

The C/EBP β transcription factor regulates epithelial cell proliferation and differentiation in the mammary gland

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Studies of C/EBP β -deficient mice have demonstrated a pivotal role for this transcription factor in hematopoiesis, adipogenesis, and ovarian function. Here we show that C/EBP β is also essential for normal development and function of the mammary gland. Ductal morphogenesis in virgin C/EBP β -deficient mice was disrupted, with ducts displaying reduced growth and branching. To distinguish whether the effect of C/EBP β deficiency on mammary epithelium is indirect or cell autonomous, we performed ovarian and mammary gland transplants. Transplants of wild-type ovaries into mutant females partially restored ductal morphogenesis during puberty but failed to support mammopoiesis during pregnancy. At term, mutant mice harboring wild-type ovaries exhibited reduced alveolar proliferation and impaired epithelial cell differentiation, including a complete absence of milk protein expression. Mammary gland transplant experiments demonstrated that development of C/EBP β -deficient epithelium was defective within a wild-type stroma and host background. Cell proliferation during pregnancy was reduced and differentiation, as measured by the activity of milk protein genes, was inhibited. However, wild-type epithelium developed in a C/EBP β -deficient stroma. Thus, C/EBP β plays an essential, cell autonomous role in the proliferation and differentiation of mammary secretory epithelial cells and is required for the activation of milk protein genes.

[Key Words: Mammopoiesis; ductal morphogenesis; alveoli; milk proteins; gene deletion]

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The C/EBPs (CCAAT/enhancer binding proteins) are a family of transcription factors with diverse regulatory functions. C/EBP α , β , and δ are expressed in partially overlapping patterns in many organs (Cao et al. 1991; Williams et al. 1991), whereas C/EBP ϵ is restricted to cells of the myeloid lineage (Yamanaka et al. 1997b). C/EBP γ (Ig/EBP) is thought to have inhibitory functions (Cooper et al. 1995) and is expressed at low levels in most tissues and cell types (Roman et al. 1990). In cell culture experiments, C/EBP α and C/EBP β are potent regulators of cell proliferation and differentiation in hepatocytes, adipocytes and myeloid cells (Johnson and Williams 1994).

The creation of mice with targeted disruptions of *c/ebp* genes has led to a more detailed understanding of the biological functions of the C/EBP proteins. Mice lacking C/EBP α die shortly after birth because of reduced glycogen storage in the liver (Wang et al. 1995) and display impaired development of granulocytes (Zhang et

al. 1997). C/EBP ϵ null animals are viable but fail to produce mature neutrophils and eosinophils, and older mice develop myelodysplasia (Yamanaka et al. 1997a). C/EBP β -deficient mice exhibit multiple immune deficiencies (Screpanti et al. 1995; Tanaka et al. 1995), as well as defective differentiation of adipocytes (Tanaka et al. 1997). In addition, C/EBP β -null females are sterile because of impaired ovulation and the absence of corpora lutea resulting from a block during granulosa cell differentiation (Sterneck et al. 1997). Fertility in these mice can be restored by implanting wild-type ovaries. However, their offspring die shortly after birth and, despite apparently normal suckling behavior, the pups do not have milk in their stomachs (Sterneck et al. 1997). These findings suggested that the C/EBP β mutation may also affect mammary gland development or function.

Several *c/ebp* genes expressed in the mammary gland are developmentally regulated during pregnancy and lactation. C/EBP α and C/EBP β mRNAs are expressed in the mammary gland of virgin animals. C/EBP α levels do not change significantly during pregnancy and lactation, whereas C/EBP β mRNA levels are induced during preg-

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nancy, decline slightly at midlactation, and increase again within 48 hr of the onset of involution. C/EBP δ mRNA expression is generally low but increases dramatically at the onset of involution (Gigliotti and DeWille 1998). C/EBP δ expression also correlates with growth arrest of mammary epithelial cells in culture (O'Rourke et al. 1997).

Development of the mammary gland proceeds in distinct stages (Imagawa et al. 1994) and is controlled by an interplay of steroid and peptide hormones (Topper and Freeman 1980). Although a rudimentary anlage is formed in the embryo, extensive morphogenesis of the gland commences with puberty and the secretion of ovarian steroid hormones. This period is characterized by rapid elongation of the epithelial ducts, which penetrate the mammary fat pad. In mature virgin mice the fat pad is filled with evenly spaced primary ducts from which secondary ducts and smaller side branches emanate. Further mammary development is induced during pregnancy under the influence of prolactin and placental hormones. During this period, rapid cell proliferation takes place that leads to formation of secretory alveoli, terminal differentiation of secretory alveolar cells, and synthesis of large amounts of milk proteins. Expression of milk proteins is initiated in a specific temporal sequence (Robinson et al. 1995). WDNM1, a secreted protein whose function is not known, is produced first and its mRNA can be detected easily at 10 days of pregnancy. Two days later mRNA for β -casein becomes detectable, whereas the whey acidic protein (WAP) gene is expressed at detectable levels from day 15 onwards. These mRNAs thus serve as markers for the differentiation state of the epithelium (Robinson et al. 1995).

In the present study we have examined mammary gland development in C/EBP β -deficient mice, and show that glandular development is impaired in virgin, pregnant, and lactating animals. We have also used ovarian and mammary gland transplantation experiments to assess whether these defects originate from the mammary epithelium or stromal compartment, or from possible effects of altered ovarian hormonal signals. Both experimental approaches mimicked unique physiological situations and revealed that functional mammary development requires the presence of C/EBP β in the mammary epithelium itself. Our studies demonstrate that mammary epithelial cells require C/EBP β for proper proliferative and morphogenic responses during mammary gland development, and for their differentiation to secretory cells during pregnancy as demonstrated by the complete absence of milk protein expression in C/EBP β -deficient mammary glands.

Results

Expression of C/EBP β during normal mammary gland development

The steady-state mRNA levels corresponding to different C/EBP isoforms were measured at various stages of mammary gland development (Fig. 1). Expression of

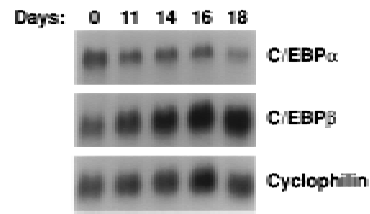


Figure 1. Expression profile of C/EBP RNA during mammary gland development. Northern blot analysis of total RNA from mammary glands of virgin and pregnant wild-type mice at various stages of gestation. The blot was hybridized sequentially with probes for the indicated genes.

C/EBP α was highest in the virgin state and was maintained at a slightly lower level throughout pregnancy. In contrast, expression of C/EBP β was lowest in the virgin and increased throughout pregnancy. Whereas C/EBP β expression has been described in several transformed or immortalized mammary epithelial cells in culture as well as in mammary epithelial tumors (Raught et al. 1995, 1996; O'Rourke et al. 1997), its role in adipose tissue development and function is also well characterized (Tanaka et al. 1997 and references therein). Therefore, the observed signal may be derived from both components of the gland. The cyclophilin signal served as a loading control.

Aberrant mammary gland development in C/EBP β -deficient virgin mice

Extensive development of the mammary gland commences at three weeks of age when mice reach puberty. In C/EBP β -deficient mice a delayed ductal outgrowth was observed as early as 3 weeks of age (Fig. 2). The extent of ductal outgrowth is best evaluated from the position of the terminal end buds in relationship to the mammary lymph node. The lymph node lies distal to the nipple from which the mammary ductal tree originates. In three-week-old wild-type and hemizygous mice the ducts had elongated close to the lymph node (Fig. 2A). In contrast, the mammary ducts were shorter in C/EBP β -deficient mice and had not reached the lymph node (Fig. 2B). The terminal end buds, which are club-shaped epithelial thickenings, are the sites of most rapid cell proliferation and ductal elongation at the distal ends of growing ducts. They have a characteristic architecture with central body cells surrounded by a distinctive layer of cap cells. In both control and C/EBP β -deficient mice the histology of the terminal end buds appeared normal (data not shown). In mature wild-type mice, extensively branched ducts occupy the mammary fat pad (Fig. 2G). In contrast, ductal morphogenesis was severely disturbed in the majority of mature mutant mice. The ducts were distended and had very few side branches. The degree of failure of normal ductal morphogenesis and branching varied among the eight animals that were examined between 2 and 6.5 months of age (Fig. 2C–E). Such variation in penetrance of the phenotype was also observed in the

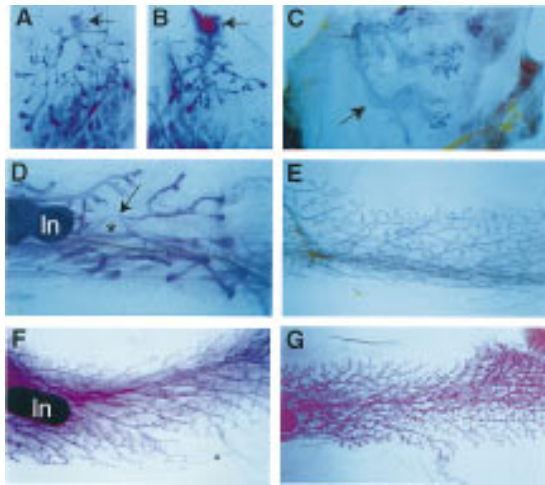


Figure 2. Whole mount analysis of ductal morphogenesis in virgin mice. A difference in growth was already detected in 3-week-old females. The ductal tree in wild-type mice (A) is larger than in C/EBP β -deficient mice (B). The arrow points to the nipple. (C) An example of extremely disturbed ductal development in a 2-month-old C/EBP β -deficient mouse. The primary ducts appear like large tubes and lack secondary branches (\rightarrow). (D) A 3.5-month-old C/EBP β -deficient mouse with very irregular ducts. Note the expanded balloon-like duct indicated by the asterisk. (E) A 5-month-old C/EBP β -deficient mouse with almost normal primary ducts that have sparse secondary branches. (F) A 4-month-old C/EBP β -deficient mouse with ovary transplant. Ovarian hormones induce formation of secondary side branches. (G) A 3-month-old wild-type mouse with fully developed ductal system. Magnification in A–D, 4 \times ; E–G, 13 \times . (In) Lymph node.

accumulation of fat and overall wasting of C/EBP β -deficient animals (Tanaka et al. 1997).

Mammary gland development in C/EBP β -deficient mice carrying wild-type ovaries

Since ovarian function is disrupted in C/EBP β -deficient mice (Sterneck et al. 1997), we asked whether the perturbation of mammary development could be rescued by wild-type ovary transplants. Ductal morphogenesis in four mature virgins, which received wild-type ovaries at the age of 3–4 weeks, was partially restored (Fig. 2F). The peripheral ducts had more side branches than the ducts in the center. This result demonstrates that the impaired development of the mammary gland in virgin C/EBP β -null mice is in part due to the lack of normal ovarian function, and confirms the importance of ovarian steroids in pubertal mammary gland development.

To evaluate whether C/EBP β only affects ductal morphogenesis or also controls alveolar development, we studied mammary structures in pregnant mice (Fig. 3). At day 18 postcoitum, wild-type glands had undergone extensive lobuloalveolar growth and were filled with enlarged alveoli that contain milk (Fig. 3B,D). In contrast, mammary gland development in C/EBP β -deficient mice

with wild-type ovaries (Fig. 3A,C) resembled that of wild-type animals at early pregnancy (not shown). Two distinct lesions were observed. Alveolar density in the fat pad was severely curbed and little or no alveolar differentiation on a histological level was observed (Fig. 3A). Specifically, secretory vesicles and fat globules were seen in the cytoplasm of wild-type secretory cells (Fig. 3D) but were absent in the null mice (Fig. 3C). These results demonstrate that alveolar development and differentiation of epithelial cells is severely impaired in pregnant C/EBP β -deficient mice with wild-type ovary transplants.

Development of C/EBP β -deficient mammary epithelium in wild-type hosts

To investigate whether the mammary gland phenotype was caused by impaired epithelial cells or indirect influences of systemic defects we performed mammary epithelial transplants. Surgical removal of the area between the nipple and the fat pad at three weeks of age leaves a fat pad free of the endogenous mammary epithelium. A small piece of mammary epithelium from another syngeneic animal can be implanted and will develop. After several weeks an entire epithelial tree will grow from the transplant and penetrate the host fat pad (DeOme et al. 1959). One fat pad per mouse received a small piece of mutant epithelium, whereas the contralateral fat pad hosted a control transplant from a wild-type or hemizygous animal. Thus, the transplanted epithelia are exposed to normal circulating hormone levels and wild-type stroma throughout puberty and pregnancy. Trans-

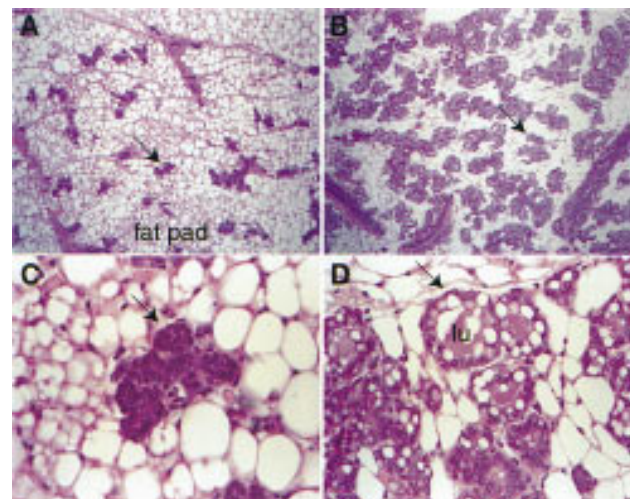


Figure 3. Alveolar development in C/EBP β -deficient mice carrying a wild-type ovary transplant. At day 18 of pregnancy the mammary gland of C/EBP β -deficient mice (A) is sparsely filled with underdeveloped alveoli (\rightarrow). No signs of secretory differentiation are visible at high magnification (C). The mammary gland of a wild-type mouse (B) is filled with differentiated alveoli (\rightarrow). At high magnification, the expanded lumen (lu) and fat globules are visible (D).

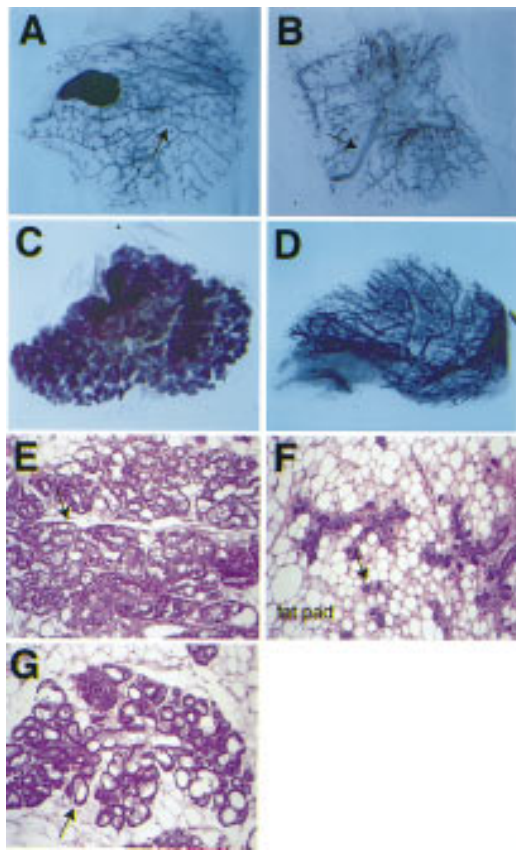


Figure 4. Development of *C/EBPβ*-deficient epithelium in wild-type stroma. Mammary epithelia from hemizygous (A,C) and *C/EBPβ*-deficient (B,D) animals were transplanted into fat pads of 3-week-old wild-type mice and harvested from virgin mice 8 weeks later (A,B) or after the mice had delivered a litter (C,D). Note the thick primary ducts (→) and irregular side branching in the *C/EBPβ*-deficient transplant in the virgin stage (B). At term, *C/EBPβ*-deficient epithelia (D) have underdeveloped small alveoli, whereas the heterozygous transplant (C) fills the stroma with fully developed alveoli. (E) Histological section of gland shown in C. (F) Histological section of gland shown in D. (G) Section through endogenous gland of host animal at term. Arrows point to alveoli.

plants were harvested from 11 mice after 4 or 8 weeks and whole mounts were prepared to evaluate the shape and extent of epithelial outgrowth. Of the seven *C/EBPβ*-deficient epithelia that were recovered from the wild-type fat pads, all but one, which looked normal, produced ducts that were thicker (Fig. 4B) than those produced by wild-type controls (Fig. 4A) and only three had filled the fat pad entirely. This result demonstrates that requirement for *C/EBPβ* in the virgin mammary gland is cell autonomous and epithelia from *C/EBPβ*-deficient mice maintain their abnormal growth characteristics even when they develop in a wild-type host under the influence of functional ovaries.

To analyze the development of the epithelial transplants in response to pregnancy hormones, 17 wild-type hosts were mated and their mammary glands were har-

vested for whole mounts and RNA isolation within 12 hr after the animals delivered a litter. Again, the developmental state of the *C/EBPβ*-deficient epithelium of term mice (Fig. 4D) resembled that of wild-type animals at early pregnancy (not shown). The *C/EBPβ*-deficient alveoli were underdeveloped as compared to control wild-type animals or transplanted wild-type epithelia (Fig. 4C). In histological sections the mutant alveoli appeared small, the lumina were not expanded, and the epithelial cells showed no signs of secretory activity (Fig. 4F), whereas fully differentiated cells were found in transplanted hemizygous epithelium (Fig. 4E) and the endogenous gland of the host (Fig. 4G). In summary, these data demonstrate that epithelia from *C/EBPβ*-deficient mice maintain their abnormal growth characteristics even when they are embedded in a wild-type fat pad and develop in a wild-type host under the influence of functional ovaries, suggesting that the *C/EBPβ* mutation perturbs epithelial cells directly.

Expression of milk protein genes in *C/EBPβ*-deficient epithelium

To characterize the state of epithelial cell differentiation and to begin to identify the deficiency of mutant epithelium at the molecular level, we analyzed the expression of several milk protein genes (Fig. 5). The β -casein and WAP genes are highly expressed in wild-type mice at day 18 of pregnancy or at day 3 of lactation. However, no

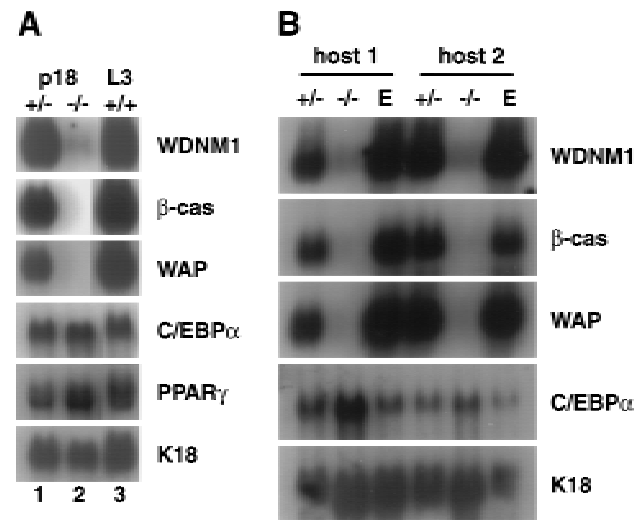


Figure 5. Gene expression in pregnant and lactating tissues. (A) Northern blot analysis of total RNA from mammary glands of hemizygous (lane 1) and *C/EBPβ* null mice with wild-type ovaries (lane 2) at day 18 of pregnancy. Lane 3 represents RNA from a wild-type mouse on day 3 of lactation. The blot was hybridized sequentially with probes for the indicated genes. (B) Northern blot analysis of total RNA derived from mammary glands of two wild-type host animals at term with epithelial transplants from a hemizygous donor (lane 1) or a *C/EBPβ* null donor (lane 2). Lane 3 represents RNA from an unoperated endogenous gland. The blot was analyzed as described in A.

expression of β -casein or WAP mRNA was detected in mammary tissue from mutant mice at day 18 of pregnancy (Fig. 5A). Expression of WDNM1 was barely detectable and at least 100-fold lower than in wild-type mice (Fig. 5A). As a control for the contribution of epithelium to the RNA samples we assessed the expression of the epithelial cell specific keratin18 gene (Singer et al. 1986) (Fig. 5A). Similar results were obtained when mammary glands from wild-type hosts with C/EBP β -deficient epithelial transplants were analyzed (Fig. 5B). Even at long exposures, no expression of β -casein and WAP was seen (not shown). Thus, the biochemically undifferentiated phenotype of mutant glands agrees with the morphologically immature appearance of C/EBP β -deficient alveoli.

We also analyzed whether the expression of C/EBP α was altered in the absence of C/EBP β . C/EBP α mRNA levels in wild-type mammary glands were significantly lower than C/EBP β and did not change significantly during pregnancy (Fig. 1; Gigliotti and DeWille 1998). The majority of the transcripts is likely derived from the adipose tissue (Tanaka et al. 1997). In the mammary glands of C/EBP β -deficient pregnant mice, as well as in glands of mutant epithelium in wild-type hosts, C/EBP α mRNA was readily detectable and comparable to the levels detected in control tissue. Similar expression levels for PPAR γ 2, a marker gene for adipocytes, were also found in wild-type and C/EBP β -deficient glands (Fig. 5A,B). Thus, the C/EBP β mutation does not affect the expression of the related C/EBP α gene or a stroma-specific gene.

Development of wild-type epithelium in C/EBP β -deficient stroma

C/EBP β is required for the differentiation of adipocytes (Tanaka et al. 1997) and it is well known that mammary epithelial development depends on interaction with appropriate mesenchymal signals (Robinson and Hennighausen 1997). Therefore, we asked whether the C/EBP β -deficient fat pad contributed to the mammary gland phenotype of mutant mice. To this aim, wild-type epithelia were transplanted into cleared fat pads of either wild-type or C/EBP β -deficient animals. The fat pads containing the epithelial implants were then transplanted into wild-type hosts to eliminate any genotype-specific systemic effects. The development of wild-type epithelium in three out of three C/EBP β -deficient fat pads harvested after 8 weeks was indistinguishable from that in wild-type fat pads (Fig. 6A,B). To evaluate the influence of the stroma on alveolar development, four hosts were mated and mammary tissues were analyzed immediately after parturition. Extensive lobuloalveolar development of wild-type epithelium took place in both wild-type and C/EBP β -deficient fat pads (Fig. 6C,D). From these observations we conclude that the absence of C/EBP β from the stromal compartment does not influence ductal elongation and branching during puberty or terminal differentiation of secretory epithelial cells.

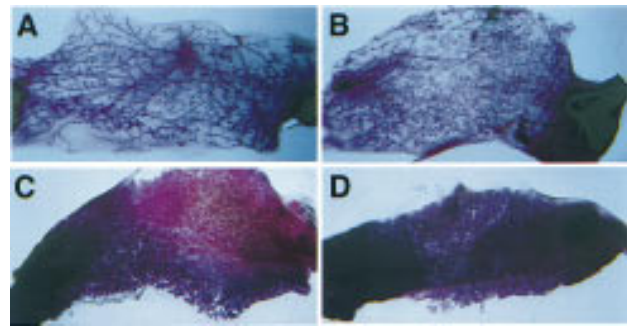


Figure 6. Development of wild-type epithelium in C/EBP β -deficient stroma. Development of transplanted wild-type epithelia at virgin stage (A,B) and at term (C,D) is the same in wild-type stroma (A,C) and in C/EBP β -deficient stroma (B,D).

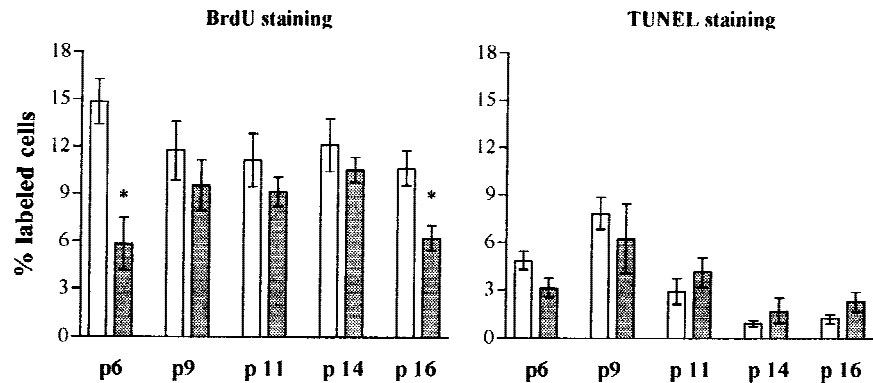
Cell proliferation is reduced in C/EBP β -deficient mammary epithelium

The impaired development of C/EBP β -deficient epithelial cells, specifically their reduced density in the mammary fat pad, could be the result of decreased proliferation rate and/or increased rate of apoptosis. To distinguish between these possibilities we injected wild-type mice carrying epithelial transplants with BrdU at different stages of pregnancy. BrdU incorporation as a measure of epithelial cell proliferation was determined at days 6, 9, 11, 14, and 16 of pregnancy (Fig. 7A). Cell proliferation was lower in C/EBP β -deficient epithelia throughout pregnancy. It was significantly reduced at day 6 of pregnancy to only 40% of the rate seen in wild-type mice. It remained lower until day 14 and was significantly reduced again on day 16. Adjacent sections were used for TUNEL analysis to determine the number of apoptotic cells. Though not statistically significant, the mammary epithelial apoptotic index in early pregnancy was lower in C/EBP β -deficient mice than in controls but increased from day 11 on (Fig. 7B). These results demonstrate that induction of increased epithelial cell proliferation in early pregnancy and proliferation at late pregnancy stages are impaired in the absence of C/EBP β .

Discussion

Mammary gland development during puberty, pregnancy, and lactation is regulated by the intricate interaction of the female endocrine system, the mammary gland epithelium, and stroma. The relevance of distinct genetic pathways for functional mammary gland development has been illustrated in mice with targeted mutations in genes from these pathways (for review, see Hennighausen and Robinson 1998). Using such mouse models it was possible to link the function of specific proteins to distinct stages of postnatal mammary development. Whereas some gene products, such as the prolactin receptor (PRLR) (Ormandy et al. 1997), the Stat5a transcription factor (Liu et al. 1997), or the LAR phosphatase (Schaapveld et al. 1997), control late stages of

Figure 7. Cell proliferation and apoptosis rates in *C/EBP β* -deficient mammary epithelium during pregnancy. At the indicated stages of pregnancy mice bearing contralateral *C/EBP β* -deficient and control mammary epithelial transplants were injected with BrdU and sacrificed 2 hr later. BrdU labeled nuclei were detected by immunostaining and apoptotic nuclei were stained by TUNEL. Between 6000 and 9000 nuclei were counted and the percentage of labeled cells was determined. The mean \pm standard deviations were calculated and compared by T test. (*) A *P* value of <0.001 . Wild-type (open bars) and *C/EBP β* -deficient (filled bars) transplants.



alveolar proliferation and differentiation during pregnancy, others, such as the progesterone receptor (PR) (Lydon et al. 1995; Humphreys et al. 1997) and the estrogen receptor- α (ER) (Boccanfuso and Korach 1997; Cunha et al. 1997), are required during puberty and early pregnancy. In addition to steroid and peptide receptors, cell cycle regulators and transcription factors control alveolar development. The absence of cyclin D1 (Fantl et al. 1995; Sicinski et al. 1995) and *a-myb* (Toscani et al. 1997) causes severe underdevelopment of alveoli at term that has histological resemblance to the defect in *C/EBP β* -deficient mice. *C/EBP β* has now emerged as a critical component in the control of both proliferation and early stages of differentiation. The requirement for *C/EBP β* is epithelial cell autonomous and in the absence of *C/EBP β* only limited proliferation of epithelial cells occurs and the differentiation program leading to a functional gland is not executed (Fig. 8). Thus, we propose that *C/EBP β* participates in hormonal signaling cascades whose endpoint is the lactating mammary gland.

Systemic vs. mammary intrinsic effects

The mammary gland defect observed in *C/EBP β* -deficient mice raised the possibility of two distinct mechanisms. In one, ductal morphogenesis depends on the presence of *C/EBP β* in mammary tissue. In the other, ductal development is dependent on secondary systemic signals that are disrupted by the lack of *C/EBP β* in endocrine tissues. Since *C/EBP β* -deficient mice display severely impaired reproductive functions (Sterneck et al. 1997), it was necessary to dissect the role(s) of *C/EBP β* within the mammary gland from those imposed indirectly through ovarian or other signals. Using ovarian transplants, with the aim of restoring physiological hormone levels, we could demonstrate that the defects in ductal morphogenesis observed in mature virgins are partly of secondary nature and caused by nonfunctional ovaries. Impaired ductal outgrowth and branching has also been observed in mice deficient in the estrogen (Boccanfuso and Korach 1997) or progesterone receptor (Lydon et al. 1995; Humphreys et al. 1997). This corroborates the ovarian component of the *C/EBP β* defect. Al-

though wild-type ovary transplants allow pregnancy to occur in otherwise sterile mutant females, the development of the mammary gland is still severely impaired with only limited growth and a complete absence of milk production. Thus, the inability of *C/EBP β* -deficient females to sustain their pups postpartum is primarily caused by mammary gland defects and not simply by impaired maternal behavior or development of the pups.

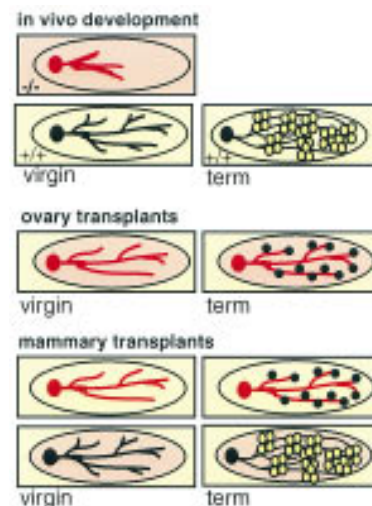


Figure 8. Summary of transplantation results. *C/EBP β* -deficient mammary glands display irregular ductal development in virgin glands. In contrast, in wild-type virgins the fat pads are interlaced with regular ducts with small side branches. At term the wild-type gland is filled with secretory alveoli that produce milk (open white circles). When *C/EBP β* -deficient mice receive an ovarian transplant ductal morphogenesis in virgins resembles more the wild-type phenotype. However, at term the *C/EBP β* -deficient glands are sparsely filled with small underdeveloped alveoli that are unable to produce milk (black full circles). Development of *C/EBP β* -deficient epithelia in wild-type fat pads is inhibited and cannot be rescued by the wild-type stroma. Wild-type epithelium is able to develop functional alveoli also in *C/EBP β* -deficient stroma. *C/EBP β* -deficient epithelium (red), *C/EBP β* -deficient stroma (pink), wild-type epithelium (black), wild-type stroma (yellow).

However, C/EBP β is also expressed in other organs, such as the adrenal gland (E. Sterneck, unpubl.), the placenta (Chen et al. 1995), and possibly the pituitary (Wegner et al. 1992), which are important for the development of the lactating mammary gland. Thus, we cannot rule out that other hormonal defects of the mutants are in part responsible for the impaired development of the mammary gland.

The development of the mammary gland depends on coordinated communication between epithelial and stromal cells (Daniel and Silberstein 1987). Because C/EBP β is critical for the development and differentiation of adipose tissue (Tanaka et al. 1997), it was important to address the contribution of the mutant stroma to the mammary gland phenotype. Our transplant experiments demonstrated that the mutant mammary gland fat pad can support normal development and differentiation of wild-type mammary epithelia (Fig. 8). Thus, we have ruled out the stromal component as a leading cause of deficient mammary gland development in C/EBP β mutant mice. All of these data establish that C/EBP β deficiency in the epithelial cell is the cause of the impaired mammary gland development in C/EBP β mutant mice. Transplant experiments established that functional C/EBP β is required in the mammary epithelial cell proper. In summary, our data show that C/EBP β deficiency in the epithelial cells is the primary cause of the perturbed mammary gland development.

Putative C/EBP β signaling pathways

Experimental evidence from a variety of cell culture systems has led to the notion that C/EBP β participates in cellular responses to many different extracellular signals (for review, see Johnson and Williams 1994). Mammary gland development is controlled by the systemic steroid hormones estrogen and progesterone in conjunction with systemic peptide hormones, including EGF and prolactin (Prl), and paracrine growth regulators (Topper and Freeman 1980; Hennighausen and Robinson 1998). Therefore, we suggest that C/EBP β is required for epithelial cells to recognize hormonal signals and/or to execute programs initiated by extracellular cues. C/EBP β may well be the mediator of any individual signaling pathway or a funnel for several pathways. In the future, cultures of epithelial cells and analyses of their responses to various signals may give insight into the specific molecular mechanisms underlying the mammary gland phenotype.

At this point, no genuine upstream signals or downstream targets of C/EBP β have been identified *in vivo*. However, a comparison of the morphological and biochemical characteristics of mammary tissue from C/EBP β -null mice with those of other engineered mouse mutants already sheds some light onto putative suspects. Disabling the Prl and JAK2/Stat5a pathway inactivates late stages of mammary gland development and terminal differentiation is not obtained (Liu et al. 1997; Ormandy et al. 1997). However, extensive alveolar structures partially fill the fat pad and the β -casein gene is

expressed at near normal levels (Liu et al. 1997). Similarly, inactivation of the inhibin β B gene inhibits late stages of functional epithelial differentiation, but the expression of β -casein is unaffected (Robinson and Hennighausen 1997). Thus, we suggest that C/EBP β is involved in stages prior to Prl or inhibin β B requirement and functions independently of the JAK2/Stat5a pathway. The absence of the receptors for estrogen and progesterone has a more profound consequence on mammary development. ER deficiency leads to severely impaired ductal development, and retarded alveolar development was observed in PR-null mice (Boccinfuso and Korach 1997; Cunha et al. 1997). It has been shown that C/EBP β can participate in estrogen signaling by mediating repression of the interleukin-6 promoter in breast tumor derived MCF-7 cells (Stein and Yang 1995; Galien et al. 1996). Thus, compromised estrogen responses may be in part responsible for the observed phenotype of C/EBP β -deficient mammary epithelial cells.

In C/EBP β mutant mice at term expression of the WAP, WDNM1, or β -casein genes was virtually undetectable, suggesting that they are targets for C/EBP β . We suggest that the β -casein gene is a direct target, and the WAP gene is controlled indirectly. The promoter of the β -casein gene, a major milk protein, contains four C/EBP binding sites that are required for basal and hormone-dependent activation of the promoter in tissue culture (Doppler et al. 1995; Raught et al. 1995). Our *in vivo* results corroborate the cell culture results. Based on a sequence analysis no C/EBP binding site has been identified in 4 kb of WAP gene promoter sequence, and it is possible that C/EBP β -deficiency imposes an early block on epithelial cell differentiation, which in turn could explain the lack of expression of WAP and WDNM1. However, because of the relaxed sequence specificity of C/EBP proteins, direct WAP and WDNM1 promoter analysis will be required to address the role of C/EBP proteins in the activation of these genes.

The true target genes that convey the transcriptional signal from C/EBP β are still elusive. We have demonstrated reduced proliferation potential of mammary epithelial cells as early as day 6 of gestation. The reduced number of epithelial cells in term mammary tissue results from an overall reduced cell proliferation, particularly in early pregnancy, the phase of most rapid growth, and slightly elevated apoptosis rates in the second half of pregnancy. This observation may reflect an inability of the C/EBP β -deficient epithelium to respond efficiently to signals mediated by the above-mentioned hormones. Many cell culture models have linked members of the C/EBP family to proliferation, growth arrest, or apoptosis (for review, see Johnson and Williams 1994). This notion raises the possibility that C/EBP β -deficiency imposes an early block on epithelial cell differentiation, such that the stage at which milk protein genes are expressed is never reached and the lack of milk production is only an indirect effect of the mutation. This question can be addressed in the future by the development of a mouse mutant with a targeted conditional deletion of the C/EBP β gene at different stages of pregnancy.

Materials and methods

Mice

The C/EBP β -deficient mice generated by homologous recombination have been described previously (Sterneck et al. 1997). Mice of mixed genetic strain background (129/Sv \times C57BL/6) or pure 129/Sv strain background were used for experiments. No strain-specific differences were noted. Therefore, the strain backgrounds are not further specified in Results. Homozygous mutants and control subjects were littermates derived from intercrossing hemizygous parents. The mice were housed and bred in a specific pathogen-free facility with a 12-hr light cycle starting at 7:00 a.m. and chow and water ad libitum. Procedures were conducted in compliance with the guidelines of the Animals Studies Committee of the National Cancer Institute and the National Institute of Diabetes, Digestive and Kidney Diseases. For pregnancy stages of mammary glands the mice were mated and inspected for the presence of vaginal plugs in the mornings. The day of the vaginal plug was counted as day 0 of pregnancy.

Whole mounts and histology

The first inguinal glands were dissected at the indicated times of development and spread on a glass slide. After fixation for 2–4 hr in Carnoy's fixative the tissues were hydrated and stained in carmine alum overnight as previously described (Robinson et al. 1995). They were dehydrated, cleared in xylene, and mounted. After pictures were taken on a stereomicroscope the samples were embedded in paraffin and sectioned at 5 μ m and stained with hematoxylin and eosin according to standard procedures.

Mammary transplants

Mammary epithelia were transplanted into cleared mammary fat pads as previously described (Robinson and Hennighausen 1997). For donors of mixed genetic background, the host animals were F₁ hybrid offspring of 129/Sv and C57BL/6 parents. For clearing of the endogenous mammary epithelium, the tissue between the nipple and the lymph node was excised from 3-week-old female mice and a piece of mammary tissue from a mature virgin donor was implanted into the center of the remaining fat pad. The transplanted mammary glands were harvested from virgin mice 4 or 8 weeks after the operation. To obtain transplanted tissues at term, the hosts were mated 8 weeks after they received the transplant and the mammary glands were harvested the morning after delivery. For transplantations of the entire fat pad 3-week-old mice were used as described before (Robinson and Hennighausen 1997). The endogenous inguinal fat pad of wild-type mice was removed. Fat pads of wild-type and C/EBP β -deficient mice with an epithelial transplant from a wild-type mouse were implanted on opposite sides and attached to the peritoneum by sutures. The transplants were harvested from virgin mice after 8 weeks or from term animals as described above.

Ovary transplants

Ovary transplants were performed as described (Sterneck et al. 1997). Briefly, 3- to 4-week-old donor and recipient littermates were anesthetized with avertin. The ovarian fat pads were moved outside through a small incision through the skin. The bursa was opened at one end and the ovary was teased out and placed into PBS at 37°C. The bursa was held open with tweezers and the replacement ovary was pushed into the bursa. The ovary, oviduct, and uterus were then placed back into the body

cavity, making sure that the ovary was fully inserted inside the bursa. The skin incisions were closed with 9-mm wound clips and the mice were given appropriate postoperative care. Two weeks postsurgery the mice were mated.

RNA isolation and Northern blots

RNA from the first inguinal glands was prepared by the acid guanidinium thiocyanate-phenol-chloroform method (Chomczynski and Sacchi 1987) or with RNA STAT-60 reagent (TELTEST "B," Friendswood, TX) according to the manufacturer's protocol. Twenty micrograms of total RNA was separated in 1.5% formaldehyde gels and blotted on GeneScreen membranes. Expression of milk protein RNA was analyzed using specific oligonucleotides as previously described (Robinson et al. 1996). Expression of C/EBP genes was analyzed with a DNA probe to the 3' UTR of the rat C/EBP β gene (Sterneck et al. 1997) and an 850-bp *Sma*I–*Hind*II fragment representing the 3' UTR of the mouse C/EBP α gene. The probe for keratin 18 was a 1.1-kb *Eco*RI cDNA fragment (Singer et al. 1986). PPAR γ 2 was detected by a 1.5-kb *Sac*I fragment spanning the coding region of the PPAR γ 2 cDNA [kindly provided by Jeffrey Gimble, Oklahoma University (Gimble et al. 1996)]. Oligonucleotides were hybridized at 55°C and cDNA probes were hybridized at 65°C.

Cell proliferation and cell death assays

Two hours before sacrifice, mice were injected with 20 μ l/g of body weight of cell labeling reagent (Amersham cell proliferation kit). Mammary glands were prepared and fixed in Tellyesniczky's solution for 4 hr. The tissues were embedded in paraffin. For detection of incorporated BrdU 5- μ m sections were processed according to the manufacturer's instructions (Amersham Life Science). On neighboring sections from the same samples, apoptotic cells were detected by TUNEL assay as described by Humphreys et al. (Humphreys et al. 1996). Between 6000 and 9000 cells were counted from six to eight different sections of a sample and the means were calculated.

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