membrane barrier (Fig. 3E). PPP resulted in a dose-dependent decrease in both variables, similar to that seen in cells attracted by serum (cf. Fig. 3A and B).

Effects of PPP on uveal melanoma xenografts. As shown in Fig. 1B, PPP efficiently reduced survival of cultured uveal melanoma cells. Next, we investigated the effects of PPP on establishment and growth of uveal melanoma in vivo. OCM-1 cell xenografts were produced s.c. in severe combined immunodeficient mice. This cell line, which has been reported

![Effects of PPP on phosphorylation of IGF-IR, Akt, and ERK1/2 as well as on MMP-2 activity and apoptotic rate.](image)

**A**

- pIGF-1R
- pERK 1/2
- pAkt

**B**

- MMP2
- Actin

**Fig. 5.** Effects of PPP on phosphorylation of IGF-IR, Akt, and ERK1/2 as well as on MMP-2 activity and apoptotic rate. A. In a separate experiment, established OCM-1-xenografted mice were treated with DMSO (control) and PPP (20 mg/kg/12 h) for 4 days. From the tumor samples, protein lysates were prepared and analyzed for expression of phosphorylated IGF-IR, Akt, and ERKs as well as gelatinolytic activity of MMP-2. B. Separate samples were prepared for histologic sections. In the control section, two typical mitoses are indicated by M. In PPP-treated tumor, several apoptotic cells are seen and indicated by A.

![Image](image)
to be tumorigenic elsewhere (29), established well in 2 weeks (Fig. 4A). One group of mice, after inoculation, was directly treated with PPP for 3 days; after which, they were kept untreated until the experiment was stopped after additional 17 days. Figure 4A shows clearly detectable tumors in control (DMSO) mice after 11 days and in the PPP-treated ones after 20 days. At this time, the tumor volumes measured 658 and 54 mm³ (in average) in DMSO- and PPP-treated mice, respectively. Thus, this early short PPP treatment dramatically reduced tumorigensis of uveal melanoma cells. We also investigated the effects of PPP on already established tumors. The tumors exhibited sizes of 100 to 150 mm³ at this time and grew steadily. The mice were treated with PPP for 12 days. As shown in Fig. 4B, the tumors immediately responded with growth inhibition and, after 7 days, they started regressing drastically. The mice did not exhibit any weight loss or other side effects during the treatment period. Similar results were also obtained in xenografts established from OCM-3 and 92-1 cells (data not shown). In a parallel experiment on established tumors treated with DMSO and PPP for 12 days, we investigated the presence of micrometastases in liver in the two groups. Figure 4C (left) shows the histology of liver tissues from a control mouse. Micrometastases can easily be detected. In samples from PPP-treated mice, extremely few tumor cell aggregates were visible in the liver. Figure 4C (right) shows the highly significant decrease in incidence of micrometastases in the PPP group of mice.

In a parallel experiment, mice with established tumors were treated with DMSO or PPP only for 4 days (before regression started; cf. Fig. 4B). Some tumor samples were snap-frozen for analysis of phosphorylation of IGF-IR, Akt, and ERK1/2 as well as for assay of MMP-2 gelatinolytic activity. Other samples were fixed in formalin for preparation of histologic sections. Figure 5A shows that treatment with PPP caused strong inhibition of IGF-IR, Akt, and ERK1/2 phosphorylation in tumor tissues. Moreover, expression of MMP-2 was decreased. Figure 4B shows histology of tumor samples. As compared with the control tumors (from mice treated with DMSO), those from PPP-treated mice exhibited a lot of apoptotic figures (Fig. 4B). Analysis of the number of apoptotic cells from several sections shows a strongly significant increase in samples isolated from PPP-treated mice compared with DMSO-treated ones (Table 1).

### Table 1. PPP induces apoptosis in uveal melanoma in vivo

<table>
<thead>
<tr>
<th>Treatment</th>
<th>No. apoptotic cells/10 high-power fields (mean ± SD)</th>
<th>Significance</th>
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<tbody>
<tr>
<td>DMSO</td>
<td>22.3 ± 1.6</td>
<td>—</td>
</tr>
<tr>
<td>PPP</td>
<td>63.7 ± 4.1</td>
<td>P &lt; 0.001</td>
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*OCM-1-xenografted mice treated with DMSO and PPP (20 mg/kg/12h) for 4 days.
†Histologic sections were prepared from tumors and analyzed for numbers of apoptotic cells per 10 high power fields (200 x). Means and SDs from 10 sections are shown.
‡Significance levels were determined by Students t test.

### Discussion

This study shows that the IGF-IR inhibitor PPP efficiently decreases survival of cultured uveal melanoma cells and causes regression of uveal melanoma tumor xenografts in mice. Moreover, PPP strongly attenuates several mechanisms involved in invasion (cell adhesion to ECM proteins, MMP-2 expression and activity, cell migration, and basement membrane and endothelial invasion) as well as reduces implantation and establishment of uveal melanoma cells. Because both invasion and tumor cell establishment comprise crucial events in metastasis, our findings raise the possibility of applying IGF-IR inhibitors as adjuvants or neoadjuvants in therapeutic management of uveal melanoma patients.

The invasive progression of tumors is a major impediment for successful treatment of neoplastic diseases in general (30) and of uveal melanoma in particular (1, 2). Today, up to 50% of the patients with operated primary uveal melanoma develop metastases and, unfortunately, most of them succumb to the disease.

Usually, ECM constitutes the first barrier against tumor spread. The term vascular mimicry describes the unique ability of highly aggressive uveal melanoma cells to form matrix-rich networks, mimicking embryonic vasculogenesis (31). Hess et al. (32) have recently suggested that the phosphoinositol-3-kinase pathway promotes vascular mimicry by regulating the activity of MMP-2 and cleavage of the laminin 5-2 chain. Interestingly, the IGF-IR signaling pathway, through the phosphoinositol-3-kinase branch, has been shown to play important roles in the interaction with ECM. IGF-IR has been found to control cell migration and integrin-mediated adhesion to ECM proteins (26). IGF-IR may influence adhesion to ECM proteins in several ways. It may affect expression of integrins and may also cause activation of specific integrins (26, 33). In this context, focal adhesion kinase, shown to be regulated by IGF-IR through the phosphoinositol-3-kinase/Akt pathway (26, 33), probably is involved.

Further, IGF-IR has been shown to affect biosynthesis and activation of MMPs (22, 34). These findings are consistent with our present results showing that inhibition of IGF-IR, using PPP, significantly decreases uveal melanoma cell adhesion to the ECM proteins fibronectin, laminin, and collagen IV, as well as down-regulates the expression and activity of MMP-2. We could also prove that MMP-2 expression is down-regulated in tumors in vivo.

The next crucial step in metastasis is the tumor cell adherence to, and digestion of, the basement membrane to enable passage into the blood stream. The main components of basement membrane are laminin and collagen IV. Both the adherence of tumor cells to the basement membrane and MMP-2-mediated basement membrane digestion have been shown to be under the control of IGF-IR (34). Using the Matrigel model, we found that PPP strongly inhibits the basement membrane invasion of uveal melanoma cells.

A third important step in metastasis is establishment of circulating tumor cells in a new tissue. This step can experimentally be investigated by determining the ability of tumor cells to produce tumors in immunosuppressive animals. We found that early short treatments with PPP drastically delayed tumor formation. This result is consistent with other studies showing
References


In conclusion, our study provides further support for an important role of IGF-IR in uveal melanoma and raises the possibility of using IGF-IR inhibitors in treatment of ongoing disease as well as in prevention of development of metastases.

that the IGF-IR is important for production of metastatic tumors (34, 35). Most importantly, we could show in the current study that the incidence of micrometastases in the liver of uveal melanoma xenografted mice was drastically reduced.
The insulin-like growth factor-I receptor inhibitor picropodophyllin causes tumor regression and attenuates mechanisms involved in invasion of uveal melanoma cells.

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