REVIEW

Wnt signaling in triple-negative breast cancer

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Wnt signaling regulates a variety of cellular processes, including cell fate, differentiation, proliferation and stem cell pluripotency. Aberrant Wnt signaling is a hallmark of many cancers. An aggressive subtype of breast cancer, known as triple-negative breast cancer (TNBC), demonstrates dysregulation in canonical and non-canonical Wnt signaling. In this review, we summarize regulators of canonical and non-canonical Wnt signaling, as well as Wnt signaling dysfunction that mediates the progression of TNBC. We review the complex molecular nature of TNBC and the emerging therapies that are currently under investigation for the treatment of this disease.

Oncogenesis (2017) 6, e310; doi:10.1038/oncsis.2017.14; published online 3 April 2017

INTRODUCTION

Breast cancer represents one of the most significant disease burdens of any cancer worldwide. Today, women have a one in eight chance of developing breast cancer over their lifetime, a risk that is significantly increased if they have inherited harmful mutations in *BRCA1* or *BRCA2*.¹ However, breast cancer is a complex, heterogeneous disease characterized by a great multitude of aberrations at the genomic and molecular level, which can manifest in dysregulated signaling pathways. A hallmark of many cancers is aberrant regulation of the Wnt signaling pathway, and breast cancer is no exception.²

Triple-negative breast cancer (TNBC), an aggressive subtype of breast cancer with a poor prognosis,³ is characterized by tumors that do not express estrogen receptors (ERs) or progesterone receptors (PRs), nor display an overexpression of human epidermal growth factor receptor 2 (HER2).⁴ Therapies targeted against HER2-positive breast cancers, such as trastuzumab (Herceptin),⁵ and those targeted against ER-positive breast cancers, such as tamoxifen,⁶ have no therapeutic benefit to individuals with the TNBC subtype. Surgical intervention and chemotherapy have been the major treatment avenues for TNBC; however, recently developed small molecules and immunotherapeutics⁷ are showing promise. In this review, we will summarize the involvement of dysregulated Wnt signaling in the progression of TNBC and TNBC stem cells, as well as the emerging therapies that are currently under investigation.

THE WNT SIGNALING PATHWAY

The Wnt/ β -catenin pathway (canonical pathway)

There are currently two models of canonical Wnt/ β -catenin signaling. In the classical model, the destruction complex remains intact in the absence of Wnt stimulation (Figure 1a). Casein kinase 1 (CK1) primes β -catenin for destruction by phosphorylating Ser45, which then allows activated Glycogen synthase kinase 3 (GSK3) to

phosphorylate β -catenin at Ser33, Ser37 and Thr41.⁸ The phosphorylated residues of β -catenin interact with the β -propeller domain of the E3 ubiquitin ligase β -TrCP, which then ubiquitinates β -catenin, thus targeting it for proteasomal degradation.⁹ Wnt/ β -catenin signaling is initiated by Wnt ligands binding to a Frizzled receptor (FZD), as well as the co-receptors low-density lipoproteins 5/6 (LRP5/6). This results in activation of FZD, permitting binding of Dishevelled (Dvl)¹⁰ and phosphorylation of one or more cytoplasmic motifs of LRP5/6. A single phosphorylated motif is sufficient to activate Wnt signaling.¹¹ Phosphorylated LRP5/6 can then interact with Axin. This interaction destabilizes the β -catenin destruction complex, which requires Axin as a scaffold and contains Dvl, the serine–threonine kinases casein kinase 1 α/β (CK1), glycogen synthase kinase 3 α/β (GSK3) and adenomatous polyposis coli (APC).¹² Destabilizing the destruction complex prevents phosphorylation of β -catenin, which then accumulates in the cytosol before translocating to the nucleus. Once there, it binds to Transcription factor/lymphoid enhancer-binding factor (TCF/LEF) transcription factors and displaces transcriptional repressor Groucho to initiate the transcription of Wnt target genes (Figure 1b).¹³ In the new model,^{14–16} the destruction complex is stabilized by Axin in both the presence and absence (Figure 1c) of Wnt activation, and β -catenin is degraded through phosphorylation-mediated recognition by β -TrCP in the intact complex. This allows newly synthesized β -catenin to accumulate in the cytosol before nuclear translocation (Figure 1d). This was demonstrated through co-immunoprecipitation, whereby β -catenin phosphorylated at Ser33/Ser37/Thr41 was shown to interact with the destruction complex upon Wnt activation, which also disrupted the interaction of β -TrCP with the Axin1– β -catenin complex.¹⁵ It has also been proposed that GSK3 inhibition, and thus β -catenin translocation after Wnt activation, is mediated through the sequestration of GSK3 inside multivesicular endosomes.¹⁷ This further demonstrates the complexity of Wnt signaling.

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Received 11 November 2016; revised 9 January 2017; accepted 24 January 2017

Figure 1. Classical and new Wnt/β-catenin pathway canonical and non-canonical pathways. (a) Overview of the ‘classical’ model of Wnt/β-catenin signaling in OFF state with no ligand bound to FZD receptor. (b) Overview of the ‘classical’ model of Wnt/β-catenin signaling pathway in ON state where Wnt ligand is bound to FZD receptor. (c) Overview of ‘new’ model of Wnt/β-catenin signaling in OFF state with no ligand bound to FZD receptor. (d) Overview of the ‘new’ model of Wnt/β-catenin signaling in ON state with Wnt ligand bound to FZD receptor. (e) Overview of Wnt planar cell polarity (PCP) pathway in ON state. Wnt binds multiple receptors including FZD and co-receptors ROR and Ryk. This activates Rac and Rho, which activate ROCK and c-Jun N-terminal kinase (JNK), respectively, leading to actin polymerization and regulates cytoskeletal arrangements. (f) Overview of Wnt/Ca²⁺ pathway in ON state. Wnt is able to bind FZD, Ryk to initiate signal transduction, which is effected through Dvl and G proteins (α, β, γ). Gene transcription is induced through proteins PKC, CaMKII and Cn (Calcineurin)-activating transcription factors.

Planar cell polarity pathway

The planar cell polarity pathway (Figure 1e) is a non-canonical, β-catenin-independent pathway that regulates cellular organization and polarity, partly through cytoskeletal organization.¹⁸ Wnt ligands, such as Wnt5a, bind to FZD receptors and co-receptors, including

ROR,¹⁹ Ryk²⁰ and PTK.²¹ Dvl interacts with Rac¹²² and Dvl-associated activator of morphogenesis 1 (DAAM1). Rac1 activates c-Jun N-terminal kinase, leading to actin polymerization,²² while DAAM1 activates Rho, which in turn activates Rho-associated kinase (ROCK) to regulate cellular cytoskeletal arrangements.²³

Figure 2. Molecular structures of the key Wnt signaling proteins and interactions. **(a)** X-ray crystal structure of the *Xenopus* Wnt8 complex with the mouse FZD8 cysteine-rich domain (PDB 4F0A). The key structural regions of the Wnt fold are highlighted, as are the major Wnt-interacting regions of the CRD. **(b)** X-ray crystal structure of the Smoothed receptor (PDB 5L7D), a Class F G protein-coupled receptor, related to FZD. The key structural regions of Smo are highlighted, as well as helix 8, which is of relevance for Dishevelled binding by FZD. **(c)** Schematic representation of the location of the DIX, PDZ and DEP domains within Dvl. **(d)** X-ray crystal structure of the DIX homodimer (PDB 4WIP). **(e)** X-ray crystal structure of the PDZ domain bound to a peptide (red; the peptide-binding site is shown in yellow). **(f)** X-ray crystal structure of a DEP homodimer (PDB 5LNP), highlighting residues known to affect Wnt signaling (shown in red). **(g)** Model of the LRP6 ectodomain generated by molecular dynamics flexible fitting of the crystal structures of the P1E1–P2E2 domains (PDB 3S94) and P3E3–P4E4 domains (PDB 4A0P), and a homology model of the LDL-R type A domains (generated in Prime, based on the crystal structure of the LDL receptor ectodomain (PDB 1N7D)) to the electron microscopy structure (EMDatabank accession 1964). Gaps in the crystal structures and between the various components modeled using Prime. **(h)** X-ray crystal structure complex of the cysteine-rich domain 2 of Dickkopf with Kremen and the LRP6 P3E4–P4E4 domains (PDB 5FWW).

The Wnt/Ca²⁺ pathway

The Wnt/Ca²⁺ pathway (Figure 1f) is activated through Wnt binding to FZD, which interacts with G proteins and Dvl.²⁴ These interactions can activate cGMP-specific phosphodiesterase or phospholipase C, resulting in a release of intracellular calcium. This results in the activation of downstream signaling proteins PKC, calcineurin and CaMKII.²⁵ CaMKII activates nuclear factor of activated T-cells, which can regulate cell adhesion and migration.²⁶ Wnt5a induces activation of CaMKII-dependent Wnt/Ca²⁺ signaling. CaMKII phosphorylates transforming growth factor β -activated kinase, which activates Nemo-like kinase.²⁷ This

cascade antagonizes canonical Wnt/ β -catenin signaling by Nemo-like kinase phosphorylation of TCF4 and prevents the β -catenin–TCF4 complex from binding to DNA.²⁸

Wnt ligands

To date, 19 members of the Wnt family have been identified in mammals, all ranging between 350 and 400 amino acids in length and characterized by a conserved fold containing a conserved motif of 24 cysteine residues (Figure 2a).²⁹ Wnt ligands are modified by lipidation, specifically, the addition of a palmitoyl group to a conserved serine by the membrane-bound

O-acyltransferase Porcupine.³⁰ Wnt lipidation is crucial for secretion from the endoplasmic reticulum³¹ and essential for Wnt function. Wnt lipidation was initially suggested to occur at Cys77 of Wnt3a (cysteine 3 of the Wnt fold);³² however, lipidation at this cysteine has been conclusively disproven by crystallographic,³³ mutational³¹ and imaging studies.³⁰

FZD receptors

FZD receptors are a group of 10 membrane proteins featuring an extracellular cysteine-rich domain (CRD) and a seven-transmembrane domain.³⁴ Along with the Smoothed receptor (Smo), the FZDs comprise the family of Class F G protein-coupled receptors. The crystal structure of XWnt8 in complex with the mouse FZD8 CRD³³ revealed an unusual interaction involving the direct binding of the Wnt lipid to a binding site on one side of the CRD (the 'thumb' region), as well as the binding of the region from cysteines 19 to 22 of XWnt8 to the other side of the CRD (the 'index finger' region; Figure 2a).³⁵ Although no complete structures are available for any FZD, several structures of Smo are known,^{36–38} most recently including both the CRD and seven-transmembrane regions (Figure 2b),³⁹ which are suggestive of the likely structure of FZD.

Disheveled

Three Dvl homologs are known (Dvl1/2/3), sharing high overall sequence similarity.⁴⁰ Dvl consists of three structurally defined domains: the DIX, PDZ and DEP domains. These three domains are separated by large insertions of unknown structure (Figure 2c); however, some functional significance has been ascribed to conserved sequences within the unstructured regions.⁴¹

Dvl polymerizes via the head-to-tail interaction of its DIX domain (Figure 2d). The DIX domain also mediates interaction with Axin.⁴² Mutations (V67A, K68A, Y27D) in the polymerization interface of the DIX domain strongly suppress Wnt signaling.⁴³ The PDZ domain of Dvl (Figure 2e) interacts with a conserved motif in the FZD C-terminal (KTxxxW).²⁴ The PDZ–FZD interaction is relatively weak, and is likely supplanted by interactions of the DEP domain with FZD. Greater insight in the role of the DEP domain in Wnt signaling was recently revealed, with this domain shown to bind as a monomer to FZD, then undergo subsequent domain swapping to assemble Wnt signalosomes. Furthermore, upon Wnt stimulation, DEP domain swapping initiates DIX-dependent Dvl and Axin polymerization, leading to the inhibition of GSK3 and Wnt signal transduction. Mutants (E499G, D460K, G436P, K438M, D449I and D452I) in the DEP domain strongly diminish Wnt signaling upon Wnt stimulation (Figure 2f).^{44,45} Dvl has also been shown to promote ubiquitination-mediated FZD degradation by RNF43.⁴⁶ This finding suggests a dual agonist/antagonist role for Dvl in Wnt signaling.

Low-density lipoprotein receptor 5/6

The extracellular domain of LRPs consists of four β -propeller repeats interspersed with epidermal growth factor repeats, followed by three LRP type A repeats (Figure 2g).¹⁰ The majority of Wnts bind to the first β -propeller/epidermal growth factor repeat (P1E1–P2E2), although Wnt3 and Wnt 3a preferentially bind to the second repeat (P3E3–P3E4).⁴⁷ Wnt3 and Wnt3a binding to LRPs is competitively inhibited by Dickkopf binding to LRP (Figure 2h).^{48–50} The intracellular action of LRP5/6 is less clearly understood, although it is known that Wnt activation initiates phosphorylation of the intracellular PPPSPxP motifs of LRP5/6 by GSK3 and CK1, allowing the recruitment of Axin.⁵¹ Importantly, it has also been shown that without the FZD–Dvl interaction, Wnt is unable to induce phosphorylation of LRP6, reinforcing the complex interplay of proteins involved in Wnt signaling.^{24,51}

ROR family receptor tyrosine kinases

The ROR family of receptor tyrosine kinases consists of two evolutionarily conserved members, ROR1 and ROR2.⁵² The ROR ectodomains feature a FZD-type CRD most closely related to that of the skeletal muscle receptor tyrosine-protein kinase.⁵³ ROR2 is involved in Wnt5a-mediated signaling; Wnt5a binding to ROR2 initiates ROR2 homodimerization, stimulating autophosphorylation at Tyr646.⁵⁴ It has been demonstrated that Wnt5a and Wnt3a bind to ROR2; however, only Wnt5a is able to initiate the activation of the ROR2 signaling cascade.⁵⁵ Recently, high expression of ROR1 has been demonstrated in TNBC cell lines, where it interacts with CK1 ϵ to promote tumor survival and growth after stimulation with Wnt5a to activate phosphoinositide 3-kinase (PI3K)/AKT signaling.⁵⁶

DEAD-box helicases

DEAD-box helicases (DDXs), named for a conserved amino-acid sequence in their ATP-binding domain (Asp-Glu-Ala-Asp), belong to a highly conserved family of ATP-dependent DNA/RNA helicases.⁵⁷ They consist of a highly conserved helicase core with two domains, displaying high similarity to the recA bacterial DNA recombination protein (Figure 3).⁵⁸ These multifunctional proteins have roles in translation initiation, pre- and post-translational modifications, DNA repair, microRNA (miR) processing, ribosome biogenesis and RNA decay.^{59–61} Furthermore, DDXs have been recently implicated in breast tumorigenesis and activation of cancer stem cell (CSC) stemness through various pathways, including Wnt.^{62,63} DDXs can be regulated by β -catenin/TCF-driven transcription and have also been shown to regulate upstream Wnt signaling. The role of DDXs is discussed in further detail later in the section titled 'DDXs, Wnt and TNBC'.

BREAST CANCER SUBTYPES

Breast cancer is a diverse and complex disease, broadly characterized by four molecularly distinct subtypes, including luminal A, luminal B, HER2-overexpressed and triple-negative breast cancer (TNBC).⁶⁴ The luminal A subtype is characterized as ER/PR-positive and HER2-negative, expressing Bcl-2, cytokeratin 8/18 and low Ki67.⁶⁵ Luminal B subtypes are more aggressive ER⁺ breast tumors, characterized as HER2[–] with high Ki67, or HER2⁺, PR[–] and ER⁺,⁶⁶ with cyclin B1 overexpression.⁶⁷ The HER2 subtype is characterized by amplification of the ERBB2/HER2 gene.⁶⁸

TNBC, including basal-like and claudin-low subtypes, accounts for 10–20% of breast cancers and is characterized by a lack of PR, ER and HER2 overexpression.⁶⁹ TNBC patients present with higher

Figure 3. X-ray crystal structure of the ATP-binding and C-terminal helicase domains of the DEAD-box helicase DDX3 (PDB 5E7M). AMP-PNP, a non-hydrolyzable ATP analog, is shown in spheres in the ATP-binding site.

Table 1. Summary of the six molecular subtypes of TNBC characterized by Lehmann et al.⁷³

TNBC subtype	Gene ontology pathway (GOP)	Genes found in GOPs with Wnt association
Luminal androgen receptor (LAR)	Steroid pathway Androgen metabolism Fatty-acid synthesis	<i>FKBP5</i> ⁷⁴ <i>SPDEF</i> ⁷⁵ <i>FASN</i> ⁷⁶
Mesenchymal (M)/mesenchymal stem-like (MSL)	EMT	<i>MMP2</i> , ⁷⁷ <i>SNAI2</i> , ⁷⁷ <i>TCF4</i> , ⁷⁷ <i>TWIST1</i> , ⁷⁷ <i>ZEB1</i> ⁷⁸
Basal 1 (BL1)/Basal 2 (BL2)	Wnt/ β -catenin signaling DNA damage	<i>CTNNB1</i> , ⁷³ <i>DKK2/3</i> , ⁷³ <i>TCF4</i> , ⁷³ <i>TCF7L2</i> , ⁷³ <i>CCND2</i> , ⁷³ <i>FZD4</i> , ⁷³ <i>CAV1</i> , ⁷³ <i>CAV2</i> ⁷³ <i>CHEK1</i> , ⁷⁹ <i>FANCA</i> , ⁸⁰ <i>FANCG</i> , ⁸⁰ <i>MSH2</i> , ⁸¹ <i>RAD21</i> ⁸²
Immunomodulatory (IM)	Proliferation/cell cycle JAK/STAT cytokine pathway IL7 pathway	<i>AURKB</i> , ⁸³ <i>PLK1</i> , ⁸⁴ <i>CENPA</i> , ⁸⁴ <i>BUB1</i> , ⁸⁴ <i>CCNA2</i> , ⁸⁵ <i>PRC1</i> , ⁸⁶ <i>MYC</i> , ⁸⁷ <i>NRAS</i> ⁸⁸ <i>CCR2</i> , ⁸⁹ <i>CCR5</i> ⁹⁰ <i>IL7</i> ⁹¹

Abbreviations: EMT, epithelial–mesenchymal transition; IL, interleukin; TNBC, triple-negative breast cancer. Various gene ontology pathways were found to be enriched in the LAR, MSL, BL1, BL2 and IM subtypes. Analysis of the genes enriched in these pathways identified genes associated with Wnt signaling.

incidence of distant disease recurrence within 3 years of diagnosis, with a high frequency of visceral metastases.⁷⁰ The prognosis for patients diagnosed with TNBC is poor, with patients who respond poorly to adjuvant treatment exhibiting worse outcomes.⁴

TNBC subtypes

TNBC has been categorized into a number of distinct molecular subtypes; however, there remains much intertumoral mutational and transcriptional heterogeneity within these subtypes. The molecular heterogeneity of TNBC confounds the clinical approach to TNBC treatment. TNBCs are characterized by high clonal frequencies of single gene mutations in the key tumorigenesis driver genes, including *TP53*, *PIK3CA* and *PTEN*, indicating that clonal evolution of these mutated genes is an early event in TNBC development.⁷¹ However, mutation frequencies within these genes are not uniform among TNBC cases.^{71,72}

Lehmann et al.⁷³ determined gene expression signatures in 587 TNBC cases from 21 breast cancer data sets and identified six molecularly distinct TNBC subtypes. These include basal-like 1, basal-like 2, immunomodulatory, mesenchymal (M), mesenchymal stem-like (MSL), and luminal androgen receptor (LAR). These subtypes, various gene ontology pathways and associated Wnt genes are described in Table 1. Recent RNA profiling performed by Burstein et al.⁹² showed overlap of LAR and MES subtypes based on Lehmann's gene expression profiling, but was unable to reproduce all observations.⁹² The findings of both of these studies indicate the presence of at least four molecularly distinct and stable TNBC subtypes, defined as LAR, mesenchymal (MES), basal-like immune-suppressed (BLIS) and basal-like immune-activated (BLIA).⁹² Furthermore, these studies suggest molecular targets for the development of therapeutics specific to the treatment of TNBC.

LAR subtype

The LAR subtype accounts for ~10% of TNBCs, whereby tumor cells exhibit positive staining for androgen receptors (ARs) and are driven by AR signaling.^{69,73} The LAR subtype of TNBC displays genomic amplification of *CCND1*, a gene regulated by the Wnt/ β -catenin pathway.⁹² There is some discordance within the literature in regards to the prognostic utility of AR status, with studies indicating no significant effect on survival rates associated with AR expression,⁹³ although AR⁺ TNBC individuals have been shown to have a positive clinical response to the nonsteroidal antiandrogen, bicalutamide.^{94,95} In a study designed to test the benefit of tamoxifen on ER[−] and TNBC patients, it was found that expression of AR⁺ versus AR[−] individuals predicted a decreased recurrence rate and treatment benefit with AR⁺ patients;⁹⁶ this is a result of tamoxifen exhibiting agonist activity on AR-expressing cells.⁹⁷

MES subtype

The MES subtype, encompassing Lehmann's M, MSL and claudin-low subtypes, is characterized by the overexpression of genes associated with cellular motility, proliferation and growth signaling pathways.^{73,92} MES subtypes have high expression of platelet-derived growth factor, insulin-like growth factor 1 and c-kit.⁹² MES tumors express mesenchymal stem cell markers, including the breast stem cell marker ALDH1A1, and are enriched in genes associated with epithelial–mesenchymal transition (EMT) and other stem-like properties.^{73,98} Within Lehmann's M and MSL subtypes, there are a number of enriched genes associated with EMT that are also modulated by Wnt signaling, including *MMP2*, *TWIST*, *SNAI2* and *TCF4*.⁷⁷ A gene set involved in Wnt/ β -catenin signaling in the M and MSL subtypes, including *CTNNB1* (β -catenin), *DKK2*, *DKK3*, *SFRP4*, *TCF4*, *TCF7L2* and *FZD4*, was also found to be enriched.⁷³ MES tumors are associated with a poorer distant metastasis-free survival at 5 years compared to other subtypes, likely associated with increased expression of cellular motility genes leading to increased metastasis.⁷³

BLIS subtype

BLIS is characterized as an immune-suppressed TNBC subtype with downregulated immune signaling pathways and reduced expression of immune function genes.^{92,99} BLIS tumors exhibit enhanced expression of mitotic and cell cycle pathway genes, with overexpression of proliferative genes, including *CENPF*, *BUB1* and *PRC1*.⁹⁹ Sry-related HMG box (SOX) transcription factors, and the immune-regulatory molecule V-domain-containing T-cell activation inhibitor.⁹² SOX transcription factors share a closely related consensus binding sequence to TCF/LEF transcription factors¹⁰⁰ and are known modulators of Wnt/ β -catenin signaling.¹⁰¹ Survival analysis shows that patients with the BLIS subtype TNBC experience lower rates of recurrence-free survival compared to other TNBC subtypes.⁹⁹

BLIA subtype

The BLIA subtype is characterized by upregulation of immune activating pathways, with overexpression of STAT transcription factors and cytotoxic T-lymphocyte-associated protein 4.⁹² Furthermore, the BLIA subtype demonstrates amplification of *CDK1*, which was recently found to phosphorylate the Wnt regulator TAZ.^{92,102} BLIA tumors have increased levels of lymphocytic infiltration and are thus associated with improved disease-free survival rates and patient outcomes compared to other TNBC subtypes, although still associated with a relatively high risk of recurrence (~20%).^{92,103}

Wnt dysregulation in TNBC and TNBC stem cells

Aberrant Wnt signaling is a characteristic of TNBC, with both canonical and non-canonical pathways implicated in TNBC tumorigenesis^{104,105} and metastasis.¹⁰⁶ Enrichment of Wnt/ β -catenin signaling is evident in TNBC and is associated with poor clinical outcomes within this subtype.^{107,108} TNBC patients displaying dysregulated Wnt/ β -catenin signaling are more likely to develop lung and brain secondary metastases.¹⁰⁶ Studies have shown that nuclear accumulation of β -catenin promotes cell migration, colony formation, stem-like features and chemoresistance of TNBC cells *in vitro* and TNBC tumorigenesis in mouse cancer models, thus suggesting that canonical Wnt signaling is a major driving force of TNBC tumorigenesis.¹⁰⁴ Although the Wnt/ β -catenin pathway is associated with the clinicopathological features of TNBC, this is not due to *CTNNB1* mutations.¹⁰⁸ Studies have also implicated dysregulation of non-canonical Wnt signaling pathways in the highly metastatic behavior of TNBC cells and CSCs, specifically through aberrant c-Jun N-terminal kinase activation.¹⁰⁹

CSCs, or cancer stem-like cells, are a small subset of cells within the heterogeneous tumor bulk that are thought to be responsible for tumor initiation.¹¹⁰ These cells also have intrinsic mechanisms for chemoresistance, such as upregulation of drug transporters, including the breast cancer resistance protein (also known as ABCG2).¹¹¹ By evading the standard chemotherapeutic treatments, it is thought that the CSCs are also responsible for the relapse experienced in many cancers, especially TNBCs.¹¹² Studies have also shown that these cells are a main contributor to metastasis, and are able to initiate solid tumor formation when xenotransplanted at low cell densities.¹¹³ TNBC stem cells are isolated from tumors as CD44⁺ (homing cell adhesion molecule), CD24⁻ (heat stable antigen), CD49f⁺ cells.¹¹⁴ CSCs also differ metabolically to other cancer cells. They are more reliant on mitochondrial respiration, which is supported by their higher mitochondrial reactive oxygen species, enhanced oxygen consumption and higher mitochondrial mass, allowing for features such as resistance to DNA damage.¹¹⁵

Wnt signaling is essential for normal breast stem cell function and mammary gland development during embryogenesis, postnatal development and pregnancy,¹¹⁶ with adult mammary glands containing Wnt-responsive stem cell populations.¹¹⁷ Studies have shown that aberrant Wnt signaling in breast cancer stem cells (BCSCs) is a key event in breast tumorigenesis.¹¹⁸ Wnt/ β -catenin signaling has been linked to TNBC tumorigenesis by regulating the key tumor-associated characteristics, including migration, stemness, proliferation and chemoresistance in TNBC cells and CSCs.¹⁰⁴ A recent study has also demonstrated that Wnt/ β -catenin signaling activity is higher in breast CSCs than the bulk tumor population, based on β -catenin, TCF4 and LEF1 expression in Aldefluor-positive cells versus Aldefluor-negative cells.¹¹⁹ Treatment with Wnt3a increased the number of ALDH⁺ breast CSCs, and knockdown of Wnt1 reduced the tumor-forming efficiency of breast CSCs *in vitro*.¹¹⁹

Furthermore, studies have shown that Wnt-derived breast tumors are maintained by clones capable of re-activating Wnt overexpression post-Wnt inhibition, indicating that aberrant Wnt activation is a key driver of breast cancer recurrence and progression.¹²⁰ A recent review highlighted the potential importance of Wnt/ β -catenin signaling, along with other developmental signaling pathways, including Cripto-1 and Notch/CSL, in the regulation of TNBC stem cells and therapy resistance in TNBC.¹²¹ An overview of Wnt signaling dysregulation is given in Figure 4.

FZDs in TNBC

FZD6. FZD6 exhibits increased gene copy number variations and overexpression in breast cancers. This is more frequent in TNBC than ER⁺ tumors. A study by Corda *et al.*¹²² determined that FZD6

Figure 4. Overview of Wnt signaling regulators contributing to TNBC progression and their targeted therapies. Canonical and non-canonical Wnt pathways are activated through Fzd, LRP and ROR receptors. Blue arrows indicate suppression/inhibition of Wnt regulators and pathways (with a net result of downregulation of Wnt target gene transcription, indicated by yellow cross); red arrows indicate activation of Wnt regulators and pathways.

was involved in the regulation of cell motility, invasion and three-dimensional (3D) growth, although it did not regulate proliferation in TNBC. This was confirmed by a significant reduction in distant metastases detected in various organs *in vivo* after the injection of MDA-MB-231 cells depleted of FZD6. Short hairpin RNA directed at *FZD6* *in vitro* was found to reduce cell invasion through a reduction in active Rho and the subsequent reduction in fibronectin fibres. This indicated that FZD6 regulates cell motility and invasion through non-canonical Wnt signaling. This study also suggests that FZD6 overexpression in TNBC has a high prognostic value in determining the risk of metastasis.¹²²

FZD7. Microarray analysis determined that FZD7 expression is upregulated in TNBC tissue and cell lines, and promotes tumorigenesis via canonical Wnt signaling pathways.¹²³ Short hairpin RNA-mediated silencing of FZD7 reduced invasiveness and colony formation in TNBC cell lines.¹²³ A recent study found that Δ Np63, an isoform of Transformation-related protein 63 (p63), enhanced FZD7 expression and increased Wnt signaling in TNBC tumor tissue and cell lines.¹²⁴

Aberrant FZD7 expression is implicated in TNBC stem cell-mediated tumorigenesis. A study recently found that knockdown of Δ Np63 in TNBC cell lines decreased FZD7 expression and tumorsphere formation, indicating that Δ Np63/FZD7 upregulation induced TNBC stem cells and promoted tumor formation in TNBC.¹²⁴ The findings of this study highlight the potential clinical importance of Δ Np63/FZD7-Wnt signaling in TNBC stem cells as a key driver of tumorigenesis and progression of TNBC.^{124,125}

FZD8. Gene expression studies have recently linked FZD8-driven Wnt signaling to chemoresistance in TNBC cell lines and TNBC stem cells. Treatment with cisplatin and tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) in TNBC cell lines

resulted in increased FZD8 expression in residual tumors of xenograft models.¹²⁶ Furthermore, FZD8 silencing led to increased Wnt pathway-driven TNBC cell apoptosis *in vitro* and *in vivo*.¹²⁶ The study showed that treatment with TRAIL/cisplatin increased expression of LEF-1 and TCF-7 in residual TNBC stem cells, thus implicating upregulation of Wnt signaling components in the development of chemoresistance.¹²⁶ An inverse correlation between FZD8 and miR-100 was shown, where decreased miR-100 expression was linked to increased FZD8 expression and Wnt signaling, resulting in increased loco-regional breast cancer metastasis.^{126,127} The role of miRs in Wnt signaling and TNBC is discussed in further detail below. c-Myc overexpression has been linked to FZD8 overexpression in TNBC cell lines, associating c-Myc-driven transcription to chemoresistance and TNBC CSC survival.¹²⁸

LRP5/6 in TNBC

LRP5/6 are essential for normal mammary development by regulating breast stem cell activity and are linked to basal-derived breast tumorigenesis.^{129–131} Studies in transgenic mice indicated that LRP5 knockdown led to resistance to Wnt1-induced tumor formation.¹³⁰

Gene expression analyses found that LRP6 is overexpressed in human TNBC.^{123,131} *In vivo* studies have shown that LRP6 silencing inhibited tumor growth in TNBC cell line-derived xenograft models.¹³² LRP6 and Wnt target gene *SOX9* have been shown to influence regulation of one another in TNBC cell lines. LRP6 overexpression led to *SOX9* upregulation, while knockdown of *SOX9* reduced LRP6 transcription and decreased cell invasion and proliferation.¹³³

LRP6 overexpression led to the upregulation of Wnt signaling and was associated with increased stemness in TNBC cells.¹³⁴ CD138 (Syndecan-1) is an EMT marker associated with both development and breast tumorigenesis,¹³⁵ and has been shown to modulate TNBC stem cell properties by targeting Wnt signaling.¹³⁴ Ibrahim *et al.*¹³⁴ showed that CD138 modulates Wnt signaling in TNBC stem cells through LRP6, whereby CD138 silencing resulted in downregulated LRP6 expression and Wnt signaling.¹³⁴

RORs in TNBC

Primary breast cancer DNA microarray data set analysis has shown that ROR1 is expressed on breast cancer cells and absent in normal breast cells, with high ROR1 expression associated with poorer survival.¹³⁶ Furthermore, the study showed that ROR1 silencing in TNBC cell lines increased apoptosis and reduced cell growth. High ROR1 expression in breast cancer cells is associated with high expression of EMT gene profiles and high incidences of disease recurrence and progression.¹³⁷ ROR1 knockdown in TNBC cell lines resulted in reduced EMT-associated protein expression, reduced cell migration and invasion *in vitro*, and inhibited metastasis in xenograft models.¹³⁷ ROR2 expression is present in both TNBC and non-TNBC, with ROR2⁺ TNBC patients exhibiting poorer survival outcomes compared to other subtypes.¹³⁸ ROR2 knockdown in TNBC cell lines inhibited Wnt signaling and reduced TCF/LEF transcription.¹³⁸ These findings indicate the potential prognostic and therapeutic significance of high ROR1/2 expression in TNBC.

DDXs, WNT and TNBC

DDX3. DDX3 is a regulator of Wnt/ β -catenin signaling, where it interacts with and increases the kinase activity of casein kinase 1 ϵ and is required for the phosphorylation of Dvl2.¹³⁹ It is known to have an oncogenic role in breast cancer, where non-tumorigenic MCF10A cell lines overexpressing DDX3 showed increased EMT, motility and invasiveness.¹⁴⁰ The same study demonstrated that DDX3 expression was positively correlated with a more aggressive

phenotype, and was highly expressed in TNBC cell lines. DDX3 overexpression resulted in E-cadherin downregulation and subsequent nuclear β -catenin translocation.¹⁴⁰ Similarly, DDX3 inhibition by NZ51, a ring-expanded nucleoside analog that is predicted to bind to the ATP-binding site of DDX3, led to decreased proliferation, motility and invasiveness in TNBC cell lines and reduced tumor load and metastatic burden in preclinical *in vivo* models.¹⁴¹

DDX5 (p68). DDX5 acts as a co-activator of Wnt/ β -catenin signaling through regulation of TCF4 expression. In turn, β -catenin/TCF4 regulates DDX5 expression, forming a positive feedback loop associated with increased EMT marker expression in TNBC cells.¹⁴² DDX5 is thought to regulate p53-mediated repair of DNA damage, and DDX5 overexpression contributes to tumorigenesis and progression in breast cancers.¹⁴³ DDX5 is highly expressed in basal-like breast cancers compared to luminal-like, and correlates with high EGFR and Ki67 expression in TNBC tissue.¹⁴⁴ Furthermore, the study found that DDX5 regulates the expression of miR-21 and miR-182 in basal breast cancers, and is associated with malignant disease.

THE ROLES OF MIRS IN WNT SIGNALING AND TNBC

miRs are endogenous, short, non-coding RNA molecules that regulate cancer-related genes at the post-transcriptional level.¹⁴⁵ miRs are differentially expressed in BCSCs and cancer cells, indicating that breast cancer-specific miRs are important in maintaining stemness and promoting tumorigenesis in BCSCs.¹⁴⁶ Twenty-seven miRs differentially expressed in locally advanced TNBC have been previously identified, with many of these predicted to be involved in regulation of Wnt signaling pathway genes.^{147,148}

miR-374a overexpression led to suppression of Wnt pathway inhibiting components (PTEN and WIF1) and ultimately increased Wnt-mediated EMT and metastasis in multiple TNBC cell lines.¹⁴⁹

miR-340 is downregulated in TNBC cell lines and has been linked with TNBC tumorigenesis regulation in multiple studies.¹⁵⁰ Induction of miR-340 resulted in downregulation of Wnt pathway target genes (*CTNNB1*, *MYC* and *ROCK1*), decreased proliferation and increased apoptosis in a metastatic TNBC cell line. The study showed that miR-340 overexpression reduced cell motility and invasiveness, indicating that miR-340 has a fundamental role in regulating breast metastases.¹⁵¹ Another study recently found that induction of miR-340 in TNBC cell lines led to reduced expression of *SOX2*, an oncogene associated with the canonical Wnt signaling pathway.¹⁵²

A study by Isobe *et al.*¹⁵³ found that miR-142 upregulation is associated with BCSCs and activates canonical Wnt signaling by promoting APC breakdown in TNBC cell lines. The study found that miR-142 expression activated canonical Wnt signaling, leading to increased miR-150 expression, thereby contributing to breast tissue hyperproliferation, BCSC proliferation and reducing apoptosis in TNBC cell lines.¹⁵³

A recent study has shown that miR-218-5p expression was significantly increased in TNBC, as well as bone metastases, from breast cancer patients.¹⁵⁴ Anti-miR-218-5p led to a reduction in cell proliferation *in vitro* and decreased tumor growth, active osteoclasts and osteolytic lesions *in vivo*, while the opposite was seen with transfection of miR-218-5p. The miR was also shown to directly modulate Wnt/ β -catenin signaling by binding to secreted FZD-related protein 1 and SOST. Anti-miR-218-5p suppressed Wnt signaling, which downregulated Parathyroid hormone-related protein expression, reducing breast cancer-induced osteolytic disease.¹⁵⁴

CURRENT AND EMERGING THERAPIES FOR TNBC AND TNBC STEM CELLS

Systemic cytotoxic chemotherapy is clinically indicated in early TNBC and is associated with a greater treatment benefit than hormone receptor-positive tumors.¹⁵⁵ Numerous early-phase clinical trials are currently underway, investigating various targeted molecules and combination therapies for the treatment of TNBC. In this section, we review current and emerging small molecule therapeutics for the treatment of TNBC (Figure 4); immunotherapeutics are reviewed elsewhere.⁷

Chemotherapy

Anthracycline/taxane-based regimens are currently the standard of care in the treatment of adjuvant and neoadjuvant TNBC. However, a recent *in vitro* study has shown that treatment with docetaxel or doxorubicin had transient and negligible impact on cell growth in two TNBC cell lines, respectively. Furthermore, the study found that docetaxel and doxorubicin treatment resulted in deregulation of genes associated with stemness in TNBC cells.¹⁵⁶ Molecular analysis found that doxorubicin treatment deregulated stem cell signaling pathways associated with cell growth, renewal and differentiation, with altered gene expression demonstrated in components of the Wnt signaling pathway, including FZD2, FZD4, FZD5, FZD6, FZD7, FZD9, Axin1, Wnt11, Wnt10a and Wnt5a. As such, the study concluded that docetaxel and doxorubicin induce stemness in differentiated TNBC cells, which likely accounts for acquired chemoresistance seen in refractory TNBC tumors.¹⁵⁶

Platinum agents

Platinum-based chemotherapeutics are a class of DNA-damaging agents, including cisplatin, carboplatin and oxaliplatin; these have established efficacy in breast cancer treatment.¹⁵⁷ *In vitro* studies have indicated that combining TRAIL and cisplatin significantly increased BCSC death compared to other standard of care treatments in TNBC cell lines.¹⁵⁸ The study showed that treatment with TRAIL and cisplatin inhibited Wnt1-mediated signaling and expression of cyclin D1, as well as the phosphorylation of β -catenin. Combination treatment with cisplatin and TRAIL also enhanced apoptosis, and inhibited proliferation and tumorsphere formation.¹⁵⁸

Wnt signaling inhibitors

Treatment with the small molecule β -catenin/TCF inhibitor CWP232228 inhibited β -catenin-mediated transcription, leading to inhibition of stem cell proliferation and reduction in tumor bulk in TNBC cell lines and TNBC patient-derived xenograft models, respectively.¹⁵⁹ PRI-724, a CREB-binding protein inhibitor, and LGK-974, a Porcupine inhibitor, are two small molecules currently undergoing clinical development. Both molecules are currently under investigation for single agent use in ongoing phase I clinical trials in TNBC patients,¹⁶⁰ with interim results yet to be released. Recent *in vitro* studies have shown that LGK-974 in combination with the PI3K/AKT/mTOR inhibitor BKM120 worked synergistically to decrease cell viability and enhance antitumor efficacy in TNBC cell lines.¹⁶¹

PARP inhibitors

Poly (ADP-ribose) polymerase (PARP) is an enzyme involved in DNA repair mechanisms necessary for maintaining BRCA-mutated cell viability.^{162,163} Included in the PARP enzyme family are tankyrase (TNKS)-1 and TNKS2. TNKS1 and TNKS2 are regulators of Wnt signaling through their interaction with Axin.^{164,165} TNBCs share phenotypic characteristics with BRCA-mutated cancers, thus providing support for the use of PARP inhibitors.¹⁶⁶ The small molecule TNKS1/2 inhibitor XAV939 showed effectiveness in the

destabilization of Axin and reduction of Wnt activity, although data suggest that a combination approach may be more beneficial.¹⁶⁵ Clinical trials evaluating the oral PARP inhibitor olaparib in BRCA1/2-positive metastatic breast cancer are currently underway, with interim results showing efficacy.¹⁶⁷ Veliparib is another PARP inhibitor currently being evaluated in combination with paclitaxel and carboplatin for metastatic TNBC.¹⁶⁸ Data from Phase I clinical trials of veliparib show acceptable safety, tolerance and good antineoplastic activity.¹⁶⁸

Histone deacetylase inhibitors

Histone deacetylase (HDAC) inhibitors are emerging as promising anti-TNBC agents because of their multifunctional capacity to regulate gene expression, cell growth and survival, as well as their ability to restore cellular aberrations due to epigenetic effects.¹⁶⁹ Entinostat is an HDAC inhibitor recently shown to have anti-CSC effects in TNBC stem cells. An *in vivo* study found that entinostat treatment reduced TNBC stem cell populations, tumorsphere formation and miR-181a expression in TNBC cell lines.¹⁷⁰ Furthermore, the study found that entinostat treatment in TNBC patient-derived xenografts reduced tumor growth and inhibited the development of lung metastases.¹⁷⁰ Further *in vivo* studies have shown that triple therapy, combining entinostat, all-*trans* retinoic acid and doxorubicin, induced apoptosis of TNBC stem cells in culture and induced differentiation of TNBC CSCs both *in vitro* and *in vivo*.¹⁷¹ Panobinostat (LBH589) decreased cell survival and cell cycle progression at the G2/M stage in TNBC cell lines and *in vivo*. It also increased acetylation of the histones H3 (Lys3) and H4 (Lys8).¹⁶⁹ Treatment with panobinostat upregulated cadherin-1 (CDH1) and reversed the M phenotype; CDH1 has been identified as a Wnt-signaling component in invasive breast carcinoma.¹⁷² An *in vivo* study found that salinomycin, a compound that selectively inhibits CSCs,¹⁷³ in combination with panobinostat, significantly inhibited the growth of TNBC stem cells in TNBC patient-derived xenografts. The study found that salinomycin and panobinostat worked synergistically to inhibit cell cycle progression, enhance apoptosis and regulate EMT in TNBC stem cells.¹⁷³

CONCLUSIONS

The dysregulation of Wnt signaling is synonymous with cancer. TNBC is an aggressive, highly proliferative phenotype, which is characteristic of overactive signaling pathways. The accelerated development of sequencing technologies has allowed us to characterize the highly heterogeneous molecular landscape of TNBC with unprecedented detail. These technologies have allowed the discovery of new potential therapeutic targets, as well as to suggest where existing drugs may be of therapeutic value, for instance, in the use of tamoxifen on AR-positive TNBC patients. Like TNBC, Wnt signaling is highly complex and not yet fully characterized. The discovery of novel regulators in TNBC, such as DDXs, adds to the complexity, but also presents exciting new opportunities for the development of potential therapeutic targets. Structural knowledge of Wnt pathway proteins and interactions has expanded in recent years, providing opportunities for rational/structure-based drug design of novel cancer therapeutics.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ACKNOWLEDGEMENTS

SP is supported by funds from the Rotary Club of Belmont, Australian Rotary Health Research Fund and Curtin University School of Biomedical Sciences. MA is a recipient

of an NHMRC Early Career Fellowship (GNT1054245), a Cancer Council of Western Australia Suzanne Cavanagh Early Career Investigator Grant and a Raine Priming Grant. AD is supported by strategic research funds from the School of Biomedical Sciences (Curtin University), Commercialization Advisory Board of Curtin University, Cancer Council of Western Australia and Actinogen Ltd, Perth, Western Australia.

REFERENCES

- Chen S, Parmigiani G. Meta-analysis of BRCA1 and BRCA2 penetrance. *J Clin Oncol* 2007; **25**: 1329–1333.
- Polakis P. Wnt signaling in cancer. *Cold Spring Harb Perspect Biol* 2012; **4**: a008052.
- Ovcaricek T, Frkovic SG, Matos E, Mozina B, Borstnar S. Triple negative breast cancer-prognostic factors and survival. *Radiol Oncol* 2011; **45**: 46–52.
- Foulkes WD, Smith IE, Reis-Filho JS. Triple-negative breast cancer. *N Engl J Med* 2010; **363**: 1938–1948.
- Hudis CA. Trastuzumab—mechanism of action and use in clinical practice. *N Engl J Med* 2007; **357**: 39–51.
- Jordan VC. Tamoxifen (ICI46,474) as a targeted therapy to treat and prevent breast cancer. *Br J Pharmacol* 2006; **147**: S269–S276.
- Pusztai L, Karn T, Safonov A, Abu-Khalaf MM, Bianchini G. New strategies in breast cancer: immunotherapy. *Clin Cancer Res* 2016; **22**: 2105–2110.
- Wu G, Huang H, Garcia Abreu J, He X. Inhibition of GSK3 phosphorylation of beta-catenin via phosphorylated PPPSPXS motifs of Wnt coreceptor LRP6. *PLoS ONE* 2009; **4**: e4926.
- Liu C, Kato Y, Zhang Z, Do VM, Yankner BA, He X. beta-Trcp couples beta-catenin phosphorylation-degradation and regulates Xenopus axis formation. *Proc Natl Acad Sci USA* 1999; **96**: 6273–6278.
- MacDonald BT, He X. Frizzled and LRP5/6 receptors for Wnt/beta-catenin signaling. *Cold Spring Harb Perspect Biol* 2012; **4**: a007880.
- Tamai K, Zeng X, Liu C, Zhang X, Harada Y, Chang Z et al. A mechanism for Wnt coreceptor activation. *Mol Cell* 2004; **13**: 149–156.
- Clevers H. Wnt/beta-catenin signaling in development and disease. *Cell* 2006; **127**: 469–480.
- Roose J, Molenaar M, Peterson J, Hurenkamp J, Brantjes H, Moerer P et al. The Xenopus Wnt effector XTcf-3 interacts with Groucho-related transcriptional repressors. *Nature* 1998; **395**: 608–612.
- Azzolin L, Panciera T, Soligo S, Enzo E, Biciatto S, Dupont S et al. YAP/TAZ incorporation in the beta-catenin destruction complex orchestrates the Wnt response. *Cell* 2014; **158**: 157–170.
- Li VS, Ng SS, Boersema PJ, Low TY, Karthaus WR, Gerlach JP et al. Wnt signaling through inhibition of beta-catenin degradation in an intact Axin1 complex. *Cell* 2012; **149**: 1245–1256.
- Azzolin L, Zanconato F, Bresolin S, Forcato M, Basso G, Biciatto S et al. Role of TAZ as mediator of Wnt signaling. *Cell* 2012; **151**: 1443–1456.
- Taelman VF, Dobrowolski R, Plouhinec JL, Fuentealba LC, Vorwald PP, Gumper I et al. Wnt signaling requires sequestration of glycogen synthase kinase 3 inside multivesicular endosomes. *Cell* 2010; **143**: 1136–1148.
- Gomez-Orte E, Saenz-Narciso B, Moreno S, Cabello J. Multiple functions of the noncanonical Wnt pathway. *Trends Genet* 2013; **29**: 545–553.
- Nishita M, Itsukushima S, Nomachi A, Endo M, Wang Z, Inaba D et al. Ror2/Frizzled complex mediates Wnt5a-induced AP-1 activation by regulating Dishevelled polymerization. *Mol Cell Biol* 2010; **30**: 3610–3619.
- Habu M, Koyama H, Kishida M, Kamino M, Iijima M, Fuchigami T et al. Ryk is essential for Wnt-5a-dependent invasiveness in human glioma. *J Biochem* 2014; **156**: 29–38.
- Golubkov VS, Chekanov AV, Cieplak P, Aleshin AE, Chernov AV, Zhu W et al. The Wnt/planar cell polarity protein-tyrosine kinase-7 (PTK7) is a highly efficient proteolytic target of membrane type-1 matrix metalloproteinase: implications in cancer and embryogenesis. *J Biol Chem* 2010; **285**: 35740–35749.
- Li L, Yuan H, Xie W, Mao J, Caruso AM, McMahon A et al. Dishevelled proteins lead to two signaling pathways. Regulation of LEF-1 and c-Jun N-terminal kinase in mammalian cells. *J Biol Chem* 1999; **274**: 129–134.
- Habas R, Kato Y, He X. Wnt/Frizzled activation of Rho regulates vertebrate gastrulation and requires a novel Formin homology protein Daam1. *Cell* 2001; **107**: 843–854.
- Wong HC, Bourdelas A, Krauss A, Lee HJ, Shao Y, Wu D et al. Direct binding of the PDZ domain of Dishevelled to a conserved internal sequence in the C-terminal region of Frizzled. *Mol Cell* 2003; **12**: 1251–1260.
- De A. Wnt/Ca²⁺ signaling pathway: a brief overview. *Acta Biochim Biophys* 2011; **43**: 745–756.
- Dejmek J, Saffholm A, Kamp Nielsen C, Andersson T, Leandersson K. Wnt-5a/Ca²⁺-induced NFAT activity is counteracted by Wnt-5a/Yes-Cdc42-casein kinase 1alpha signaling in human mammary epithelial cells. *Mol Cell Biol* 2006; **26**: 6024–6036.
- Ishitani T, Kishida S, Hyodo-Miura J, Ueno N, Yasuda J, Waterman M et al. The TAK1-NLK mitogen-activated protein kinase cascade functions in the Wnt-5a/Ca(2+) pathway to antagonize Wnt/beta-catenin signaling. *Mol Cell Biol* 2003; **23**: 131–139.
- Ishitani T, Ninomiya-Tsuji J, Matsumoto K. Regulation of lymphoid enhancer factor 1/T-cell factor by mitogen-activated protein kinase-related Nemo-like kinase-dependent phosphorylation in Wnt/beta-catenin signaling. *Mol Cell Biol* 2003; **23**: 1379–1389.
- Cadigan KM, Nusse R. Wnt signaling: a common theme in animal development. *Genes Dev* 1997; **11**: 3286–3305.
- Gao X, Hannoush RN. Single-cell imaging of Wnt palmitoylation by the acyltransferase porcupine. *Nat Chem Biol* 2014; **10**: 61–68.
- MacDonald BT, Hien A, Zhang X, Iranloye O, Virshup DM, Waterman ML et al. Disulfide bond requirements for active Wnt ligands. *J Biol Chem* 2014; **289**: 18122–18136.
- Willert K, Brown JD, Danenberg E, Duncan AW, Weissman IL, Reya T et al. Wnt proteins are lipid-modified and can act as stem cell growth factors. *Nature* 2003; **423**: 448–452.
- Janda CY, Waghray D, Levin AM, Thomas C, Garcia KC. Structural basis of Wnt recognition by Frizzled. *Science* 2012; **337**: 59–64.
- Dijksterhuis JP, Petersen J, Schulte G. WNT/Frizzled signalling: receptor-ligand selectivity with focus on FZD-G protein signalling and its physiological relevance: IUPHAR Review 3. *Br J Pharmacol* 2014; **171**: 1195–1209.
- Takada R, Satomi Y, Kurata T, Ueno N, Norioka S, Kondoh H et al. Mono-unsaturated fatty acid modification of Wnt protein: its role in Wnt secretion. *Dev Cell* 2006; **11**: 791–801.
- Wang C, Wu H, Katritch V, Han GW, Huang XP, Liu W et al. Structure of the human smoothened receptor bound to an antitumour agent. *Nature* 2013; **497**: 338–343.
- Wang C, Wu H, Evron T, Vardy E, Han GW, Huang XP et al. Structural basis for Smoothened receptor modulation and chemoresistance to anticancer drugs. *Nat Commun* 2014; **5**: 4355.
- Weierstall U, James D, Wang C, White TA, Wang D, Liu W et al. Lipidic cubic phase injector facilitates membrane protein serial femtosecond crystallography. *Nat Commun* 2014; **5**: 3309.
- Byrne EF, Sircar R, Miller PS, Hedger G, Luchetti G, Nachtergaele S et al. Structural basis of Smoothened regulation by its extracellular domains. *Nature* 2016; **535**: 517–522.
- Sussman DJ, Klingensmith J, Salinas P, Adams PS, Nusse R, Perrimon N. Isolation and characterization of a mouse homolog of the Drosophila segment polarity gene dishevelled. *Dev Biol* 1994; **166**: 73–86.
- Gao C, Chen YG. Dishevelled: the hub of Wnt signaling. *Cell Signal* 2010; **22**: 717–727.
- Fiedler M, Mendoza-Topaz C, Rutherford TJ, Mieszczynek J, Bienz M. Dishevelled interacts with the DIX domain polymerization interface of Axin to interfere with its function in down-regulating beta-catenin. *Proc Natl Acad Sci USA* 2011; **108**: 1937–1942.
- Schwarz-Romond T, Fiedler M, Shibata N, Butler PJ, Kikuchi A, Higuchi Y et al. The DIX domain of Dishevelled confers Wnt signaling by dynamic polymerization. *Nat Struct Mol Biol* 2007; **14**: 484–492.
- Gammons MV, Renko M, Johnson CM, Rutherford TJ, Bienz M. Wnt signalosome assembly by DEP domain swapping of Dishevelled. *Mol Cell* 2016; **64**: 92–104.
- Tauriello DV, Jordens I, Kirchner K, Slootstra JW, Kruiwagen T, Bouwman BA et al. Wnt/beta-catenin signaling requires interaction of the Dishevelled DEP domain and C terminus with a discontinuous motif in Frizzled. *Proc Natl Acad Sci USA* 2012; **109**: E812–E820.
- Jiang X, Charlat O, Zamponi R, Yang Y, Cong F. Dishevelled promotes Wnt receptor degradation through recruitment of ZNRF3/RNF43 E3 ubiquitin ligases. *Mol Cell* 2015; **58**: 522–533.
- Gong Y, Bourhis E, Chiu C, Stawicki S, DeAlmeida VI, Liu BY et al. Wnt isoform-specific interactions with coreceptor specify inhibition or potentiation of signaling by LRP6 antibodies. *PLoS ONE* 2010; **5**: e12682.
- Cheng Z, Biechele T, Wei Z, Morrone S, Moon RT, Wang L et al. Crystal structures of the extracellular domain of LRP6 and its complex with DKK1. *Nat Struct Mol Biol* 2011; **18**: 1204–1210.
- Ahn VE, Chu ML, Choi HJ, Tran D, Abo A, Weis WI. Structural basis of Wnt signaling inhibition by Dickkopf binding to LRP5/6. *Dev Cell* 2011; **21**: 862–873.
- Bourhis E, Wang W, Tam C, Hwang J, Zhang Y, Spittler D et al. Wnt antagonists bind through a short peptide to the first beta-propeller domain of LRP5/6. *Structure* 2011; **19**: 1433–1442.
- Zeng X, Huang H, Tamai K, Zhang X, Harada Y, Yokota C et al. Initiation of Wnt signaling: control of Wnt coreceptor Lrp6 phosphorylation/activation via frizzled, dishevelled and axin functions. *Development* 2008; **135**: 367–375.
- Minami Y, Oishi I, Endo M, Nishita M. Ror-family receptor tyrosine kinases in noncanonical Wnt signaling: their implications in developmental morphogenesis and human diseases. *Dev Dyn* 2010; **239**: 1–15.

- 53 Stiegler AL, Burden SJ, Hubbard SR. Crystal structure of the frizzled-like cysteine-rich domain of the receptor tyrosine kinase MusK. *J Mol Biol* 2009; **393**: 1–9.
- 54 Schlessinger J. Cell signaling by receptor tyrosine kinases. *Cell* 2000; **103**: 211–225.
- 55 Liu Y, Rubin B, Bodine PV, Billiard J. Wnt5a induces homodimerization and activation of Ror2 receptor tyrosine kinase. *J Cell Biochem* 2008; **105**: 497–502.
- 56 Zhang S, Chen L, Cui B, Chuang HY, Yu J, Wang-Rodriguez J et al. ROR1 is expressed in human breast cancer and associated with enhanced tumor-cell growth. *PLoS ONE* 2012; **7**: e31127.
- 57 Jarmoskaite I, Russell R. DEAD-box proteins as RNA helicases and chaperones. *Wiley Interdiscip Rev RNA* 2011; **2**: 135–152.
- 58 Samatanga B, Klostermeier D. DEAD-box RNA helicase domains exhibit a continuum between complete functional independence and high thermodynamic coupling in nucleotide and RNA duplex recognition. *Nucleic Acids Res* 2014; **42**: 10644–10654.
- 59 Rocak S, Linder P. DEAD-box proteins: the driving forces behind RNA metabolism. *Nat Rev Mol Cell Biol* 2004; **5**: 232–241.
- 60 Li L, Monckton EA, Godbout R. A role for DEAD box 1 at DNA double-strand breaks. *Mol Cell Biol* 2008; **28**: 6413–6425.
- 61 Linder P, Jankowsky E. From unwinding to clamping-the DEAD box RNA helicase family. *Nat Rev Mol Cell Biol* 2011; **12**: 505–516.
- 62 Fuller-Pace FV. DEAD box RNA helicase functions in cancer. *RNA Biol* 2013; **10**: 121–132.
- 63 Shin EM, Hay HS, Lee MH, Goh JN, Tan TZ, Sen YP et al. DEAD-box helicase DP103 defines metastatic potential of human breast cancers. *J Clin Invest* 2014; **124**: 3807–3824.
- 64 Perou CM, Sørlie T, Eisen MB, van de Rijn M, Jeffrey SS, Rees CA et al. Molecular portraits of human breast tumours. *Nature* 2000; **406**: 747–752.
- 65 Eroles P, Bosch A, Perez-Fidalgo JA, Lluch A. Molecular biology in breast cancer: intrinsic subtypes and signaling pathways. *Cancer Treat Rev* 2012; **38**: 698–707.
- 66 Creighton CJ. The molecular profile of luminal B breast cancer. *Biologics* 2012; **6**: 289.
- 67 Eroles P, Bosch A, Pérez-Fidalgo JA, Lluch A. Molecular biology in breast cancer: intrinsic subtypes and signaling pathways. *Cancer Treat Rev* 2012; **38**: 698–707.
- 68 Prat A, Carey LA, Adamo B, Vidal M, Tabernero J, Cortés J et al. Molecular features and survival outcomes of the intrinsic subtypes within HER2-positive breast cancer. *J Natl Cancer Inst* 2014; **106**: dju152.
- 69 Abramson VG, Lehmann BD, Ballinger TJ, Pietenpol JA. Subtyping of triple-negative breast cancer: implications for therapy. *Cancer* 2015; **121**: 8–16.
- 70 Kast K, Link T, Friedrich K, Petzold A, Niedostatek A, Schoffer O et al. Impact of breast cancer subtypes and patterns of metastasis on outcome. *Breast Cancer Res Treat* 2015; **150**: 621–629.
- 71 Shah SP, Roth A, Goya R, Oloumi A, Ha G, Zhao Y et al. The clonal and mutational evolution spectrum of primary triple-negative breast cancers. *Nature* 2012; **486**: 395–399.
- 72 Weisman PS, Ng CK, Brogi E, Eisenberg RE, Won HH, Piscuoglio S et al. Genetic alterations of triple negative breast cancer by targeted next-generation sequencing and correlation with tumor morphology. *Mod Pathol* 2016; **29**: 476–488.
- 73 Lehmann BD, Bauer JA, Chen X, Sanders ME, Chakravarthy AB, Shyr Y et al. Identification of human triple-negative breast cancer subtypes and preclinical models for selection of targeted therapies. *J Clin Invest* 2011; **121**: 2750–2767.
- 74 Wang G, Wang J, Sadar MD. Crosstalk between the androgen receptor and beta-catenin in castrate-resistant prostate cancer. *Cancer Res* 2008; **68**: 9918–9927.
- 75 Noah TK, Lo YH, Price A, Chen G, King E, Washington MK et al. SPDEF functions as a colorectal tumor suppressor by inhibiting beta-catenin activity. *Gastroenterology* 2013; **144**: 1012–23 e6.
- 76 Wang H, Xi Q, Wu G. Fatty acid synthase regulates invasion and metastasis of colorectal cancer via Wnt signaling pathway. *Cancer Med* 2016; **5**: 1599–1606.
- 77 Laezza C, D'Alessandro A, Paladino S, Maria Malfitano A, Chiara Proto M, Gazzerri P et al. Anandamide inhibits the Wnt/beta-catenin signalling pathway in human breast cancer MDA MB 231 cells. *Eur J Cancer* 2012; **48**: 3112–3122.
- 78 Kahlert UD, Maciaczyk D, Doostkam S, Orr BA, Simons B, Bogiel T et al. Activation of canonical WNT/beta-catenin signaling enhances in vitro motility of glioblastoma cells by activation of ZEB1 and other activators of epithelial-to-mesenchymal transition. *Cancer Lett* 2012; **325**: 42–53.
- 79 Greenow KR, Clarke AR, Williams GT, Jones R. Wnt-driven intestinal tumorigenesis is suppressed by Chk1 deficiency but enhanced by conditional haploinsufficiency. *Oncogene* 2014; **33**: 4089–4096.
- 80 Huard CC, Tremblay CS, Magron A, Levesque G, Carreau M. The Fanconi anemia pathway has a dual function in Dickkopf-1 transcriptional repression. *Proc Natl Acad Sci USA* 2014; **111**: 2152–2157.
- 81 Castiglia D, Bernardini S, Alvino E, Pagani E, De Luca N, Falcinelli S et al. Concomitant activation of Wnt pathway and loss of mismatch repair function in human melanoma. *Genes Chromosomes Cancer* 2008; **47**: 614–624.
- 82 Xu H, Yan Y, Deb S, Rangasamy D, Germann M, Malaterre J et al. Cohesin Rad21 mediates loss of heterozygosity and is upregulated via Wnt promoting transcriptional dysregulation in gastrointestinal tumors. *Cell Rep* 2014; **9**: 1781–1797.
- 83 Huang YL, Anvarian Z, Doderlein G, Acebron SP, Niehrs C. Maternal Wnt/STOP signaling promotes cell division during early *Xenopus* embryogenesis. *Proc Natl Acad Sci USA* 2015; **112**: 5732–5737.
- 84 Kikuchi K, Niikura Y, Kitagawa K, Kikuchi A. Dishevelled, a Wnt signalling component, is involved in mitotic progression in cooperation with Plk1. *EMBO J* 2010; **29**: 3470–3483.
- 85 Suzuki A, Pelikan RC, Iwata J. WNT/beta-catenin signaling regulates multiple steps of myogenesis by regulating step-specific targets. *Mol Cell Biol* 2015; **35**: 1763–1776.
- 86 Chiacchiera F, Rossi A, Jammula S, Pianti A, Scelfo A, Ordóñez-Moran P et al. Polycomb complex PRC1 preserves intestinal stem cell identity by sustaining Wnt/beta-catenin transcriptional activity. *Cell Stem Cell* 2016; **18**: 91–103.
- 87 He TC, Sparks AB, Rago C, Hermeking H, Zawel L, da Costa LT et al. Identification of c-MYC as a target of the APC pathway. *Science* 1998; **281**: 1509–1512.
- 88 Conrad WH, Swift RD, Biechele TL, Kulikauskas RM, Moon RT, Chien AJ. Regulating the response to targeted MEK inhibition in melanoma: enhancing apoptosis in NRAS- and BRAF-mutant melanoma cells with Wnt/beta-catenin activation. *Cell Cycle* 2012; **11**: 3724–3730.
- 89 Xu M, Wang S, Qi Y, Chen L, Frank JA, Yang XH et al. Role of MCP-1 in alcohol-induced aggressiveness of colorectal cancer cells. *Mol Carcinog* 2016; **55**: 1002–1011.
- 90 Yasuhara R, Irie T, Suzuki K, Sawada T, Miwa N, Sasaki A et al. The beta-catenin signaling pathway induces aggressive potential in breast cancer by up-regulating the chemokine CCL5. *Exp Cell Res* 2015; **338**: 22–31.
- 91 Yu Q, Xu M, Sen JM. Beta-catenin expression enhances IL-7 receptor signaling in thymocytes during positive selection. *J Immunol* 2007; **179**: 126–131.
- 92 Burstein MD, Tsimelzon A, Poage GM, Covington KR, Contreras A, Fuqua SA et al. Comprehensive genomic analysis identifies novel subtypes and targets of triple-negative breast cancer. *Clin Cancer Res* 2015; **21**: 1688–1698.
- 93 Pistelli M, Caramanti M, Biscotti T, Santinelli A, Pagliacci A, De Lisa M et al. Androgen receptor expression in early triple-negative breast cancer: clinical significance and prognostic associations. *Cancers* 2014; **6**: 1351–1362.
- 94 Arce-Salinas C, Riesco-Martinez MC, Hanna W, Bedard P, Warner E. Complete response of metastatic androgen receptor-positive breast cancer to bicalutamide: case report and review of the literature. *J Clin Oncol* 2016; **34**: e21–e24.
- 95 Gucalp A, Tolane S, Isakoff SJ, Ingle JN, Liu MC, Carey LA et al. Phase II trial of bicalutamide in patients with androgen receptor-positive, estrogen receptor-negative metastatic breast cancer. *Clin Cancer Res* 2013; **19**: 5505–5512.
- 96 Hilborn E, Gacic J, Fornander T, Nordenskjöld B, Stal O, Jansson A. Androgen receptor expression predicts beneficial tamoxifen response in oestrogen receptor-alpha-negative breast cancer. *Br J Cancer* 2016; **114**: 248–255.
- 97 Ciupek A, Rechoum Y, Gu G, Gelsomino L, Beyer AR, Brusco L et al. Androgen receptor promotes tamoxifen agonist activity by activation of EGFR in ERalpha-positive breast cancer. *Breast Cancer Res Treat* 2015; **154**: 225–237.
- 98 Prat A, Parker JS, Karginova O, Fan C, Livasy C, Herschkowitz JI et al. Phenotypic and molecular characterization of the claudin-low intrinsic subtype of breast cancer. *Breast Cancer Res* 2010; **12**: 1.
- 99 Liu Y-R, Jiang Y-Z, Xu X-E, Yu K-D, Jin X, Hu X et al. Comprehensive transcriptome analysis identifies novel molecular subtypes and subtype-specific RNAs of triple-negative breast cancer. *Breast Cancer Res* 2016; **18**: 1.
- 100 Lefebvre V, Dumitriu B, Penzo-Mendez A, Han Y, Pallavi B. Control of cell fate and differentiation by Sry-related high-mobility-group box (Sox) transcription factors. *Int J Biochem Cell Biol* 2007; **39**: 2195–2214.
- 101 Kormish JD, Sinner D, Zorn AM. Interactions between SOX factors and Wnt/beta-catenin signaling in development and disease. *Dev Dyn* 2010; **239**: 56–68.
- 102 Zhang L, Chen X, Stauffer S, Yang S, Chen Y, Dong J. CDK1 phosphorylation of TAZ in mitosis inhibits its oncogenic activity. *Oncotarget* 2015; **6**: 31399–31412.
- 103 Prat A, Pineda E, Adamo B, Galván P, Fernández A, Gaba L et al. Clinical implications of the intrinsic molecular subtypes of breast cancer. *Breast* 2015; **24**: S26–S35.
- 104 Xu J, Prospero JR, Choudhury N, Olopade OI, Goss KH. β -Catenin is required for the tumorigenic behavior of triple-negative breast cancer cells. *PLoS ONE* 2015; **10**: e0117097.
- 105 Mohammed MK, Shao C, Wang J, Wei Q, Wang X, Collier Z et al. Wnt/ β -catenin signaling plays an ever-expanding role in stem cell self-renewal, tumorigenesis and cancer chemoresistance. *Genes Dis* 2016; **3**: 11–40.
- 106 Dey N, Barwick BG, Moreno CS, Ordanic-Kodani M, Chen Z, Oprea-Ilie G et al. Wnt signaling in triple negative breast cancer is associated with metastasis. *BMC Cancer* 2013; **13**: 1.
- 107 Khrantsov AI, Khrantsova GF, Tretiakova M, Huo D, Olopade OI, Goss KH. Wnt/ β -catenin pathway activation is enriched in basal-like breast cancers and predicts poor outcome. *Am J Pathol* 2010; **176**: 2911–2920.

- 108 Geyer FC, Lacroix-Triki M, Savage K, Arnedos M, Lambros MB, MacKay A et al. beta-Catenin pathway activation in breast cancer is associated with triple-negative phenotype but not with CTNNB1 mutation. *Mod Pathol* 2011; **24**: 209–231.
- 109 Borg J-P, Belotti E, Daulat A, Lembo F, Bertucci F, Charafe-Jauffret E et al. Deregulation of the non-canonical pathway in triple-negative breast cancer. *FASEB J* 2013; **27**: 610.1–1.
- 110 Medema JP. Cancer stem cells: the challenges ahead. *Nat Cell Biol* 2013; **15**: 338–344.
- 111 Sicchieri RD, da Silva WA, Mandarano LR, de Oliveira TM, Carrara HH, Muglia VF et al. ABCG2 is a potential marker of tumor-initiating cells in breast cancer. *Tumour Biol* 2015; **36**: 9233–9243.
- 112 Pogoda K, Niwinska A, Murawska M, Pienkowski T. Analysis of pattern, time and risk factors influencing recurrence in triple-negative breast cancer patients. *Med Oncol* 2013; **30**: 388.
- 113 Cheng L, Ramesh AV, Flesken-Nikitin A, Choi J, Nikitin AY. Mouse models for cancer stem cell research. *Toxicol Pathol* 2010; **38**: 62–71.
- 114 Atkinson RL, Yang WT, Rosen DG, Landis MD, Wong H, Lewis MT et al. Cancer stem cell markers are enriched in normal tissue adjacent to triple negative breast cancer and inversely correlated with DNA repair deficiency. *Breast Cancer Res* 2013; **15**: R77.
- 115 Peiris-Pages M, Martinez-Outschoorn UE, Pestell RG, Sotgia F, Lisanti MP. Cancer stem cell metabolism. *Breast Cancer Res* 2016; **18**: 55.
- 116 van Amerongen R, Bowman AN, Nusse R. Developmental stage and time dictate the fate of Wnt/ β -catenin-responsive stem cells in the mammary gland. *Cell Stem Cell* 2012; **11**: 387–400.
- 117 Zeng YA, Nusse R. Wnt proteins are self-renewal factors for mammary stem cells and promote their long-term expansion in culture. *Cell Stem Cell* 2010; **6**: 568–577.
- 118 Howard B, Ashworth A. Signalling pathways implicated in early mammary gland morphogenesis and breast cancer. *PLoS Genet* 2006; **2**: e112.
- 119 Jang GB, Kim JY, Cho SD, Park KS, Jung JY, Lee HY et al. Blockade of Wnt/ β -catenin signaling suppresses breast cancer metastasis by inhibiting CSC-like phenotype. *Sci Rep* 2015; **5**: 12465.
- 120 Cleary AS, Leonard TL, Gestl SA, Gunther EJ. Tumour cell heterogeneity maintained by cooperating subclones in Wnt-driven mammary cancers. *Nature* 2014; **508**: 113–117.
- 121 Rangel MC, Bertolette D, Castro NP, Klauzinska M, Cuttitta F, Salomon DS. Developmental signaling pathways regulating mammary stem cells and contributing to the etiology of triple-negative breast cancer. *Breast Cancer Res Treat* 2016; **156**: 211–226.
- 122 Corda G, Sala G, Lattanzio R, Iezzi M, Salles M, Fragassi G et al. Functional and prognostic significance of the genomic amplification of frizzled 6 (FZD6) in breast cancer. *J Pathol* 2016; **241**: 350–361.
- 123 Yang L, Wu X, Wang Y, Zhang K, Wu J, Yuan Y et al. FZD7 has a critical role in cell proliferation in triple negative breast cancer. *Oncogene* 2011; **30**: 4437–4446.
- 124 Chakrabarti R, Wei Y, Hwang J, Hang X, Blanco MA, Choudhury A et al. Δ Np63 promotes stem cell activity in mammary gland development and basal-like breast cancer by enhancing Fzd7 expression and Wnt signalling. *Nat Cell Biol* 2014; **16**: 1004–1015.
- 125 Phesse T, Flanagan D, Vincan E. Frizzled7: a promising Achilles' heel for targeting the Wnt receptor complex to treat cancer. *Cancers* 2016; **8**: 50.
- 126 Yin S, Xu L, Bonfil RD, Banerjee S, Sarkar FH, Sethi S et al. Tumor-initiating cells and FZD8 play a major role in drug resistance in triple-negative breast cancer. *Mol Cancer Ther* 2013; **12**: 491–498.
- 127 Jiang Q, He M, Guan S, Ma M, Wu H, Yu Z et al. MicroRNA-100 suppresses the migration and invasion of breast cancer cells by targeting FZD-8 and inhibiting Wnt/ β -catenin signaling pathway. *Tumour Biol* 2016; **37**: 5001–5011.
- 128 Yin S, Cheriyan VT, Rishi AK, Reddy KB. c-Myc and Frizzled 8 play a major role in the regulation of cancer stem cells and drug resistance in triple-negative breast cancer. *Cancer Res* 2015; **75**: 2225.
- 129 Badders NM, Goel S, Clark RJ, Klos KS, Kim S, Bafico A et al. The Wnt receptor, Lrp5, is expressed by mouse mammary stem cells and is required to maintain the basal lineage. *PLoS ONE* 2009; **4**: e6594.
- 130 Lindvall C, Evans NC, Zylstra CR, Li Y, Alexander CM, Williams BO. The Wnt signaling receptor Lrp5 is required for mammary ductal stem cell activity and Wnt1-induced tumorigenesis. *Journal of Biological Chemistry* 2006; **281**: 35081–35087.
- 131 Lindvall C, Zylstra CR, Evans N, West RA, Dykema K, Furge KA et al. The Wnt co-receptor Lrp6 is required for normal mouse mammary gland development. *PLoS ONE* 2009; **4**: e5813.
- 132 Liu C-C, Prior J, Piwnicka-Worms D, Bu G. LRP6 overexpression defines a class of breast cancer subtype and is a target for therapy. *Proc Natl Acad Sci* 2010; **107**: 5136–5141.
- 133 Wang H, He L, Ma F, Regan MM, Balk SP, Richardson AL et al. SOX9 regulates low density lipoprotein receptor-related protein 6 (LRP6) and T-cell factor 4 (TCF4) expression and Wnt/ β -catenin activation in breast cancer. *J Biol Chem* 2013; **288**: 6478–6487.
- 134 Ibrahim SA, Hassan H, Vilardo L, Kumar SK, Kumar AV, Kelsch R et al. Syndecan-1 (CD138) modulates triple-negative breast cancer stem cell properties via regulation of LRP-6 and IL-6-mediated STAT3 signaling. *PLoS ONE* 2013; **8**: e85737.
- 135 Nikolova V, Koo C-Y, Ibrahim SA, Wang Z, Spillmann D, Dreier R et al. Differential roles for membrane-bound and soluble syndecan-1 (CD138) in breast cancer progression. *Carcinogenesis* 2009; **30**: 397–407.
- 136 Zhang S, Chen L, Cui B, Chuang H-Y, Yu J, Wang-Rodriguez J et al. ROR1 is expressed in human breast cancer and associated with enhanced tumor-cell growth. *PLoS ONE* 2012; **7**: e31127.
- 137 Cui B, Zhang S, Chen L, Yu J, Widhopf GF, Fecteau J-F et al. Targeting ROR1 inhibits epithelial–mesenchymal transition and metastasis. *Cancer Res* 2013; **73**: 3649–3660.
- 138 Henry C, Quadir A, Hawkins N, Jary E, Llamas E, Kumar D et al. Expression of the novel Wnt receptor ROR2 is increased in breast cancer and may regulate both β -catenin dependent and independent Wnt signalling. *J Cancer Res Clin Oncol* 2015; **141**: 243–254.
- 139 Cruciat C-M, Dolde C, de Groot RE, Ohkawara B, Reinhard C, Korswagen HC et al. RNA helicase DDX3 is a regulatory subunit of casein kinase 1 in Wnt- β -catenin signaling. *Science* 2013; **339**: 1436–1441.
- 140 Botlagunta M, Vesuna F, Mironchik Y, Raman A, Lisok A, Winnard P et al. Oncogenic role of DDX3 in breast cancer biogenesis. *Oncogene* 2008; **27**: 3912–3922.
- 141 Xie M, Vesuna F, Botlagunta M, Bol GM, Irving A, Bergman Y et al. NZ51, a ring-expanded nucleoside analog, inhibits motility and viability of breast cancer cells by targeting the RNA helicase DDX3. *Oncotarget* 2015; **6**: 29901–29913.
- 142 Guturi KKN, Sarkar M, Bhowmik A, Das N, Ghosh MK. DEAD-box protein p68 is regulated by β -catenin/transcription factor 4 to maintain a positive feedback loop in control of breast cancer progression. *Breast Cancer Res* 2014; **16**: 1.
- 143 Moore HC, Jordan LB, Bray SE, Baker L, Quinlan PR, Purdie CA et al. The RNA helicase p68 modulates expression and function of the Δ 133 isoform (s) of p53, and is inversely associated with Δ 133p53 expression in breast cancer. *Oncogene* 2010; **29**: 6475–6484.
- 144 Wang D, Huang J, Hu Z. RNA helicase DDX5 regulates microRNA expression and contributes to cytoskeletal reorganization in basal breast cancer cells. *Mol Cell Proteomics* 2012; **11**: M111. 011932.
- 145 Esquela-Kerscher A, Slack FJ. Oncomirs—microRNAs with a role in cancer. *Nat Rev Cancer* 2006; **6**: 259–269.
- 146 Shimono Y, Zabala M, Cho RW, Lobo N, Dalerba P, Qian D et al. Downregulation of miRNA-200c links breast cancer stem cells with normal stem cells. *Cell* 2009; **138**: 592–603.
- 147 Avery-Kiejda KA, Braye SG, Mathe A, Forbes JF, Scott RJ. Decreased expression of key tumour suppressor microRNAs is associated with lymph node metastases in triple negative breast cancer. *BMC Cancer* 2014; **14**: 1.
- 148 Goh JN, Loo SY, Datta A, Siveen KS, Yap WN, Cai W et al. microRNAs in breast cancer: regulatory roles governing the hallmarks of cancer. *Biol Rev Camb Philos Soc* 2016; **91**: 409–428.
- 149 Cai J, Guan H, Fang L, Yang Y, Zhu X, Yuan J et al. MicroRNA-374a activates Wnt/ β -catenin signaling to promote breast cancer metastasis. *J Clin Invest* 2013; **123**: 566–579.
- 150 Wu ZS, Wu Q, Wang CQ, Wang XN, Huang J, Zhao JJ et al. miR-340 inhibition of breast cancer cell migration and invasion through targeting of oncoprotein c-Met. *Cancer* 2011; **117**: 2842–2852.
- 151 Mohammadi-Yeganeh S, Paryan M, Arefian E, Vasei M, Ghanbarian H, Mahdian R et al. MicroRNA-340 inhibits the migration, invasion, and metastasis of breast cancer cells by targeting Wnt pathway. *Tumour Biol* 2016; **37**: 8993–9000.
- 152 Mohammadi Yeganeh S, Vasei M, Tavakoli R, Kia V, Paryan M. The effect of miR-340 over-expression on cell-cycle-related genes in triple-negative breast cancer cells. *Eur J Cancer Care* 2016; **7**: 79032–79046.
- 153 Isobe T, Hisamori S, Hogan DJ, Zabala M, Hendrickson DG, Dalerba P et al. miR-142 regulates the tumorigenicity of human breast cancer stem cells through the canonical WNT signaling pathway. *Elife* 2014; **3**: e01977.
- 154 Taipaleenmaki H, Farina NH, van Wijnen AJ, Stein JL, Hesse E, Stein GS et al. Antagonizing miR-218-5p attenuates Wnt signaling and reduces metastatic bone disease of triple negative breast cancer cells. *Oncotarget* 2016; **7**: 79032–79046.
- 155 Group EBCTC. Adjuvant chemotherapy in oestrogen-receptor-poor breast cancer: patient-level meta-analysis of randomised trials. *Lancet* 2008; **371**: 29–40.
- 156 Tudoran O, Soritau O, Balacescu L, Visan S, Barbos O, Cojocneanu-Petric R et al. Regulation of stem cells-related signaling pathways in response to doxorubicin treatment in Hs578T triple-negative breast cancer cells. *Mol Cell Biochem* 2015; **409**: 163–176.
- 157 Sparano JA. Defining a role and predicting benefit from platinum-based therapy in breast cancer: an evolving story. *J Clin Oncol* 2015; **33**: 1–3.
- 158 Yin S, Xu L, Bandyopadhyay S, Sethi S, Reddy KB. Cisplatin and TRAIL enhance breast cancer stem cell death. *Int J Oncol* 2011; **39**: 891.

- 159 Jang G-B, Hong I-S, Kim R-J, Lee S-Y, Park S-J, Lee E-S *et al*. Wnt/ β -catenin small-molecule inhibitor CWP232228 preferentially inhibits the growth of breast cancer stem-like cells. *Cancer Res* 2015; **75**: 1691–1702.
- 160 Takebe N, Miele L, Harris PJ, Jeong W, Bando H, Kahn M *et al*. Targeting Notch, Hedgehog, and Wnt pathways in cancer stem cells: clinical update. *Nat Rev Clin Oncol* 2015; **12**: 445–464.
- 161 Solzak JP, Atale R, Hancock B, Radovich M. Dual PI3K and Wnt pathway inhibition is a synergistic combination against triple-negative breast cancer. *Cancer Res* 2015; **75**: 5340.
- 162 Farmer H, McCabe N, Lord CJ, Tutt AN, Johnson DA, Richardson TB *et al*. Targeting the DNA repair defect in BRCA mutant cells as a therapeutic strategy. *Nature* 2005; **434**: 917–921.
- 163 Bryant HE, Schultz N, Thomas HD, Parker KM, Flower D, Lopez E *et al*. Specific killing of BRCA2-deficient tumours with inhibitors of poly (ADP-ribose) polymerase. *Nature* 2005; **434**: 913–917.
- 164 Huang SM, Mishina YM, Liu S, Cheung A, Stegmeier F, Michaud GA *et al*. Tankyrase inhibition stabilizes axin and antagonizes Wnt signalling. *Nature* 2009; **461**: 614–620.
- 165 Bao R, Christova T, Song S, Angers S, Yan X, Attisano L. Inhibition of tankyrases induces Axin stabilization and blocks Wnt signalling in breast cancer cells. *PLoS ONE* 2012; **7**: e48670.
- 166 Livraghi L, Garber JE. PARP inhibitors in the management of breast cancer: current data and future prospects. *BMC Med* 2015; **13**: 1.
- 167 Kaufman B, Shapira-Frommer R, Schmutzler RK, Audeh MW, Friedlander M, Balmaña J *et al*. Olaparib monotherapy in patients with advanced cancer and a germline BRCA1/2 mutation. *J Clin Oncol* 2015; **33**: 244–250.
- 168 Pahuja S, Beumer JH, Appleman LJ, HA-H Tawbi, Stoller RG, Lee JJ *et al*. (eds). in *ASCO Annual Meeting Proceedings*, Chicago, IL, USA; 29 May–2 June, 2015.
- 169 Tate CR, Rhodes LV, Segar HC, Driver JL, Pounder FN, Burow ME *et al*. Targeting triple-negative breast cancer cells with the histone deacetylase inhibitor panobinostat. *Breast Cancer Res* 2012; **14**: R79.
- 170 Schech A, Kazi A, Yu S, Shah P, Sabnis G. Histone deacetylase inhibitor entinostat inhibits tumor-initiating cells in triple-negative breast cancer cells. *Mol Cancer Ther* 2015; **14**: 1848–1857.
- 171 Merino VF, Nguyen N, Jin K, Sadik H, Cho S, Korangath P *et al*. Combined treatment with epigenetic, differentiating, and chemotherapeutic agents cooperatively targets tumor-initiating cells in triple-negative breast cancer. *Cancer Res* 2016; **76**: 2013–2024.
- 172 Prasad CP, Mirza S, Sharma G, Prashad R, DattaGupta S, Rath G *et al*. Epigenetic alterations of CDH1 and APC genes: relationship with activation of Wnt/ β -catenin pathway in invasive ductal carcinoma of breast. *Life Sci* 2008; **83**: 318–325.
- 173 Kai M, Kanaya N, Wu SV, Mendez C, Nguyen D, Luu T *et al*. Targeting breast cancer stem cells in triple-negative breast cancer using a combination of LBH589 and salinomycin. *Breast Cancer Res Treat* 2015; **151**: 281–294.

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