Biological Function and Prognostic Significance of Peroxisome Proliferator-Activated Receptor δ in Rectal Cancer

Lie Yang¹,², Hong Zhang³, Zong-Guang Zhou¹, Hui Yan¹, G. Adell⁴, and Xiao-Feng Sun¹,²

Abstract

Purpose: To investigate the expression significance of PPAR β/δ in relation to radiotherapy (RT), clinicopathologic, and prognostic variables of rectal cancer patients.

Experimental Design: We included 141 primary rectal cancer patients who participated in a Swedish clinical trial of preoperative RT. Tissue microarray samples from the excised rectal cancers and the adjacent or distant normal mucosa and lymph node metastases were stained with PPAR δ antibody. Survival probability was computed by the Kaplan–Meier method and Cox regression model. The proliferation of colon cancer cell lines KM12C, KM12SM, and KM12L4a was assayed after PPAR δ knockdown.

Results: PPAR δ was increased from adjacent or distant normal mucosa to primary cancers, whereas it decreased from primary cancers to lymph node metastases. After RT, PPAR δ was increased in normal mucosa, whereas it decreased in primary cancers and lymph node metastases. In primary cancers, the high expression of PPAR δ was related to higher frequency of stage I cases, lower lymph node metastasis rate, and low expression of Ki-67 in the unirradiated cases, and related to favorable survival in the cases either with or without RT. The proliferation of the KM12C, KM12SM, or KM12L4a cells was significantly accelerated after PPAR δ knockdown.

Conclusions: RT decreases the PPAR δ expression in primary rectal cancers and lymph node metastases. PPAR δ is related to the early development of rectal cancer and inhibits the proliferation of colorectal cancer cells. Increase of PPAR δ predicts favorable survival in the rectal cancer patients either with or without preoperative RT. Clin Cancer Res; 17(11); 3760–70. ©2011 AACR.

Introduction

PPARs are ligand-activated transcription factors that are members of the nuclear hormone receptor superfamily, containing 3 isoforms, α, β/δ, and γ (1). PPAR α and γ have been well characterized as central regulators of lipid and glucose homeostasis (2), but less is known about the biological roles of PPAR δ. Available studies indicate that the major functions of PPAR δ are associated with its regulator roles in multiple biological processes such as lipid metabolism and energy homeostasis, embryo implantation, wound healing, inflammatory response, cell proliferation, and carcinogenesis (3, 4). PPAR δ therefore represents a potential drug target for the treatment of some diseases, such as obesity and metabolic syndrome, which has led to the development of several synthetic drug agonists (5).

Recent studies have implicated PPAR δ in the pathogenesis of colorectal cancer. Inactivation of adenomatous polyposis coli (APC) upregulates PPAR δ expression in colorectal cancer cells (6). PPAR δ is overexpressed in both human and rat colorectal cancers (7–9). Loss of PPAR δ expression in colon cancer cells results in decreased growth of xenografts (10). Ligand activation of PPAR δ potentiates colon tumorigenesis in mice (4, 11, 12). These studies support a promotive role of PPAR δ in colorectal carcinogenesis, but other studies conflict with these reports. Targeted deletion of APC alleles decreases PPAR δ expression in mouse intestine (13). PPAR δ expression is decreased in human colon cancers as compared with the matched normal mucosa (14, 15). Ligand activation of PPAR δ attenuates colon tumorigenesis in mice (4, 16, 17). Experiments that examined polyp formation in PPAR δ–null APCmin mice show either no effect (18), or paradoxically, an increase in polyp number and size compared with wild-type mice (13, 16, 17, 19). Our recent study shows that PPAR δ may facilitate the differentiation of colon cancer cells (20). Collectively, the role of PPAR δ in colorectal carcinogenesis remains highly controversial.

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Note: Supplementary data for this article are available at Clinical Cancer Research Online (http://clincancerres.aacrjournals.org/).

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doi: 10.1158/1078-0432.CCR-10-2779

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Translational Relevance

PPAR δ/β has been implicated in the pathogenesis of colorectal cancer, but its exact role remains controversial. This study shows that PPAR δ is related to the early development of rectal cancer, and involved in the inhibition of the proliferation of colorectal cancer cells. These findings support the role of PPAR δ as a tumor suppressor and thus may be a good therapeutic target for colorectal cancer. In this study, we show that pre-operative radiotherapy (RT) increases the PPAR δ expression in normal rectal mucosa, but decreases it in primary rectal cancers and lymph node metastases. Therefore, application of a PPAR δ agonist or regulator together with preoperative RT in PPAR δ—low expression tumors may enhance the efficacy of RT. We show that RT decreases the PPAR δ expression in primary rectal cancers and lymph node metastases, and the increase of PPAR δ in primary rectal cancers is related to favorable survival of the patients, indicating that PPAR δ is a useful prognostic factor for rectal cancer patients.

Current studies on the role of PPAR δ in colorectal cancer are mostly based on cell lines or animal models, lacking clinical study with long-term observation data of patients. Little is known about the relationship of PPAR δ expression with radiotherapy (RT), clinicopathologic, and prognostic factors in colorectal cancer patients. In the present study, we examined the expression of PPAR δ in human rectal cancers and the matched adjacent or distant normal mucosa and lymph node metastases, with or without preoperative RT, by using immunohistochemical assay (IHC). We analyzed the relationship between PPAR δ expression and preoperative RT, clinicopathologic factors including survival of rectal cancer patients, and the proliferation of 3 colon cancer cell lines with different metastatic potentials. To our knowledge, this is the first study correlating PPAR δ expression with RT and prognosis in the rectal cancer patients who participated in a Swedish clinical trial of preoperative RT.

Materials and Methods

Patients
This study included 141 primary rectal cancer patients from the Southeast Swedish Health Care region who participated in a randomized Swedish rectal cancer trial of preoperative RT between 1987 and 1990 (Swedish Rectal Cancer Trial, 1997; ref. 21). The written informed consent was given by each participant. Seventy-seven patients received tumor resection alone, and 64 received preoperative RT followed by tumor resection. Locally curative resection was carried out in all patients. RT was given with 25 Gy in 5 fractions over a median of 6 days (range, 5–12 days), delivered with 6–10 MV photons. Surgery was then carried out in a median of 3 days (range, 1–13 days) after RT. None of the patients received adjuvant chemotherapy. Table 1 presents the characteristics of the patients and tumors including gender, age, TNM stage [classified according to the Cancer Staging Manual of American Joint Committee on Cancer, 7th edition, 2010 (22)], grade of differentiation, number of other tumors, surgical type, resection margin, and mean distance to the anal verge. There was no statistical difference between the non-RT and RT groups regarding these characteristics (P > 0.05, Table 1).

Follow-up was carried out by matching all patients against the Swedish Cancer Register and the Cause of Death Register until 2004. The median follow-up period was 84 months (range, 0–193 months). Information about local or distant recurrence, disease-free survival (DFS) and overall survival (OS) were obtained from patient medical records. We conducted our study after approval by the Institutional Review Board of the Linköping University, Sweden.

Tissue collection and tissue microarray
Specimens were collected from primary rectal cancers (n = 141) and the matched adjacent normal mucosa (n = 81), distant normal mucosa (n = 115), and metastases in the regional lymph nodes (n = 36). Distant normal mucosa was taken from proximal or distal margin (4–35 cm from the primary tumor) of the resected rectum, and adjacent

<table>
<thead>
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<td>Characteristics</td>
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<td>To anal verge, cm</td>
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²Pearson χ² test.

bThere had been previous colorectal cancer and/or other types of tumors before the present rectal cancer.

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normal mucosa was from mucosa adjacent to the primary tumor, both were histologically free from pretumor and tumor. All the specimens including normal mucosa, primary cancers and lymph node metastases were paraffin embedded for tissue microarray (TMA).

Representative paraffin-embedded tissue blocks were selected as donor blocks for the TMA. Three morphologically representative regions were chosen in each block and 3 cylindrical core tissue specimens (0.6 mm in diameter) were taken from these areas, inserted in a recipient paraffin block. Sections from this block were cut into 5 μm chips using a microtome, mounted on microscope slides. The tissue microarrays were constructed using a manual arrayer (Beecher Inc.).

**Immunohistochemical assay**

IHC staining for PPAR δ expression was done on 5-μm TMA sections from paraffin-embedded surgical specimens. The sections were deparaffinized in xylene, rehydrated in ethanol, and rinsed in distilled water. Masked epitope retrieval was done by boiling the sections in 1× DIVA buffer (Biocare Medical) at 125°C for 30 seconds in a high-pressure cooker. Then, the sections were stored at room temperature for 20 minutes, followed by rinsing with PBS containing 0.5% bovine serum albumin. The sections were incubated in 3% H2O2-methanol for 5 minutes to block the activity of endogenous peroxidase. After being washed in PBS, the sections were incubated with power block (Spring Bioscience) for 10 minutes to reduce nonspecific background staining. The sections were incubated with the PPAR δ polyclonal rabbit anti-Human IgG (ARP37889, against N terminal; Aviva Systems Biology) in a 1:400 dilution with 5% skimmed milk PBS buffer over night. After being washed in PBS, the sections were incubated with a secondary antibody, Envision System Labelled Polymer-HRP Anti-Rabbit (Dakocytomation) for 25 minutes. The sections were rinsed in PBS before reacting with 3, 3-diaminobenzidine tetrahydrochloride (ChemMate) for 8 minutes to produce coloration. Finally, the sections were counterstained with hematoxylin, followed by dehydration with ethanol. All steps were done at room temperature. Sections known to show positive staining for PPAR δ were included in each run, receiving either the primary antibody or PBS, as positive or negative controls. In all staining procedures, the positive controls showed clear staining, whereas there was no staining in the negative controls.

**Measurements of PPAR δ expression by IHC**

The IHC slides were examined independently in a blinded fashion by 2 investigators (L. Yang and H. Zhang) without knowledge of clinicopathologic or biological information. Each investigator estimated the proportion of cells stained and the intensity of staining in the whole section. The intensity in epithelial cells or tumor cells was scored as 0 (negative staining), 1 (weak staining exhibited as light yellow), 2 (moderate staining exhibited as yellow brown), and 3 (strong staining exhibited as brown). The proportion of cells stained was accessed using a 5 scoring system: 0 (no positive cells), 1 (<10% positive cells), 2 (10%-40% positive cells), 3 (40%-70% positive cells), and 4 (>70% positive cells). The percentage of cells at each intensity was multiplied by the corresponding intensity value to obtain an immunostaining score ranging from 0 to 12. The scores were combined to obtain an overall mean score. The cutoff value for expression levels from weak to strong was based on a measurement of heterogeneity using log-rank test statistical analysis with respect to OS. Using this assessment system, optimal cutoff values were identified by the mean score as follows: 0 (negative), 1–3 (weak), 4–9 (moderate), and 9–12 (strong). If there was a discrepancy in individual scores, then both investigators reevaluated the slides together to reach a consensus before combining the individual scores. To avoid an artificial effect, the cells on the margins of the sections and in areas with poor morphology were not counted. For statistical analysis, the negative or weakly stained cases were considered low expression and the moderate or strongly stained cases were considered high expression.

**Evaluation of apoptosis, proliferation, and phosphatase of regenerating liver-3, cyclooxygenase-2, and survivin**

The data for apoptosis, detected by terminal deoxynucleotidetransferase–mediated deoxyuridine triphosphate—biotin nick end labeling (TUNEL) assay (n = 136), proliferation (n = 126), phosphatase of regenerating liver-3 (PRL-3; n = 125), cyclooxygenase-2 (COX-2; n = 138), and survivin (n = 98) of primary rectal cancers determined by IHC, were taken from previous studies conducted with the same cases used in the present study at our laboratory (23–27). The percentage of apoptotic cancer cells was determined by counting approximately 1,000 tumor cells. Cases were considered as negative if apoptotic cells constitute less than 5% of tumor cells. Proliferation in the cancer cells was measured using IHC for Ki-67 as an indicator. Low proliferation was defined in sections where less than 30% of cancer cells expressed Ki-67 and high proliferation in section where Ki-67 was expressed in 30% or more of cancer cells.

**Western blot assay**

Total proteins from the normal rectal mucosa samples, primary rectal cancers, or lymph node metastases were extracted using 1× RIPA lysis buffer (Santa Cruz) and the concentrations were assayed by the Bicinchoninic acid Protein Assay (Pierce). Each 20-μg aliquot of total proteins was loaded in duplicate in 10% SDS-PAGE gel, and then transferred onto a 0.2 μm polyvinylidene difluoride membrane (Bio-Rad). After completing protein transfer, the membrane was blocked in 5% (w/v) skimmed milk in PBS and incubated overnight with the rabbit polyclonal antibody (IgG) against PPAR δ (0.3 μg/mL; Aviva Systems Biology), COX-2 (0.5 μg/mL; catalogue no. ab52237, Abcam), or survivin (1.0 μg/mL; catalogue no. ab8228, Abcam), respectively. The blots were detected by the
secondary antibody, horseradish peroxidase (HRP)-linked polyclonal goat anti-rabbit IgG (0.1 μg/mL; DakoCytomation) and visualized with Amersham ECL plus Western blot detection system (Amersham Biosciences/GE Healthcare). Protein expression was normalized to the level of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) detected by the rabbit monoclonal antibody (HRP conjugated, 0.04 μg/mL; Cell Signaling).

Cell culture
Three human colon cancer cell lines, KM12C, KM12SM, and KM12L4a (from Prof. I.J. Fidler, Anderson Cancer Center, Houston, TX), untreated or treated by lentivirus targeting PPAR δ gene (Lenti-PPAR δ) or by lentivirus without RNA interference effect (Lenti-control) from our previous study (20), were used. All the cell lines were maintained in Eagle's minimal essential medium (MEM) with Earle's salts, l-glutamine and nonessential amino acids (Sigma-Aldrich), supplemented with 1.5% NaHCO3, 1 mm Na-Pyruvate (Invitrogen), 1× MEM Vitamin Solution (Invitrogen), 1% penicillin–streptomycin (Invitrogen), 1 μg/mL Puromycin (only for the treated cells to maintain the purity) and 10% FBS (Invitrogen). The expression of PPAR δ had been stably silenced in the cells treated by Lenti-PPAR δ while remaining unaffected in the Lenti-control treated cells, as confirmed in our previous study (ref. 20; see Supplementary Data). Of the 3 cell lines, KM12C has poor metastatic potential, whereas KM12SM and KM12L4a have high metastatic potentials (28).

Cell proliferation assay
The effect of PPAR δ knockdown on the proliferation of KM12C, KM12SM, and KM12L4a cells was assayed with MTT (methyl thiazolyl tetrazolium). The log-phase cells, untreated or treated with Lenti-PPAR δ or Lenti-control, were seeded in 96-well plates (4 x 103/well) for 24 to 96 hours. Each group was analyzed every 24 hours in triplicate by the following method: each well had 20 μL MTT (5 mg/mL; Sigma-Aldrich) added and were incubated for a further 4 hours at 37°C; subsequently the formazan crystals were solubilized with 150 μL dimethyl sulfoxide (DMSO; Sigma-Aldrich). The absorbance (A) value was measured at 570 nm wavelength on an automatic Microplate Reader (model 550; Bio-Rad) with DMSO as the blank. The growth curve of cells was drawn using growth date as x-axis and A value as y-axis.

Statistical analysis
All statistical analyses were carried out using STATISTICA software package (version 7.0; STATSOFT Inc., Tulsa, OK). McNemar’s or Pearson χ2 methods were used to test the significance of the differences in PPAR δ expression between the adjacent or distant normal mucosa, primary cancers and lymph node metastases, and the association of PPAR δ expression with clinicopathologic variables. The relationship between PPAR δ expression and survival, local recurrence or distant recurrence was tested using Kaplan–Meier analysis (Log rank test) and Cox proportional hazards regression analysis (likelihood ratio test). ANOVA and post hoc multiple comparison (LSD, 2-tailed) was applied to test the quantitative analysis in Western blot assay and the difference of absorbance values in MTT assay. The test was 2-sided and a P value of less than 0.05 was considered statistically significant.

Results
PPAR δ was increased from normal mucosa to primary cancers, whereas it decreased from primary cancers to lymph node metastases
By IHC, PPAR δ was detected predominantly in the cytoplasm of epithelial cells in normal mucosa, and tumor cells of primary cancers and lymph node metastases, with a little staining in the nuclei (Fig. 1A). For the further analyses of this study, only the staining of cytoplasmic PPAR δ was measured and presented. In the cases without RT, the frequency of high expression (moderate or strong staining) of PPAR δ was significantly increased from adjacent normal mucosa (2%, 1 of 44) to distant normal mucosa (10%, 6 of 61) to primary cancers (61%, 47 of 77), whereas expression decreased from primary cancers to lymph node metastases (32%, 9 of 28; P = 0.008; Fig. 1B). There was not a significant difference between adjacent and distant normal mucosa (P = 0.23).

The specificity of the PPAR δ antibody used in IHC was examined by Western blot. As shown in Figure 1C, each of the electrophoretic lanes had a clear band at the expected position for PPAR δ (48 kDa) as well as the band for GAPDH (37 kDa). Quantitative analysis showed that the expression of PPAR δ protein normalized by GAPDH was significantly increased by 64.0% ± 2.6% from normal mucosa to primary cancers, and decreased by 10.8% ± 1.3% from primary cancers to lymph node metastases (P = 0.036; Fig. 1C), which was identical to the results of IHC.

PPAR δ was increased in normal mucosa, whereas it decreased in primary cancers and lymph node metastases after RT
Compared with the unirradiated cases, the frequency of highly expressed PPAR δ in irradiated cases was significantly increased in both adjacent normal mucosa (2% vs. 16%; P < 0.001) and distant normal mucosa (10% vs. 33%; P = 0.014), whereas it decreased in primary cancers (61% vs. 41%; P = 0.031) and in lymph node metastases (32% vs. 22%; P = 0.043; Fig. 2).
the relationship between PPAR δ expression and TNM stage ($P = 0.43$) or lymph node metastasis disappeared ($P = 0.27$; data not shown).

In the cases either with or without RT, the high expression of PPAR δ in primary cancers was related to low expression (<30% of cancer cells) of Ki-67 ($P = 0.016$, 0.025; Fig. 3C). The non-RT cancers with better differentiation showed a significantly higher frequency of highly expressed PPAR δ than did those with poor differentiation ($P = 0.004$; data shown in our previous study [20]). There was no significant relationship between PPAR δ expression and age, gender, growth pattern, apoptosis, PRL-3, COX-2,
or survivin ($P > 0.05$; data not shown). The Western blot assay confirmed that the expression of PPAR $\delta$ was not significantly different from the expression of COX-2 or survivin in primary rectal cancers ($P = 0.43$; see Supplementary Data).

High expression of PPAR $\delta$ in primary cancers was related to favorable survival, without relation to local/distant recurrence

Kaplan–Meier analysis showed that the high expression of PPAR $\delta$ in the unirradiated primary cancers was related to better DFS ($P = 0.003$; Fig. 4A) but not OS ($P = 0.13$; data not shown). In the irradiated cases, the high expression of PPAR $\delta$ was related to better OS ($P = 0.032$; Fig. 4B) but not DFS ($P = 0.10$; data not shown). In multivariate analysis, the above significances still remained, independent of age, gender, tumor differentiation, growth pattern, and TNM stage (Table 2). As shown in Table 2, the unirradiated patients with low expression of PPAR $\delta$ were 3.6 times more likely to die of rectal cancer than those with high expression of PPAR $\delta$ (HR, 3.6; $P = 0.042$; 95% CI, 1.1–14.7). The irradiated patients with low expression of PPAR $\delta$ were 4.9 times more likely to die than those with high expression of PPAR $\delta$ (HR, 4.9; $P = 0.029$; 95% CI, 1.8–12.1).

![Figure 2](image-url)  
**Figure 2.** The influence of RT on PPAR $\delta$ expression. After RT, the frequency of highly expressed PPAR $\delta$ was significantly increased from 2% to 16% in adjacent normal mucosa ($P < 0.001$; 95% CI, 0.14–0.26) and from 10% to 33% in distant normal mucosa ($P = 0.014$; 95% CI, 0.30–0.47), whereas it decreased in primary cancers from 61% to 41% ($P = 0.031$; 95% CI, 0.35–0.58) and from 32% to 22% in lymph node metastases ($P = 0.043$; 95% CI, 0.19–0.30).

![Figure 3](image-url)  
**Figure 3.** The relationship between PPAR $\delta$ expression in primary rectal cancers and TNM stage, lymph node metastasis and Ki-67 expression. The cases were unirradiated. The cases with high expression (moderate or strong staining) of PPAR $\delta$ exhibited (A) higher frequency of stage I cases and lower frequency of stage II, III, and IV cases ($P = 0.015$) and (B) lower rate of lymph node metastasis ($P = 0.046$), compared with those with low PPAR $\delta$ expression (negative or weak staining). C, both the non-RT and RT cases showed a negative correlation of PPAR $\delta$ expression with the expression of Ki-67 ($P = 0.016$, 0.025).
In the cases either with or without RT, the expression of PPAR δ in primary cancers was not significantly related to local recurrence or distant recurrence, although the cases with high expression of PPAR δ tended to have delayed time to local or distant recurrence ($P > 0.05$; see Supplementary Data).

Knockdown of PPAR δ promoted the proliferation of KM12C, KM12SM, and KM12L4a cells

By MTT assay, the colon cancer cells, KM12C, KM12SM, and KM12L4a with silenced PPAR δ, exhibited a significant increase in proliferation ($A$ value) at each time point, compared with untreated cells and the control cells treated
Discussion

In the present study, we found that PPARδ was predominantly located in the cytoplasm of the epithelial cells in normal mucosa, and tumor cells of primary cancers and lymph node metastases, with little expression in the nuclei. This finding is consistent with the studies by Takayama and colleagues (8) and Yoshinaga and colleagues (29). The specificity of the PPARδ antibody in the present study was verified by Western blot, which showed a clear band at the expected position of PPARδ protein as shown in Figure 1C. The cytoplasmic accumulation of PPARδ in human rectal tissue may be necessary for the proteins to be available for their nuclear role whenever required. Known as a nuclear receptor, PPARδ was found to be located mainly in the nuclei of colorectal cancer cells (7, 30). However, these studies could not be reproduced by Western blot as reported by Foreman and colleagues (31). Future studies need to confirm the location of PPARδ protein in colorectal cancer cells, by Western blot comparing nuclear extracts with cytoplasmic fractions.

The expression patterns of PPARδ in colorectal cancers have been reported, but the majority of available data are not supported by Western blot. For example, He and colleagues (6) and Gupta and colleagues (32) observed the overexpression of PPARδ mRNA in human colorectal cancer tissues by Northern blots. IHC analysis showed the increase of PPARδ expression in human colon cancers but no quantified analysis of Western blot was provided (8). Another report suggested that expression of PPARδ was higher in flat dysplastic adenomas from APCmin heterozygous mice while Western blot analysis showed no change in the expression of PPARδ in adenomas as compared with normal mucosa (30). To date, few studies have reported the differential expression of PPARδ between primary cancers and lymph node metastases. In the present study, we used both IHC and Western blot to quantitate the expression of PPARδ in normal rectal mucosa, primary cancers and lymph node metastases. Both IHC and Western blot showed that PPARδ was increased in primary cancers compared with adjacent or distant normal mucosa, and then deceased from primary cancers to lymph node metastases. Our finding confirms that PPARδ may be involved in the pathogenesis of rectal cancer, and indicates that PPARδ is related to the lymph node metastasis.

We analyzed the influence of RT on the PPARδ expression and found that PPARδ was increased in the adjacent or distant normal mucosa samples, and decreased in primary cancers and lymph node metastases after RT. Increase of PPARδ has recently been reported to protect cells from stress-induced injury, oxidation and DNA damage (33, 34). In this regard, the increase of PPARδ in the normal mucosa samples may be a protective response to RT, and its

with the Lenti-control (P < 0.05). There was no significant difference between untreated cells and control cells (P > 0.05; Fig. 5). The comparison of the A values at each time point among the 3 cell lines did not show significant difference (P > 0.05; data not shown).

Table 2. Cox multivariate analysis assessing the prognostic significance of cytoplasmic PPARδ expression in primary tumor

<table>
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<th>Indicator of poor survivala</th>
<th>HR (95% CI)</th>
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<td>PPARδ expression</td>
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<tr>
<td>Non-RT High vs. low</td>
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<td>Non-RT (I + II) vs. (III + IV)</td>
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<td>RT</td>
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<td>24.2 (4.6–23.7)</td>
<td>&lt;0.001</td>
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aThe analysis of DFS is presented for the cases without RT, whereas OS is for the cases with RT.
bSignificance was analyzed by Cox regression model; a P value of < 0.05 was considered significant.
stage I cases and lower frequency of lymph node metastasis. These findings indicate that PPAR \( \delta \) may be an early event in the development of rectal cancer and may be involved in the lymph node metastasis. Therefore, the examination of PPAR \( \delta \) in rectal cancer may be valuable for the preoperative evaluation, which may help in deciding treatment protocol for patients.

In the present study, we observed a significant association between increased PPAR \( \delta \) and favorable patient survival in primary rectal cancers. The increase of PPAR \( \delta \) in primary cancers was significantly related to better DFS in the cases without RT and better OS in those with RT. Ishizuka and colleagues reported that increased PPAR \( \delta \) was associated with favorable postoperative OS in the patients with colorectal cancers (35). However, that study used small samples (26 cases) in a short observation period (mean 32.3 months) and had no the information about RT. The present finding suggests that high expression of PPAR \( \delta \) is an independent indicator of a good prognosis for the rectal cancer patients undergoing surgery alone or surgery plus preoperative RT. We observed that the cases with highly expressed PPAR \( \delta \) tended to have delayed time to local/distant recurrence, although the differences did not reach statistical significance. This result seems inconsistent with the relationship between increased PPAR \( \delta \) and favorable patient survival observed in this study. The nonsignificance between the PPAR \( \delta \) expression and local/distant recurrence may be explained by the relatively small sample size in this study, which has confined the statistical significance. In addition, the expression of PPAR \( \delta \) may be associated with some unknown factors, which affected the survival of the patients more strongly than local/distant recurrence.

To explore the mechanisms underlying our findings, we further analyzed the relationships of PPAR \( \delta \) with apoptosis, Ki-67, PRL-3, COX-2, and survivin in the primary cancers. Ki-67 is a proven indicator of cell proliferation (36). COX-2 has been shown to participate in the apoptosis inhibition, angiogenesis, cell invasion, and metastasis of colorectal cancer (37). Survivin is a member of the apoptosis inhibitors (38). PRL-3 has been identified as an important protein in the metastatic process of colorectal cancer (26). We found that increased PPAR \( \delta \) was significantly related to the decreased Ki-67 in either non-RT or RT cases, but not to apoptosis, PRL-3, COX-2, or survivin. This finding indicates that PPAR \( \delta \) may be involved in the cell proliferation of rectal cancer, without participation in the physiological process of apoptosis, PRL-3, COX-2, and survivin.

Further assays showed that all 3 cell lines—KM12C, KM12SM, and KM12L4a—exhibited increased proliferation after PPAR \( \delta \) knockdown, indicating an inhibiting role of PPAR \( \delta \) on the proliferation of colon cancer cells. This finding confirms our previous observation that the silencing of PPAR \( \delta \) significantly promoted the proliferation of HCT-116 cells (39). We have recently shown the promotional role of PPAR \( \delta \) in the differentiation of both colon cancer cell lines and tissue (20). Taken together, the
present findings show that PPAR δ plays an inhibiting role in the progression of colorectal cancer, the mechanism underlying which is associated with its functions of inhibiting the proliferation and promoting the differentiation of cancer cells. Consistent with our findings, recent studies have provided evidence to support the inhibiting role of PPAR δ in colorectal carcinogenesis (13, 16, 17, 19). The inhibiting role of PPAR δ in colorectal cancer is consistent with the other findings in the present study, where increased PPAR δ is related to a higher frequency of early stage tumors, a lower rate of lymph node metastasis and favorable survival of rectal cancer patients. We did not find a significant difference in cellular proliferation among the KM12C, KM12SM, and KM12La cells after PPAR δ knockdown. This indicates that the effect of PPAR δ on the proliferation of colon cancer cells was not associated with the cell metastatic potentials.

Compared with ordinary IHC sections, TMA technology greatly improves internal experimental control as it allows simultaneous staining of massive sections. The present study benefits from the advantages of TMA; however the use of only 3 core tissue specimens per block when preparing TMA might lead to a limitation of reproductive samples due to tumor heterogeneity.

In conclusion, increased PPAR δ in primary rectal cancers is related to higher frequency of early stage tumors and lower rate of lymph node metastasis. PPAR δ plays a role in inhibiting the proliferation of colorectal cancer cells. These findings show that PPAR δ is related to the early development of rectal cancer and support the role of PPAR δ as a tumor suppressor in colorectal carcinogenesis. We show that preoperative RT increases the PPAR δ expression in normal mucosa while decreasing it in primary rectal cancers and lymph node metastases. Therefore, the application of a PPAR δ agonist or regulator together with preoperative RT in PPAR δ-low expression tumors may enhance the effect of RT. We show that increased PPAR δ in primary rectal cancers is independently related to a favorable survival of patients, showing that PPAR δ is a useful prognostic factor for rectal cancer patients.

Disclosure of Potential Conflict of Interest

No potential conflicts of interest were disclosed.

Acknowledgments

The authors thank Prof. I.J. Fidler (Anderson Cancer Center, Houston, TX) for offering KM12 cell lines and Dr. David Hinselwood (a Scottish doctor) at the Department of Oncology, Linköping University, Sweden, for revising this article.

Grant Support

This study was financially supported by grants from the Swedish Cancer Foundation, Swedish Research Council and the Health Research Council in the South-East of Sweden, and the Natural Science Foundation of China (grant no. 30801332).

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Received October 16, 2010; revised March 26, 2011; accepted March 29, 2011; published OnlineFirst April 29, 2011.

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Clinical Cancer Research

Biological Function and Prognostic Significance of Peroxisome Proliferator-Activated Receptor δ in Rectal Cancer

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Clin Cancer Res 2011;17:3760-3770. Published OnlineFirst April 29, 2011.

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