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<td>Author(s)</td>
<td>Shen, B; Chu, ESH; Zhao, G; Man, K; Wu, CW; Cheng, JTY; Li, G; Nie, Y; Lo, CM; Teoh, N; Farrell, GC; Sung, JJY; Yu, J</td>
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PPARgamma inhibits hepatocellular carcinoma metastases
in vitro and in mice

B Shen1,2,8, ESH Chu1,8, G Zhao1,3,8, K Man4, C-W Wu1, JTY Cheng5, G Li5, Y Nie2, CM Lo6, N Teoh6, GC Farrell6, JFY Sung1 and J Yu*,1,7

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BACKGROUND: We have previously demonstrated that peroxisome proliferator-activated receptor (PPARγ) activation inhibits hepatocarcinogenesis. We aim to investigate the effect of PPARγ on hepatocellular carcinoma (HCC) metastatic potential and explore its underlying mechanisms.

METHODS: Human HCC cells (MHCC97L, BEL-7404) were infected with adenovirus-expressing PPARγ (Ad-PPARγ) or Ad-LacZ and treated with or without PPARγ agonist (rosiglitazone). The effects of PPARγ on cell migration and invasive activity were determined by wound healing assay and Matrigel invasive model in vitro, and in an orthotopic liver tumour metastatic model in mice.

RESULTS: Pronounced expression of PPARγ was demonstrated in HCC cells (MHCC97L, BEL-7404) treated with Ad-PPARγ and rosiglitazone and showed an additive effect. Activation of PPARγ by rosiglitazone significantly reduced the incidence and severity of lung metastasis in an orthotopic HCC mouse model. Key mechanisms underlying the effect of PPARγ in HCC include upregulation of cell adhesion genes, E-cadherin and SYK (spleen tyrosine kinase), extracellular matrix regulator tissue inhibitors of metalloproteinase (TIMP) 3, tumour suppressor gene retinoblastoma 1, and downregulation of pro-metastatic genes MMP9 (matrix metallopeptidase 9), MMP13, HPSE (heparanase), and Hepatocyte growth factor (HGF). Direct transcriptional regulation of TIMP3, MMP9, MMP13, and HPSE by PPARγ was shown by ChIP-PCR.

CONCLUSION: Peroxisome proliferator-activated receptor-gamma exerts an inhibitory effect on the invasive and metastatic potential of HCC in vitro and in vivo, and is thus a target for the prevention and treatment of HCC metastases.


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Keywords: PPARγ; hepatocellular carcinoma metastasis; heparanase; matrix metallopeptidase; tissue inhibitors of metalloproteinase

Despite significant advances in early detection and therapy, hepatocellular carcinoma (HCC) still remains the third leading cause of cancer-related deaths worldwide (Bosh et al, 2004). The high mortality rate of HCC is mainly attributable to late presentation at advanced stage, where curative surgical resection is no longer feasible or of limited efficacy (Mann et al, 2007). Tumour recurrence in HCC can occur as metastases, whereas more than 90% of HCC-related deaths are the result of secondary local or distant disease. However, efficacious or curative drug therapy for HCC and its metastases remains elusive.

Peroxisome proliferator-activated receptor-gamma (PPARγ) is a ligand-activated transcription factor that belongs to the nuclear hormone receptor super family; its roles include control of several biological processes related to growth, differentiation, cell cycle, and apoptosis (Koeffler 2003). Activation of PPARγ has been shown to inhibit proliferation in several cancers in vitro and in vivo (Koeffler 2003; Grommes et al, 2004). Our group has recently reported that PPARγ activation by its agonist (Yu et al, 2006) or ectopic expression of PPARγ by Ad-PPARγ transfection (Yu et al, 2010) inhibits HCC growth and progression by suppressing cell proliferation, inducing cell apoptosis, and causing cell cycle arrest (Yu et al, 2006, 2010). The PPARγ expression in HCC is significantly reduced in tumour tissues compared with surrounding non-tumourous liver, especially in poorly differentiated tumour than in well-differentiated tumour (Yu et al, 2006). Peroxisome proliferator-activated receptor-gamma also has a role in inhibiting tumour growth and metastatic spread in colon (Takano et al, 2008) and thyroid cancers (Ohta et al, 2001; Chen et al, 2006) as well as lung carcinoma (Panigrahy et al, 2002). However, PPARγ’s effect on invasive and metastatic potential of HCC has yet to be defined.

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†These authors contributed equally to this work.
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The aim of the present study was to determine the effect and the underlying molecular mechanism of PPAR\(\gamma\) on HCC cell migration and invasion using HCC cell lines and formation of distant metastases in vivo in an orthotopic murine liver tumour model.

**MATERIALS AND METHODS**

**Human HCC cell lines and culture**

The human HCC cell line MHCC97L, stably labelled with luciferase, was a gift from K Man, Department of Surgery, The University of Hong Kong (Man et al., 2010). BEL-7404 was obtained from the Institute of Biochemistry and Cell Biology (SIBS, Shanghai, China) (Chen et al., 1980). Cells were cultured in DMEM (Dulbecco’s modified Eagle medium) supplemented with 10% foetal bovine serum (Invitrogen, Carlsbad, CA, USA) and incubated at 37 °C and 5% CO\(_2\).

**Adenovirus-mediated PPAR\(\gamma\) gene transfer**

Recombinant adenovirus expressing the mouse PPAR\(\gamma\)-1 cDNA (Ad-PPAR\(\gamma\)) or E. coli \(\beta\)-galactosidase gene (Ad-LacZ, control adenovirus vector) (gift from JK Reddy, Department of Pathology, Feinberg School of Medicine, Northwestern University, Chicago) was propagated, isolated in human embryonic kidney 293 (HEK293) cells, then purified with Adeno-X Maxi Purification Kit (Clontech, Mountain View, CA, USA); the adenovirus with infectious titre range from 1.0 \(\times\) 10\(^{5}\) to 10\(^{6}\) pfu (plaque-forming unit) ml\(^{-1}\) was stored at −80 °C until use.

**RNA extraction, cDNA synthesis, and RT–PCR**

Total RNA was extracted from cell pellets by Trizol (Invitrogen) and reverse transcribed into cDNA using MultiScribe Reverse Transcriptase (Applied Biosystems). Samples were amplified using the ABI Prism 7700 Sequence Detection System (Applied Biosystems).

**Tumour cell migration assay**

Wound healing assay was performed for analysis of cell migration in vitro. Briefly, MHCC97L (5 \(\times\) 10\(^{5}\) cells per well) or BEL-7404 (5 \(\times\) 10\(^{5}\) cells per well) cells were seeded in 12-well plates and infected with Ad-LacZ (70 multiplicities of infection, MOI) or Ad-PPAR\(\gamma\) (70 MOI), and treated with or without rosiglitazone (50 \(\mu\)M) at 37°C until 90% confluent (Yu et al., 2010). Sterile tips were used to scratch cell layers, which were subsequently washed with PBS, and cultured with DMEM media and 1% FBS. Cells were photographed (phase-contrast microscope) at 0, 24, 36, and 48 h after incubation. The distance travelled by cells was measured between the two boundaries of an acellular area and results of the wound healing assay were expressed as a ratio to Ad-LacZ-treated cells. The distance travelled by cells was measured between the two boundaries of an acellular area and results of the wound healing assay were expressed as a ratio to Ad-LacZ-treated cells.

**RNA extraction, cDNA synthesis, and RT–PCR**

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**Chromatin immunoprecipitation (ChIP) assay**

Chromatin immunoprecipitation analysis was performed using the human HCC cell line MHCC97L, which has metastatic potential to lung (Man et al., 2010). MHCC97L cells (2 \(\times\) 10\(^{6}\) cells in 0.1 ml PBS) were infected subcutaneously into the left dorsal flank of 4-week-old male Balb/c nude mice. Subcutaneous tumours were harvested once the subcutaneous tumours reached about 10 mm\(^3\) and cut into 1.0 mm\(^3\) pieces. One piece of tumour was then implanted into the left liver lobes in a separate group of nude mice (6-week-old) (10 per group) (Man et al., 2010). After tumour implantation, mice were randomly treated with or without rosiglitazone (200 p.p.m. in chow), a dosage was selected base on our previous experiments (Yu et al., 2010). Liver tumour growth and lung metastasis were monitored by Xenogen IVIS-200, an optical in vivo imaging system (Caliper Life Science, Hopkinton, MA, USA) weekly. Mice were euthanised at week 7 after tumour implantation (Man et al., 2010), and tumours and lung nodules were analysed histologically. Signal intensity of tumours detected by Xenogen IVIS was expressed as Radiant Efficiency (radiance/illumination power density = ps\(^{-1}\) cm\(^{-2}\) sr\(^{-1}\)). All experimental procedures were approved by the Animal Ethics Committee of the Chinese University of Hong Kong.

**Characterisation of PPAR\(\gamma\)-binding region and semi-quantititation of ChIP-PCR**

To evaluate the direct modulation of transcriptional activity on the promoters of metastasis-related genes by PPAR\(\gamma\), the Genomatix tool, ModellInspector, was used to scan all the known PPAR\(\gamma\)-binding regions linked to metastasis-related genes. Human promoter regions (size between 2 kb) in the Genomatix promoter database were scanned. Matched binding sites were selected with core and matrix similarity of > 0.8 (http://www.genomatix.de/ online_help/help_matinspector/matinspector_help.html). The predicted PPAR\(\gamma\)-binding sites on the promoter of target genes were validated by ChIP-PCR (specific primers listed in Table 1).
**Western blot analysis**

Total protein was extracted from cell pellets and protein concentration measured by Bradford assay (Bio-Rad, Hercules, CA, USA). A total of 30 mg of protein was separated by 12% sodium dodecyl sulphate–polyacrylamide gel electrophoresis, then transferred onto equilibrated polyvinylidene difluoride membrane (Amersham Biosciences, Buckinghamshire, UK). Membranes were probed with primary antibodies for tissue inhibitors of metalloproteinase 3 (TIMP3), heparanase (HPSE), and E-cadherin (Santa Cruz Biotechnology).

**Statistical analysis**

Data were presented as means ± s.d. Multiple group comparisons were analysed by one way ANOVA after Bonferroni’s correction. Non-parametric data between two groups was computed by Chi-square test or Fisher Exact test. The difference for two different groups was determined by Student’s t test. A P-value of less than 0.05 was considered statistically significant.

**RESULTS**

**Peroxisome proliferator-activated receptor-gamma suppresses HCC cell migration and invasiveness of MHCC97L and BEL-7404 human HCC cell lines**

The MHCC97L and BEL-7404 cells were infected with Ad-PPARγ or Ad-LacZ control in the presence or absence of rosiglitazone for 48 h and induction of PPARγ was confirmed by Western blot (Figure 1A). Enhanced PPARγ expression by Ad-PPARγ or rosiglitazone, markedly slowed cell migration scratchy ‘wound’ at edges of MHCC97L and BEL-7404 HCC cells (Figure 1B). Quantitative analyses at 36 h confirmed a significant reduction in wound closure in Ad-PPARγ or rosiglitazone-treated cells compared with Ad-LacZ-infected control cells (Figure 1C). There appeared to be an additive effect of Ad-PPARγ plus rosiglitazone compared with Ad-PPARγ or rosiglitazone only in BEL-7404 cell line (Figure 1C).

To study the effect of PPARγ conferred on the invasiveness of HCC, MHCC97L and BEL-7404 cells were infected with Ad-PPARγ or treated with rosiglitazone using a Matrigel model (Figure 2A). *In vitro* invasively growing HCC cells were significantly impaired by up to 60% when infected with Ad-PPARγ or primed by rosiglitazone at 48 h (Figure 2B). Moreover, the combination of Ad-PPARγ and rosiglitazone incrementally suppressed cell invasion compared with Ad-PPARγ or rosiglitazone alone (Figure 2B).

**Activation of PPARγ by rosiglitazone inhibits HCC metastases to the lung in vivo**

In light of the observed anti-migration and anti-invasion effects of PPARγ on HCC cell lines *in vitro*, we tested whether activation of PPARγ by rosiglitazone could alter metastatic potential of MHCC97L *in vivo* in an orthotopic metastasis mouse model, where subcutaneously grown tumours derived from MHCC97L cells expressing luciferase were implanted into the livers of nude mice; small successful transplantation of tumours was confirmed by xenogen imaging 2 weeks after surgery (Figure 3A). Mice were randomly treated with rosiglitazone or vehicle for 7 weeks and then re-imaged *in vivo*. By week 7, 87.5% (7 out of 8) of the vehicle-treated control mice demonstrated lung metastases after orthotopic implantation. In contrast, only 33% (3 out of 9) of rosiglitazone-treated animals developed lung metastases (P < 0.05) (Figure 3B). The luciferase signals emanating from the lungs originally from the MHCC97L cells expressing luciferase in the rosiglitazone-treated mice were significantly lower than in the vehicle-treated group (1.7 × 10^5 vs 3.14 × 10^6 p s ^−1 cm ^−2 sr ^−1, P < 0.05) (Figure 3C). Subsequent histology confirmed that lung nodules were secondary metastatic deposits from liver (Figure 3B). Collectively, these results provide clear evidence that PPARγ activation inhibits lung metastasis in an orthotopic HCC model *in vivo*.

**Peroxisome proliferator-activated receptor-gamma modulates the expression profiles of metastasis-related genes in MHCC97L cells**

To elucidate the molecular mechanisms underlying the inhibitory effect of PPARγ on HCC cell invasiveness, gene expression profiles in Ad-PPARγ- or Ad-LacZ-infected MHCC97L were analysed using a human tumour metastasis pathway PCR array. When compared with control Ad-LacZ-infected cells, PPARγ altered downstream targets involved in cell adhesion, extracellular matrix (ECM) proteins, cell growth, and cell motility (Table 2), all of which are critical to the regulation of cancer cell invasiveness and metastasis. Peroxisome proliferator-activated receptor-gamma exerted its anti-metastatic effects by increasing the expression of cell adhesion genes, E-cadherin (5.2-fold), spleen tyrosine kinase (SYK) (1.7-fold), and ECM regulator metallopeptidase inhibitor 3 (TIMP3) (7.2-fold), a physiological inhibitor of matrix metallopeptidases (MMPs). The PPARγ also suppressed expression of pro-metastatic genes, such as MMP9 (~1.7-fold), MMP13 (~2.0-fold), HPSE (~6.5-fold), and significantly diminished hepatocyte growth factor (HGF) (~2.2-fold), a cellular growth and motility regulator. Further, PPARγ-induced retinoblastoma 1 (RBI) expression, a potent tumour suppressor gene by four-fold (Table 2).

To validate these changes on expression profiling, western blot and semi-quantitative RT–PCR were performed on MHCC97L and BEL-7404 cells infected with Ad-PPARγ or Ad-LacZ in the presence or absence of rosiglitazone. Ectopic expression of PPARγ by Ad-PPARγ or activation of PPARγ by rosiglitazone, increased E-cadherin and TIMP3 protein expression with a concomitant diminution of HPSE in both HCC cell lines (Figure 4A1). Combination of Ad-PPARγ and rosiglitazone exerted an additive effect on the induction of E-cadherin and TIMP3 and suppression of PPARγ expression compared with Ad-PPARγ alone (Figure 4A2).

**Table 1 Primer sequences for semi-quantitative PCR detection**

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<td>MMP13 promoter 2</td>
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<td>TCCCCGCGAGAAATGAATTG</td>
</tr>
<tr>
<td>HPSE promoter</td>
<td>GGTTGGTTGATCTCCTTCCA</td>
<td>CTTTCTCTCCACCTGATCAG</td>
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</table>

**Gene Forward primer (5’ to 3’) Reverse primer (5’ to 3’)**

**Results**

**Activation of PPARγ by rosiglitazone inhibits HCC metastases to the lung in vivo**

In light of the observed anti-migration and anti-invasion effects of PPARγ on HCC cell lines *in vitro*, we tested whether activation of PPARγ by rosiglitazone could alter metastatic potential of MHCC97L *in vivo* in an orthotopic metastasis mouse model, where subcutaneously grown tumours derived from MHCC97L cells expressing luciferase were implanted into the livers of nude mice; small successful transplantation of tumours was confirmed by xenogen imaging 2 weeks after surgery (Figure 3A). Mice were randomly treated with rosiglitazone or vehicle for 7 weeks and then re-imaged *in vivo*. By week 7, 87.5% (7 out of 8) of the vehicle-treated control mice demonstrated lung metastases after orthotopic implantation. In contrast, only 33% (3 out of 9) of rosiglitazone-treated animals developed lung metastases (P < 0.05) (Figure 3B). The luciferase signals emanating from the lungs originally from the MHCC97L cells expressing luciferase in the rosiglitazone-treated mice were significantly lower than in the vehicle-treated group (1.7 × 10^5 vs 3.14 × 10^6 p s ^−1 cm ^−2 sr ^−1, P < 0.05) (Figure 3C). Subsequent histology confirmed that lung nodules were secondary metastatic deposits from liver (Figure 3B). Collectively, these results provide clear evidence that PPARγ activation inhibits lung metastasis in an orthotopic HCC model *in vivo*.

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of HPSE (Figure 4A1). A similar enhanced suppressive effect of the combination of Ad-PPAR\(_g\) and rosiglitazone was observed on MMP9 and MMP13 mRNA expression by RT–PCR (Figure 4A2).

**Direct transcriptional regulation of TIMP3, MMP9, MMP13, and HPSE by PPAR\(_g\)**

To further determine whether PPAR\(_g\)-mediated downstream gene expression changes were associated with direct promoter binding, PPAR\(_g\)-binding sites in the promoters of identified targets, TIMP3, MMP9, MMP13, and HPSE, were mined using ModellInspector software (Genomatix Software GmbH, Munich, Germany). Chromatin immunoprecipitation assay with PPAR\(_g\) antibody was performed in MHCC97L cells followed by PCR confirmation (Figure 4B). By ChIP-PCR assay, PPAR\(_g\) binds to the promoters of TIMP3, MMP9, MMP13, and HPSE in MHCC97L cells (Figure 4B). These findings suggest that TIMP3, MMP9, MMP13, and HPSE are direct targets of PPAR\(_g\) in liver cancer cells.
DISCUSSION

The significance of PPARγ on the process of metastasis is not well studied in contrast to the effect of PPARγ on tumour growth. In this study, we show that ectopic expression of PPARγ by Ad-PPARγ or its agonist, rosiglitazone, in two HCC cell lines (MHCC97L, BEL-7404) inhibits metastatic activity in vitro, in particular in wound healing, cell migration, and invasion. Moreover, the combination of Ad-PPARγ and rosiglitazone results in an enhanced anti-metastatic effect. Together, these results indicate that restoration of PPARγ allow for interactions with endogenous or exogenous agonists that activate the anti-metastatic processes associated with PPARγ. The role of PPARγ in inhibiting HCC metastases in vitro was further elucidated in an orthotopic HCC xenograft model. In this murine model, activation of PPARγ suppressed HCC lung metastasis in nude mice. Surgical resection or liver transplantation is the curative mainstays of HCC management, however, post-operative recurrence predominantly related to metastasis remains a challenge. Our finding was supported by recent studies, which indicated that activation of PPARγ inhibits metastasis of lung cancer (Reka et al, 2010) and non-small cell lung cancer (Choudhary et al, 2010) in vitro and in vivo. Collectively, our finding that activation of PPARγ by rosiglitazone significantly suppresses metastatic potential could have a beneficial impact on the clinical practice and in adjuvant therapy of HCC after surgical resection and transplantation.

The molecular mechanisms by which PPARγ exerts its anti-invasive and anti-metastases functions in HCC have not yet been defined. To identify key regulators of PPARγ-mediated anti-metastatic effect in HCC, we utilised cDNA microarray and ChIP-PCR to study MHCC97L cells infected with Ad-PPARγ; genes that were significantly altered in expression levels were then validated by RT–PCR and immunoblotting. We report that the suppression of cell invasion and migration mediated by PPARγ was mediated via downregulation of MMPs (MMP9, MMP13), increased expression of TIMP3 and E-cadherin. Of which, MMP9 and MMP13 are members of extracellular proteinases with key functions in the formation and remodelling of tumour invasion (Khasigov et al, 2003). Matrix metallopeptidase 9 has been described to promote tumour malignant progression, invasion, and metastatic spread by activating tumour growth factor-β (TGF-β) (Yu and Stamenkovic, 2000). Whereas, MMP13 is a central role in the modulation of other MMPs (Leeman et al, 2002) such as stimulating pro-MMP9 activation (Knäuper et al, 1997). Elevated MMP13 has been shown in various human malignancies including breast cancer, colorectal neoplasms, melanoma, and squamous head and neck tumours,
particularly at the invading edge of such cancers; increased MMP13 expression was associated with poor prognosis, tumour aggressiveness, and metastases (Leeman et al., 2002; Luukkaa et al., 2006; Kondratiev et al., 2008; Chang et al., 2009).

Tissue inhibitors of metallo proteinase 3 is an important endogenous inhibitor of MMPs. Upregulation of TIMP3 expression by PPARγ suggested its suppressive functions in tumour invasion and metastasis via suppressing of MMPs (Bachman et al., 1999;...
Table 2. Effect of PPARγ on its downstream gene expression profiles of cancer metastasis pathways in HCC cell lines.

<table>
<thead>
<tr>
<th>Full name</th>
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Figure 4. (A1) Western blots were performed to confirm the downstream gene expression regulated by PPARγ in MHCC97L and BEL-7404. GAPDH was used as an internal control. The relevant band densitometry analysis was performed and displayed in the lower panel. (A2) Semi-quantitative RT–PCR and real-time quantitative PCR analyses were performed to validate the candidate genes expression. The data are expressed as means ± s.d., *P<0.001, compared with the Ad-LacZ. (B) Chromatin Immunoprecipitation (ChIP-qPCR) was performed to identify direct targets of PPARγ protein. Input (2%) represents the genomic DNA. (C) Schematic diagram for the mechanisms of anti-metastasis function of PPARγ derived from cDNA array, western blot, and ChIP-qPCR. PPARγ-mediated suppression of cell invasion and migration was associated with several biological effects: (1) Directly upregulating MMP9 (MMP9, MMP13) and downregulating their inhibitor (TIMP3) by direct binding to the promoter of each targets, and subsequent modulation of cell–cell adhesion molecule E-cadherin, which in turn inhibited the cell–cell adhesion and extracellular matrix (ECM) turnover; (2) Directly inhibiting the transcription of HPSE gene, which contributes to the suppression of ECM turnover and distant metastasis; (3) Downregulation of HGF to suppress the invasive potential; (4) Upregulation of tumour suppressors (RB1 and SYK), which protects against tumourigenesis through suppressing ECM turnover, cell proliferation, and causing cell cycle arrest. Blue colour indicates the direct targets, and green colour indicates second targets of PPARγ. The colour reproduction of this figure is available at the British Journal of Cancer online.
Tumour suppressor genes, RB1 and SYK, have been reported to be dysfunctional in several cancers where aberrant expression levels appear to correlate with poor prognosis (Bailet et al., 2009; Kouraklis et al., 2009). In contrast, overexpression of RB1 or SYK in several cancer cell types can inhibit tumour growth and reduce metastasis in mouse xenografts (Valente et al., 1996; Coopman et al., 2000). Spleen tyrosine kinase also inhibits the motility of human breast cancer cells (Zhang et al., 2009) and promotes the formation of cell–cell contact through mediation of E-cadherin activity (Larive et al., 2009). In the present study, we describe a near two-fold induction of SYK and impressive upregulation by four-fold of RB1 by PPARg; in keeping with its proposed anti-metastatic and -invasive effects.

In conclusion, activation of PPARg has demonstrated efficacy in suppressing HCC cell migration and invasion in vitro, and in inhibiting distant metastases from liver in an orthotopic HCC model in vivo. Peroxisome proliferator-activated receptor-gamma may provide a potential target for the prevention and treatment of metastatic HCC.

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Conflict of interest
The authors declare no conflict of interest.

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Bachman KE, Herman JG, Corn PG, Merlo A, Costello JF, Cavenee WK, Bos FX, Ribes J, Montesano R, Bottaro DP (1998) Induction of mesenchymal transition, a plausible mechanism of driving cancer cell migration (Cowden Dahl et al., 2008). Cell–cell adhesion molecule, E-cadherin, is also a key component of epithelial adherent junctions. Loss of E-cadherin expression is a hallmark of epithelial-to-mesenchymal transition, a plausible mechanism of driving cancer progression and metastasis (Onder et al., 2008; Makrilia et al., 2009). We demonstrate further that PPARg inhibits transcription of HPSE by direct binding to its promoter. Upregulation of HPSE has been associated with increased lymph node, as well as distant metastases (Sanderson et al., 2005) and reduced post-operative survival of cancer patients (Sato et al., 2009).

The anti-metastasis function of PPARg in vitro also appeared to be associated with the downregulation of HGF by cDNA array. HGF is known to act as a multifunctional growth factor and is upregulated in many human cancers including HCC (Ljubimova et al., 1997; Maulik et al., 2002). Hepocyte growth factor drives epithelial cells to undergo EMT and can downregulate EMT-associated E-cadherin expression in murine liver tumour cells (Ding et al., 2010). Also, HGF has been reported to stimulate MMP9 expression (Mizuno et al., 2005) and increase MMP3 promoter activity in HCC (Ozaki et al., 2003), which can lead to increased cancer cell invasiveness (Reboul et al., 2001; Wang et al., 2007; Lee et al., 2010). Notably, HGF activity may be mediated by MMPs (including MMP9 and MMP13) in the extracellular matrix, and thus induce HCC cells to proliferate and invade (Monvoisin et al., 2002; Mohammed et al., 2005). In contrast, HGF may down-modulate TIMP-3 expression resulting in increased MMP accumulation, hence contributing to the invasiveness and aggressiveness in cancer cells (Castagnino et al., 1998). Thus, the anti-metastasis effect of PPARg in HCC may be in part related to the inhibition of HGF expression (Figure 4C).

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