

Impaired involution of mammary glands in the absence of milk fat globule EGF factor 8

Rikinari Hanayama*[†] and Shigekazu Nagata*^{†‡§}

*Department of Genetics, Osaka University Medical School, 2-2 Yamada-oka, Suita, Osaka 565-0871, Japan; [†]Laboratory of Genetics, Integrated Biology Laboratories, Graduate School of Frontier Biosciences, Osaka University, Osaka 565-0871, Japan; and [‡]Solution Oriented Research for Science and Technology, Japan Science and Technology Corporation, Osaka 565-0871, Japan

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During the involution of mammary glands, epithelial cells undergo apoptosis and are cleared for the next cycle of lactation. The clearance of apoptotic epithelial cells is mediated by neighboring epithelial cells and by macrophages that migrate into the mammary glands. Here, we report that milk fat globule EGF factor 8 (MFG-E8), a secreted glycoprotein that binds to apoptotic cells by recognizing phosphatidylserine, was expressed by epithelial cells and macrophages in mammary glands and was involved in engulfment of apoptotic cells. A deficiency of *MFG-E8* caused the accumulation of a large number of milk fat globules (MFGs) in the mammary ducts during involution, indicating that the excess MFGs were cleared by an *MFG-E8*-dependent mechanism. The *MFG-E8*^{-/-} mice developed mammary duct ectasia with periductal mastitis, and the redevelopment of the mammary gland for their second litter was impaired. These results demonstrate that *MFG-E8*-mediated phagocytosis of apoptotic epithelial cells and MFGs is important for efficient involution of mammary glands.

exosomes | mastitis | phosphatidylserine

Mammary glands are unique mammalian organs that are developed to nurse offspring. For each pregnancy, the glands undergo a cycle of development, lactation, and involution. In each cycle, mammalian epithelial cells proliferate, differentiate, and die. The involution of mammary glands is triggered when the suckling stimulus is lost, usually upon weaning, and milk accumulates in the glands. The involution takes ≈10 days in mice and can be divided into two phases (1–3). In the first phase, which is reversible and lasts 48 h after weaning, mammary epithelial cells lose their differentiated function. In the second phase, the basement membranes and extracellular matrix in the mammary glands are degraded by proteases, leading to destruction of the lobular-alveolar architecture of the mammary glands. As involution progresses, mammary epithelial cells are removed and adipocytes concomitantly reappear, and the lobular-alveolar structure is reorganized to approach that of virgin glands (4).

During the involution process, epithelial cells undergo apoptosis and are shed into the lumen (5). At the initial stage of involution, these apoptotic cells are thought to be engulfed mainly by neighboring epithelial cells; in the later stage, they seem to be cleared by macrophages that migrate into the gland (3, 6). Milk fat globules (MFGs) are minute globules carrying fat that are secreted from the epithelial cells into the lumen during lactation (7). Upon weaning, many MFGs remain in the lumen, and they seem to be engulfed by epithelial cells lining the lobules in the mammary gland (8). How these epithelial cells and macrophages recognize and engulf apoptotic cells and MFGs has not been well elucidated.

Many molecules expressed on the surface of apoptotic cells or phagocytes have been proposed as ligands and receptors for the engulfment of apoptotic cells (9–11). Mouse MFG EGF factor 8 (*MFG-E8*), called lactadherin in humans, is a 72-kDa glycoprotein secreted from mammary epithelial cells, macrophages, and immature dendritic cells (12–15). We recently showed that *MFG-E8* specifically binds to apoptotic cells by

recognizing phosphatidylserine (PS) exposed on the surface of the apoptotic cells and promotes the engulfment of apoptotic cells by phagocytes (12). *MFG-E8*-deficient mice are incapable of removing the apoptotic cells in the germinal centers of the secondary lymphoid tissues and develop lupus-like autoimmune diseases (16).

In this report, we show that the expression of *MFG-E8* was strongly up-regulated when mammary glands underwent involution. Primary epithelial cells prepared from involuting mammary glands expressed *MFG-E8*, and they engulfed apoptotic cells in an *MFG-E8*-dependent manner. Macrophages present in the mammary gland at the late stage of involution also expressed *MFG-E8* and engulfed apoptotic cells. The mammary glands of *MFG-E8*-deficient mice developed normally in the first pregnancy and could provide sufficient milk for pups. However, the involution of the *MFG-E8*^{-/-} glands was impaired, and a large quantity of MFG was left uncleared in the mammary ducts. These glands showed ectasia with periductal mastitis, and their redevelopment for the second litter was poor. These results indicate that *MFG-E8* plays an important role in removing apoptotic epithelial cells and MFGs during mammary gland involution and that this process is essential for the redevelopment of mammary glands.

Materials and Methods

Mice. *MFG-E8*^{-/-} mice (16) and *caspase-activated DNase* null mice (17) were described previously. The *MFG-E8*^{-/-} mice were backcrossed 10 times to C57BL/6 mice and used in this study, except in Fig. 4 and also in Fig. 5, which is published as supporting information on the PNAS web site, in which the littermates between *MFG-E8*^{+/-} and *MFG-E8*^{-/-} mice carrying the mixed background of C57BL/6 and 129/Sv were used. All mice were housed in a specific pathogen-free facility at Osaka University Medical School, and all animal experiments were carried out in accordance with protocols approved by the Osaka University Medical School Animal Care and Use Committee.

Induction of Involution. Pregnant mice delivered their young at 12–16 weeks of age, and their litter sizes were normalized to 8–10 pups. After full lactation was established (5 days of nursing), dams were separated from their pups to initiate involution. The glands were harvested at 0- to 10-day time points after forced weaning. For a second pregnancy, the dams were separated from pups 21 days after parturition and immediately mated for the next pregnancy. The pregnancy was confirmed by observing the vaginal plug, and the mothers who mated within 7 days after weaning were further analyzed.

Conflict of interest statement: No conflicts declared.

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Abbreviations: MFG, milk fat globule; *MFG-E8*, MFG EGF factor 8; PS, phosphatidylserine; H&E, hematoxylin/eosin.

[§]To whom correspondence should be addressed. E-mail: nagata@genetic.med.osaka-u.ac.jp.

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Histochemical Analysis. For hematoxylin/eosin (H&E) staining, mammary glands were fixed in 4% paraformaldehyde/4% sucrose in 0.1 M phosphate buffer (pH 7.2), embedded in paraffin, and sectioned at 4 μ m. For immunohistochemical analysis, glands were embedded in OCT compound and frozen in liquid nitrogen. Frozen sections (4 μ m) were fixed in cold acetone and incubated with an avidin/biotin blocking kit (Vector Laboratories), followed by incubation with 5% goat serum/1% BSA. The sections were stained in PBS containing 5% goat serum/1% BSA with biotinylated anti-CD68 (Serotec) or anti-CD45.2 (BD Biosciences), which was followed by staining with Alexa Fluor 488-conjugated (Molecular Probes) or Cy3-conjugated (Sigma) streptavidin. The sections were also stained with Cy3-conjugated anti-MFG-E8 (clone 18A2) (15). TUNEL staining was performed by using an Apoptag kit (Chemicon). After staining, the sections were mounted with FluorSave mounting reagent (Calbiochem) containing 1 μ g/ml DAPI (Dojindo Laboratories, Kumamoto, Japan) and observed by fluorescence microscopy (IX-70, Olympus, Melville, NY).

The whole-mount analysis of mammary glands was carried out according to the method of Thompson *et al.* (18) (<http://labs.amc.org/learn/nec>). In brief, glands were excised on day 10 of involution, dehydrated, stained with alum carmine, and stored in methyl salicylate (Sigma). The glands were then observed with a stereomicroscope (Stemi DV4, Zeiss) and photographed with a digital camera (DSC-F505V, Sony, Tokyo).

Cell Culture. Primary mammary epithelial cells were prepared according to the method described by Ehmann *et al.* (19), with some modifications. Briefly, mice at the age of 12 weeks were subjected to normal parturition and allowed to lactate for 5 days. Two days after removal of the pups, the mammary glands were excised, minced into small pieces, and treated at 37°C for 2 h with liver digest medium (Invitrogen). The suspension was poured through a 133-mm nylon mesh (NBC, Tokyo) to capture the epithelial clumps. The clumps were placed on a 10-cm culture dish in a 1:1 mixture of DMEM and F12 medium containing 10% FBS and 1% insulin-transferrin-selenium-X supplement (Invitrogen), 10 ng/ml TGF- α (Sigma), and 50 ng/ml basic FGF (Sigma) and cultured for 3 days. Cells were harvested with trypsin/EDTA, stained with phycoerythrin-conjugated anti-Mac-1 (BD Biosciences), and subjected to cell sorting (FACSARIA, BD Biosciences). Mac-1-negative cells grown on eight-well Lab-tek II chamber slides (Nalge Nunc) showed an epithelium-like morphology and produced β -casein. Mac-1-positive cells were cultured for 1 week in DMEM containing 10% FBS and macrophage colony-stimulating factor (20) and used as mammary gland macrophages.

In Vitro Phagocytosis Assay. Phagocytosis of apoptotic cells was assayed as described in ref. 12. In brief, thymocytes from 6-week-old *caspase-activated DNase* null mice (17) were treated with 10 μ M dexamethasone to induce apoptosis. Apoptotic thymocytes (1×10^6 cells) were added to 2.0×10^4 epithelial cells or macrophages that were cultured in eight-well Lab-tek II chamber slides, and phagocytosis was allowed to proceed for 1.5 h. The cells were fixed with 1% paraformaldehyde, subjected to TUNEL, and observed by light microscopy. The TUNEL-positive thymocytes were counted, and the number of TUNEL-positive apoptotic cells per phagocyte (phagocytosis index) in a total of 150 phagocytes was determined.

Analysis of Milk and MFGs. After parturition, dams at the age of 12 weeks for the first pregnancy or 20 weeks for the second pregnancy were allowed to lactate for 5 days, and the pups were removed. To analyze milk, the mammary glands were excised 12 h after weaning and homogenized by using a Polytron in 50 mM Tris-HCl buffer (pH 8.0) containing 150 mM NaCl, 1%

Nonidet P-40, 1 mM (*p*-amidinophenyl) methanesulfonyl fluoride hydrochloride, 1 μ g/ml leupeptin, and 1 μ g/ml pepstatin. The cell lysates were cleared by centrifugation at $30,000 \times g$ for 15 min, and aliquots (2 μ g of protein) were subjected to 10–20% gradient SDS/PAGE.

To analyze MFGs, the mammary glands were excised 2 days after weaning and minced in PBS. Suspension of the minced glands was centrifuged at $3,000 \times g$ for 15 min, and the floating MFGs were collected. After washing four times with PBS, MFGs were suspended in PBS, stained with biotinylated anti-MFG-E8 (clone 18A2) and then phycoerythrin (PE)-conjugated streptavidin (BD Biosciences), and analyzed by flow cytometry (FACSCalibur, BD Biosciences). MFGs were also stained with PE-conjugated annexin V (BD Biosciences) in 10 mM Hepes buffer (pH 7.4) containing 140 mM NaCl and 2.5 mM CaCl₂.

Results

Expression of MFG-E8 in Mammary Epithelial Cells During Involution. MFG-E8, which is expressed in macrophages for the engulfment of apoptotic cells (12), is expressed in mammary glands (21). To investigate the role of MFG-E8 in the development and involution of mammary glands, the expression of MFG-E8 in the gland at various developmental stages was examined by Northern hybridization. As shown in Fig. 1A, MFG-E8 was weakly expressed in the mammary glands of virgin animals and in early pregnancy. Its expression was up-regulated in late pregnancy and increased further soon after the glands underwent involution. Real-time PCR analysis with MFG-E8-specific primers (15) indicated that the expression levels of MFG-E8 mRNA in the mammary gland on day 5 of lactation and days 4, 7, and 10 of involution were 16.5, 30.5, 13.4, and 4.8 times higher, respectively, than the MFG-E8 mRNA level in virgin glands. Staining with an anti-MFG-E8 mAb showed that MFG-E8 was expressed by epithelial cells of the involuting mammary glands (Fig. 1B). There were no CD68-positive cells in the lactating mammary glands. However, they could be detected on day 2 of involution, and their number increased as involution progressed, up to 14.2% of the total cell number in the mammary glands on day 4 of involution. These CD68-positive cells expressed MFG-E8 (Fig. 1C) as well as other macrophage antigens such as Mac-1 and F4/80 (data not shown).

Epithelial cells prepared from mammary glands on day 2 of involution efficiently engulfed apoptotic cells (Fig. 1D), and the phagocytosis index (the number of engulfed apoptotic cells per epithelial cell) was 1.83 (Fig. 1E). The ability of *MFG-E8*^{-/-} epithelial cells to engulf apoptotic cells was significantly reduced, and their phagocytosis index was 0.63. Adding recombinant MFG-E8 (0.1 μ g/ml) to *MFG-E8*^{-/-} epithelial cells caused significant recovery, to 1.55 (Fig. 1E), indicating that the mammary epithelial cells could engulf apoptotic cells by means of an MFG-E8-dependent mechanism. Similarly, the CD68-positive macrophages from the involuting mammary glands efficiently engulfed apoptotic cells in an MFG-E8-dependent manner (Fig. 1E). The phagocytosis index of the macrophages was 2.1, which was similar to that of thioglycollate-elicited peritoneal macrophages (12).

Requirement for MFG-E8 in the Clearance of Apoptotic Cells and MFGs.

Apoptotic cells can be recognized *in situ* by TUNEL staining. When apoptotic cells are engulfed, the DNA of the dying cells is further digested and cannot be stained by TUNEL (22). Using this technique, we examined the effect of MFG-E8 on the clearance of apoptotic cells during involution of the mammary gland. As shown in Fig. 2A, $\approx 2.3\%$ of the epithelial cells in WT mammary glands were TUNEL-positive on day 2 of involution and increased to 6.3% on day 3 of involution. These TUNEL-positive cells could be stained with mAb against the activated caspase 3 (data not shown), confirming that these cells were

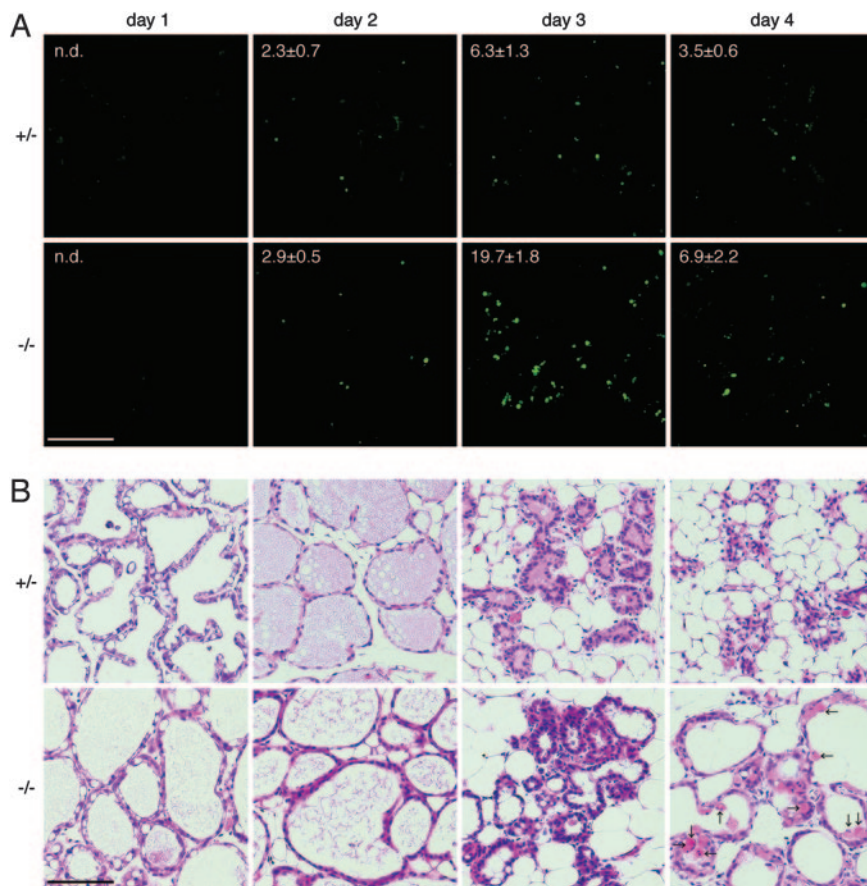


Fig. 2. Impaired clearance of apoptotic mammary epithelial cells in *MFG-E8*-deficient mice. (A) TUNEL staining. The mammary glands were prepared from *MFG-E8*^{+/-} or *MFG-E8*^{-/-} mice on the indicated days of involution and stained for TUNEL. The number of TUNEL-positive cells in the mammary glands on the indicated days of involution was determined and is shown as the percentage of DAPI-positive cells. At least 1,800 cells for each gland (300 cells per field) were analyzed. The analyses were performed with three mice for each genotype, and the average values are shown with SD. n.d., not detected. (Scale bar, 100 μ m.) (B) Staining with H&E. Sections of mammary glands from *MFG-E8*^{+/-} or *MFG-E8*^{-/-} mice at the indicated day of involution were stained with H&E. Amorphous materials, which could be debris of dead cells, are indicated by arrows. (Scale bar, 100 μ m.)

many cells with a plasma cell-like appearance invaded the mammary tissues of the *MFG-E8*^{-/-} dams.

The aberrant redevelopment of the mammary glands caused a decrease in milk production. As shown in Fig. 4C, extracts of mammary glands from *MFG-E8*^{-/-} mice at the first lactation contained the major milk proteins (α -, β -, and γ -caseins and whey acidic protein) at a level comparable to that of WT mammary glands. At the second lactation, WT mammary glands expressed a similar level of the major milk proteins, as at the first lactation. In contrast, the expression level of these proteins in *MFG-E8*^{-/-} glands was significantly reduced at the second lactation. *MFG-E8*^{-/-} dams could feed their first litter normally (Fig. 5), but the rate of weight gain of the second litter nursed by the *MFG-E8*^{-/-} dams was significantly reduced, and the average weight of the second litter at age 20 days was \approx 20% less than that of the second litter nursed by the WT dams. In addition, \approx 20–30% of the pups of the second and third litters of the *MFG-E8*^{-/-} dams died within 10 days of birth. When these pups were nursed by WT dams, they grew normally, indicating that the ateliosis of the pups nursed by *MFG-E8*^{-/-} dams was caused by the difficulty in lactation.

Discussion

Unlike most organs, the development and involution of mammary glands occurs postnatally, providing an excellent system for studying the molecular mechanisms of organ development. The

development of the mammary gland is hormonally regulated by pregnancy, during which the gland grows greatly and differentiates in preparation for secreting milk (25). Upon weaning, the entire alveolar epithelium involutes, accompanied by the apoptotic cell death of epithelial cells (5). In this process, the remaining MFGs are also cleared from the gland (8). The involution process is regulated at the transcriptional level and is associated with the up- or down-regulation of a set of genes (26, 27). In this report, we showed that the *MFG-E8* gene was strongly expressed in the mammary gland during the lactation and involution periods. The epithelial cells that comprise the gland expressed *MFG-E8*. In addition, macrophages that migrated into the mammary gland during the late stage of involution also expressed *MFG-E8*. It will be interesting to study whether similar transcription factors are involved in the *MFG-E8* gene expression in mammary epithelial cells and macrophages.

As with professional phagocytes of thioglycollate-elicited peritoneal macrophages and immature dendritic cells (12, 15), the engulfment of apoptotic cells by mammary epithelial cells, which are nonprofessional phagocytes, depended on *MFG-E8*. It is likely that *MFG-E8* secreted from epithelial cells binds to PS exposed on apoptotic cells and promotes their engulfment by means of integrins on epithelial cells (28). In addition, we found that epithelial cells cleared MFGs in an *MFG-E8*-dependent manner. In fact, uncleared MFGs were left in the *MFG-E8*^{-/-} mammary glands in longer periods than apoptotic cells. It is

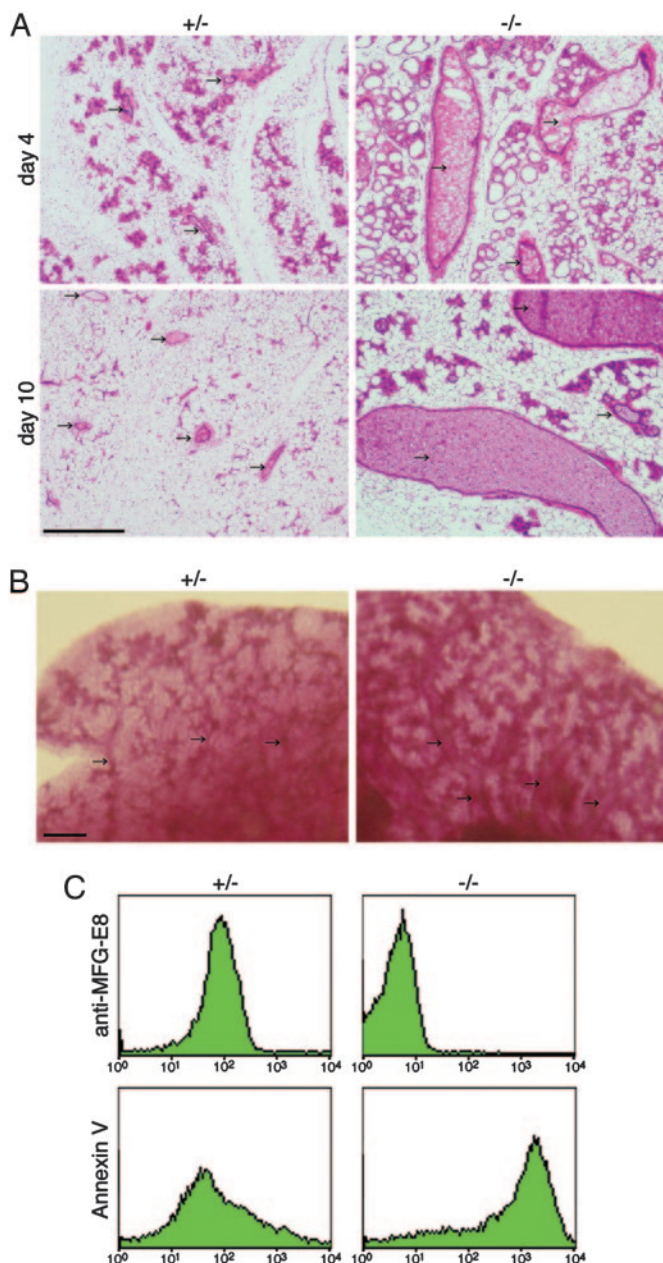


Fig. 3. Accumulation of MFGs in *MFG-E8*-deficient mice. (A) Accumulation of MFGs in *MFG-E8*^{-/-} mammary glands. *MFG-E8*^{+/-} or *MFG-E8*^{-/-} dams were allowed to nurse pups for 5 days and were separated from the pups. The mammary glands were excised 4 and 10 days after weaning. The paraffin sections of the mammary glands were stained with H&E. Mammary ducts are indicated by arrows. (Scale bar, 0.5 mm.) (B) Whole-mount staining of mammary glands. Mammary glands of *MFG-E8*^{+/-} or *MFG-E8*^{-/-} mice were excised 10 days after weaning and stained with alum carmine. Arrows indicate mammary ducts. (Scale bar, 1 mm.) (C) Staining of MFGs for MFG-E8 and PS. MFGs prepared from involuting mammary glands of *MFG-E8*^{+/-} or *MFG-E8*^{-/-} mice were stained with anti-MFG-E8 antibody (Upper) or annexin V (Lower).

likely that the unengulfed apoptotic cells undergo the secondary necrosis and that cellular contents are released. However, MFGs are rather stable and left in the mammary ducts.

MFGs are a type of exosome carrying fat and are secreted from mammary epithelial cells by budding (7). They are produced by the lactating epithelial cells that line the lumens of the mammary gland. In this process, fat droplets formed inside the cells move to the apical region, where they become enveloped in

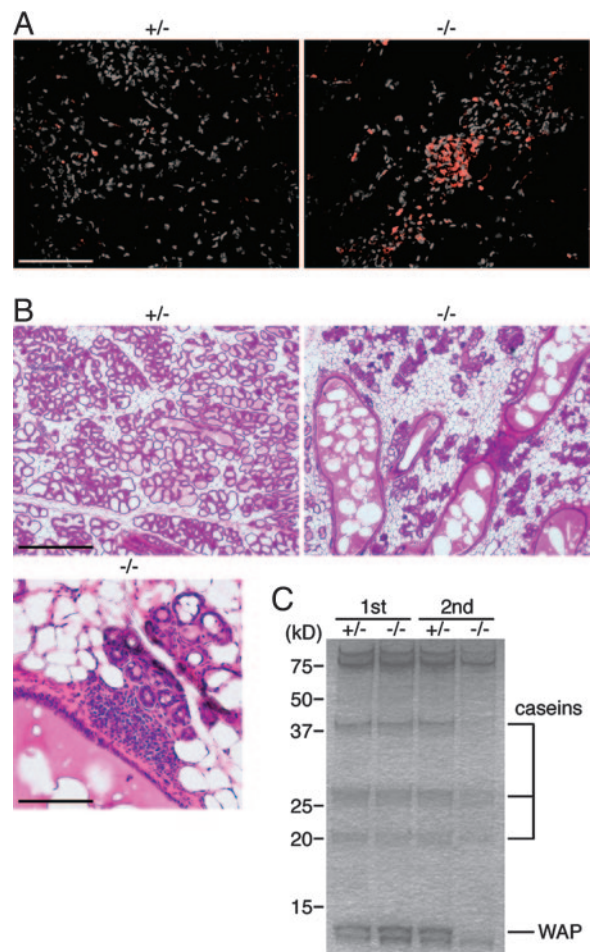


Fig. 4. Mastitis in mammary glands in *MFG-E8*-deficient mice and abnormal mammary gland redevelopment. (A) Inflammation in the *MFG-E8*^{-/-} mammary gland at late involution. *MFG-E8*^{+/-} or *MFG-E8*^{-/-} dams were allowed to lactate pups for 5 days and then forced to wean. At day 6 after weaning, sections of mammary glands were stained with anti-CD45 (red) and counterstained with DAPI (gray). (Scale bar, 0.1 mm.) (B) Abnormal redevelopment of the *MFG-E8*^{-/-} mammary gland. *MFG-E8*^{+/-} or *MFG-E8*^{-/-} mice underwent a second pregnancy and were allowed to nurse the pups. At day 5 of lactation, sections of the mammary glands were stained with H&E. In Lower, the sections from a *MFG-E8*^{-/-} mammary gland were enlarged. (Scale bars: Upper, 0.5 mm; Lower, 0.1 mm.) (C) Reduced synthesis of milk proteins in *MFG-E8*^{-/-} mice at the second lactation. *MFG-E8*^{+/-} and *MFG-E8*^{-/-} mice underwent one (1st) or two (2nd) cycles of pregnancy and lactation. On day 5 of the first or second lactation, the pups were removed. Twelve hours later, protein extracts were prepared from the mammary glands, and 2- μ g aliquots were separated by SDS/PAGE and stained with Coomassie brilliant blue. Molecular mass standard proteins were run in parallel, and their molecular masses are indicated at the left. The proteins seen in the mammary gland extracts were α -casein (43 kDa), β -casein (29 kDa), γ -casein (21 kDa), and whey acidic protein (WAP, 14 kDa). The total protein obtained from each gland was 7.0 and 7.2 mg for *MFG-E8*^{+/-} at the first and second lactation and 8.2 and 2.9 mg for *MFG-E8*^{-/-} at the first and second lactation, respectively.

plasma membrane and are expelled from the cell into the lumen (7). MFGs prepared from the milk of *MFG-E8*^{-/-} mice exposed PS on their surface. PS is usually localized to the inner surface of plasma membranes; this localization is maintained by a putative ATP-dependent translocase (29). We recently found that the nuclei expelled from erythroid precursor cells are deprived of ATP and expose PS on their surface (24). Similarly, MFGs carry few organelles such as mitochondria; therefore, ATP would be depleted in MFGs, thus inactivating the translocase and leading to the exposure of PS on the outer surface of

the globules. MFGs prepared from WT mice carried MFG-E8 on their surface, suggesting that as soon as MFG-E8 is secreted from mammary epithelial cells, it binds to MFGs. Thus, similar to apoptotic cells, MFG-E8-bound MFGs would be efficiently engulfed by epithelial cells, leading to the clearance of MFGs from the lumen during mammary gland involution. MFG-E8 was strongly expressed in the mammary gland also during the lactation period. The quantity of milk produced by a mother often exceeds that necessary for nursing babies. Thus, during the lactation period, the excess MFGs may be absorbed by epithelial cells in an MFG-E8-dependent manner, to be reused.

The involution of the mammary gland was severely impaired in *MFG-E8*^{-/-} mice. Even after the mice started to nurse the second litter, the lobular-alveolar structure was not properly developed in the *MFG-E8*^{-/-} mammary glands. This defect is partly because the mammary gland was largely occupied by the enlarged ducts that were filled with MFGs. Accordingly, when pups started to suckle, the number of MFGs left in the glands was reduced, and the mammary gland started to redevelop (unpublished data). Another possible reason for the poor development of the *MFG-E8*^{-/-} mammary gland is the mastitis that was observed in almost all *MFG-E8*^{-/-} mice soon after the first involution. Mastitis is often caused by bacterial infection, which occurs most frequently during early involution (30). However, our mice were maintained in specific pathogen-free conditions, and mastitis was never found in the WT mice, suggesting that the mastitis in the *MFG-E8*^{-/-} mice was induced without bacterial

infection. The involution of the mammary gland is regulated by the STAT3 transcription factor (31, 32) and is accompanied by the activation of various inflammatory genes such as CXCL1 and CXCL10 (27), which recruit inflammatory cells into the mammary gland. It is possible that the prolonged involution process in *MFG-E8*^{-/-} mice causes the extended activation of inflammatory genes, leading to severe inflammation. The engulfment of apoptotic cells by macrophages and epithelial cells is reported to inhibit inflammation by producing antiinflammatory factors such as TGF- β (33). Moreover, the dead cells that were not efficiently engulfed by phagocytes may undergo necrotic cell death, which is believed to induce inflammation (34). Thus, it is also possible that the dead epithelial cells that escaped from being engulfed may activate inflammation to induce the mastitis. Neutrophils and lymphocytes that were recruited in mastitis will produce cytotoxic substances, which could block the development of the mammary gland (35). In any case, our results indicate that if the mammary glands do not properly undergo involution, they will have problems in redeveloping for the second litter.

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