What Signals Operate in the Mammary Niche?

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Abstract. Adult stem cells reside in a specialized microenvironment, the niche, which controls their behavior. As mammary stem cells, and consequently their niches, are still poorly defined, we look at better-characterized adult mammalian stem cell niches in the hematopoietic system and the skin. We attempt to define the mammary stem cell niche functionally, based on the widely used mammary fat pad reconstitution assay.

We note that the concept of the niche needs to be extended from the specialized microenvironment described in the hematopoietic system, to a model that takes into account the macroenvironment, as recently shown in the skin, and systemic clues as we will illustrate for the mammary gland where the reproductive hormones are major determinants of stem cell activation. In fact, in the mammary gland a special type of stem cells is determined only during pregnancy.

Reproductive hormones act on hormone receptor positive cells, sensor cells, in the mammary epithelium to induce paracrine signaling that leads to activation of stem cells. Some of the downstream mediators are in common with other niches such as Wnt and possibly Notch signaling. Other signals are specific to the mammary gland such as amphiregulin and RANKL.

Keywords: Estrogens, progesterone, mammary gland, stem cell niche, microenvironment, macroenvironment

INTRODUCTION

The breast undergoes most of its development after birth under the control of systemic hormones. Reproductive hormones are messengers of systemic requirements to the mammary gland tissue and orchestrate major changes in this organ by modulating local signaling pathways [15,41].

The rudimentary ductal system present at birth begins to expand during puberty. With repeated menstrual cycles, its complexity increases. The most dramatic increase in epithelial cell number occurs during pregnancy. Throughout lactation there is little cell proliferation; in the absence of breastfeeding or after weaning, the breast involutes and returns to a state that morphologically resembles a pre-pregnancy gland. With every new pregnancy this cycle is repeated; this seemingly unlimited regeneration capacity is attributed to the presence of breast epithelial stem cells.

Adult stem cells are capable of self-renewal and can give rise to differentiated cell lineages. In the breast two cell lineages are distinguished; the luminal cells that line the lumen and thereby form a tube and the contractile myoepithelial cells that form a meshwork around the luminal cells.

Studying the hematopoietic system, Schofield first recognized that stem cells do not operate in isolation but depend on their microenvironment; he coined the term “niche” for the microanatomical space that is formed by neighboring cells that allows only one stem cell to fit in and maintains the self-renewal capacity of this cell [94]. As a stem cell divides, one cell has to leave the niche; it may still be multipotent but has less self-renewal capacity than the cell that remains in the niche [37,133].

A cell that has left the niche but finds an empty niche and homes to it can reacquire stem cell properties [45, 82].

The concept of the niche has since been confirmed experimentally in Drosophila melanogaster and...
stem cells and their niches have been well characterized for instance in the drosophila gonads and midgut [26,51,70,135].

In mammals, adult stem cells have been mapped in few tissues. Schofield had hypothesized that the niche for hematopoietic stem cells is at the interface with the bone in the bone marrow and indeed, hematopoietic stem cells (HSCs) have been identified there, either next to immature osteoblasts or next to fenestrated endothelial cells [20,139]. Epidermal stem cells are found in the bulge area of hair follicles [25], the crypt was identified as the location for intestinal stem cells [6], and in the brain, stem cells are in the subventricular zone [55].

The identity and localization of mammalian epithelial stem cells is still debated, consequently, the niche components remain to be defined. Here, we take a look at what is known about mammalian adult stem cell niches in different tissues and the signaling that occurs within them. We will speculate about the composition of the mammary stem cell niche and discuss potential signals operating between niche and stem cells.

STEM CELL NICHEL: DIFFERENT LEVELS OF CONTROL

Hematopoietic niche

The best-characterized adult stem cell is the hematopoietic stem cell [127,133]; murine HSCs were identified based on their ability to provide reconstitution of all blood cell lineages in lethally irradiated mice [7, 117,134]. Adult HSCs home to two niches in the bone marrow, the endosteal one, in which they physically interact with a subset of immature osteoblasts [3,79], and the vascular one (Fig. 1) [59,118], in which they attach to the fenestrated endothelial cells of bone marrow sinusoids [113,114]. The endosteal zone favors quiescence, whereas the vascular niche is thought to allow differentiation and ultimately mobilization to the circulation [54].

Experiments with genetically engineered mice showed that activation of parathyroid receptor 1 (PTH1R) signaling as well as ablation of the bone morphogenetic protein (BMP) receptor (BMPR1A) signaling in the bone-marrow stroma resulted in an increase in both the number of osteoblasts and HSCs in the bone marrow [20,139] and revealed the role of immature osteoblasts as central components of the niche.

Osteoblasts secrete and/or activate factors (Fig. 2) such as angiopoietin, stem cell factor (SCF) and the matrix acid glycoprotein, osteopontin [1,80] which activate their respective receptors Tie2, c-Kit, and CD44, on HSCs. Tie2/Ang-1 and KIT signaling maintain HSCs quiescent ensuring their self-renewal capacity [1, 66,116]. Osteopontin, recognized by β1-integrin on the HSCs’ membrane physically sequesters HSCs in the niche thereby indirectly inhibiting HSC proliferation [80,105].

Reticular cells are specialized fibroblasts that constitute the vascular niche and contribute to the endosteal niche; they express CXCL12/SDF-1 (stromal cell-derived factor-1) that binds to the chemokine receptor CXCR4 on the HSCs membrane. This interactions and the resulting signaling are required to maintain HSCs in vascular niches [110].

A central signaling pathway in self-renewal and differentiation of HSCs is canonical Wnt signaling. Constitutive activation of this signaling cascade blocks differentiation and leads to accumulation of functionally defective HSCs [53,92]. Ectopic expression of the secreted Wnt antagonist dickkopf-1 (Dkk-1) in niche cells results in a failure to maintain HSCs quiescent and consequent loss of self-renewal ability [109]. On the other
Fig. 2. Hematopoietic Stem Cell Niche. The scheme shows the signaling molecules and their receptors in the HSC that regulate HSC behaviour; SCF/KIT, Jagged1/Notch and Wnt/Fz are involved in osteoblast induced cell renewal; osteopontin/CD44; angiopoietin/Tie2 and also Wnt/Fz are involved in osteoblast induced quiescence of the stem cells and CXCL-12/CXCR4 in reticulocyte induced quiescence. Both PTH/PTH1R stimulation and BMP/BMPR1A downregulation leads to an increase in osteoblast number and consequently in the HSC number.

Hand activation of the Wnt signaling pathway increases the ability of HSCs to reconstitute the hematopoietic system of irradiated mice [87]. Wnt-5a which activates noncanonical and inhibits canonical Wnt signaling maintaining HSCs in a quiescent state [76]. Taken together, these somewhat contradictory data suggest that canonical Wnt signaling has an important role in controlling the stem cell pool, however, the biological consequences of activating this signaling cascade are dose and stage-dependent.

Notch signaling has been implicated in the control of the HSC pool as ectopic activation of this pathway results in an increased number of HSCs [49,104]. Consistent with a role of Notch signaling in hematopoiesis, both receptors and ligands are expressed in the bone marrow [85]. However, loss-of-function experiments suggest that the pathway is not essential [68].

The transcription factor c-myc promotes differentiation of stem cells [2,126] suggesting that its role in the skin is different from its role in the hematopoietic system. However, loss-of-function experiments are required to draw this conclusion.

Epidermal niche(s)

Another organ system amenable to functional studies is the skin; it comprises multilayered interfollicular epidermis (IFE) and different appendages, such as hair follicles, sweat glands and sebaceous glands [9] (Fig. 3). Epidermal stem cells are found in the IFE, in the sebaceous glands and in the bulge region of the hair follicle; stem cells of the bulge region have been isolated and have been shown to reconstitute all epidermal compartments [46,115] and will be discussed here. In the bulge, stem cells can interact with surrounding epithelial cells, the inner and outer root sheath cells, as well as with the basement membrane and fibroblasts (Fig. 3). To what extent these different interactions are important to stem cell function is currently a matter of debate.

The surface protein, β1-integrin, used by HSCs to bind osteopontin, mediates binding to the basal membrane component laminin and collagen in epithelial cells and is enriched in epidermal stem cells versus more differentiated populations [48,65]. It is functionally important as blocking β1-integrin by ectopic expression of a dominant negative mutant results in differentiation [140].

The transcription factor p63 is essential for the establishment of stratified epithelia [72,136]. Whether it has a role in stem cell maintenance like in the thymus [21,95] or whether it is required for the commitment to stratification is debated [56,57]. Interestingly, p63 is downregulated by activation of Notch signaling in suprabasal cells [78]. In contrast to its stimulatory effects in the stem cell compartment of the hematopoietic system, Notch stimulates differentiation in the skin [86].

Overexpression of the transcription factor c-myc promotes differentiation of stem cells [2,126] suggesting that its role in the skin is different from its role in the hematopoietic system. However, loss-of-function experiments are required to draw this conclusion.

Central to activation of bulge stem cells is periodic activation of canonical Wnt signaling [42,111,120]. Upon elevation of the levels of β-catenin hair follicles enter anagen (phase of rapid proliferation) [64,121] whereas Wnt signaling is suppressed during telogen (phase of mitotic quiescence, regression phase) [31,84]. Expression of the bone morphogenetic proteins (BMPs)-2 and -4 in the dermis and subcutaneous fat cycles out of phase with respect to this recurrent Wnt signaling activation [84]. High BMP signaling activity keeps the hair follicle refractory whereas low BMP signaling activity renders it competent for regeneration [84]. Thus, stem cell activity is not only influenced by the microenvironment but ultimately controlled by signaling from another organ; Plikus and colleagues introduced the term “macroenvironment” to describe this long distance signaling.
BREAST EPITHELIAL STEM CELLS AND THEIR NICHE

As breast epithelial stem cells can give rise to myoepithelial and luminal cells, it has been surmised that stem cells express markers of both the luminal and myoepithelial cell lineage [83,106,107]. Cells co-expressing both type of markers have been identified; they are located in an intermediate position between the myoepithelial and luminal cells in the human breast [39]. These cells are epithelial and hence express the panepithelial marker EpCAM but they do not express proteins found on the apical membrane of differentiated luminal cells, such as MUC1. Such cells were purified from reduction mammoplasty specimens. Consistent with being bipotential progenitors, the MUC−/ESA+ cells gave rise to both luminal and myoepithelial cell types in 2D and 3D cultures [39]. However, in vitro cultures have limitations; for example hormone receptor expression does not occur and it remains to be shown that these populations reconstitute in vivo. More recently, a stem cell zone has been localized in the ducts based on immunohistochemistry assays for putative stem cell and niche markers [123].

In rodents, reconstitution of epithelium-divested mammary fat pads provides a functional assay for stem cells [27,32,137]. In 3-week-old prepubertal mice the inguinal mammary gland can be surgically cleared of the endogenous epithelium. Any part of the mammary gland is able to give rise to a new ductal system when grafted to such a cleared fat pad [81,137] independently of age, reproductive history or reproductive state of the donor mice [30]. The reconstitution ability is lost over seven consecutive transplant generations [27].

The reconstitution of the cleared fat pad by engrafted mammary epithelial cells (MECs) has been...
very successfully used as an assay for stem cell activity combined with immunophenotyping and fluorescence activated cell sorting (FACS) of MECs [69, 96,108,128]. This line of work revealed subpopulations of MECs with distinct reconstitution potential. Cells with high reconstitution potential are enriched in a population expressing the glycosyl phosphatidylinositol (GPI)-anchored protein CD24 and high levels of β1 (CD29) and α6 (CD49d) integrin (CD24[+]/CD29[high]CD49d[high]) [96,108].

Consistent with the role of integrins in binding ECM components, these surface molecules are highly expressed in cells that interact with the basal lamina [93]. Indeed, cells in the CD24[+]/CD49d[high]CD29[high] compartment bear myoepithelial characteristics; they express Keratin 14 [96,108], high levels of the epidermal growth factor receptor (EGFR), and do not express the estrogen receptor α (ERα) [4]. Taken together, these observations suggest that the mammary stem cells localize basally within the mammary epithelium, possibly with extensive direct contact to the basal lamina.

Previous studies based on morphological characteristics had attributed a particular phenotype to stem cells. Based on their appearance by electron microscopy or by toluidine blue stain on ultrathin sections, they were called “small light cells” (SLC) [22,99]. SLCs are found dispersed in an intermediate position between luminal and myoepithelial cells like the bipotential progenitor cells in the human breast described above [22,99]. The SLCs were shown to be the only cells that enter mitosis when mammary explants were cultured [99]. Consistent with these cells being stem cells, their numbers decrease as the reconstitution potential of mammary epithelium diminishes upon serial transplantation [99].

These two lines of work point to two different scenarios for the localization of mammary stem cells (Fig. 5): in the first, the mammary stem cell is part of the myoepithelial sheet; in the second scenario, the mammary stem cell is located between the luminal and myoepithelial layers. These two models may not be mutually exclusive; the morphologically defined SLCs may express the same surface markers as the myoepithelial cells and therefore part of the CD24low/CD29high and CD49dhigh population in FACS sorting. Alternatively, light cells may simply contaminate the FACS sorted population; obviously, even a single contaminating stem cell should, by definition, reconstitute a mammary gland.

While SLCs represent around 3% of the epithelial cell population [22] estimates of stem cell frequency based on FACS sorting vary between different groups from 1 in 200 [73] to 1 in 5000 MECs [96].

A FUNCTIONAL DEFINITION OF THE MAMMARY STEM CELL NICHE

In both the scenarios presented above (Fig. 5), the microenvironment of the putative stem cell comprises neighboring luminal and myoepithelial cells as well as more or less extensive contact with the basal lamina. Based on the functional definition of a stem cell by the reconstitution assay, various other factors that determine the stem cell’s repopulation ability can be discerned. These may directly or indirectly affect signaling in the mammary stem cell niche and may or may not be considered part of the niche.

First of all, MECs only give rise to a ductal tree when they are grafted to the mammary fat pad; when they are transplanted to other adipose tissues they yield only very limited outgrowth [77]. Thus, similar to the skin, signals from cells not in direct contact with the stem cell are important for the stem cell function. The contribution of different cell types within the fat pad, such as fibroblasts, adipocytes, endothelial, perivascular and immune cells, remains to be determined.

An additional level of complexity stems from the fact that the gland has to coordinate all its activities with systemic requirements conveyed, above all, by the female reproductive hormones estrogens, progesterone and prolactin. Indeed, MECs that are grafted
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Fig. 5. Localization of the mammary gland stem cell. A: Schematic diagram illustrating the cellular composition of a milk duct in the mouse mammary gland: LC: luminal cell; MC: myoepithelial cell; MSC: mammary stem cell AD: adipocyte; FB: fibroblast. B: Localization of the mammary gland stem cells based on functional studies of different FACS sorted populations. The population enriched for cells of high reconstitution potential expresses myoepithelial markers. C: localization of the mammary stem cells based on morphological characteristics. Small light cells (SLCs) localize in an intermediate position between myoepithelial and luminal layer and do not reach the lumen.

to a cleared fat pad will only grow out when estrogens are present; in ovariectomized females no ductal outgrowth occurs; local estrogen administration restores it [29]. The systemic hormonal milieu translates into local cues via the estrogen and progesterone receptors present on a subset of luminal epithelial cells, that we call the sensor cells [15]. These sensor cells are either part of the niche or near the niche, they may directly or indirectly signal to the stem cells. As the hormonal milieu changes, the niche and it’s signaling also change.

THE NICHE DURING PUBERTY

The fat pad reconstitution assay reflects growth as it occurs physiologically during puberty. Pubertal ductal outgrowth is driven by ovarian estrogens [29,75] that act through ERα in the mammary epithelium [67], which is expressed in about 30% of the luminal epithelial cells at this stage [90,138]. When ERα−/− MECs are grafted to cleared fat pads no outgrowth occurs; however, when ERα deficient cells are mingled with WT cells, they contribute extensively to the ductal outgrowth and give rise to both myoepithelial and luminal cells [67]. Thus, ERα deficient mammary epithelia contain stem cells; a finding that is consistent with the observation that the cell fraction enriched for stem cells is hormone receptor negative [4]. The ER negative stem cell can only unfold its potential when it is in vicinity of an ERα positive cells that translate estrogen action into a paracrine signal. Whether the hormone receptor positive sensor cell is actually part of the niche and interacts directly with the stem cell or whether it may be a few cell diameters away from the stem cell remains unclear.

Which are the downstream mediators of estrogens? The EGFR ligands epidermal growth factor (EGF), transforming growth factor α (TGFα), and amphiregulin administered locally, can restore ductal elongation in ovariectomized mice arguing that EGFR signaling is important downstream of ERα [24,50,100]. The physiologically relevant EGF family member is amphiregulin; its transcription is specifically and selectively induced by 17β-estradiol in the puberal mammary epithelium [23]. The membrane-bound protease TACE/ADAM17 cleaves and thereby activates
Fig. 6. Model of pubertal mammary stem cell niche: Estrogens from the systemic environment act on ERα receptor positive sensor cell, induce expression of amphiregulin that is activated by TACE and acts via the EGFR on stromal cells. The basal membrane is disrupted in this phase of the development. In response to this stimulation, the stromal cells release factors like FGFs and MMPs that in turn impinge directly or indirectly on the stem cell to modulate its behavior. See text for further explanations.

MSC: mammary stem cell; LC: luminal cell; MC: myoepithelial cell; FB: fibroblast, AD: adipocyte; HR: hormone receptor, here: ERα.

The nature of the stromal factors released in response to amphiregulin that talk back to the mammary epithelium remains to be elucidated as well as the cell type within the stroma that is responsible for secreting/activating these factors. Furthermore, it is unclear whether these stromal factors act directly on the stem cell or impinge on niche cells.

Candidate stromal factors are the matrix metalloproteinases (MMP) −14 and −2 [58], previously reported as EGFR targets which may activate membrane bound/ECM-sequestered growth factors and/or components of ECM such as collagens, laminin, fibronectin, proteoglycans, the TGF-β inhibitor decorin, and FasL [101]. Other attractive contenders for the stromal mediators are the fibroblast growth factors (FGFs) −7 and −2 [102], they elicit ductal morphogenesis in a 3D matrigel assay [97] and stimulate growth and branching of EGFR−/− mammary organoids in culture [103]. Interestingly, FGF-2 (basic FGF) is a critical component in the medium for mammospheres [33, 35] suggesting that activation of FGF signaling is important to stem cells [33]. Whether these FGFs are induced in the stroma in response to amphiregulin and whether they act directly on the stem cell or indirectly via niche cells both in vivo and in mammospheres remains to be determined.

amphiregulin [61,112]. Both amphiregulin and TACE are required in the mammary epithelium for ductal outgrowth [23,103]. In chimeric epithelia formed by amphiregulin−/− and WT cells, the mutant cells proliferate and contribute to cell both compartments consistent with amphiregulin acting as a paracrine mediator of estrogen action [23]. The mutant cells never give rise to an entire ductal segment but are always observed within a few cell diameters of WT cells [23] indicating that close interactions are required during ductal outgrowth to relegate the estrogen-induced signals to ERα negative stem and/or progenitor cells.

In the absence of EGFR, the cognate receptor of amphiregulin, ductal outgrowth fails to occur. Tissue recombination studies revealed that EGFR deficient stroma abolishes the growth of a wild type epithelium whereas EGFR deficient epithelium grows in a WT fat pad [131] indicating that the prime target of amphiregulin are stromal cells, which may in turn release factors to stimulate either the niche or the stem cells (Fig. 6).

During estrogen-driven ductal outgrowth, epithelial cells at the tip of the growing ducts, are in direct contact with the stroma because the basal lamina is disrupted in this zone of active proliferation [88]. Thus, it is conceivable that stromal cells may be part of the niche (Fig. 6).
Fig. 7. Model of pregnancy-associated stem cell niche. Systemic progesterone and prolactin, in the presence of estrogens, act together in the development of the mammary gland during pregnancy. These hormones reach their receptors in luminal cells belonging to the stem cell microenvironment. In response, the sensor cells release several factors like IGF-2, Wnt4 and RANKL. These influence the surrounding cells, and may directly or indirectly influence the stem cells and parity induced-stem cells. MSC: mammary stem cell; PI-MSC: parity identified-mammary stem cell; LC: luminal cell; MC: myoepithelial cell; FB: fibroblast; AD: adipocyte; HR: hormone receptor; BM: basal membrane.

It is also possible that default inhibitory influences exerted on the niche are relieved by a stroma-derived factor. TGF-β acts as a key negative regulator of proliferation on many epithelial cell types including MECs [28,36]. Intriguingly, decorin is a TGF-β inhibitor highly expressed in mammospheres [33] and could be secreted by stromal cells in response to EGFR stimulation.

TGF-β regulates the expression of Wnt-5a [89], a Wnt that activates non canonical wnt signaling [52,60,122,130] and can inhibit canonical wnt signaling [71,119,129]. Wnt-5a mediates a subset of TGF-β inhibitory actions and in Wnt-5a−/− mammary epithelia the highly proliferative ductal tips, terminal endbuds (TEBs) are enlarged [89] suggesting that down modulation of TGF-β/Wnt-5a signaling, may indeed promote activation of stem cells. However, it is difficult to reconcile these functional data with the observation that expression of Wnt-5a mRNA is enriched in the TEBs compared to subtending ducts [58] when TGF/β activity and hence Wnt-5a expression should be down modulated at the active growth front.

In the mammary gland, as in the hematopoietic system and in the epidermis, Wnt signaling may be important in determining the stem cell compartment. Mice that ectopically express Wnt-1 in the mammary epithelium have an increase in the CD29high CD49high compartment [96] and tumors arising in the mammary glands of these mice express progenitor markers [63]. Moreover, microRNAs, which have been shown to be enriched in stem cell populations, are increased when Wnt-1 is overexpressed in MECs [43]. Whether wnt signaling is required in the stem cell itself, in niche cells, or exerts more indirect effects remains to be addressed.

Expression of Notch-3, SCF and the Wnt receptor Frizzled 2 has been found increased when expression profiles of mammospheres and of cells grown on adherent plates were compared [33] suggesting that Notch, Wnt and Kit signaling all of which are important in the hematopoietic niche may also have a role in the mammary gland niche. Consistent with such a scenario, inhibition of Notch signaling interferes with growth of mammospheres [34]. In vivo data, however, suggest that regulation of Notch signaling is complex and cell type dependent. Both ectopic activation [47,98] as well as loss of function [91] can interfere with ductal outgrowth arguing that the timing and the level of expression as well as the targeted cell type will need to be considered. BMP signaling function has not been addressed in the mammary gland.
STEM CELL NICHE DURING PREGNANCY

Once the ducts reach the edges of the fat pad, they cease to grow; this usually coincides with the onset of sexual maturity. With recurrent estrous cycles the complexity of the ductal system increases through lateral branching driven by cyclic increases in serum progesterone levels [88]. During early pregnancy, progesterone levels increase and sidebranching is further enhanced; later, prolactin drives alveologenesis. It is not clear to what extent this proliferation requires stem cell activity or merely reflects an expansion of transient amplifying cells.

During pregnancy a novel stem cell population becomes apparent, the parity-identified stem cells, (PI-SCs). These cells were tracked using double transgenic mice, which express Cre driven by the milk protein, whey acidic protein (WAP) promoter and a LacZ gene that is preceded by a floxed stop codon (WAP-Cre/Rosa26-flox-stop-flox-lacZ). In these double transgenic mice, MECs that differentiate and hence express Cre/Rosa26-flox-stop-flox-lacZ). In these double transgenic mice, which express Cre driven by the milk protein promoter, WAP will induce Cre recombinase resulting in the expression of β-galactosidase. Subsequently, the LacZ positive cells contribute substantially to the outgrowth and are found in all cellular compartments [69]. Interestingly, PI-SCs are grafted on their own, as achieved by injecting them into cleared fat pads in limiting dilutions, they do not give rise to complete ductal outgrowths but form lobules indicating that their intrinsic potential is more restricted than that of the common mammary SCs [12,124,125].

A few PI-SCs develop during estrous [10] but the bulk of PI-SCs is induced during pregnancy [124]. This points to the existence of a pregnancy-specific stem cell niche. Intriguingly, the pregnancy stem cell niche can induce adult testicular cells into milk-producing MECs [11]. The male germ cells are capable of self-renewal upon transplantation and contribute to both luminal and myoepithelial cell lineages in the reconstituted glands [11].

What distinguishes the pregnancy niche from the niche in the virgin female? During pregnancy the same cell types will be interacting with stem cells as in the virgin animal (Figs 6 and 7). In contrast to the “pubertal niche” the basal membrane remains intact during pregnancy suggesting that interactions with the stroma may be less extensive at this stage. At this stage, the myoepithelial layer is less continuous probably allowing luminal cells to contact the basal membrane [88].

Progesterone and prolactin act in the presence of estrogen, that are required for progesterone receptor (PR) expression, to induce side branching and alveologenesis [17,18] (for review see [13]) by acting on their respective sensor cells. As a result local signals are induced that may directly impinge on stem cells or act on neighboring luminal or myoepithelial cells, possibly stromal cells that are part of the niche (Fig. 6).

Receptor Activator for Nuclear Factor κB Ligand (RANKL) [14,74], Wnt-4 [16], calcitonin [44] and insulin like growth factor-2 (IGF-2) [14] (Fig. 7) have been implicated downstream of the pregnancy hormones. We speculate that Wnt-4 mobilizes stem cells either acting on them directly or indirectly through activation of niche cells. The offspring is induced to proliferate by RANKL. The functional importance of calcitonin induction by progesterone in this context is unclear. IGF-2 is an important factor for maintaining human embryonic stem cells in culture [8]; whether its role in alveologenesis in the mammary gland is linked to effects on stem cells remains to be tested.

A better understanding of PI-SCs and their niche is very important in light of the observation that early pregnancies have a protective effect against cancer. This is evident from epidemiologic studies on humans [40] and can be reproduced in rodents [19,38]. Indeed, Wagner and Smith argue that the specific properties of the PI-SC population that set them apart from MSCs may underly the protective effect of pregnancy.

CONCLUSIONS

Thinking about stem cells and their niches has been largely influenced by the pioneering work performed in the readily accessible hematopoietic system. In the mammary gland, in addition to the immediate microenvironment, various stromal components and above all the systemic hormonal milieu are crucial control elements of stem cell function. The female reproductive hormones acting through sensor cells within the luminal epithelium, trigger a cascade of paracrine signal-
ing events that impinge on stem cells and their niches. Some signaling pathways have been implicated that appear to be of general importance to stem cells and their niches such as Wnt, Notch, BMP signaling. In the mammary gland, their exact role remains to be established. In addition a number of pathways with organ-specific functions, such as amphiregulin in the hematopoietic niche versus osteopontin in the hematopoietic niche have been identified.

The challenge is to reconcile morphological hallmarks of stem cells in situ and FACS phenotyping ex vivo so that the stem cells and their niches can eventually be defined and characterized in vivo. We need to integrate the models about stemness and differentiation as in the hematopoietic system where cells are single and live in suspension, with the complex 3D structure of the mammary gland that changes substantially during development, as well as the mesenchymal interactions that influence it.

A better understanding of the mammary niche and its dependence on developmental stage and hormonal milieu will provide novel insights into how breast cancer arises.

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