Targeting cancer stem cells: Modulating apoptosis and stemness
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Citation for published version (APA):
Çolak, S. (2016). Targeting cancer stem cells: Modulating apoptosis and stemness
Chapter 5

Decreased mitochondrial priming determines chemoresistance of colon cancer stem cells

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Published in *Cell Death and Differentiation*, 2014
Abstract
Tumor heterogeneity is in part determined by the existence of cancer stem cells (CSCs) and more differentiated tumor cells. CSCs are considered to be the tumorigenic root of cancers and suggested to be chemotherapy resistant. Here we exploited an assay that allowed us to measure chemotherapy-induced cell death in CSCs and differentiated tumor cells simultaneously. This confirmed that CSCs are selectively resistant to conventional chemotherapy, which we revealed is determined by decreased mitochondrial priming. In agreement, lowering the anti-apoptotic threshold using ABT-737 and WEHI-539 was sufficient to enhance chemotherapy efficacy, while ABT-199 failed to sensitize CSCs. Our data therefore point to a crucial role of BCLXL in protecting CSCs from chemotherapy and suggest that BH3 mimetics, in combination with chemotherapy, can be an efficient way to target chemotherapy-resistant CSCs.
Introduction

Colorectal cancer is the third most common cancer worldwide\(^1\),\(^2\). Patients with advanced stage colorectal cancer are routinely treated with either 5-fluorouracil (5-FU), leucovorin and oxaliplatin (FOLFOX), or with 5-fluorouracil, leucovorin and irinotecan (FOLFIRI), often in combination with targeted agents such as anti-VEGF or anti-EGFR at metastatic disease\(^5\),\(^6\). Despite this intensive treatment, therapy is still insufficiently effective and chemotheraphy resistance occurs frequently. Although still speculative, it has been suggested that unequal sensitivity to chemotherapy is due to an intra-tumoral heterogeneity that is orchestrated by cancer stem cells (CSCs) that can self renew and give rise to more differentiated progeny\(^7\),\(^8\). When isolated from patients, CSCs efficiently form tumors upon xenotransplantation into mice which resemble the primary tumor from which they originated\(^9\)-\(^11\). Additionally, many xenotransplantation studies have emphasized the importance of CSCs for tumor growth\(^5\),\(^9\)-\(^12\). Colon CSCs were originally isolated from primary human colorectal tumor specimens using CD133 as cell surface marker and shown to be highly tumorigenic when compared to the non-CSCs population within a tumor\(^9\),\(^10\). Later, other cell surface markers as well as the activity of the Wnt pathway have been used to isolate colon CSCs from tumors\(^12\),\(^13\). Spheroid cultures have been established from human primary colorectal tumors that selectively enrich for the growth of colon CSCs\(^11\),\(^12\), although it is important to realize that these spheres also contain more differentiated tumor cells\(^12\). In agreement, we have shown that the Wnt activity reporter that directs the expression of enhanced green fluorescent protein (TOP-GFP) allows for the separation of CSCs from more differentiated progeny in the spheroid cultures\(^12\).

CSCs are suggested to be responsible for tumor recurrence after initial therapy, as they are considered to be selectively resistant to therapy\(^11\),\(^14\). Conventional chemotherapy induces, among others, DNA damage and subsequent activation of the mitochondrial cell death pathway, which is regulated by a balance between pro- and anti-apoptotic BCL2 family members\(^15\). Upon activation of apoptosis, pro-apoptotic BH3 molecules are activated and these may perturb the balance in favor of apoptosis initiated by mitochondrial outer membrane polarization (MOMP), release of cytochrome c and subsequent activation of a caspase cascade.

The apoptotic balance of cancer cells can be measured with the use of a functional assay called BH3 profiling\(^16\). BH3 profiling is a method to determine the apoptotic “priming” level of a cell by exposing mitochondria to standardized amounts of roughly 20-mer peptides derived from the alpha-helical BH3 domains of BH3-only proteins and determining the rate of mitochondrial depolarization. Using this approach, priming was meas-
ured in various cancers and compared to normal tissues\textsuperscript{17, 18}. In all cancer types tested, the mitochondrial priming correlated well with the observed clinical response to chemotherapy. That is, cancers that are highly primed are more chemosensitive, while chemoresistant tumor cells as well as normal tissues are poorly primed\textsuperscript{17, 18}. This suggests that increasing mitochondrial priming can potentially increase chemosensitivity, which can be achieved by directly inhibiting the anti-apoptotic BCL2 family members\textsuperscript{18}. To this end small molecule inhibitors, so called BH3 mimetics, have been developed. ABT-737 and the highly related ABT-263 both inhibit BCL2, BCLXL and BCLW\textsuperscript{19-21} and were shown to be effective in killing cancer cells \textit{in vitro} and \textit{in vivo}\textsuperscript{21} with a preference for BCL2\textsuperscript{19, 22}. As BCL2 protein expression is often upregulated in hematopoietic cancers, it represents a promising target, which was supported by high efficacy of these BH3 mimetics in animal experiments\textsuperscript{21}. However, \textit{in vivo} efficacy is limited due to thrombocytopenia, which relates to a dependence of platelets on BCLXL for survival\textsuperscript{23, 24}. To overcome this toxicity, a BCL2-specific compound, ABT-199, was developed\textsuperscript{25}. Souers \textit{et al.} showed that inhibition of BCL2 using ABT-199 blocks tumor growth of acute lymphoblastic leukemia cells in xenografts\textsuperscript{25}. In addition to the single compound effects of ABT-199, combination with rituximab inhibited growth of non-Hodgkin’s lymphoma, mantle cell lymphoma, and acute lymphoblastic leukemia tumor cells growth \textit{in vivo}\textsuperscript{25}. In addition, highly effective tumor lysis was observed in all three patients with chronic lymphocytic leukemia that were treated with ABT-199\textsuperscript{25}. More recently, a BCLXL specific-compound, WEHI-539, was discovered using high-throughput chemical screening\textsuperscript{26}. As the apoptotic balance appears a useful target for the treatment of cancers and CSCs have been suggested to resist therapy selectively, we set out to analyze whether specifically colon CSCs are resistant to therapy and whether this is due to an enhanced anti-apoptotic threshold, specific to CSCs. To study chemosensitivity, we developed a robust single cell-based analysis in which we can measure apoptosis simultaneously in CSCs and their differentiated progeny. Utilizing this system we showed that colon CSCs and not their differentiated progeny are resistant to chemotherapeutic compounds and that this was due to the fact that these cells are less primed to mitochondrial death. Furthermore, inhibition of anti-apoptotic BCLXL molecule with either ABT-737 or WEHI-539, but not ABT-199, breaks this resistance and sensitizes the CSCs to chemotherapy.
Results

Colon cancer stem cells are resistant to various drugs.

Previously, cell death in colon CSCs and differentiated tumor cells has been compared by utilizing spheroid cultures enriched for CSCs in comparison with such cultures that were forced to undergo differentiation in vitro. In this approach differentiated tumor cells, also called sphere-derived adherent cultures (SDAC), were generated by placing spheroids in adherent cultures using medium containing Foetal Calf Serum. Chemotherapy-induced cell death was subsequently compared between spheroids and SDAC cultures. This approach therefore compares mixtures of cells maintained under different conditions and observed difference may not necessarily relate to intrinsic resistance of CSCs. Here we developed a robust assay, which allows treatment and cell death measurement of colon CSCs and their differentiated progeny under the exact same conditions. As we have shown that Wnt activity discriminates between CSCs and differentiated progeny within one sphere, we used this segregation to measure cell death in spheroid cultures after chemotherapy treatment with CaspGlow, a FACS-based assay that measures caspase-3 activity at the single cell level (Figure 1a).

This assay revealed that CSCs were more resistant to conventional chemotherapeutic compounds, including oxaliplatin (Figure 1a and b), cisplatin, 5-FU, etoposide, the chemotherapy regimens FOLFIRI and FOLFOX and also to the death receptor ligand TRAIL (Figure 1b). In addition to caspase-3 activity, other apoptotic features were measured at the single cell level using a similar FACS-based set-up. Phosphatidylserine exposure (Figure 2a) as well as loss of mitochondrial potential (Figure 2b) confirmed that colon CSCs were more resistant to chemotherapy when compared to their differentiated progeny. To show that resistance of colon CSCs towards chemotherapy is a common phenomenon, various spheroid cultures derived from different patients with colorectal cancer (Co108 and Co123) were transduced with TOP-GFP. As shown before, also in these spheroid cultures, WNT reporter activity correlated with CD133 expression and clonogenic potential (Supplementary Figure 1), pointing to the presence of CSCs in the CD133+ and TOP-GFP high fraction. Treatment of these distinct colorectal spheroid cultures with oxaliplatin also revealed a differential resistance to oxaliplatin, in which CSCs (TOP-GFP high) are more resistant when compared to differentiated tumor cells (TOP-GFP low) (Figure 2c).

Colon CSCs are less primed to death.

Several distinct mechanisms have been described that can lead to resistance towards chemotherapy. One of these involves a block in apoptosis induction, which can be
Figure 1: Colon cancer stem cells are resistant towards chemotherapy. A) Spheroid primary cultures were derived from primary human colorectal tumors and transduced with TOP-GFP construct. In these cultures Wnt pathway activity drives the expression of GFP (TOP-GFP). These spheroid cultures with TOP-GFP (Co100) were treated with chemotherapy e.g. oxaliplatin for 24 hr. Subsequently, spheroid cultures were stained with CaspGlow to measure caspase-3 activity. Cells are first gated on TOP-GFP high (Blue bar) and TOP-GFP low (Red bar), differentiated tumor cells and cancer stem cells, respectively. Caspase-3 activity is then separately analyzed for the differentiated tumor cells and cancer stem cells. B) This method was used to test various chemotherapeutic therapies. Spheroid cultures treated for 24 hr with oxaliplatin, cisplatin, 5-FU, etoposide, FOLFIRI, and FOLFOX or 4 hr with TRAIL are shown. Caspase-3 activity shows more cell death in differentiated cells compared to CSCs. Significance comparing differentiated versus CSCs is indicated (* p<0.05, ** p<0.01, *** p<0.001).
BCLXL is required for colon CSCs chemoresistance

Figure 2: Colon cancer stem cell resistance can be observed with multiple techniques in various patient derived spheroid cultures. A) Spheroid cultures were treated with oxaliplatin for 24 hr and (A) Phosphatidylserine (PS) exposure or (B) loss of mitochondrial activity were measured. C) Different spheroid cultures derived from different patients (Co108 and Co123), were treated with oxaliplatin and PS exposure was measured in differentiated tumor cells (TOP-GFP<sup>lo</sup>) and CSCs (TOP-GFP<sup>hi</sup>). All spheroid cultures showed cell-death in the differentiated tumor cells and not in the CSCs cells. Significance comparing differentiated versus CSCs is indicated (* p<0,05, ** p<0,01).
caused by a shift in the apoptotic balance, where a deregulation of anti- and pro-apoptotic signals is tilted in favour of anti-apoptotic proteins. In agreement, colon cancer spheroids over-expressing the anti-apoptotic BCL2 members BCL2, BCLW and BCLXL (Figure 3a and Supplementary Figure 2a and c) were completely resistant to oxaliplatin when analysed in the CaspGlow assay (Figure 3b and Supplementary Figure 2c). Ectopic overexpression of the other BCL2 proteins BFL1, BCLB and MCL1 was not sufficient to overcome oxaliplatin-induced cell death in spheroid cultures (Supplementary Figure 2), suggesting that changing the apoptotic balance using the previously reported stable anti-apoptotic BCL2 family members is indeed a possible means to escape from chemotherapy.

To directly evaluate whether the resistance of CSCs is due to a difference in this apoptotic balance, BH3 profiling experiments were used to measure the level of mitochondrial priming in colon CSCs and differentiated tumor cells (Figure 3c and d). Spheroid cultures were treated with BH3 peptides and mitochondrial depolarization was measured. Importantly, we have shown before that these BH3 peptides have differential binding to anti-apoptotic BCL2 family proteins. For instance, HRK peptide has a much greater affinity for BCLXL than BCL2 or BCLW (Figure 3c). For all BH3 peptides, less depolarization was measured in CSCs when compared to differentiated cells (Figure 3d and Supplementary Figure 3). This profiling indicates that colon CSCs display decreased mitochondrial priming in comparison to differentiated tumor cells and thereby can resist chemotherapy.

**BCLXL specific inhibitor WEHI-539 sensitizes colon cancer CSCs towards chemotherapy.**

In some cancers expression of anti-apoptotic molecules is highly enhanced when compared to the untransformed cells from the same patient. As overexpression of BCL2, BCLXL or BCLW could enhance chemotherapy resistance (Supplementary Figure 2c), we analyzed whether the small molecule inhibitor ABT-737, which binds to all these anti-apoptotic family members, could target CSCs. In addition, we utilized the more selective compounds ABT-199, which solely targets BCL2, and WEHI-539, which solely targets BCLXL. First we tested whether the mitochondrial pathway regulates clonogenic potential of CSCs. Limiting dilution analysis with CSCs that were pre-treated with ABT-737, ABT-199 or WEHI-539 revealed that ABT-737 and WEHI-539 both were sufficient to decrease clonogenic capacity, whereas ABT-199 did not affect clonogenic growth (Figure 4a). As WEHI-539 is selective for BCLXL, this points to a dependency of CSCs on BCLXL for survival. Importantly, ABT-737 or
BCLXL is required for colon CSCs chemoresistance

**Figure 3: Colon CSCs and differentiated cells are differentially mitochondrial primed.**

A) Spheroid culture (Co100) ectopically overexpressing BCLXL protein were generated, as shown by western blot BCLXL (upper panel). Lower panel shows a control western for ERK1/2. B) After oxaliplatin treatment, differentiated tumor cells (TOP-GFP<sup>low</sup>) overexpressing BCLXL cells are no longer sensitive and show significantly reduced cell death, by caspase-3 measurement. C) Figure showing specificity of BH3 peptides for binding to anti-apoptotic BCL2 family protein members. Red indicates high affinity and green low affinity. D) Spheroid cultures (Co100) were subjected to BH3 profiling using various BH3 peptides. Cells were stained for CD133, treated with indicated BH3 peptides and mitochondrial depolarization was measured with JC1. Mitochondrial depolarization in differentiated tumor cells (CD133<sup>low</sup>) and CSCs (CD133<sup>high</sup>) is shown relative to complete depolarization obtained with CCCP. Significance comparing oxaliplatin-treated control versus BCLXL overexpressing differentiated tumor cells is indicated in Figure 3B and comparing differentiated versus CSCs in Figure 3D (* p<0.05, ** p<0.01, *** p<0.001).
WEHI-539-induced loss of clonogenicity could be restored when BCLXL was ectopically overexpressed (Figure 4b). To confirm that CSCs are dependent on BCLXL and not BCL2, western blot analysis were performed for BCL2, BCLXL and BCLW proteins and indeed all spheroid cultures expressed BCLXL and small amounts of BCLW, but no BCL2 protein levels could be detected (Figure 4c). Next, we evaluated whether decreased mitochondrial priming was responsible for chemotherapy resistance of CSCs. We therefore set out to perturb the apoptotic balance with sub-lethal amounts of ABT-737, ABT-199 or WEHI-539 to determine whether this would be a means to sensitize colon CSCs towards chemotherapy. As expected, colon CSCs were resistant to single treatments with low levels of BH3 mimetics and also against oxaliplatin. However when spheroid cultures were treated with ABT-737 or WEHI-539 compounds, CSCs were effectively sensitized towards oxaliplatin (Figure 4d) and other chemotherapeutic agents (Supplementary Figure 4). Similar to our observations on clonogenicity, the BCL2 specific inhibitor ABT-199 did not sensitize CSCs toward chemotherapy (Figure 4d). Finally, to determine whether combination treatment affected clonogenicity of CSCs, we used low doses of oxaliplatin that slightly reduced CSC clonogenicity (Figure 4e). However, combination of oxaliplatin with sub-lethal doses of ABT-737, but not ABT-199, resulted in further loss of clonogenicity of CSCs (Figure 4e). In agreement, when spheroid cultures were treated with ABT-737, ABT-199 or WEHI-539 in combination with oxaliplatin and growth of the cultures was measured over time (Figure 4f), we observed a strong initial decrease in cell viability in the first 4 days. However, spheroid cultures treated with oxaliplatin alone or in combination

Figure 4: ABT-737 and WEHI-539 sensitize colon CSCs towards chemotherapy. A) Untransduced or B) BCLXL-transduced spheroid cultures were treated with 1 µM of ABT-737, ABT-199 or WEHI-539 for 24 hr and limiting dilution assay was performed on CSCs (TOP-GFPhigh). ABT-737 and WEHI-539 decreased clonogenic capacity in CSCs (A), but not when BCLXL is overexpressed (B). Significance is indicated (** p<0.01) and is related to vehicle (DMSO) alone treatment. C) Western blot analysis on the different spheroid cultures for BCL2 (upper panel) and BCLXL (middle panel) and BCLW (lower panel) proteins. Control westerns for the kinase ERK1/2, which is stably expressed in the spheroids, are shown. D) Spheroid cultures were treated with 100 nM ABT-737 or WEHI-539 in combination with 50 µM oxaliplatin for 24 hr and subsequently analyzed for caspase-3 activity or E) Clonogenic capacity using limiting dilution on CSCs (TOP-GFPhigh). Significance is indicated (** p<0.01 and *** p<0.001) and is related to oxaliplatin alone treatments. F) Spheroid cultures were treated with 100 nM ABT-737, ABT-199 or WEHI-539 in combination with 0.5 µM oxaliplatin for 24 hr and cell numbers were measured at various time points using cell titer blue.
BCLXL is required for colon CSCs chemoresistance

A BCLXL is required for colon CSCs chemoresistance

**Clonogenic capacity (%)**

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**% Casp3-active cells**

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**Fluorescence (530/590nm)**

- Oxaliplatin
- ABT-199 + Oxaliplatin
- ABT-737 + Oxaliplatin
- WEHI-539 + Oxaliplatin

**Days**

0 5 10 15 20
with ABT-199 regained proliferative potential in the days after, while combination treatment with ABT-737 or WEHI-539 strongly inhibited this revival (Figure 4f). This indicates that decreased mitochondrial priming of CSCs is an important aspect of CSCs that directs enhanced chemotherapy resistance of these cells and should thus be targeted to optimize chemotherapy.

Discussion

In the last decade it has become increasingly clear that many tumors are heterogeneous, consisting of CSCs and more differentiated tumor cells. Like other CSCs, colon CSCs are highly tumorigenic and therefore targeting these cells is suggested to enhance the success of cancer therapy. However, colon CSCs are resistant to chemotherapy and the molecular mechanisms that orchestrate this are still poorly understood. Here we demonstrate that colon CSCs are less mitochondrial primed and increasing mitochondrial priming by ABT-737 sensitizes colon CSCs towards therapy.

In a non-cancerous cell there is a balance between anti-apoptotic molecules and pro-apoptotic BH3 molecules. This balance determines the activation of BAX and BAK proteins and thereby mitochondrial integrity in a cell. There are two main models for activation of BAX and BAK molecules. In the first model, called the direct activation model, BH3 proteins are divided in activators (e.g. BIM and BID) and sensitizers (e.g. BAD and HRK)\(^3\). In this model sensitizers can bind to anti-apoptotic molecules (e.g. BCL2, MCL1) and can inhibit them. This inhibition of anti-apoptotic molecules results in an indirect activation of BAX and BAK molecules, while the activators are able to activate BAX and BAK molecules directly, resulting in MOMP.

In the second model, called the neutralization model, BAX and BAK activation is a spontaneous event, which is prevented by the anti-apoptotic molecules. Anti-apoptotic BCL2 proteins need to be neutralized by BH3-only proteins to allow BAX and BAK oligomerization and MOMP\(^3\). Recently, a unified model has been proposed that incorporates both models suggesting the existence of two modes of inhibition in which the BCL2 family prevents apoptosis\(^3\).

Similar to earlier studies\(^16,18,28\), we used BH3 profiling to determine the level of mitochondrial priming in CSCs and differentiated cells (Figure 3d). In this assay BH3 peptides were used that mimic the function of BH3 molecules. Using the differential specificities of the various BH3 molecules, it is possible to probe the mitochondrial resistance of a cell. As described before, BID and BIM are very potent in inducing mitochondrial depolarization\(^17\). However, lower concentrations of BIM revealed lower
BCLXL is required for colon CSCs chemoresistance

In both models, BIM and BID proteins can be described as potent apoptosis activators either as they represent direct inducers or because they are promiscuous in inhibiting a wide-range of anti-apoptotic BCL2 members. Interestingly, CSCs were selectively more resistant to all other BH3 peptides that were tested. This can be due to the higher affinity of these peptides or may be explained by a loss of activators (like BID and BIM) in CSCs. Alternatively, it may point to the expression of higher levels of anti-apoptotic BCL2 proteins. The latter possibility is supported by the observation that simple overexpression of BCLXL, BCL2 or BCLW is sufficient to prevent chemotherapy-induced apoptosis in the differentiated tumor cells, while the BH3 mimetic ABT-737 and WEHI-539 sensitizes both CSCs and the more differentiated cells. Further work is needed to determine the mechanism behind the decreased mitochondrial priming, but our findings strongly argue for a BCLXL-dependent anti-apoptotic threshold in colon CSCs.

In various tumors it has been shown that the apoptotic balance is disturbed in favor of anti-apoptotic molecules. In many lymphomas, BCL2 protein expression is increased. Moreover, also in solid tumors various anti-apoptotic proteins are shown to be highly expressed. In line with this, colorectal cancers have been reported to express enhanced levels of BCL2 and BCLXL and this correlates with clinical outcome of the patient. Similarly, microsatellite instable colorectal cancers frequently display loss of BAX due to a frameshift mutation and this correlates with decreased 5-FU sensitivity. Previously, we have shown that colon CSCs express IL-4 and this can signal in an autocrine fashion. In line with our current findings, we showed that treatment of colon CSCs with neutralizing antibody against IL-4, decreased protein levels of the anti-apoptotic proteins c-FLIP, BCLXL and PED and sensitized colon CSCs to oxaliplatin treatment.

Although our data strongly argue for a role of decreased mitochondrial priming of CSCs in chemotherapy resistance, it is important to note that therapy resistance of CSCs may have other underlying reasons. For instance, Kranenburg and colleagues have shown that colon CSCs highly express BIRC6 and knock-down of BIRC6 sensitized colon CSCs towards oxaliplatin. Similarly slow cell cycling and high drug transporters activity have been connected to resistance. In agreement, ABCB5 was shown to be expressed in CSCs and render them drug resistant. The latter two mechanisms are less likely to be responsible for therapy resistance in our model as we have previously shown that cell cycle activity is not different between CSCs and differentiated tumor cells. In addition, CSCs were also resistant to TRAIL, which is not dependent on either cell cycle or drug efflux. Moreover,
experiments performed with doxorubicin showed no difference in uptake of the drug (not shown), indicating that drug efflux is not different in colon CSCs. Here we show that chemoresistance of CSCs is instead due to differential mitochondrial priming. Previously, we have used an inducible caspase-9 to target colon CSCs. Upon activation of caspase-9, colon CSCs were killed efficiently \textit{in vitro} and \textit{in vivo} \cite{43}. In line with this study this indicates that the apoptotic block in colon CSCs is upstream of MOMP.

For many years, there is an effort to target the anti-apoptotic molecules in cancer. ABT-737, and its orally bioavailable variant ABT-263, target BCL2, BCLXL and BCLW \cite{21}, while ABT-199 and WEHI-539 specifically target BCL2 and BCLXL respectively \cite{25,44,45,26}. Here, these three inhibitors were tested in combination with chemotherapy. Interestingly, only ABT-737 and WEHI-539 were effective in sensitizing colon CSCs towards chemotherapy, while ABT-199 failed to kill or sensitize CSCs. In agreement, BCL2 protein expression was hardly detected in the CSCs analysed in contrast to BCLXL. Importantly, also the more differentiated tumor cells can be sensitized by BH3 mimetics, which indicates that a BCLXL-dependent anti-apoptotic threshold is present in differentiated cancer cells as well, but that this threshold is simply elevated in the CSCs and sufficient to block chemotherapy-induced death. As relatively low doses of BH3 mimetics are sufficient to sensitize CSCs, it may be feasible to design combination therapies that spare thrombocytes while effectively targeting colorectal cancer. Whether such combination therapies pose a threat to normal intestinal stem cells remains to be established. We have previously shown that hematopoietic stem cells are relatively unprimed \cite{18}, but whether these normal stem cells or intestinal stem cells are also sensitized by BCLXL-targeting BH3 mimetics remains to be determined. Taken together, we have shown that colon CSCs are resistant to various drugs due to a decreased mitochondrial priming and that exogenous priming of CSCs with ABT-737 or WEHI-539 is sufficient to sensitize colon CSCs towards chemotherapy.

\textbf{Materials and Methods}

\textbf{Cell culture}

Colon spheroid cultures were derived from different patients with colorectal cancer in accordance with the rules of the medical ethical committee of the AMC. Briefly, primary resected human colon carcinomas were digested enzymatically for 1 hr with collagenase II (1.5 mg/ml; C6885 Sigma-Aldrich, Zwijndrecht, The Netherlands) at 37°C. The dissociated samples were strained
BCLXL is required for colon CSCs chemoresistance

(70 μm pore size) and washed in CSC medium and subsequently cultured. Spheroid cultures were cultured in stem cell medium (advanced DMEM/F12 (Gibco, Bleiswijk, The Netherlands)) supplemented with 6 mg/ml Glucose, 5 mM HEPES, 2 mM L-Glutamine, 4 μg/ml Heparin, epidermal growth factor (EGF; 50 ng/ml) and basic fibroblast growth factor (bFGF; 10 ng/ml). Spheroid cultures were maintained in ultra low adherent flasks (Corning, Amsterdam, The Netherlands). The vector for the TCF/LEF reporter driving expression of GFP (TOP-GFP) was a gift from Laurie Ailles and was described previously. Spheroid cultures were transduced lentivirally with TCF/LEF reporter and single cell cloned by single-cell plating in 96-well ultralow adhesion plates (Corning) with FACSaria (BD Biosciences, Breda, The Netherlands). To generate spheroid cultures that ectopically overexpress BCLXL, spheroid cultures were transduced with pHEFTIR-BCLXL. The lentiviral construct pHEFTIG was a kind gift from Arjen Bakker and Dr. Bianca Blom (AMC, Amsterdam). First GFP within pHEFTIG was replaced by RFP to generate pHEFTIR. Next, BCLXL was cloned into pHEFTIR to generate a pHEFTIR-BCLXL. Spheroid cultures were transduced with pHEFTIR (Control) or pHEFTIR-BCLXL (BCLXL). To generate spheroid culture expressing BCL2, BCLW, BCLXL, BFL1, BCL proteins, spheroid cultures were retroviral transduced with pMX-IRES-Blasticidin with the cDNA of the particular BCL2 protein and selected for at least two weeks. These constructs were kind gift from prof. Jannie Borst (The Netherlands Cancer Institute, Amsterdam) and were described previously. Spheroid culture ectopically expressing MCL1-3a mutant, was generated using LZRS-MCL1-3A, which was a gift from Dr. Mien-Chie Hung (The University of Texas M.D. Anderson Cancer Center, Texas). This MCL1-3A mutant construct was previously described and is was shown that it can not be degraded by the E3 ligase β-TrCP.

Reagents

The following chemotherapeutic compounds were tested; 50 μM Oxaliplatin, 10 μg/ml etoposide, 200 μg/ml 5-FU (all from Sigma-Aldrich), 50 μM Cisplatin (Platosin, Pharmachemie, Haarlem, The Netherlands) all for 24 hr, and 100 ng/ml rhTRAIL (Enzo Life Sciences, Raamsdonkveer, The Netherlands) in combination with 1.5 μg/ml anti-FLAG antibody (Sigma-Aldrich) for 4 hr. FOLFOX and FOLFIRI treatments consisted of 1.25 μM Oxaliplatin (FOLFIRI) or 1 μM Irinotecan (Sigma-Aldrich) treatment followed after 90 min by 50 μM 5-FU, cells were analyzed after 24 hr. Spheroid cultures were treated for 24 hr with ABT-199, ABT-737 (Selleck Chemicals, Huissen, The Netherlands) or WEHI-539 (ChemScene, NJ, USA) in combination with a chemotherapeutic compound or only vehicle.
Immunoblotting and antibodies

Cells were lysed in 1x RIPA lysis and extraction buffer (Thermo Fisher Scientific, Etten-Leur, The Netherlands) containing complete protease inhibitor (Roche, Woerden, The Netherlands). After clearing the lysate by high speed centrifugation (14,000 rpm, 10 min, 4°C), the protein concentration was determined using BCA protein assay (Thermo Scientific, Breda, The Netherlands). 20 μg extracted proteins were separated by 12% precast gels (Biorad, Veenendaal, The Netherlands) and transferred to on Hybond-P (Amersham, Upsalla, Sweden). Membranes were blocked with 5% milk in Phosphate Buffered Saline solution containing 0.2% TWEEN (PBS-T) for 1 hr. The membrane was incubated with the appropriate amount of antibody. Primary antibody incubations were carried out in 2.5% milk/PBS-T overnight at 4°C and subsequently washed 3 times with PBS-T. Membranes were then incubated with anti-mouse IgG horseradish peroxidase conjugates for 1 hr, washed 3 times and detection of bound antibody was performed with ECLplus reagents (Amersham). Western blots were analysed by LAS4000. The antibodies used were anti-BCLXL (Santa-Cruz, clone S18, 1:500, Heidelberg, Germany), anti-BCL2 (Santa-Cruz, clone N19 1:500), anti-BCLW (Abcam, clone 31H4, 1:500, Cambridge, UK), anti-BFL1 (Santa-Cruz, 1:250), anti-MCL1 (Cell Signaling, clone 4572, 1:1000, Leiden, The Netherlands) and anti-ERK1/2 (Cell Signaling, 1:10,000).

RNA extraction, cDNA synthesis and RT-PCR

RNA was extracted from cells using Trizol reagent (Invitrogen, Leek, The Netherlands) in accordance with the manufacturer’s protocol. RNA concentration was determined with NanoDrop ND-2000 (Thermo Scientific) and 1 μg of RNA was used to synthesis cDNA using SuperScript III accordance with the manufacturer’s protocol (invitrogen). RT-PCR was performed with LC480 SYBR green (Roche) in accordance with the manufacturer’s instructions on a LC480. The following primers were used: 18S sense: AGACAAACAAGCTCCGTGAAGA, 18S antisense: CAGAAGTGACGCAGCTCTCA, BCLB sense: CTAAGGAGCAGAGGGCC, and BCLB anti-sense: GTGGAAAGGGGGTCCTGAAG.

Limiting-dilution assay

Spheroid cultures were dissociated and the 10% TOP-GFP**high** and 10% TOP-GFP**low** cells were FACS deposited using FACSaria (BD Biosciences) in a limiting dilution fashion at 1, 2, 4, 8, 16, 32, 64, 128, and 256 cells per well in ultra-low 96 wells plates (Corning). Clonal frequency was evaluated with the Extreme Limiting Dilution Analysis (ELDA) ‘limdil’ function as described 48.
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**FACS staining**
Various colon spheroid cultures transduced with TOP-GFP were dissociated and were stained with AC133/CD133 – APC antibody (1:25) (Miltenyi Biotec, Paris, France) in PBS containing 1% bovine serum albumin (PBS-B) for 30 min at 4°C. Subsequently, cells were washed with PBS-B and resuspended in PBS-B. 7-AAD (BD Biosciences) was used to exclude death cells. Staining was analysed on a FASCanto (BD Biosciences).

**Cell death assays**
Primary colon cancer spheroid cultures were dissociated with trypsin-EDTA and seeded as single cells on adherent cell culture 12 wells plate (Greiner, Alphen a/d Rijn, The Netherlands) overnight. The next day, adherent cells were treated with chemotherapeutic drugs or targeted compounds for the indicated times. After treatment, cells were harvested and identification of the CSCs was performed by gating on the TOP-GFP<sup>high</sup> cells in the spheroid cultures, while the differentiated cells were identified in the same cultures by gating on the TOP-GFP<sup>low</sup> cells. To measure apoptosis at the single cell level in both populations three distinct FACS-based measurements of cell death were used. The first, Caspase 3 activity was measured with CaspGlow active staining kit (Red-DEVD-FMK) according to the manufacturer’s instructions (BioVision, California, USA). In short, after treatment, spheroid cultures were made single cells using trypsin-EDTA. 50.000 cells were washed with stem cell medium and stained with RED-DEVD-FMK for 1 hr at 37 °C. Subsequently, cells were washed twice with wash-buffer. The second FACS-based measurement of cell death was based on the exposure of phosphatidylinerine. Spheroid cultured cells were stained with Annexin V-APC (BD biosciences) and 7-AAD (BD biosciences) for 15 min at RT. The third assay involved mitochondrial activity, which was measured with Mitotracker Orange CMTMRos (Molecular Probes, Bleiswijk, The Netherlands). Spheroid cultures were incubated for 30 min with 25 nM Mitotracker Orange at 37 °C. All three apoptosis staining methods were followed by flow cytometry performed with FACSCanto (BD biosciences). Cell death was measured in CSCs by gating on TOP-GFP<sup>high</sup> cells and in differentiated tumor cells by gating on TOP-GFP<sup>low</sup> cells.

**Cell survival assay**
Primary colon cancer spheroid cultures were dissociated and seeded as in cell death assays. Adherent cells were treated with oxaliplatin in combination with ABT-737, ABT-199, or WEHI-539. After 24 hr treatment, cells were harvested and 2000 cells were transferred into ultra low adherent 96 wells plates (Corning). Cell survival was
measure at different time points by adding 20μl/well of cell titer blue reagent (Promega, Leiden, The Netherlands). Subsequently, cells were incubated for 4 hr and fluorescence was measured on a Biotek HT synergy plate reader (BioTek, Potton, UK)

**BH3 profiling**
Mitochondrial priming was measured as described before 16. Briefly, spheroid cultures were stained with AC133/CD133 - APC antibody (1:25) (Miltenyi Biotec). Subsequently cells were washed in T-EB (500 mM Trehalose, 10 mM Hepes-KOH pH 7.7, 80 mM KCl, 1 mM EGTA, 1 mM EDTA, 0.1% BSA, 5 mM succinate) and 50.000 cells resuspended in 100 μl T-EB were added to 100 μl of T-EB containing 20 μg/ml digitonin, 20 μg/ml oligomycin, and peptides. Cells were incubated for 90 min with various 100 μM BH3 peptides at RT. As shown before, BIM and BID peptides were very potent and therefore 6 μm and 12 μm of these peptide were tested, respectively. More concentrations of BIM peptides are tested and shown in Supplementary Figure 4. JC-1 staining was performed for 30 min at RT (1 μM) and MOMP was measured in CSCs (10 % CD133^high^ cells) and more differentiated cells (10 % CD133^low^ cells) on a FACSCanto (BD Bioscience).

The following BH3 peptides (Eurogentec, Maastricht, The Netherlands) were used;
BIM: MRPEIWIAQELRRIGDEFNA;
BID: EDIIRNIAHRHLAQGDSMDR;
PUMA: EQWAREIGAQLRRMADDLNA;
BIK: MEGSDALALRLACIGDEMDV;
BNIP3: VVEGEKEVEALKKSADWVSD;
HRK: SSAAQLTAARLKALGDELHQ;
NOXA A: AELPPEAAQLRKIGDKVYC;
BMF: HQAEVQIARKLQLIADQFHR

**Acknowledgements**
We would like to thank Louis Vermeulen, Felipe de Sousa e Melo, Michael Bots, Maarten Bijlsma and Jeremy Ryan for constructive discussion, and the surgeons Prof. Bemelman, Prof. Gulik, Dr. Tanis, Dr. Buskens and Dr. Van de Ven (AMC Amsterdam) for providing colon cancer samples. We also want to thank Dr. Mien-Chie Hung for providing MCL1-3a mutant construct and Dr. Rogier Rooswinkel and prof. Jannie Borst (The Netherlands Cancer Institute, Amsterdam) for providing all other pro-survival BCL2 family members, Dr. Laurie Ailles (University of Toronto) for providing Tcf/Lef reporter construct and Arjen Bakker and Dr. Bianca Blom
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(AMC Amsterdam) for providing pHEFTIG construct. Finally, we thank Berend Hooibrink, Toni van Capel en Kate Cameron for assistance with fluorescence-activated cell sorting experiments. This work was supported by a VICI grant from the Netherlands Organisation for Scientific Research and a Dutch Cancer Society (KWF Kankerbestrijding) grant (2009-4416 and 2012-5612) (to J.P.M.)

Conflict of Interest
The authors declare no conflict of interest.
CHAPTER 5

References


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**Supplementary figures**

Supplementary Figure 1: TOP-GFP activity correlates with CD133 and clonogenicity. TOP-GFP transduced spheroid cultures were stained for CD133-APC (A) and the lowest (blue) and highest (red) CD133 expressing cells were analyzed for their TOP-GFP activity (B). C). 10% TOP-GFP<sup>high</sup> and 10% TOP-GFP<sup>low</sup> cells were deposited in ultra-low 96 wells plates in a limiting dilution fashion. After two weeks colonies were scored and variation in clonogenic capacity between differentiated and CSCs is shown. Significance comparing differentiated versus CSCs is indicated (*** p<0,001).
Supplementary Figure 2: Ectopic overexpression of BCL2, BCLXL and BCLW, not BCLB, BFL1 and MCL1, block oxaliplatin induced cell death in spheroid cultures. A) western blot analysis showing overexpression of the various BCL2 family members in spheroid cultures. B) As a reliable antibody for BCLB was not available, BCLB-transduced and control spheroids were analyzed by qRT-PCR analysis for BCLB. C) Spheroid cultures expressing various BCL2 proteins were treated with 50μM oxaliplatin for 24 hr and caspase-3 activity was measured with CaspGlow in differentiated cells (TOP-GFP<sup>low</sup>) and CSCs (TOP-GFP<sup>high</sup>). Significance comparing oxaliplatin-treated control versus various BCL2 family member overexpressing differentiated tumor cells is indicated (** p<0.01).
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Supplementary Figure 3: Differential mitochondrial priming measured with BIM peptide in differentiated tumor cells and CSCs. Spheroid culture (Co100) was subjected to BH3 profiling using decreasing concentration of BIM BH3 peptide. Cells were stained for CD133, treated with BIM BH3 peptides and mitochondrial depolarization was measured with JC1. Mitochondrial depolarization in differentiated tumor cells (CD133\textsuperscript{low}) and CSCs (CD133\textsuperscript{high}) is shown relative to complete depolarization obtained with CCCP. Significance comparing differentiated versus CSCs is indicated (* p<0.05, ** p<0.01, *** p<0.001).
Supplementary Figure 4: ABT-737 and WEHI-539 sensitizes colon CSCs towards various chemotherapies. Spheroid cultures were treated with 100nM ABT-737 or WEHI-539 in combination with (A) 10 mg/ml etoposide, (B) 200 mg/ml 5-FU or (C) 50 mM cisplatin. Caspase-3 activity was measured with CaspGlow in CSCs (TOP-GFP_high). Significance is indicated (* p<0.05, ** p<0.01, *** p<0.001) and is related to oxaliplatin alone treatments.