LEF-1 drives aberrant β-catenin nuclear localization in myeloid leukemia cells

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ABSTRACT

Canonical Wnt/β-catenin signaling is frequently dysregulated in myeloid leukemias and is implicated in leukemogenesis. Nuclear-localized β-catenin is indicative of active Wnt signaling and is frequently observed in acute myeloid leukemia (AML) patients; however, some patients exhibit little or no nuclear β-catenin even where cytosolic β-catenin is abundant. Control of the subcellular localization of β-catenin therefore represents an additional mechanism regulating Wnt signaling in hematopoietic cells. To investigate the factors mediating the nuclear-localization of β-catenin, we carried out the first nuclear/cytoplasmic proteomic analysis of the β-catenin interactome in myeloid leukemia cells and identified putative novel β-catenin interactors. Comparison of interacting factors between Wnt-responsive cells (high nuclear β-catenin) versus Wnt-unresponsive cells (low nuclear β-catenin) suggested the transcriptional partner, LEF-1, could direct the nuclear-localization of β-catenin. The relative levels of nuclear LEF-1 and β-catenin were tightly correlated in both cell lines and in primary AML blasts. Furthermore, LEF-1 knockdown perturbed β-catenin nuclear-localization and transcriptional activation in Wnt-responsive cells. Conversely, LEF-1 overexpression was able to promote both nuclear-localization and β-catenin-dependent transcriptional responses in previously Wnt-unresponsive cells. This is the first β-catenin interactome study in hematopoietic cells and reveals LEF-1 as a mediator of nuclear β-catenin level in human myeloid leukemia.

Introduction

Canonical Wnt signaling is an evolutionary conserved signal transduction pathway strictly controlled during normal development but frequently dysregulated in cancer.¹ In the absence of a Wnt ligand, the central mediator of this signaling pathway, β-catenin, is constitutively phosphorylated by a destruction complex (DC) consisting of GSK3β, CK1, Axin and APC, priming it for subsequent degradation by the proteasome. Upon Wnt ligand binding to the Wnt receptors (Frizzled and LRP5/6), the DC becomes saturated with phosphorylated β-catenin (which cannot be degraded) resulting in cytosolic accumulation of non-phosphorylated β-catenin.² Following nuclear translocation, β-catenin complexes with the T-cell factor (TCF)/lymphoid enhancer factor (LEF) transcriptional regulators and promotes activation of proto-oncogenic Wnt target genes, like c-myc, cyclinD1 and survivin (http://web.stanford.edu/group/nusselab/cgi-bin/wnt/target_genes). Thus, Wnt signaling activation is dependent on the movement of β-catenin into the nucleus, yet this remains a poorly understood process in blood cells. β-Catenin lacks canonical nuclear-localization or -export sequences and its subcellular distribution has instead been associated with multiple factors in context-dependent settings.³ β-Catenin is frequently over-expressed in acute myeloid leukemia (AML)⁴ where its expression correlates with inferior patient survival.⁵ β-Catenin has been shown...
to play a key role in the initiation of AML and chronic myeloid leukemia (CML)." Furthermore, frequent chromosomal aberrations driving AML and CML are known to co-operate with β-catenin. Key to the activation of Wnt signaling is the movement of β-catenin into the nucleus and this is frequently observed in AML. We have previously demonstrated that approximately 10% of primary AML patient blast samples exhibit little nuclear β-catenin expression, despite substantial cytosolic levels, a phenomenon replicated in 10-20% of myeloid leukemia cell lines upon Wnt stimulation. In fact, this is characteristic of normal human hematopoietic stem/progenitor cells (HSPC) which similarly limit β-catenin nuclear-localization, possibly to protect normal HSC from detrimental levels of Wnt signaling. The permissive nuclear-localization of β-catenin observed in myeloid leukemias is therefore aberrant and warrants further investigation.

To better understand β-catenin nuclear-localization mechanisms in myeloid leukemia cells, we generated the first β-catenin interactomes in hematopoietic cells. These analyses have shown that LEF-1, a β-catenin-dependent transcription factor, can also regulate the level of nuclear β-catenin in myeloid leukemia cells. The relative level of nuclear LEF-1 expression correlates with relative nuclear levels of β-catenin in primary AML patient blasts indicating this axis has clinical relevance. Furthermore, the nuclear-localization of β-catenin can be promoted by LEF-1 overexpression and conversely is reduced by LEF-1 knockdown. Finally, we demonstrate LEF-1 expression is suppressed in Wnt-unresponsive cells through rapid proteolytic degradation that is not observed in Wnt-responsive cells. Overall, this study characterizes β-catenin interactions within a hematopoietic context and identifies LEF-1 as a regulator of nuclear β-catenin localization in human leukemia.

Methods

Patient samples, cell culture and β-catenin stabilization
Bone marrow, peripheral blood or leukapheresis samples from patients diagnosed with AML/myelodysplastic syndromes (MDS) (for clinical information see Online Supplementary Table S1) were collected in accordance with the Declaration of Helsinki and with approval of University Hospitals Bristol NHS Trust and London Brent Research Ethics Committee. Mononuclear cells were separated using Ficoll-Hypaque (Sigma-Aldrich, Poole, UK) and samples with ≥80% viability included in the study. K562, HEL, ML-1, U937, THP1 and PLB-985 cell lines (ECACC, Salisbury, UK) were cultured as previously described. For proliferation assays, cell lines were treated with 5 mM MgCl₂, 0.5% Igepal-CA630/NP40) containing complete medium. For co-immunoprecipitation (co-IP), 8 μg of crosslinked β-catenin (Clone-14) or IgG (Clone MOPC-31C) antibody (Becton Dickinson, Oxford, UK) were incubated with 1 mg of either preclotted cytoplasmic or nuclear lysate overnight at 4°C (Online Supplementary Methods). Post-acquisition analyses were performed using FlowJo.

Mass spectrometry and data analyses
Cytosolic or nuclear β-Catenin/IgG co-IPs were prepared and analyzed by mass spectrometry (MS) as detailed in the Online Supplementary Methods. Post-acquisition, duplicate values and proteins detected by only a single peptide were first removed. Tandem Mass Tag (TMT) ratios were imported into Perseus v.1.5.6.0 (Max Planck Institute of Biochemistry, Munich, Germany) followed by logarithmic transformation, normalization (through median subtraction) and removal of proteins not present in at least two of three replicates. A one-sample t-test was performed with significance of protein binding (Log₂ P-value) plotted versus fold change in protein binding (Log₂). The MS proteomics data have been deposited with the ProteomeXchange Consortium (http://proteomecentral.proteomexchange.org) via the PRIDE partner repository with the dataset identifier PXD009305. Interaction specificity was assessed using the publicly available CRAPoMe database (Contaminant Repository for Affinity Purification: http://www.crapome.org).

Assessment of T-cell factor reporter and flow cytometry
Activity of BAR lentiviral construct was performed as previously described. Multi-parameter flow cytometric measurements were acquired using a MACSQuant® Analyzer 10 in conjunction with MACSQuantTyler v.2.8 (Miltenyi Biotec, Bisley, UK) or an Accuri C6 in conjunction with C sampler software v.1.0.264.21 (BD). Post-acquisition analyses were performed using FlowJo.
v.10.5.3 (Tree Star Inc., Ashland, OR, USA). Threshold for TCF reporter fluorescence was set using matched-controls expressing mutant ‘found unresponsive’ fuBAR. Cell viability was assessed using 2 μg/mL propidium iodide (Miltenyi Biotech).

**Statistical analysis**

Statistical analyses were performed using GraphPad Prism v.7.0 (GraphPad Software Inc., San Diego, CA, USA) and Perseus. Correlation was assessed using a Spearman Rank correlation coefficient (R). Significance of difference was assessed using a one-sample or Student t-test and data represents mean±one Standard Deviation (SD) derived from three biological replicates.

**Results**

**Myeloid leukemia cell lines exhibit heterogeneous nuclear β-catenin localization and Wnt activation**

Previously we showed that myeloid leukemia cell lines vary markedly in their capacity for nuclear β-catenin localization upon Wnt stimulation, mimicking the heterogeneity of nuclear β-catenin translocation in AML patients. To investigate the mechanistic basis for this, we selected two sets of myeloid leukemia lines which differed markedly in Wnt signaling output in response to agonist. K562 and HEL were Wnt-responsive cell lines that localized high levels of β-catenin into the nucleus upon treatment with the Wnt agonist, CHIR99021, a GSK3β inhibitor (Figure 1A). Cell viability was not significantly affected by 16 h CHIR99021 treatment ([Online Supplementary Figure S1A and B]). Similar patterns of subcellular localization were observed for the active (non-phosphorylated) form of β-catenin in response to CHIR99021 ([Online Supplementary Figure S1C]). As expected, the phosphorylated forms of β-catenin (Ser33/37/Thr41) were reduced upon CHIR99021 treatment ([Online Supplementary Figure S1C]). Correspondingly, these lines showed robust induction of a TCF reporter (a measure of β-catenin-dependent transcription), whilst cells expressing reporter with a mutated TCF

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**Figure 1.** Myeloid leukemia cell lines exhibit a heterogeneous response to Wnt stimulation. (A) Representative immunoblots showing total β-catenin subcellular localization in myeloid cells following CHIR99021 treatment (GSK3β inhibitor). Lamin A/C and α-tubulin indicate the purity/loading of the nuclear (N) and cytosol (C) fractions, respectively. (B) Representative flow cytometric histograms showing intensity of the TCF-dependent expression of YFP from the ‘β-catenin activated reporter’ (BAR) reporter, or negative control ‘found unresponsive β-catenin activated reporter’ (fuBAR) control (containing mutated promoter binding sites) following treatment with CHIR99021/vehicle control (dimethyl sulfoxide (DMSO)) as above. (C) Summary showing the relative percentage nuclear β-catenin localization (as a proportion of the total) induced in myeloid cell lines upon CHIR99021 treatment. (D) Summary showing the median fluorescence intensity generated from the BAR/fuBAR reporters in myeloid cell lines treated ± CHIR99021. *P<0.05; ****P<0.0001; ns: not significant. MFI: mean fluorescence intensity.
binding site showed no induction (Figure 1B). In contrast, ML-1 and U937 cells had a highly restricted capacity for nuclear β-catenin localization (vs. K562 and HEL cells) despite substantial cytosolic stabilization of the protein (Figure 1B and C). Consistent with this, CHIR99021 treatment of ML-1 or U937 cells caused no detectable activation of the TCF reporter (Figure 1B and 1D). The same patterns of Wnt-responsiveness were observed when cell lines were treated with tWnt3a (Online Supplementary Figure S2). These findings demonstrate that the ability of some leukemia cells to drive a transcriptional response to Wnt agonist is limited by their capacity to permit the nuclear accumulation of β-catenin; we termed these cells Wnt-unresponsive.

**β-catenin interactome analyses reveal contrasting protein interactions between Wnt-responsive and Wnt-unresponsive leukemia cell lines**

Given the wealth of factors previously reported to regulate the nuclear localization of β-catenin,3 we designed an interactome screen of cytosolic and nuclear β-catenin interacting partners from representative Wnt-responsive (K562 and HEL), and Wnt-unresponsive (ML1) cell lines so that we could shortlist candidate factors involved in this process (Figure 2A). Prior to mass spectrometry, we validated the efficiency of β-catenin co-IP (Figure 2B) from both a positive control for high β-catenin expression (SW620 colorectal cells containing mutated APC: demonstrating a 4.0- and 4.3-fold enrichment of β-catenin in cytosol and nuclear fractions, respectively) and in the context of agonist-stabilized β-catenin (HEL: showing a 6.3- and 9.1-fold enrichment). We also confirmed co-IP of a known interactor, TCF-4 (TCF7L2), from the nuclear compartment.

Following MS, raw tandem mass tag (TMT) ratios were processed to generate a set of statistically ranked interactions based on significance of fold-change in protein binding (raw and processed MS data available in Online Supplementary MS data sheets). An extensive profile of β-catenin interactions were observed in K562 cells (225 significantly enriched cytosolic interactions, 118 significantly enriched nuclear interactions) (Figure 3A and B). In contrast, a comparatively sparse interaction profile was observed in ML1 cells (38 significantly enriched cytosolic interactions, 26 significantly enriched nuclear interactions) (Figure 3C and D). An extensive repertoire of β-catenin interactions was also detected in the other Wnt-responsive cell line analyzed, (HEL; 154 significantly enriched cytosolic and 9.1-fold enrichment). We also confirmed co-IP of a known interactor, TCF-4 (TCF7L2), from the nuclear compartment.

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**Nuclear LEF-1 expression correlates with nuclear β-catenin localization in cell lines and primary acute myeloid leukemia patient cells**

We next examined the data from the fractions of Wnt-responsive cells for candidate proteins that could promote the nuclear localization of β-catenin. Several had previously been implicated in negative regulation of β-catenin nuclear localization, GSK3β,\(^{31}\) α-catenin,\(^{32}\) Axin1/2,\(^{33}\) and APC,\(^{34}\) whilst TCF-4\(^{35}\) and LEF-1\(^{36}\) are nuclear localized transcription factors that bind β-catenin. To validate the MS data, we examined the protein expression of these candidates in a panel of myeloid cell lines by immunoblotting. With the exception of APC, which is problematic to blot,\(^{37}\) immunoblotting confirmed MS analysis in that expression of the negative regulators was mostly limited to Wnt-responsive cells making it unlikely they were responsible for restricted nuclear β-catenin in Wnt-unresponsive cells (Figure 5A).

We next examined the expression of two known nuclear β-catenin interactors, TCF-4 and LEF-1, which are ubiquitously expressed in multiple tissues. Both of these proteins are predominantly nuclear in Wnt-responsive cell lines and were absent from the Wnt-unresponsive lines (Figure 5A; matching the proteomics data in Figure 3 and Online Supplementary Figure S3). Of these two proteins, LEF-1 bound β-catenin with higher significance than TCF4 in both K562 and HEL nuclei. This, together with our previous observation that overexpression of TCF-4 actually suppressed β-catenin-dependent transcription\(^{38}\) (and thus is unlikely to promote nuclear β-catenin level), led us to focus our investigation on LEF-1. This protein is known to be dysregulated in AML\(^{39}\) so we examined the clinical relevance of this by correlating relative LEF-1 nuclear localization with that of β-catenin in primary AML blasts. In our cohort of 23 nuclear/cytosol fractionated AML patient samples, we observed a highly significant degree of correlation (Spearman Rank R=0.63, P<0.005) between the rel-
ative nuclear-localized (as a proportion of total) β-catenin protein and relative nuclear-localized (as a proportion of total) LEF-1 protein (Figure 5B and C). Interestingly, the overall frequency of β-catenin protein overexpression (approx. 80%) observed in this largely pediatric AML cohort was higher than previous reports (Online Supplementary Figure S6). Taken together, these data indicate that nuclear β-catenin translocation capacity is linked to LEF-1 expression in myeloid cell lines, and nuclear LEF-1 is a strong clinical predictor of nuclear β-catenin levels in patient-derived AML blasts.

Modulation of LEF-1 expression in myeloid cells regulates nuclear-localization of β-catenin and alters Wnt-responsiveness

The above data indicated a correlative relationship between nuclear LEF-1 level and capacity for nuclear β-catenin localization. To demonstrate causation, we examined the effect of modulating LEF-1 expression on β-catenin localization. Initially, we tested multiple LEF1 shRNA sequences (Online Supplementary Figure S7A) and selected the best (TRCN0000428355) for knockdown of nuclear LEF-1 in the Wnt-responsive K562 and LEF-1 regulates β-catenin nuclear localization

![Figure 2: Experimental strategy for analysis of β-catenin interaction partners in myeloid leukemia cell lines. (A) Wnt-responsive K562/HEL cells and Wnt-unresponsive cells ML-1 cells were treated with CHIR99021 to stabilize β-catenin prior to cytosolic/nuclear fractionation. From these fractions, either an IgG or β-catenin (β-cat) co-immunoprecipitation (co-IP) was performed generating eight samples which were each TMT labeled with a unique isobaric mass tag. All samples were pooled, fractionated, cleaned and analyzed by mass spectrometry. Mass intensities from each tag report the relative peptide abundance in each sample. Quantitative fold-enrichment of β-catenin co-IP was obtained by comparison with fraction-matched IgG co-IP control. Three biological replicates were performed on each cell line. (B) Representative immunoblots showing the efficiency of total β-catenin co-IP performed from the cytosol or nuclear fractions of SW620 cells and HEL cells. Detection of the known interaction partner TCF-4 was used to assess binding partner efficiency. ID: immunodepleted lysate. β-catenin arrows represent full-length/degradation intermediates; TCF-4 arrows indicate 58/79kDa transcriptional isoforms.]
HEL cell lines. This approach resulted in a 72%±7% knockdown in LEF-1 protein in the nuclei of K562 cells and an 89%±4% knockdown in HEL cells (Figure 6A). Somewhat lower levels of LEF-1 knockdown were observed in CHIR99021-treated cells (65%±19% and 83%±7%, respectively) probably a result of LEF1 being a Wnt target gene and thus being induced through Wnt agonist treatment. LEF-1 knockdown perturbed nuclear localization of β-catenin by approximately one-third (28%) in K562 following CHIR99021 treatment, proportionate to control cells. This reduction was accentuated in HEL cells (41%) which corresponded to the greater degree of LEF-1 knockdown in these cells (Figure 6B). The knockdown of LEF-1 protein resulted in significantly reduced growth of both K562 and HEL cells at multiple time points across a range of serum concentrations (Figure 6C). Use of a second LEF1 shRNA and a different method of Wnt stimulation (rWnt3a) resulted in a similar finding (Online Supplementary Figure S7B and C). These data suggest LEF-1 promotes the optimal translocation of β-catenin into the nucleus of Wnt-responsive cells and partly contributes to their growth.

Next, we examined whether LEF-1 expression was sufficient to permit nuclear-localization of β-catenin. To establish this, we stably over-expressed LEF1 in the Wnt-unresponsive (and LEF-1 negative) U937 and ML1 cells. Overexpression of LEF-1 resulted in substantial cytosolic expression of the full-length LEF-1 protein (50kDa) but weak nuclear expression; despite this, we observed a dramatic increase in nuclear localized β-catenin in both ML1 (4-fold) and U937 (2.3-fold) cells over-expressing LEF1 following CHIR99021 treatment (Figure 6D and E). This disparity may be explained by the abundant expression of a short-form of LEF-1 in the nucleus (25-30kDa) that was absent in Wnt-responsive lines (discussed below). These effects were mirrored using Wnt5a treatment (Online Supplementary Figure S7C) and we also showed that LEF-1 overexpression was able to facilitate nuclear localization of β-catenin in two further AML cell lines (PLB-958 and THP1) (Online Supplementary Figure S7D). These data demonstrate that overexpression of LEF-1 can significantly increase the capacity for nuclear β-catenin localization in Wnt-unresponsive cell lines.

To assess the impact of LEF-1 knockdown (and subsequent nuclear β-catenin reduction) on Wnt signaling, we measured TCF reporter activity. As predicted, Wnt signaling induction in Wnt-responsive K562 and HEL cells following CHIR99021 treatment was severely diminished.
following LEF-1 knockdown (Figure 7A and B). A significant reduction in Wnt signaling output was also observed following use of an alternative LEF1 shRNA in response to CHIR99021 or Wnt3a stimulation (Online Supplementary Figure S8). Assessment of the TCF reporter activity in the LEF1 over-expressing lines was not possible due to the confounding expression of the GFP selectable marker; instead, we examined the protein expression of the classic downstream Wnt target proteins survivin, c-MYC and cyclinD1. We found that LEF1 overexpression significantly enhanced the expression of these proteins following CHIR99021 treatment (Figure 7C and D), indicating that

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**Figure 4.** Mass spectrometric analyses identify putative novel interaction partners for β-catenin in myeloid leukemia cell lines. Bar graphs summarizing the average fold change in protein binding (relative to matched IgG co-immunoprecipitation) for novel, significant β-catenin interactions observed in (A) K562 cytosolic, (B) K562 nuclear, (C) ML1 cytosolic, and (D) ML1 nuclear fractions. Significant proteins only are shown (red dots in Figure 3 panels) with a frequency on the CRAPome database of ≤10% and known interactions were removed. Red asterisks represent proteins of particular significance to myeloid leukemias and/or Wnt signaling (see Results). Proteins are ranked along x-axis according to statistical significance (most significant on the left) (see also Online Supplementary Table S2).
the observed translocation of β-catenin was sufficient to generate a transcriptional response. A caveat to these data is that these observations cannot be linked exclusively to the modulation of β-catenin nuclear localization since LEF-1 expression is a co-variable in these experiments. Taken together, these data indicate LEF-1 can regulate the nuclear level of β-catenin and our observations are also consistent with a concomitant regulation of Wnt signaling activity.

**LEF-1 protein is proteolytically cleaved in Wnt-unresponsive cells**

In the experiments above, overexpression of full-length LEF-1 in Wnt-unresponsive cell lines resulted in the emergence of a 25-30kDa species of LEF-1 protein. To evaluate the contribution of this short LEF-1 form with the increased nuclear β-catenin level observed in these cell lines we examined the β-catenin binding capacity of this species. We performed β-catenin co-IP from both the cytosol and nucleus of CHIR99021 treated LEF1-overexpressing U937 cells and immunoblotted for LEF-1 protein. Both the full-length and short-forms of LEF-1 protein co-immunoprecipitated with β-catenin from the cytosol implying both forms bind β-catenin in this cell line. In the nuclear fraction, β-catenin preferentially co-immunoprecipitated with the short LEF-1 form, though the proportion of short-form was highly enriched in this fraction (Figure 8A; input lane). These data confirm that the short-form LEF-1 has β-catenin binding ability and could mediate the increased nuclear β-catenin translocation observed in Wnt-unresponsive cell lines above upon LEF-1 overexpression with CHIR99021 treatment.

Finally, we examined the origin of the short LEF-1 polypeptide. This variant was unlikely to have derived from alternative splicing since the ectopic expression of LEF-1 was driven from cDNA. Furthermore, these short forms of LEF-1 were also observed endogenously in colorectal cell lines (Online Supplementary Figure S9A) and in primary AML samples featured in Figure 5 (full blots in Online Supplementary Figure S9B). Therefore, the presence of short-form LEF-1 was most consistent with a proteolytic cleavage mechanism. To investigate this, nuclear lysates from both K562 control and U937-LEF1 cells ± protease inhibitor cocktail (PIC) were incubated at 37°C and the relative proportions of full-length to short LEF-1 forms in each cell line observed over 0-60 min. In K562 control cells, full-length LEF-1 protein was stable with no detectable breakdown of full-length LEF-1 protein even in the absence of PIC (Figure 8C). In contrast, the full-length LEF-1 band present in U937-LEF-1 cells was reduced with concomitant enrichment of the short-form polypeptide after a 10 min incubation at 37°C (Figure 8C). The removal of the PIC reduced the half-life of full-length LEF-1 by approximately 50% with degradation occurring within 5 min. Together these data are consistent with the short-form of LEF-1 arising through proteolytic processing.
Figure 6. Modulation of LEF-1 expression affects relative nuclear β-catenin level. (A) Representative immunoblots showing the level and subcellular localization of total β-catenin and LEF-1 protein in K562 and HEL cells in response to control/LEF1 shRNA±CHIR99021 treatment. Shorter 6-hour (hrs) CHIR99021 treatments were used for these cells to minimize the LEF-1 expression induced through Wnt stimulation. (B) Summary graph showing the relative percentage nuclear β-catenin localization induced upon 6-hr CHIR99021 treatment of K562 or HEL cells±control/LEF1 shRNA. (C) Effect of LEF1 shRNA on the proliferation of K562 and HEL at different concentrations of serum after 24, 48 and 72 hrs. (D) Representative immunoblots showing the level and subcellular localization of total β-catenin and LEF-1 protein in ML-1 and U937 cells in response to control/LEF1 overexpression (o/e) ± 16-hrs CHIR99021 treatment. The positions of full-length (FL) and short-forms (SF) of LEF-1 protein on the blot are indicated by arrows. Lamin A/C and α-tubulin were used to assess fraction purity and protein loading. (E) Summary graph showing the relative percentage nuclear β-catenin localization induced upon CHIR99021 treatment of ML-1 and U937 cells for 16 hrs ± control/LEF1 overexpression. *P<0.05; **P<0.01.
Figure 7. Modulation of LEF-1 expression affects downstream Wnt signaling. (A) Representative flow cytometric histograms showing intensity of the TCF reporter (BAR) in K562 and HEL cells treated with control/LEF1 shRNA ± CHIR99021. (B) Summary data showing the median fluorescence intensity generated from the BAR reporter in K562 and HEL cells treated with control/LEF1 shRNA ± CHIR99021. (C) Representative immunoblots showing expression of known Wnt target proteins survivin, c-MYC and cyclinD1 in ML-1 and U937 cells in response to control/LEF1 overexpression ± CHIR99021. Lamin A/C and α-tubulin were used to assess fraction purity and protein loading. (D) Summary data showing the relative fold-change in nuclear protein expression of classic Wnt targets survivin, c-MYC and cyclinD1 in CHIR99021 treated ML1 and U937 cells over-expressing LEF1. Dashed line represents relative level (=1) present in nuclei of CHIR99021-treated ML1 and U937 cells expressing control plasmid. *P<0.05; **P<0.01; ***P<0.001. ns: not significant.

Figure 8. Short-form LEF-1 is proteolytically-derived and is capable of β-catenin binding. (A) Representative immunoblot showing total β-catenin co-immunoprecipitation (Co-IP) from cytosolic and nuclear fraction of LEF1-over-expressing U937 cells. Total β-catenin and LEF-1 protein levels are shown. ID: immunodepleted lysate. Representative immunoblot showing LEF-1 protein levels in (B) K562 control and (C) U937-LEF1 nuclear lysates ± protease inhibitor cocktail (PIC) during time course incubation at 37°C. (D) Representative immunoblots showing LEF-1 protein levels in K562 control nuclear lysate after mixing with U937 control whole cell lysate (1:1 protein concentration) ± PIC during time course incubation at 37°C. Nuclear lamin A/C detection was used to assess protein loading.
These data suggest that LEF-1-targeted proteases are active in Wnt-unresponsive U937 cells, but inactive/absent in Wnt-responsive K562 cells. To test this, we mixed nuclear lysates from K562 cells directly with whole cell lysates from U937 control cells (which are LEF-1 negative). Remarkably, the previously stable full-length LEF-1 band present in K562 cells exhibited marked and rapid reduction with a concomitant increase in the short-form polypeptide, a process exacerbated by PIC removal (Figure 8D). These data confirm that constitutive LEF-1 degradation mechanisms are active in Wnt-unresponsive cells but are absent/inhibited in Wnt-responsive cells.

Discussion

Canonical Wnt signaling has emerged as one of the most frequently disregulated signaling pathways in myeloid neoplasms which has led to considerable interest in targeting this pathway. The central mediator β-catenin represents an appealing therapeutic target because of its leukemogenic role,7,8 prognostic influence,9 and functional redundancy in normal hematopoietic development.10 The characterization of the hematopoietic interactome of β-catenin is, therefore, of considerable interest, and the experimental approach adopted in this study was validated by the identification of multiple known β-catenin partners, such as Axin, TCF-4, α-catenin and APC. Associations with key hematopoietic transcription factors, C/EBPζ and GATA-1, were identified and these are known to co-occupy genomic sites with TCF-4 (a known β-catenin partner) during hematopoietic development.41 We confirmed the previously reported interaction of β-catenin with the oncogenic fusion protein BCR-ABL in CML, which is present in K562 cells.9 In addition to known partners, we identified putative novel interactions which have known relevance to leukemia and/or Wnt signaling (MBD3,14,16 PRC1,15,17 MSI2,18-20 LIN28B,21-23 and WT131). To investigate the nuclear localization mechanisms of β-catenin, we focused on LEF-1, given that it was one of the most significantly enriched interactors present in Wnt-responsive cell nuclei, and its nuclear expression was highly predictive of β-catenin nuclear-localization in both myeloid cell lines and primary AML blasts. A role for LEF-1 in mediating nuclear-localization of β-catenin was confirmed using both knockdown and overexpression approaches. LEF-1 contains a nuclear localization sequence and is known to shuttle between cytoplasmic and nuclear compartments.42 This suggests LEF-1 could serve as a cytosolic-nuclear chaperone for β-catenin. However, the high nuclear:cytosol ratio of LEF-1 expression observed in Wnt-responsive cells (K562 and HEL) would be more consistent with LEF-1 serving as a nuclear retention factor for β-catenin, as has previously been demonstrated for other Wnt signaling components.43 This is the first evidence of LEF-1 contributing to β-catenin nuclear-localization capacity in human myeloid leukemia cells; however, our data cannot exclude the role of other factors in this process. Nuclear export mechanisms may also be influential in limiting nuclear accumulation of β-catenin in Wnt-unresponsive cell lines. In particular, RanBP1 was an interacting partner detected for β-catenin in ML-1 nuclei but absent in K562/HEL nuclei, and may warrant further study given the documented role for RanBP1 in mediating β-catenin export.43 Two factors recently implicated in nuclear β-catenin regulation including RAPGEF544 and Twa145 were not detected in our analyses, suggesting their interaction with β-catenin is context-dependent.

A strong correlation between the relative levels of nuclear localized β-catenin and LEF-1 was identified in primary AML samples suggesting this relationship may have clinical relevance. Indeed, expression of constitutively active LEF-1 in HSPC induced AML in mice,46 and LEF-1 expression promotes the survival of myeloid leukemia cell lines.47 In our study, we observed significantly inhibited growth in K562 and HEL cells harboring LEF-1 knockdown (Figure 6C). Given the adverse prognosis of nuclear β-catenin in AML,48 and recent studies demonstrating its therapeutic merit in AML models,49-51 LEF-1 represents an attractive target in myeloid leukemia. Small molecule inhibitors of β-catenin: TCF/LEF interaction are under development and have shown efficacy in leukemia treatment.52-54 It would be interesting to observe if this efficacy is partly driven by reducing the level of nuclear β-catenin in leukemia cells. This study focused on the role of LEF-1 driving aberrant β-catenin nuclear-localization in myeloid leukemia; however, given the frequency of Wnt/β-catenin dysregulation in human cancer, this axis could be active in other malignancies.

We also observed that LEF-1 is proteolytically degraded in Wnt-unresponsive leukemia cells resulting in the emergence of short-form LEF-1 proteins (25-30kDa). These were smaller than the 38kDa transcriptional isoforms previously reported to derive from alternative splicing or alternative promoter usage55,56 which serve as dominant-negative inhibitors of Wnt signaling because they lack the β-catenin binding domain necessary to initiate transcription.57 In contrast, the proteolytic fragments observed in this study retained β-catenin binding capacity (Figure 7A) and did not appear to have dominant-negative function (Wnt targets were still activated) (Figure 6C and D). Under the conditions of the experiment, however, where LEF-1 is being over-expressed, the abundance of these fragments may have a dominant effect on β-catenin retention in the nucleus, whereas under normal circumstances, it is likely that degradation serves to remove LEF-1 and suppress the nuclear retention of β-catenin. This post-translational regulation of LEF-1 has parallels with embryonic stem cells where proteolytic cleavage of TCF-3 (TCF7L1), a closely related family member, removes it from target genes when differentiation signals trigger the suppression of Wnt signaling.58 In leukemia cells, this proteolytic cleavage could be mediated by NLK (Nemo-like kinase) which binds the E3 ubiquitin-ligase NARF (NLK associated RING finger protein) and reportedly induces ubiquitylation (and proteasomal degradation) of LEF-1 in co-operation with the E2 conjugating enzyme E2-25K,59 a mechanism previously reported to be active in leukemia cells.58

In summary, our study has made three key findings: firstly, the generation of the first β-catenin interactomes in leukemia cells; secondly, the characterization of LEF-1 as a regulator of nuclear β-catenin localization in leukemia; and, finally, the demonstration of post-transcriptional proteolytic degradation mechanisms for controlling LEF-1 expression in myeloid leukemia cells.
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