Chemoradiotherapy Resistance in Colorectal Cancer Cells is Mediated by Wnt/β-catenin Signaling

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Abstract

Activation of Wnt/β-catenin signaling plays a central role in the development and progression of colorectal cancer. The Wnt-transcription factor, TCF7L2, is overexpressed in primary rectal cancers that are resistant to chemoradiotherapy and TCF7L2 mediates resistance to chemoradiotherapy. However, it is unclear whether the resistance is mediated by a TCF7L2 inherent mechanism or Wnt/β-catenin signaling in general. Here, inhibition of β-catenin by siRNAs or a small-molecule inhibitor (XAV-939) resulted in sensitization of colorectal cancer cells to chemoradiotherapy. To investigate the potential role of Wnt/β-catenin signaling in controlling therapeutic responsiveness, nontumorigenic RPE-1 cells were stimulated with Wnt-3a, a physiologic ligand of Frizzled receptors, which increased resistance to chemoradiotherapy. This effect could be recapitulated by overexpression of a degradation-resistant mutant of β-catenin (S33Y), also boosting resistance of RPE-1 cells to chemoradiotherapy, which was, conversely, abrogated by siRNA-mediated silencing of β-catenin. Consistent with these findings, higher expression levels of active β-catenin were observed as well as increased TCF/LEF reporter activity in SW1463 cells that evolved radiation resistance due to repeated radiation treatment. Global gene expression profiling identified several altered pathways, including PPAR signaling and other metabolic pathways, associated with cellular response to radiation. In summary, aberrant activation of Wnt/β-catenin signaling not only regulates the development and progression of colorectal cancer, but also mediates resistance of rectal cancers to chemoradiotherapy.

Implications: Targeting Wnt/β-catenin signaling or one of the downstream pathways represents a promising strategy to increase response to chemoradiotherapy. Mol Cancer Res; 15(11): 1481–90. ©2017 AACR.

Introduction

Wnt/β-catenin signaling plays a central role during development and in maintaining homeostasis of multiple tissues throughout the body (1–3). Under physiologic conditions, Wnt ligands such as Wnt-1 or Wnt-3a associate with transmembrane Wnt receptors of the Frizzled family. This leads to an activation of dishevelled (DVL2), which in turn inhibits the so-called destruction complex. This multiprotein complex consists of adenomatous polyposis coli (APC), glycogen synthase kinase 3β (GSK3β), as well as Axin, which, in the absence of Wnt signaling, promote ubiquitylation and proteasomal degradation of β-catenin. Inhibition of the destruction complex results in an accumulation of β-catenin, which subsequently enters the nucleus and binds to members of the T-cell factor/lymphoid enhancer factor (TCF/LEF) family of high-mobility group transcription factors. Interaction with TCF/LEF factors promotes their ability to induce or repress the transcription of a plethora of target genes (1–3).

Aberrant Wnt/β-catenin signaling plays a critical role in the development and progression of various human malignancies. Importantly, mutations in and deregulation of components of this pathway are defining features of colorectal cancer (4, 5), which is the second leading cause of cancer-related death in Europe and the United States (6). According to recent data from the Cancer Genome Atlas Network, more than 90% of colorectal cancers show alterations of the Wnt signaling pathway, including biallelic inactivation of APC or activating mutations of CTNNB1 (β-catenin) in approximately 80% (7).

In previous studies, we demonstrated that the Wnt transcription factor TCF7L2, formerly known as TCF4, was overexpressed in primary rectal cancers that were resistant to preoperative 5-fluorouracil (5-FU)-based long-term chemoradiotherapy...
with antibodies and detected using an ImageQuant LAS 4000 technology (Lonza). Additional information regarding trans-Western blot analyses, cells were transfected using nucleofector technology or HiPerFect (Qiagen). For colony formation assays and polyvinylidene difluoride membrane (GE Healthcare), probed with antibodies and detected using an ImageQuant LAS 4000 mini CCD camera system (GE Healthcare). The respective antibodies and experimental conditions are listed in Supplementary Table S4.

Materials and Methods

Cell lines and cell culture

Human colorectal cancer cell lines SW480, SW837, SW1463, and LS1034 were obtained in 2006 from the ATCC. In 2009, directly prior to freezing the cell stocks used in this study, cell line authenticity was reconfirmed using short tandem repeat profiling (12). Cells were cultured in their recommended media (Invitrogen), supplemented with 10% FBS (Pan) and 2 mMol/L L-glutamine (BioWhittaker), and discarded no later than after 15 passages. Periodically, mycoplasma contamination was determined using the MycoAlert Mycoplasma Detection Kit (Lonza). Wild-type L cells and L-Wnt-3a cells (13), as well as hTERT-immortalized retinal pigment epithelial (RPE-1) cells (14), were kindly provided by Holger Bastians (Institute for Molecular Oncology, University Medical Center Goettingen, Goettingen, Germany). Cells were authenticated and stored and subsequently propagated as explained above. All cancer cell lines used in this study were characterized previously in regards to their radiation sensitivity and gene expression profiles (15).

siRNA transfection

Transfections with synthetic siRNA duplexes were performed as described previously (16, 17). Briefly, for cellular viability assays, cells were reverse transfected with siRNAs (Qiagen; Dharmacon/Thermo Fisher Scientific) using RNAiMAX (Invitrogen) or HiPerFect (Qiagen). For colony formation assays and Western blot analyses, cells were transfected using nucleofector technology (Lonza). Additional information regarding transfection conditions and siRNA sequences can be found in Supplementary Tables S1–S3.

Small-molecule inhibitor XAV-939

The tankyrase inhibitor XAV-939 (Tocris Bioscience) was used to inhibit tankyrases 1 and 2 (18). Cells were treated with different inhibitor concentrations for various time spans, followed by a medium exchange to remove the inhibitor. DMSO-treated cells served as a negative control. Details can be found in Supplementary Tables S1 and S2.

Western blot analysis

Western blot analysis was performed as described previously (9, 17). Briefly, 20 μg of whole-cell protein lysate was loaded and resolved on an 8% or 10% Bis-Tris polyacrylamide gel. Proteins were transferred by semi-dry blotting onto a polyvinylidene difluoride membrane (GE Healthcare), probed with antibodies and detected using an ImageQuant LAS 4000 mini CCD camera system (GE Healthcare). The respective antibodies and experimental conditions are listed in Supplementary Table S4.

Cellular viability assay

Cellular viability following siRNA transfection was assessed using the CellTiter-Blue reagent (Promega). Reduction of resazurin to resorufin was measured at various time points after the respective treatment using a plate reader (VICTOR X4, PerkinElmer) as per the manufacturer’s instructions. Cellular viability of siRNA-transfected cells was compared with cells transfected with a nonsilencing control siRNA (siNEG). Detailed information can be found in Supplementary Table S1.

Dual luciferase reporter assay

Plasmid transfections were performed as follows: cells were seeded into 12-well plates and allowed to adhere overnight. The next day, cells were forward transfected with the reporter plasmids SuperTopFlash or SuperFopFlash (Addgene), and, to normalize for transfection efficiency, cotransfected with Renilla luciferase reporter plasmid (Promega) using X-tremeGENE HP DNA Transfection Reagent (Roche). Twenty-four hours after transfection, passive lysis buffer (Promega) was added, and both Firefly and Renilla luciferase activity was measured in a microplate reader (Mithras LB940, Berthold Technologies). Relative transcription factor activity was calculated by dividing Renilla-normalized values of SuperTopFlash reporter plasmid and SuperFopFlash control reporter plasmid. Details on experimental conditions are provided in Supplementary Table S1.

(Chemo-)radiotherapy and determination of cell survival

To determine the respective surviving fractions after radiation treatment and chemoradiotherapy, a standard colony formation assay was performed. Defined numbers of cells growing in log-phase were seeded as single-cell suspensions into 6-well plates (Supplementary Table S2), and subsequently radiated with a single dose of 1, 2, 4, 6, and 8 Gy of X-rays (Gulmay Medical). For chemoradiotherapy, cells were exposed to 3 μmol/L of 5-FU (Sigma-Aldrich) for 16 hours prior to radiation. After defined time periods (Supplementary Table S2), cells were fixed with 70% ethanol and stained. Colonies with more than 50 cells were scored as survivors, and nonradiated cultures were used for data normalization. Resulting plating efficiencies for all colony formation assays can be found in Supplementary Table S5. Experiments were performed in triplicate (technical replicates) and independently repeated three times (biological replicates).

Exogenous stimulation of Wnt/β-catenin signaling using Wnt-3a

Wnt-3a-containing supernatant was produced from low-density seeded L-Wnt-3a cells, which were grown to confluence in DMEM/F12 medium. After approximately three days, medium was collected and stored at 4°C (batch I). Subsequently, fresh DMEM/F12 medium was added and cells were grown for another three days, until medium was collected (batch II). Batches I and II were pooled and sterile filtered. The resulting medium was diluted 1:1 using fresh DMEM/F12 medium supplemented with 0.35% sodium bicarbonate (Biochrom). As a negative control, wild-type L cells were treated similarly to produce a supernatant without Wnt-3a. For colony formation assays, cells were seeded in 6-well plates in medium with or
without Wnt-3a. 5-FU was added 16 hours prior to irradiation and removed by a media exchange, with or without Wnt-3a, 12 hours after irradiation (Supplementary Fig. S4).

**Overexpression of mutated β-catenin (S33Y)**

RPE-1 cells were seeded into 6-well plates and allowed to adhere overnight. Using 2.5 Gy of X-ray for 34 cycles, equaling to a cumulative dose of 68 Gy (Supplementary Fig. S5A). Specifically, cells were seeded into 6-well plates, radiated the next day, and cultured in fresh medium. At 70%–80% confluence, they were exposed to the next cycle of radiation.

**RNA isolation**

For each condition (i.e., no irradiation (SW1463, SW1463RES) or 6 hours after exposure to 4 Gy of X-rays (SW1463-4 Gy and SW1463RES-4 Gy), total RNA was isolated from logarithmic growing cells using the RNeasy Mini kit (Qiagen), per the manufacturer’s instructions, including the optional on-column DNAse digestion (Supplementary Fig. S5A). Quantity and purity of isolated RNA was assessed using a NanoDrop ND-1000 spectrophotometer (Thermo Fisher Scientific). RNA integrity was assessed by Agilent 2100 Bioanalyzer electrophoresis (Agilent Technologies). Only samples with an RNA integrity number > 9.5 were considered for additional experiments. Experiments were independently repeated three times (biological replicates).

**Microarray-based global gene expression analyses**

Gene expression profiling was performed as described previously (15, 16, 19). In brief, 200 ng of total RNA was reverse transcribed, amplified, and labeled with Cy3 using the Low RNA Input Linear Amplification Kit PLUS (Agilent Technologies). Subsequently, 1.65 μg of labeled cRNA was fragmented and hybridized overnight to a 44 k gene expression microarray (Agilent Technologies). After a washing step, the arrays were scanned on an Agilent DNA microarray scanner G2505B (Agilent Technologies) at 5 μm resolution. The respective gene expression data has been deposited in the National Center for Biotechnology Information Gene Expression Omnibus (GSE97543).

**Statistical analysis**

For analyses of the irradiation data, a multiple linear regression model was used to describe the normalized surviving fraction as a dependent variable, given the independent variables of irradiation dose, treatment group (control group or treated group), and biological replicates. Two different models, including either only irradiation dose and replicates or, additionally, treatment effect and treatment and irradiation dose interaction term were compared using ANOVA. All analyses were performed using the R statistical computing software R (version 3.1.0) using the R packages nlme and survival. For visualization, irradiation data are presented as mean and SEM from at least three independent experiments using the software KaleidaGraph (version 4.1.0). P values < 0.05 were considered significant, suggesting an effect on the treatment on the dose response.

Statistical analyses of cellular viability and luciferase reporter activity experiments were performed using an unpaired two-tailed Student t test in Microsoft Excel and visualized in Grapher (version 8.2.460). Again, P values < 0.05 were scored as significant.

To identify genes that are significantly altered in response to irradiation for each condition, we performed a one-sided, paired-sample Student t test to test whether the expression of each gene is increased or decreased as a response to irradiation. We then used false discovery rate (FDR) correction for multiple hypotheses with an FDR-level q < 0.05. Over-represented pathways were identified by using WikiPathway enrichment applying the hypergeometric enrichment test P < 0.05 (20).

**Results**

**Inhibition of β-catenin increases treatment sensitivity of colorectal cancer cells**

We previously reported a novel role for TCF7L2 in mediating responsiveness of colorectal cancer to chemoradiotherapy (8, 9). To test whether the observed effect of radiosensitization is TCF7L2-specific or dependent on the Wnt/β-catenin pathway more generally, β-catenin expression was silenced in two colon cancer cell lines, SW480 and LS1034, and in the rectal cancer cell line SW837, using RNA interference. Successful silencing was confirmed for all cell lines at 24, 48, 72, and 96 hours posttransfection by Western blot analysis (Supplementary Fig. S1). Of note, we detected a pronounced reduction of nuclear, cytosolic and total expression levels of both active and total β-catenin (Fig. 1A). siRNA-mediated silencing of β-catenin led to a mild reduction of cellular viability 48 hours after transfection, which was significant in LS1034 (Fig. 1B, left), and was accompanied by a significantly decreased TCF/LEF reporter activity in all three cell lines (LS1034 si#1: P = 4.196e-08, si#2: P = 4.317e-11; SW480 si#1: P = 1.058e-05, si#2: P = 2.646e-08; SW837 si#1: P = 0.0004, si#2: P = 5.131e-05; Fig. 1B, right). To assess the consequences of β-catenin inhibition on cellular sensitivity to chemoradiotherapy, we determined the respective surviving fractions following (chemo-) radiotherapy using a colony formation assay, as is standard in the field. Compared with the nonsilencing control siRNA, silencing of β-catenin significantly increased the sensitivity of LS1034 (P = 0.00019), SW480 (P = 2.73e-05), and SW837 (P = 2.76e-06) cells to irradiation (Fig. 1C, left). A similar effect was observed for a combination of 5-FU and irradiation (LS1034: P = 0.00186; SW480: P = 0.000272; SW837: P = 2.71e-08; Fig. 1C, right). Hereby, the addition of 5-FU only slightly increased the overall sensitivity to irradiation.

Next, we aimed to test whether we can recapitulate this sensitization effect using a small-molecule inhibitor of Wnt/β-catenin signaling. Toward this goal, we used the potent and clinically promising tankyrase inhibitor XAV-939 (21). Tankyrase inhibitors have been previously identified to inhibit β-catenin-mediated transcription by blocking Tankyrase isoforms 1 and 2 thereby stabilizing Axin through preventing its degradation (18). SW837 and SW480 cells were incubated with varying doses of XAV-939 for multiple time periods (Supplementary Information Gene Expression Omnibus (GSE97543).
siRNA-mediated silencing of β-catenin sensitizes colorectal cancer cells to (chemo-) radiotherapy. A, Active β-catenin and total β-catenin protein levels decreased 48 hours after transfection with siRNAs targeting β-catenin compared with a nonspecific negative control (siNEG) in LS1034, SW480, and SW837. Proteins were isolated as cytosolic and nuclear fractions (left) and as whole protein lysates (right). B, Cellular viability was measured 48 hours after transfection using a CellTiter-Blue assay (left), and transcriptional activity of the TCF/LEF complex was determined using a dual luciferase reporter assay (right). While silencing of β-catenin resulted in a mild reduction of cellular viability, the TCF/LEF reporter activity decreased significantly (LS1034: P = 4.96e-08 (si#1); P = 4.37e-11 (si#2); SW480: P = 1.058e-05 (si#1); P = 2.646e-08 (si#2); SW837: P = 0.0004 (si#1); P = 5.33e-05 (si#2)). C, Cell lines were irradiated 48 hours after transfection at various doses of X-ray (radiation treatment, RT, left). For chemoradiotherapy, cells were preincubated, 24 hours after transfection, with 3 μmol/L of 5-FU for 16 hours, and subsequently irradiated (right). Silencing of β-catenin significantly increased the sensitivity of LS1034 (radiation treatment: P = 0.00019, chemoradiotherapy: P = 0.00186, ANOVA model), SW480 (radiation treatment: P = 2.73e-05, chemoradiotherapy: P = 0.000272, ANOVA model), and SW837 (radiation treatment: P = 2.76e-06, chemoradiotherapy: P = 2.77e-08) to (chemo-) radiotherapy. Each experiment was repeated three times. Data are displayed as mean values, n = 2, error bars ± SEM.

**Figure 1.**

- **A**: Active β-catenin and total β-catenin protein levels were validated by Western blot analysis. For each cell line, two doses were established, one inducing Axin2 and consequently reducing (active) β-catenin as much as possible without decreasing cellular viability (Supplementary Fig. S3A and S3B). As described previously (18), treatment with XAV-939 resulted in a viability reduction of up to 20% (Supplementary Fig. S3B). Importantly, treatment with XAV-939 led to a significant radiosensitization of SW480 (1 μmol/L P = 0.016; 4 μmol/L P = 0.0112) and SW837 (5 μmol/L P = 0.0223; 10 μmol/L P = 0.000264). The sensitization effect to radiation treatment was stronger with higher doses of XAV-939, but less prominent as observed with siRNAs targeting β-catenin (Supplementary Fig. S3C).

- **B**: These results suggest that both TCF7L2, as previously demonstrated (8, 9), and β-catenin influence responsiveness to chemoradiation.

- **C**: Wnt-3a-stimulation leads to treatment resistance in nontumorigenic epithelial cells

To further investigate whether responsiveness to chemoradiation is not only β-catenin- and TCF7L2-dependent, but also Wnt/β-catenin-dependent, we used RPE-1 cells, which are highly sensitive to irradiation. RPE-1 cells represent a nontumorigenic epithelial cell model system with a stable karyotype (13, 22). Incubation with Wnt-3a, a physiologic ligand of the Frizzled receptor family, induced protein expression of Axin2 (13, 22). Incubation with Wnt-3a significantly increased the sensitivity of LS1034 (radiation treatment: P = 0.00019, chemoradiotherapy: P = 0.00186, ANOVA model), SW480 (radiation treatment: P = 2.73e-05, chemoradiotherapy: P = 0.000272, ANOVA model), and SW837 (radiation treatment: P = 2.76e-06, chemoradiotherapy: P = 2.77e-08) to (chemo-) radiotherapy. Each experiment was repeated three times. Data are displayed as mean values, n = 2, error bars ± SEM.

- **Figure 2.**

- **A**: Incubation with Wnt-3a, a physiologic ligand of the Frizzled receptor family, induced protein expression of Axin2 (13, 22). Importantly, external stimulation of Wnt/β-catenin signaling with Wnt-3a significantly increased the resistance to both irradiation (24 hours: P = 0.000185; 144 hours: P = 0.0102; Fig. 2C) and chemoradiotherapy (24 hours: P = 0.0000233; 144 hours: P = 0.00211; Fig. 2D). Conversely, exposure to XAV-939 resulted in a significantly increased sensitivity of RPE-1 cells to irradiation (P = 0.0232; Fig. 2E).

- **B**: This indicates that responsiveness to (chemo-)radiotherapy is Wnt/β-catenin dependent, and that the mechanism of Wnt/β-catenin–controlled responsiveness is not restricted to malignant cells but points to a universal phenomenon.

- **C**: Overexpression of mutated β-catenin (S33Y) leads to treatment resistance in nontumorigenic epithelial cells

To obtain further evidence that (chemo-)radiotherapy responsiveness is Wnt/β-catenin dependent, and to rule out potential Wnt- or β-catenin–independent effects of Wnt-3a (3, 23), we stimulated the pathway through overexpression of a constitutively active β-catenin (S33Y mutated) protein, which
Figure 2.
Wnt-3a stimulation induces treatment resistance in nontumorigenic epithelial cells. A, Incubation of RPE-1 cells with Wnt-3a for 24 hours (h) resulted in increased protein levels of Axin2 and active β-catenin. Proteins were isolated as cytosolic and nuclear fractions (left), and as whole protein lysates (right). B, Similarly, stimulation with Wnt-3a resulted in an approximately 800-fold increased TCF/LEF reporter activity ($P = 0.0105$). C and D, Treatment with Wnt-3a significantly increased resistance to both irradiation and chemoradiotherapy after 24 hours (radiation treatment (RT): $P = 0.000185$, chemoradiotherapy: $P = 0.0000233$; ANOVA model) and 144 hours (radiation treatment: $P = 0.0102$, chemoradiotherapy: $P = 0.00211$; ANOVA model). E, Exposure to 4 μmol/L of XAV-939 for 24 hours significantly sensitized RPE-1 cells to irradiation ($P = 0.0232$). Each experiment was repeated three times. Data are displayed as mean values, $n = 3$, error bars ± SEM.
Overexpression of mutated β-catenin (S33Y) induces treatment resistance in nontumorigenic epithelial cells. **A**, Transient transfection of mutated β-catenin resulted in increased protein levels of active and total β-catenin (left), and in elevated TCF/LEF reporter activity (P = 0.0002, right). **B** and **C**, Stable overexpression of mutated β-catenin resulted in a approximately 250-fold increased TCF/LEF reporter activity (P = 0.0017), and significantly decreased the sensitivity of RPE-1 cells to both irradiation (P = 0.0098; ANOVA model) and chemoradiotherapy (P = 0.0087; ANOVA model). **D**, Silencing of β-catenin in RPE-1 cells that stably express mutated β-catenin (S33Y) significantly decreased β-catenin protein expression and TCF/LEF reporter activity (si#1: P = 0.010, si#2: P = 0.0063; left panels), and abrogated the resistance to radiotherapy (P = 6.61e-06; middle) and chemoradiotherapy (P = 4.66e-06; right). **E**, In control RPE-1 cells that stably express the empty vector, silencing of β-catenin did neither affect TCF/LEF reporter activity (left) nor treatment sensitivity (radiation treatment: P = 0.557; chemoradiotherapy P = 0.637). Each experiment was repeated three times. Data are displayed as mean values, n = 3, error bars ± SEM.

Figure 3.

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Radioresistant colorectal cancer cells show deregulated RNA expression profiles and activate different pathways in response to irradiation.

To elucidate how Wnt/β-catenin signaling mediates resistance, we established gene expression profiles of SW1463 and radioresistant SW1463 (SW1463RES), both prior to and 6 hours after exposure to a single dose of 4 Gy (Supplementary Fig. S5B). As expected, we found a significant \( P = 0.05 \) overexpression of Wnt/β-catenin signature genes in SW1463RES compared with SW1463, including increased expression of LEF1, part of the TCF/LEF1 transcription factor complex, and WNT5B (Supplementary Table S6A).
To now assess transcriptional changes in response to radiation and to detect different activation of signaling pathways, we first identified genes differentially expressed between SW1463 cells and SW1463 cells 6 hours after exposure to 4 Gy (SW1463-4 Gy; Supplementary Table S6B). Next, we profiled our radiation-resistant SW1463 cells, prior to (SW1463RES) and 6 hours after exposure to 4 Gy (SW1463RES-4 Gy; Supplementary Table S6C). Both populations responded to radiation with a significant up- and downregulation of numerous genes and pathways, which can be found in Fig. 4D; Supplementary Tables S6B and S6C. In contrast to SW1463 cells, which responded to radiation with 4 Gy with significantly decreased expression of genes belonging to the PPAR signaling, we observed increased expression of PPAR pathway genes in SW1463RES in response to radiation with 4 Gy (Fig. 4D). In addition, SW1463RES cells responded with overexpression of genes belonging to metabolic pathways, such as AMPK signaling or the citric acid cycle (Fig. 4D).

**Discussion**

Preoperative chemoradiotherapy followed by radical surgical resection represents the standard of care for patients with locally advanced rectal cancer (4, 11, 26). However, the response of individual tumors to preoperative multimodal treatment is highly heterogeneous and ranges from complete clinical response (complete resistance). This poses a clinical dilemma, because patients with resistant tumors are exposed to the potential side effects of chemotherapy and irradiation with no clear benefit. It is therefore critical to uncover mechanisms and pathways of treatment resistance for the identification of strategies to increase the fraction of patients with rectal cancer who benefit from multimodal neoadjuvant treatment (10).

In an attempt to identify novel molecular targets and pathways that may be manipulated to sensitize tumors to chemoradiotherapy, we previously demonstrated that the Wnt transcription factor TCF7L2 is overexpressed in chemoradiotherapy-resistant rectal cancers (8). Subsequently, we showed that RNAi-mediated silencing of TCF7L2 sensitizes colon and rectal cancer cell lines to chemoradiotherapy (9). This radiosensitization was the consequence of a transcriptional deregulation of Wnt/TCF7L2 target genes, and a compromised DNA double strand break repair. In addition, silencing of TCF7L2 resulted in an increased fraction of cells in the G2-M phase of the cell cycle, which is known for increased vulnerability to radiation-induced DNA damage (27). However, it remained unclear whether this effect was a TCF7L2-inherent function or Wnt/β-catenin signaling-dependent. In this study, we confirm that the Wnt/β-catenin pathway mediates resistance of colorectal cancer cells to irradiation and 5-FU–based chemoradiotherapy.

We demonstrated that inhibition of β-catenin, either mediated through siRNAs or the small-molecule inhibitor XAV-939, sensitizes colorectal cancer cells to irradiation. This adds weight to the growing body of evidence suggesting that Wnt/β-catenin signaling mediates treatment responsiveness, in addition to its central role in tumor development and progression (1, 2). Recently, Cojoc and colleagues established gene expression profiles of prostate cancer cell lines and discovered that β-catenin regulates aldehyde dehydrogenase (ALDH1A3; ref. 28). RNAi-mediated silencing of β-catenin and ALDH1A3 led to a pronounced radiosensitization of a priori resistant cells (28). Fitting, in our experiment, radiosensitive SW1463 cells responded with a significant downregulation of ALDH1A3 to radiation. Dong and colleagues demonstrated a role for Wnt/β-catenin signaling in radiation-induced invasion of glioblastoma cells (29). In their model, radiation mediated nuclear accumulation of β-catenin and an upregulation of Wnt/β-catenin downstream genes. Pathway inhibition abrogated the proinvasion effects of radiotherapy (29). Similar to our approach, Ahn and colleagues repeatedly irradiated lung cancer cells to obtain resistant cell populations. Using gene expression profiling, they identified multiple genes that were differentially expressed between resistant cells and their parental cell lines. Wnt/β-catenin pathway genes were the most frequently altered (30). In our model, the increased resistance of SW1463RES was accompanied by elevated levels of active and total β-catenin, and increased TCF/LEF transcriptional activity. Interestingly, several of the genes described by Ahn and colleagues (24) were differentially deregulated in a similar fashion in our model, including many Wnt pathway genes. This points to an involvement of Wnt/β-catenin signaling in radiation resistance in multiple tumor entities. Of note, we previously characterized the chemoradiotherapy sensitivity of 12 colorectal cancer cell lines, which we correlated with gene expression profiles. Importantly, and nicely fitting with the observations reported here, we detected an overrepresentation of Wnt-pathway and Wnt-target genes within this signature of chemoradiosensitivity (15).

From a clinical point of view, the goal is to improve sensitivity to chemoradiotherapy. Of equal importance is to decrease potential side effects of irradiation, specifically, to decrease normal tissue toxicity (31). In this respect, Hai and colleagues demonstrated that transient activation of Wnt/β-catenin signaling prevents radiation-induced damage to salivary glands (32). Zhao and colleagues observed that an upregulation of the Wnt/β-catenin pathway accelerates mucosal repair following radiotherapy-induced oral mucositis (33). Very recently, Chandra and colleagues demonstrated that an activated Wnt/β-catenin pathway blocks radiation-induced apoptosis in osteoblasts through enhanced DNA repair, suggesting that Wnt agonists may be clinically used to block radiation-induced osteoporosis (34). In a similar attempt, we stimulated ontogenetically “normal” RPE-1 cells, which are highly sensitive to irradiation, with Wnt-3a, a physiologic ligand of the Frizzled receptor family. Treatment with Wnt-3a resulted in a strong activation of Wnt/β-catenin signaling, with increased expression levels of both Axin2 and β-catenin, accompanied by a approximately 800-fold increase of TCF/LEF reporter activity. Importantly, however, Wnt-3a stimulation resulted in significantly increased resistance to irradiation, while inhibition of β-catenin by XAV-939 sensitized RPE-1 to irradiation. A similar effect was observed through overexpression of constitutively active (S33Y-mutated) β-catenin in RPE-1 cells, which also resulted in a strong activation of Wnt/β-catenin signaling and which increased radiation resistance. This effect could be rescued by siRNA-mediated silencing of β-catenin. These results further support the notion that Wnt/β-catenin signaling controls responsiveness to chemoradiotherapy, and that this effect is not due to a β-catenin-independent branch of Wnt signaling (23).

However, open questions remain: Firstly, the underlying molecular mechanisms through which the Wnt/β-catenin signaling...
pathway functionally mediates responsiveness to chemoradiotherapy are still not fully understood. Preliminary evidence indicates that Wnt/β-catenin signaling may trigger epithelial to mesenchymal transition (EMT), which has been implicated in increased resistance to radiotherapy (28, 35–38). In our gene expression data, we observed an overexpression of genes associated with EMT in resistant cell populations, underscoring the importance of the Wnt/β-catenin-EMT-axis (data not shown). Recently, Jun and colleagues suggested a role for nonhomologous end joining (NHEJ) in Wnt/β-catenin-mediated radiation resistance (39). By screening DNA repair genes and β-catenin targets, they identified LIG4, whose expression was directly regulated by β-catenin and observed that deregulation of LIG4 mediates resistance of colon cancer cell lines to radiation. However, we did not observe changes in LIG4 expression when comparing SW1463 and Wnt-active radiation-resistant SW1463RES. In addition, LIG4 expression was not associated with treatment response in a set of 161 primary rectal cancers treated with 5-FU-based chemoradiotherapy (Emons and colleagues, unpublished data). This leads to the conclusion that, while the activation of LIG4 may represent one possible mechanism through which Wnt/β-catenin signaling controls responsiveness in colon cancer cells, there likely exist other resistance mechanisms in rectal cancer.

Of note, our data suggest a role for PPAR signaling in mediating radiation resistance. We observed increased expression of genes from the PPAR pathway in response to 4 Gy in SW1463RES, while SW1463 cells reacted in a completely opposite manner, that is, with a decrease in expression of these genes. It has been previously shown that PPAR is a downstream pathway of Wnt/β-catenin/TCF7L2 signaling (40), and that PPAR signaling, besides its prominent role in providing metabolic advantages for cancer cells (41), prevents radiation-induced cellular damage. For example, a PPAR-gamma agonist has been recently shown to protect normal tissue from radiation injury (42). Finally, the pathway is associated with a poor prognosis in a variety of human carcinomas (43, 44). Therefore, we speculate that PPAR signaling might be a novel mechanism through which Wnt signaling mediates radioresistance.

Our study did not address the question whether Wnt/β-catenin pathway inhibition in vivo sensitizes to chemoradiotherapy. However, several studies reported an association between treatment resistance and the expression of members of the Wnt/β-catenin pathway (45, 46). As described above, we previously used pretherapeutic gene expression profiling of tumor biopsies to demonstrate that TCF7L2 was expressed at higher levels in resistant compared with responsive primary rectal cancers (8). Using IHC analyses, Krigel and colleagues reported that TCF7L2 expression was a negative prognostic factor associated with a shorter overall survival of colorectal cancer patients (47). Similarly, Gomez-Martín and colleagues demonstrated that overexpression of β-catenin following chemoradiotherapy was associated with a decreased disease-free survival and a poor prognosis of rectal cancer patients (45).

In summary, our data demonstrate that Wnt/β-catenin signaling mediates responsiveness of rectal cancers to chemoradiotherapy. We suggest that targeting Wnt/β-catenin signaling or one of the downstream pathways represent a promising strategy to increase therapeutic responsiveness of rectal cancers to chemoradiotherapy, the standard treatment for patients with locally advanced stages of this disease (4, 11, 26).

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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