A novel role of microRNA146b in promoting mammary alveolar progenitor cell maintenance

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Summary

In this report, we have shown that miR146b promotes the maintenance of pregnancy-derived mammary luminal alveolar progenitors. MiR146b expression was significantly higher in the mammary glands of pregnant and lactating mice than in virgin mice. Furthermore, miR146b levels were significantly higher in mouse mammary glands exposed to the sex hormones, estrogen and progesterone, compared with those of untreated control animals. Pregnancy-derived primary mouse mammary epithelial cells in which miR146b was knocked down showed a significant reduction in the number of hollow acinar organoid structures formed on three-dimensional Matrigel and in β-casein expression. This demonstrates that miR146b promotes the maintenance of pregnancy-derived mammary luminal alveolar progenitors. It has been shown that mouse mammary luminal progenitors give rise to hollow organoid structures, whereas solid organoid structures are derived from stem cells. Among several miR146b targets, miR146b knockdown resulted in preferential STAT3 isoform caused mammary epithelial cell death and a significant reduction in β-casein mRNA expression. Therefore, we conclude that during pregnancy miR146b is involved in luminal alveolar progenitor cell maintenance, at least partially, by regulating STAT3β.

Key words: MicroRNA-146b, Alveolar Progenitors, STAT3

Introduction

Development of the mammary gland is a highly dynamic process and involves distinct embryonic, pubertal, pregnancy, lactation and involution stages. Hormones and growth factors undoubtedly play key roles in all stages of mammary development. The essential role of micro-RNAs (miRNAs) in the various stages of mammary development is being recognized. Not surprisingly, a disruption in miRNAs, has been implicated in human breast cancers (Watson and Khaled, 2008).

miRNAs have been linked to both normal physiology and pathology, including regulation of developmental processes, proliferation, differentiation and apoptosis. Loss of mature miRNAs resulted in a failure of the cells to differentiate (Andl et al., 2006; Bernstein et al., 2003; Kanellopoulou et al., 2005; Muljo et al., 2005), providing evidence that miRNAs are involved in stem cell maintenance, self-renewal, and/or differentiation (Greene et al., 2010). MiRNAs have been found in co-expressed clusters during mammary gland development, suggesting co-regulation of miRNA groups (Avril-Sassen et al., 2009; Piao and Ma, 2012). Furthermore, it has been shown that miRNAs encoding receptors of endocrine hormones are direct targets of specific miRNAs, which also indicates the role of miRNAs in mammary gland development (Cui et al., 2011).

Mouse microRNA-146b-5p (miR146b) is located on chromosome 19 at position 19q13, whereas miR146a is located on chromosome 11 at position 11q15 (Fujita et al., 2011). Structurally, the mature miR146a and miR146b differ by only two nucleotides at the 3’ end, a region that is believed to play a minor role in target recognition, and in many instances miR146a and miR146b have demonstrated redundant activities (Lewis et al., 2003). The role of miR146b in cancer progression is currently controversial. MiR146b overexpression has been shown to be associated with aggressive behavior in papillary thyroid carcinomas (PTC) (He et al., 2005) and triple negative breast cancers (Garcia et al., 2011). Conversely, miR146b has also been shown to inhibit carcinogenesis. Welch and colleagues recently classified miR146a/b as a ‘metastasis-suppressing metastamir’ (Hurst et al., 2009). The term ‘metastamir’ refers to metastasis regulatory miRNAs that have an impact on critical steps in the metastatic cascade, such as epithelial–mesenchymal transition (EMT), apoptosis and angiogenesis (Hurst et al., 2009). For example, miR146b was shown to inhibit glioblastoma cell migration (Xia et al., 2009) and to suppress breast cancer metastasis (Bhaumik et al., 2008; Hurst et al., 2009). These findings suggest that miR146b may play a dual role in cancer and that much remains to be learned about the mechanism by which this miRNA may participate during carcinogenesis.
Recently, our group isolated distinct populations of ductal-limited, alveolar-limited and multipotent mammary progenitors by single-cell cloning of the COMMA-D (CD) cell line (Kittrell et al., 2011). Transplantation of the alveolar-limited clone, referred to herein as an ‘alveolar progenitor cells/clone’, generated functional alveolar structures in vivo. Transplantation of ductal-limited clone, referred to hereinafter as a ‘ductal progenitor clone’, generated only ductal structures devoid of functional alveolar differentiation potential. Transplantation of two of the clones, referred to hereinafter as ‘multipotent progenitor clones’, generated ducts and alveolar structures (Kittrell et al., 2011). Using this model system, we identified a hormonally regulated miRNA, miR146b, that plays a pivotal role in alveolar progenitor cell maintenance.

As mentioned above, altered expression of miR146b has been linked to an invasive and metastatic capacity in diverse cancers. Finding a hormonally regulated miRNA that regulates both the processes of alveolar cell maintenance and breast cancer may provide one missing link in the molecular pathways implicated in hormonal regulation of human breast cancer.

Results
MiR146b is highly expressed in the alveolar progenitor cells
The distinct mammary alveolar, ductal and multipotent progenitor clones were used to screen for changes in the expression of 84 unique miRNAs (Fig. 1A) by an RT2 miRNA PCR array. This screening showed a set of 74 upregulated miRNAs in the clones with in vivo morphogenic potential (including the alveolar progenitor, ductal progenitor and multipotent progenitors) when compared with clones that exhibited no in vivo morphogenic potential. Twenty miRNAs showed a 10 to 20-fold increase, and 15 miRNAs showed >20-fold increase (Fig. 1A). Among those upregulated, miR146b showed >72-fold increase, and miR-203 showed >77-fold increase. To examine the expression of upregulated miRNAs in the specific progenitor clones, RT-qPCR was used on RNA derived from progenitor clones grown on Matrigel in three dimensions (3D). MiR146b was the only miRNA that showed differential expression as it was highly expressed by the alveolar progenitor clone with a 10-fold increase compared with the level in the ductal and the multipotent progenitor clones (10±1.8 versus 1.29±0.7 and 1±0.23, respectively; Fig. 1B). Others such as miR203 expression showed no difference between the distinct progenitor clones (data not shown).

To confirm these findings, miR146b expression levels were examined in the primary mammary epithelial cells (PMECs) derived from virgin, pregnant, lactating and 10 days post-weaning (involuting) female BALB/c mice (Fig. 1C). These studies showed a 6.3-fold increase in miR146b expression levels in the PMECs derived from pregnant mice compared with those from virgin and post-weaning mice (6.3±0.83 versus 1.0±0.44 and 1.3±1.17, respectively, P<0.05) and a 12-fold increase in PMEC derived from lactating compared with virgin and post-weaning mice (12.03±1.45 versus 1.0±0.44 and 1.3±1.17, respectively, P<0.05). MiR146b expression levels were normalized to keratin-18, in order to exclude the contribution of stromal cells. Because the pattern of expression of miR146a and miR146b are similar and the isoforms are reported to be co-regulated (Garcia et al., 2011), miR146a levels were similarly examined by RT-qPCR. The results showed that miR146a levels increased similarly during pregnancy and lactation. However, the increase in miR146a expression levels, were not as significant as miR146b (Fig. 1C).

Recently, the progenitor and differentiated cells in the mouse mammary glands and human breast were phenotypically and functionally characterized by fluorescence-activated cell sorting (FACS) followed by transplantation. These studies defined the subpopulations enriched in stem cells by the expression of unique surface markers, lineage-negative (lin−) CD24+CD29hi and lin−CD24+CD49fhi, luminal progenitors by lin−CD29hiCD24+CD61+, differentiated estrogen receptor-positive (ER+) luminal cells by CD24+CD133+ and myoepithelial progenitors by CD29hiCD24lo (Asselin-Labat et al., 2007; Shackleton et al., 2006; Stingl et al., 2006).

To find the mammary epithelial sub-population with the highest level of miR146b expression, the luminal progenitors (lin−CD29hiCD24+CD61+), basal/stem cells (lin−CD29hiCD24-) and differentiated luminal cells (lin−CD29hiCD24+CD61+) were FACS sorted from the mammary glands of virgin and pregnant BALB/c mouse, followed by RNA extraction and RT-qPCR. The results revealed that during pregnancy miR146b expression levels were ~2-fold higher in the luminal progenitors when compared with the basal/stem cells (10.6±2.17 versus 4.6±0.19; P<0.05) and 1.6-fold higher when compared with the differentiated luminal cells (10.6±2.17 versus 6.3±0.68; P<0.05; Fig. 1D,E). However, there were no differences in miR146b expression levels between these epithelial subpopulations derived from virgin mouse mammary glands (Fig. 1E). Interestingly, miR146b was 10.5-fold higher in the luminal progenitors (alveolar) of pregnant compared with virgin (ductal) mice (10.6±2.17 versus 1.01±0.38; P<0.05). Since expression was significantly higher in the luminal alveolar progenitors than in stem cells and differentiated luminal cells, we focused our studies on the functional role of miR146b in the luminal alveolar progenitor cells.

Sex hormones and prolactin result in the upregulation of miR146b levels in primary mammary epithelial cells
Since miR146b expression levels were significantly higher in all epithelial subpopulations from pregnant compared with virgin mice, we hypothesized that pregnancy hormones may participate in its upregulation. To investigate the influence of hormones on miR146b, we exposed glands from virgins to estrogen (E) and progesterone (P) in vivo. E plus P hormonal pellets were placed subcutaneously in virgin BALB/c mice for 3 weeks. Following treatment, the mammary glands were removed followed by PMEC isolation. RT-qPCR revealed that E plus P treatment resulted in a 2.7-fold increase in miR146b expression levels, normalized to keratin-18, compared with the level in the non-treated controls (2.7±0.08 versus 1.0±0.54; P<0.05; Fig. 2A). To rule out the contribution of stroma, PMECs derived from virgin mouse mammary glands were treated with E plus P, prolactin alone, or all three hormones for 10 days in vitro. A similar rise in miR146b levels was found upon exposure to E plus P, prolactin alone, or all three hormones compared with the level in the untreated cells (1.46±0.09 and 1.38±0.09, respectively, versus 1±0.26; P<0.05; Fig. 2C). These results suggested that sex hormones and or prolactin, directly or indirectly, mediate the upregulation in miR146b levels.

MiR146b mediates the maintenance of alveolar luminal progenitor cells
To assess the functional role of miR146b, ‘GFP’-labeled Locked Nucleic Acid oligonucleotides (LNA™, Exiqon) complementary
to miR146b were used to knockdown miR146b expression. For these studies, PMECs derived from BALB/c mouse mammary glands and/or CD-derived alveolar progenitor cells were used. In order to examine the specificity of the miR146b LNA, RT-qPCR was used to examine miR146a levels following miR146b knockdown. The data showed a non-significant reduction in miR146a, confirming that the knockdown was selective against miR146b (Fig. 3A). Furthermore, time-course experiments using
Role of MiR146b in alveolar progenitor cell survival

The CD-derived alveolar progenitor cells followed by RT-qPCR showed stable miR146b knockdown up to 72 hours post-transfection (supplementary material Fig. S1A). To assess effects on alveolar progenitor cell survival, the CD-derived alveolar progenitor cells were collected at 48, 60 and 72 hours post-transfection followed by staining with the LIVE/DEAD® Fixable Dead Cell Staining Kit. The fluorescent dye can permeate the compromised membranes of dead cells and react with free amines both in the interior and on the cell surface. A significant reduction in the alveolar progenitor cells was seen at 72 hours post-transfection (Fig. 3C,D; supplementary material Fig. S1B). Effects on cell death was confirmed by a western blot analysis, which showed an increase in cleaved caspase 3 in the alveolar progenitor cells knocked down of miR146b compared with the control groups (Fig. 3B). Our data showed selective effect on cell survival in the alveolar progenitor cells when compared with the CD-derived ductal limited and multipotent progenitor cells (Fig. 3C,D; supplementary material Fig. S1B). As seen in supplementary material Fig. S1C, >80% of the total population of CD cells were transfected with the miR146b LNA. Even though ~60% of the PMECs were transfected with the miR146b LNA, there was not a significant effect on survival of the total population of PMECs derived from pregnant mouse mammary glands (data not shown). However, as seen in Fig. 3E,F, there was a significant reduction in the percentage of alveolar luminal progenitor cells knocked down of miR146b (lin’CD29loCD24hiCD61’) compared with the control (0.55±0.097% versus 1.0±0.18%; means ± s.e.m., P<0.05).

To further examine the functional role of miR146b in the luminal alveolar progenitors, an assay involving three-dimensional (3D) hollow or solid acinar organoid formation efficiency on Matrigel was performed. Based on previously published studies, the mammary stem cells give rise to solid acinar organoids, whereas luminal progenitors form hollow acinar organoid structures when grown in 3D Matrigel cultures (Guo et al., 2012; Lim et al., 2009; Shackleton et al., 2006; Stingl et al., 2006). For these studies, PMECs were isolated from pregnant mouse mammary glands followed by transfection with miR146b LNA. At 24 hours post-transfection, the cells were transferred from tissue culture plates to Matrigel and treated with prolactin for 7 days. The epithelial organoids were counted and classified into solid or hollow acinar organoid structures. In addition, some cells were recovered from each group to test for knockdown efficiency using RT-qPCR. As seen in Fig. 4A, the PMECs were efficiently knocked down of miR146b up to 7 days post-LNA transfection. Furthermore, there was a 5.3-fold reduction in the number of hollow acinar organoid structures upon miR146b knockdown compared with the control groups (2±0 versus 10.6±2.4; means ± s.e.m., P<0.05), reflecting a reduction either in the number of luminal alveolar progenitors or their ability to give rise to the differentiated luminal structures on Matrigel (Fig. 4B,C). However, the number of the solid structures remained constant in all of the treatment groups. The number of hollow acinar structures were higher with prolactin treatment (16±4 versus 7±1.15; means ± s.e.m., P<0.05). However, the effect of miR146b knockdown was similar with or without treatment with prolactin.

To further investigate the effect of miR146b knockdown on the alveolar progenitor cell function we tested the ability of the CD-derived alveolar progenitor cell line to express β-casein milk protein following miR146b knockdown. β-casein is endogenously expressed in the CD-cell line and can be rapidly induced by prolactin in vitro (Ball et al., 1988). The CD-derived alveolar progenitor cells and primary mouse epithelial cells were transfected with miR146b LNA inhibitor in 2D culture. At 24 hours post-transfection the cells were transferred to Matrigel and treated with prolactin (PRL) for another 72 hours. The cells were then recovered from the Matrigel and an RT-qPCR for β-casein mRNA levels was performed. MiR146b knockdown produced a significant reduction in β-casein expression compared with the level in the control CD-derived alveolar progenitor cells treated with PRL (2.3E±0.1E± versus 3.7E±0.2E±; mean normalized expression ± s.e.m., P<0.05; Fig. 4D), as well as in the primary mouse mammary epithelial cells (0.1±0.03 versus 1±0.04; mean normalized expression ± s.e.m., P<0.05; Fig. 4E). These data show that miR146b regulated β-casein expression, a marker of alveolar cell viability and function.
MiR146b promotes alveolar progenitor cell maintenance, at least partially, through downregulation of STAT3

We hypothesized that miR146b may promote alveolar progenitor cell survival and function by suppressing transcription factors with key roles during mammary gland development during pregnancy and or involutions. Previous studies have shown miR146b targets include interleukin (IL)-1 receptor associated kinase (IRAK1), TNF receptor-associated factor 6 (TRAF6), nuclear factor kappa-light-chain-enhancer of activated B cells (NFkB) and SMAD family member 4 (SMAD4) (Bhaumik et al., 2008; Geraldo et al., 2012). Since some miRNAs might also upregulate expression (Ghosh et al., 2008; Vasudevan et al., 2007), we also examined signal transducer and activator of transcription 5α (STAT5a) and Elf5, two transcription factors that promote alveologenesis (Bouras et al., 2008; Siegel and Muller, 2010). For these experiments, the CD-derived alveolar progenitor cells were knocked down of miR146b to screen for targets. As shown in Fig. 5A, knockdown of miR146b by LNA inhibitors decreased the expression of SMAD4, IRAK1 and NFkB, but had no effects on STAT5a, TRAF6 and Elf5. These data suggest that the previously identified targets of miR146b may not be directly regulated in the mammary glands.

STAT3 is a transcription factor that has been implicated previously in promoting mammary epithelium cell apoptosis at the onset of involution (Chapman et al., 1999). This finding prompted us to examine whether miR146b might target STAT3. The 3’UTR of the STAT3 gene was analyzed with Ensembl 2012 (Flicek et al., 2012) and RNA22 target prediction program (Miranda et al., 2006), and two putative miR146b binding sites were identified in the STAT3a 3’UTR (1.9 kb), while three miR146b binding sites were identified in STAT3b 3’UTR (3.6 kb; Fig. 6A). The two binding sites in STAT3a 3’UTR are similar to STAT3β 3’UTR.

Knockdown of miR146b in the CD-derived alveolar progenitor cells, resulted in a significant increase in STAT3β expression compared with STAT3α (Fig. 5B; supplementary material Fig. S1E). A western blot using STAT3β-specific antibody (generated
in the laboratory of D. J. T.) showed that the upregulated lower band in total STAT3 western blot was indeed STAT3\(b\) and not a product of STAT3\(a\) proteolytic degradation (Fig. 5B; middle panel). The STAT3\(b\)-specific antibody was raised against the unique C-terminal 7 residues in STAT3\(b\) protein. The increase in STAT3\(b\) expression upon miR146b knockdown in PMECs derived from pregnant mouse mammary glands was up to 2-fold compared with the controls (Fig. 5C, and densitometry data not shown). These data suggest that miR146b preferentially regulates STAT3\(b\) protein expression.

**MiR146b binds to the 3′ UTR of STAT3\(a\) and STAT3\(b\)**

To determine if the putative miR146b binding sites resided within the 3′UTR of STAT3, we utilized luciferase constructs in which the 3′UTRs of STAT3\(a\) or STAT3\(b\) were inserted downstream of the firefly luciferase reporter gene (driven by SV40 promoter). A chimeric mRNA is transcribed consisting of the firefly luciferase and the STAT3\(a\) or 3′UTR sequence. The vector also contains a Renilla luciferase as an internal control downstream of the CMV promoter (Fig. 6B). To examine whether miR146b directly regulates STAT3\(a\) and \(b\), 293T cells were transfected either with the STAT3\(a\) or \(b\) 3′UTR reporter construct along with the miR146b precursor and the Pre-miR\(^{\text{TM}}\) negative control. Firefly luciferase activity was then measured and normalized to Renilla luciferase activities in the same wells.

The results showed a statistically significant reduction in luciferase reporter activity of STAT3\(a\) (70.2±1.7% versus 100±2.4%; mean normalized luciferase units ± s.e.m., \(P<0.0001\)) and in luciferase reporter activity of STAT3\(b\) (72.9±3.2% versus 100±3.1%; mean normalized luciferase units ± s.e.m., \(P<0.0001\)) when cells overexpressed miR146b compared with cells that expressed the Pre-miR\(^{\text{TM}}\) negative control (Fig. 6C). Luciferase reporter activity of the STAT3\(a\) (data not shown) and STAT3\(b\) 3′UTR was also significantly lower in PMECs from pregnant mice, which had significantly higher miR146b expression, compared with luciferase activity in PMECs from virgin mice (71±9.8% versus 100±5.6%; mean normalized luciferase units ± s.e.m., \(P<0.0001\); Fig. 6D). These data suggest that miR146b directly binds the 3′UTR of STAT3\(a\) and STAT3\(b\). In order to examine effects on STAT3 splicing, real time RT-PCR using specific primers to STAT3\(a\) and STAT3\(b\) was performed. The results showed that, upon miR146b knockdown, STAT3\(a\) and STAT3\(b\) mRNA levels were unchanged (Fig. 6E). Therefore, most likely, miR146b does not affect STAT3 splicing.

**STAT3\(b\) may play a functional role during mammary gland involution**

The contribution of specific STAT3 isoforms during mammary epithelial cell apoptosis at the onset of involution is not known at
this time. To correlate miR146b expression with STAT3α/β levels during mammary development, we performed total STAT3 western blots using CD-derived progenitor clones as well as BALB/c-derived mammary glands from virgin, pregnant, lactating, and 1, 3 and 6 days post-weaning (involution) mice. Interestingly, STAT3β levels were inversely correlated with miR146b expression in the CD-derived progenitor clones (compare Fig. 7A and Fig. 1B). Furthermore, STAT3β levels were inversely correlated with miR146b levels at each physiological stage, while STAT3α was equally expressed throughout the distinct mammary developmental stages (compare Fig. 7B and Fig. 1C). As shown previously (Chapman et al., 1999) and in this study (Fig. 7C,D), there is an increase in STAT3α and STAT3β tyrosine phosphorylation (phosphor-Y705-STAT3, present in both STAT3α and STAT3β) prior to a significant rise in STAT3α serine phosphorylation (phosphor-S727-STAT3 present only in STAT3α) beginning at day 2 post-weaning. These findings suggest that STAT3β, either as homodimers or heterodimer with STAT3α or other transcription factors, may contribute to the onset of mammary involution. The specific STAT3 isoform knockout mice have been generated, but the contribution of each isoform to mammary involution has not been reported (Maritano et al., 2004). In order to relate the role of miR146b to STAT3β, we studied the effect of STAT3β overexpression on alveolar luminal cell viability and function. A cDNA overexpressing STAT3β was transfected into the CD-derived alveolar progenitor cells. The cells were collected at 24 hours post-transfection and stained with the LIVE/DEAD® Fixable Dead Cell Staining Kit. As expected, similar to the effects seen with miR146b knockdown, STAT3β overexpression reduced cell viability in the CD-derived alveolar progenitors by 20% compared with the controls (76.6±1.04% versus 95.7±1.5%; means ± s.e.m.; Fig. 7E,F). Furthermore, we examined whether overexpression of STAT3 isoforms resulted in a similar reduction in β-casein mRNA levels as seen with miR146b knockdown. For these experiments, virgin-derived mouse mammary epithelial cells were transfected with the STAT3β- and STAT3α-overexpressing cDNA clones,
transferred onto Matrigel, and treated with PRL for 72 hours. The cells were then recovered for RT-qPCR. Fig. 7G, shows that similar to miR146b knockdown, overexpression of STAT3a and to a higher extent STAT3b resulted in a significant reduction in β-casein mRNA levels compared with the control (0.2±0.01 and 0.06±0.02 versus 1±0.13; mean normalized expression ± s.e.m., P<0.05). These data suggest that miR146b may regulate alveolar progenitor cells, at least partially, through regulation of STAT3b. Since we did not find a significant regulation of STAT3a protein expression by miR146b, we believe that miR146b effects are at least partially, through regulation of STAT3b.

Discussion
In this study, it is demonstrated that miR146b expression is upregulated in the luminal alveolar progenitor cells during pregnancy and lactation and upon hormonal stimulation with (E plus P) and prolactin. Furthermore, transient knockdown of miR146b reduced survival of luminal alveolar progenitor cells. These data suggest that miR146b participates in alveologenesis under the influence of sex hormones and prolactin. The effects of miR146b may, at least partially, be through suppression of key transcription factors involved in alveolar cell death during involution, such as STAT3b.
Although many studies report the role of miR146b in cancer, its role in mammary gland development is still unknown. Bockmeyer et al. studied the miRNA profiles of human basal and luminal epithelial cells as well as basal and luminal breast cancers (Bockmeyer et al., 2011). These studies showed miR146b to be a basal-specific miRNA and to be upregulated in basal-like breast cancers when compared with luminal A and B breast cancer subtypes (Bockmeyer et al., 2011). Other than in this report, expression of miR146b and its role in mammary epithelial cell development have not been described.

We reasoned that miR146b might play a role in alveolar cell maintenance by regulating a known target such as IRAK1/TRAF6, NF-κB, or SMAD4 or by regulating transcription factors with known roles during alveologenesis such as STAT5a and E74-like factor 5 (ELF5). A review of the literature showed miR146b to regulate TGF-β signaling by repressing SMAD4 in thyroid cancers (Geraldo et al., 2012). TGF-β has been shown to play an important role during mammary involution by inducing cellular apoptosis and alveolar collapse in the early stages of involution (Flanders and Wakefield, 2009). Furthermore, miR146b has been shown to inhibit breast cancer cell invasion and migration by targeting IRAK1/TRAF6 and by downregulating NF-κB signaling (Bhaumik et al., 2008). NF-κB has also been shown to destabilize STAT5a, a key transcription factor that promotes alveologenesis (Floyd et al., 2007). Interestingly, miR146b knockdown did not result in the upregulation of any of the known suspected targets. Therefore, the previously identified miR146b targets may not be similarly regulated in normal mammary glands.

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**Fig. 7. STAT3β is activated in the mammary epithelial cells during involution.**

(A) Western blot analysis of total STAT3α/β in the CDβ cell line and the CD-derived clones. (B–D) Western blot analysis of total STAT3α/β (B); phosphorylated tyrosine 705 (phospho-Tyr-STAT3; C); phosphorylated serine 727 (phosphor-ser-STAT3; D) in total glands from virgin (V), pregnant (P) and lactating (L) mice and from mice at different physiological stages of involution: days 1, 3 and 6 (B–D) and 10 (C) days post-weaning (D-PW). (E) Representative histogram of flow cytometry analysis of the alveolar progenitor cell line stained with the LIVE/DEAD® Fixable Dead Cell Staining Kit at 24 hours post-transfection with STAT3β overexpression clone. (F) Quantification of the percentage of live cells (n=3, *P<0.05). (G) RT-qPCR of mouse mammary epithelial cells transfected with STAT3α/STAT3α-OE- or STAT3β (STAT3β-OE)-overexpression clone compared with cells transfected with the empty vector (control). The cells were cultured on Matrigel for 72 hours, with prolactin treatment. Cells were recovered and analyzed for β-casein expression by qPCR (n=3, *P<0.05 compared with other cells). Data are means ± s.e.m.
Since STAT3 transcription factors were reported to play key roles in mammary epithelial cell apoptosis at the onset of involution, we explored the role of miR146b in regulating STAT3 isoforms. STAT3 has two main isoforms: the full-length STAT3\(^a\) and the truncated STAT3\(^b\), by alternative mRNA splicing of exon 23 (Huang et al., 2007; Zammarchi et al., 2011). In STAT3\(^b\), the 55-amino acid C-terminal acidic transactivation domain of STAT3\(^a\) is replaced by seven unique amino acid residues (Zammarchi et al., 2011). An important factor in STAT3 functional heterogeneity may be the existence of these two alternatively spliced isoforms. As suspected, miR146b knockdown resulted in upregulation of STAT3 expression in both PMECs derived from pregnant mammary glands and in the alveolar progenitor cells. However, the upregulation of the STAT3\(^b\) isoform was higher than that of STAT3\(^a\). There are several possibilities for preferential regulation of STAT3\(^b\) protein expression by miR146b. One is the existence of an additional miR146b binding site on the predicted STAT3\(^b\) 3′UTR, resulting in a different secondary structure and higher efficiency of mir146b binding (Flicek et al., 2012). Another mechanism, may be the regulation of alternative splicing events. The interplay between the miRNA and splicing regulation has been demonstrated previously. These studies showed that miR124 and miR133 regulate the RNA-binding protein, poly(A) tract-binding protein 1 (PTBP1). PTBP1 is a global repressor of alternative pre-mRNA splicing in non-neuronal cells (Boutz et al., 2007; Makeyev et al., 2007). Additionally, ESEfinder and PESX algorithms predicted a putative exonic splicing enhancer (ESE), which overlaps with one of the predicted miR146b binding sites. MiR146b binding to this overlapped region may regulate ESE. MiR146b could also indirectly regulate STAT3 alternative splicing by targeting key splicing factors. Real-time RT-PCR results showed that STAT3\(^a\) and STAT3\(^b\) mRNA levels remain constant in miR146b knockdown cells despite striking changes in protein expression, suggesting that miR146b most likely affects STAT3 protein translation.

STAT3 is characterized by its capacity to activate different sets of genes in different cell types (Levy and Lee, 2002), and the different STAT3 isoforms seem to regulate cell survival and maintenance in a cell-type- and context-specific manner by suppressing or activating a variety of target genes (Pensa et al., 2009). STAT3\(^b\) has been considered a dominant-negative factor by inhibiting STAT3\(^a\) gene transactivation (Caldenhoven et al., 1996; Schaefer et al., 1995). More recent studies have demonstrated that STAT3\(^b\) may activate a specific set of genes and possess non-redundant functions to STAT3\(^a\). Zammarchi et al. used morpholino oligomers to redirect endogenous STAT3 alternative splicing from STAT3\(^a\) to STAT3\(^b\) (Zammarchi et al., 2011). This study showed that inducing the STAT3\(^b\) isoform in MDA-MB-435 cells caused cell death and tumor regression in the xenograft models. The same study showed that STAT3 canonical
target genes were not affected by the switch from STAT3α to STAT3β. Rather, a unique expression signature seems to be associated with the physiological α-to-β splicing shift (Zammarchi et al., 2011). In mammary gland development, it has become clear that STAT3 regulates a complex series of processes involving alveolar epithelial cell death during involution by maintaining the balance of inflammatory and anti-inflammatory signaling (Pensa et al., 2009). The contribution of specific STAT3 isoforms to the mammary involution process is still not known. Knockout mice of specific STAT3 isoforms have been generated (Maritano et al., 2004). It is reported that STAT3β knockout mice are viable, fertile and capable of nursing their pups (Maritano et al., 2004). However, a detailed characterization of the mammary glands during involution has not been reported in these mice. Our study clearly demonstrates that both isoforms of STAT3 are upregulated during involution, and overexpression of STAT3β in the alveolar luminal progenitor cells reduced their survival. It would be interesting to decipher the contribution of specific STAT3 isoforms to the mammary involution process in specific STAT3-isoform knockout mice.

In conclusion, we have demonstrated that estrogen plus progesterone and prolactin result in the upregulation miR146b. Furthermore, miR146b levels rise in the mouse mammary glands during lactation and pregnancy. MiR146b upregulation during pregnancy and lactation may mediate the maintenance of alveolar luminal progenitor cells by selectively suppressing STAT3β and to a lesser extent STAT3α. Following weaning and during the process of involution, miR146b levels drop to basal levels, which result in the upregulation of STAT3 isoforms and induction of alveolar cell death (Fig. 8). The LNA inhibitors induce a transient knockdown in miR146b; thus they are not suitable for long term in vivo transplantation studies. It will be essential to confirm our results by more stable miR146b knockdown and overexpression in mammary epithelial cells followed by in vivo transplantation studies.

### Material and Methods

#### Cells

CommaDβ-derived clones (SP-1, SP-3, SP-4, NSP-1, NSP-2, NSP-3, NSP-4 and NSP-5) were used. The alveolar progenitor (SP-3), ductal progenitor (NSP-2) and multipotent progenitor clones (NSP-5) have previously been described and characterized (Kittel et al., 2011). 293T cells (Clontech, cat. no. 632180) were used for lucerase reporter assays.

#### Animals

We used female BALB/c mice which were either bred or purchased from Harlan Laboratories, Inc., IN, USA. We used mice in late pregnancy, as well as mice 24 hours, 3 days and 6 days post-weaning (involving mammary glands), and virgin mice in the metestrus phase of the estrous cycle as confirmed by vaginal cytology (Caligioni, 2009). Animal experiments were conducted following protocols approved by the University of Kansas Medical Center Institutional Animal Care and Use Committee.

#### Mouse surgeries

Slow-releasing pellets containing 50 μg of estradiol and 20 mg of progesterone for hormonal treatment were placed subcutaneously on the lateral side of the neck between the ear and shoulder. After three weeks, mice were sacrificed, and the mammary glands were excised. One mammary gland was processed for embedding, and the other mammary gland was used to recover the epithelial cells according to established protocol in our laboratory (Valdez et al., 2011).

#### Cell culture: 2D/3D (MatrigelTM)

Primary mouse cells were grown and maintained in F12 (Invitrogen, cat. no. 11765) supplemented with 5% fetal bovine serum (Invitrogen, cat. no. 10438026), 1% antibiotic/antimycotic reagents (AA; Invitrogen, cat. no. 15240), 50 μg/ml gentamicin (Sigma-Aldrich cat. no. G1272), 1 mg/ml hydrocortisone (Stem Cell Technologies, cat. no. 07904), and 5 ng/ml epidermal growth factor. The CD-9 derived clones (i.e. SP-3, NSP-2 and NSP-5) were grown in DMEM/F12 (Invitrogen, cat. no. 11320) containing 2% fetal bovine serum, 5 ng/ml epidermal growth factor, 10 mM HEPES (Invitrogen, cat. no.15630), 5 μg/ml insulin, 50 μg/ml gentamicin and 1% antibiotic/antimycotic.

293T cells were grown in DMEM (Sigma-Aldrich, cat. no. D5796) containing 10% fetal bovine serum, and 1% antibiotic/antimycotic. Hormonal treatment included: 100 μM progesterone (Sigma, cat. no. P0319-25G), 100 μM β-estradiol (Sigma, cat. no. E8875-1G) and 3 μg/ml ovine prolactin (Harbor-UCLA Research and education institute). The cells were grown in humidified incubator at 37°C and 5% CO2. For 3D culture, cells were plated in media containing 2% Basement Membrane Matrix MatrigelTM (BD Biosciences, cat. no. 356234) on a thin film of MatrigelTM. Cell recovery was achieved using BD Cell Recovery Solution (BD Biosciences, cat. no. 354253).

#### Hematoxylin and Eosin staining

Excised mammary glands were fixed in 4% paraformaldehyde for 2 hours on ice, then in 70% ethanol at 4°C until processing, and then embedded in paraffin according to established protocols. The slides containing 5-μm sections were deparaffinized in xylene (X3P-1GAL, Fisher scientific) and hydrated through graded ethanol series (100%, 95%, 80% and 70%), followed by hematoxylin staining (S121A-32OZ Harris hematoxylin with glacial acetic acid, Poly Scientific) for 30 seconds and destaining in acid ethanol (0.3% of concentrated HCl in 70% ethanol) by eosin staining (S176 Eosin Phthalo Color, Poly Scientific) for 30 seconds and dehydration in graded ethanol (95% and 100%) and xylene three times for 15 minutes each, followed by placement of a coverslip over the slide with xylene-based permount (SP15-100 histological mounting medium, Fisher Scientific) (Valdez et al., 2011). Light microscopy was performed by using an Axios Imager.M2 microscope (Carl Zeiss MicroImaging, Inc., Thornwood, NY, USA). Pictures were taken at 10× magnification (objective N-Achroplan; 10×/0.25 ×0.17) with an AxioCam MRc5 High Resolution Camera (Carl Zeiss MicroImaging, Inc., Thornwood, NY, USA), and the acquisition software used was ZEISS Axiovision GmbH (Carl Zeiss MicroImaging, Inc.).

#### RNA isolation and Quantitative PCR (qPCR)

Total RNA was isolated with miRNesy Mini Kit (Qiagen cat. no. 217004) using the manufacturer’s protocol, and cDNA was synthesized from 250 ng of total RNA with miScript Reverse Transcription Kit (Qiagen, cat. no. 218061). MiRNA and mRNA were measured using miScript SYBR Green PCR Kit (Qiagen, cat. no. 218073), power SYBR® green PCR Master Mix (Applied Biosystems, cat. no. 4367659), TaqMan® Gene Expression Master Mix (Applied Biosystems, cat. no. 4369016) and primers specific for hsa-miR-146b-5p (IDT, Ref. no. 52053397), hsa-miR-144a (IDT, Ref. no. 51289005), STAT3a/b reverse (IDT, Ref. no. 86187028), STAT3a forward (IDT, Ref. no. 86187029), STAT3b reverse (IDT, Ref. no. 86187030), β-actin (Applied Biosystems, cat. no. Mm00479024m1). Reactions were performed in the StepOnePlus™ Real-Time PCR System and software (Applied Biosystems) in 96-well plates. Target gene expression was normalized to snl6b/snl6 forward (IDT, Ref. no. 50315659) and snl6 reverse (IDT, Ref. no. 50315660), actin forward (IDT, Ref. no. 86146439), actin reverse (IDT, Ref. no. 86146440) keratin 18 forward (IDT, Ref. no. 58927035), keratin 18 reverse (IDT, Ref. no. 89270346). The standard curve method was used for quantification.

#### Statistical analysis

Data are presented as mean normalized expression ± s.e.m. One-way analysis of variance (ANOVA) was used for statistical comparisons. A value of P<0.05 was considered significant.

#### Microarray studies

RNA from CDβ clones with growth potential (SP-3, SP-1, NSP-2, NSP-5) and those without growth potential (NSP-3, NSP-4, SP-1, SP-4) were combined. RNA quantity and quality (RIN >8) was checked using the Agilent Bioanalyzer. Reverse transcription was done using SA Biosciences RT™ miRNA First Strand Kit, RT™ miRNA PCR Array System (human cancer array, MAH-102E). PCR array was loaded according to SA Biosciences protocol for loading map. The data were analyzed using the ΔΔCT method. (www.sabiosciences.com/pcrarraydataanalysis.php).

#### FACS analysis and sorting

Primary antibodies used were anti-mouse CD29 (APC/CY7)-conjugated antibody (Biolegend, cat. no. 102226), anti-mouse CD24 (PE)-conjugated antibody (BD Biosciences, cat. no. 553262), anti-mouse CD61 (Alexa Fluor 647)-conjugated (Biolegend, cat. no. 104314), biotin-conjugated anti-mouse CD31 (Biolegend, cat. no. 102404), biotin-conjugated anti-mouse CD41a (Biolegend, cat. no. 135910), biotin-conjugated Rat anti-mouse CD45 (BD Pharmingen, cat. no. 553077), biotin-conjugated rat anti-mouse Ter-119 (BD Pharmingen, cat. no. 559971). The biotin-conjugated antibodies were labeled using streptavidin-V450 (BD Biosciences, cat. no. 560797). Isotype control staining was performed using PE-conjugated anti-rat.
immunoglobulin G2b (IgG2b) antibody (Biolegend, cat. no. 400635), APC/Cy7-conjugated mouse IgG2b antibody (BD Pharmingen, cat. no. 558061) and biotin-conjugated rat IgG2ak (IgG2aK) antibody (BD Pharmingen, cat. no. 553928). The cells were stained at a final concentration of 1:200 for 30 minutes on ice followed by washes in Hanks’ balanced salt solution (Invitrogen, Carlsbad, CA, USA) containing 2% fetal bovine serum. For the live–dead assay, LIVE/DEAD® Fixable Dead Cell Stain Kit (Invitrogen, cat. no. L34955) was used according to the manufacturer’s protocol. FACS and data analysis were performed using the BD LSR II flow cytometer and FlowJo software (Tree Star, Inc., Ashland, OR, USA).

**MiR146b knockdown and overexpression in CDj cells and in PMECs**

To overexpress miR-146b, 40 nM of hsa-miR-146b-5p synthetic pre-miR precursors (Ambion, ReN0. AM17100) were transfected into PMECs, along with the Pre-miRTMmiRNA Precursor Negative Control #1 (Ambion ReN0. AM17110). To knockdown miR-146b, 40 nM of miRCURY LNA microRNA inhibitor (Exiqon, cat. no. MT01) miRNA target sequence expression clone in pEZX-MT01 vector with fLuc along with 40 nM of miR-146b precursor miRNA (Ambion, ReN0. AM17100) and miRNA control (Ambion ReN0. AM17110). The cells were transfected to a 96-well plate 18 hours after transfection and cultured for another 24 hours. Both firefly luciferase and Renilla luciferase activities were measured using GeneCopoeia Luc-Pair™3 miR Luciferase Assay kit (GeneCopoeia, cat. no. LPRF-M030) according to the manufacturer’s protocol, and data were recorded on a BioTek Microplate Reader using Gen5™ software. Firefly luciferase activity was normalized with Renilla luciferase activities in the same well.

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**Author contributions**

H.E. participated in the design of the study, carried out the transfections, animal surgeries, FACS sorting and analysis, western blots, imaging–RT-PCR and drafted the manuscript; Y.H. participated in animal hormonal treatment and surgeries, western blots and H&E staining; K.V. participated in imaging and manuscript figures; M.C. performed the miRNA microarrays; U.B. generated and validated the STAT3b-specific antibody. S.S., M.M., D.T.J., P.P., D.T. and L.K.C. participated in the conception and design of the study; F.B. conceived the study, and participated in its design and coordination. All authors read and approved the final manuscript.

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**Supplemental material available online at**


**References**


Fig. S1. (A) RT-PCR of miR-146b in CD-alveolar progenitor cell line at 24, 48 and 72 hours after transfection. Data are presented as the mean ± s.e.m. (n=3). (B) Representative histogram of flow cytometry analysis for cells stained with the LIVE/DEAD® Fixable Dead Cell Staining Kit. Live-dead analysis was done in the alveolar progenitor cell line at 48, 60 and 72 hours post-transfection. (C,D) Representative histograms of flow cytometry analysis for cells transfected with the GFP-labeled LNA inhibitors and their controls, showing the percentage of GFP positive cells in the CD-derived distinct progenitor cell lines (C) and in the PMECs (D). (E) Representative bar graphs of densitometry for STAT3α and STAT3β levels normalized to β-actin in CD-derived alveolar progenitor cells (n=4, *indicates P<0.05 compared to other cells). Data are presented as the mean ± s.e.m.