Metastatic tumor antigen 3 is a direct corepressor of the Wnt4 pathway

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Here we show that expression of MTA3 inhibits ductal branching in virgin and pregnant murine transgenic mammary glands. MTA3 also suppresses the Wnt4 pathway and, thus, these findings parallel phenotypic changes in Wnt4-null mice. MTA3 represses Wnt4 transcription and Wnt4 secretion, inhibiting Wnt-target genes in mammary epithelial cells. Accordingly, knockdown of endogenous MTA3 stimulates Wnt4 expression and Wnt cellular targets. The MTA3–NuRD (nucleosome remodeling and deacetylase) complex physically interacts with the Wnt4 chromatin in an HDAC-dependent manner, leading to suppression of the Wnt4 gene and Wnt4-dependent morphogenesis. These findings identify MTA3 as an upstream physiologic repressor of Wnt4 in mammary epithelial cells.

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Our understanding of MTA3 functions is primarily derived from the cancer cell line-based models. For example, MTA3 interacts with the promoter of the transcription repressor Snail and down-regulates its expression in a HDAC-dependent manner in cancer cell lines. Since Snail is known to promote epithelial-to-mesenchymal transitions primarily via down-regulating the expression of E-cadherin, MTA3-induced repression of Snail has been linked to the differentiation and increased expression of E-cadherin in human breast cancer cell lines (Fujita et al. 2003). The second known target of MTA3 was BCL6. MTA3 Mi2/NuRD complex deacylates BCL6, and regulates differentiation of B lymphocytic cell lines (Fujita et al. 2004). Although cell line-based findings have provided important insights into the biological activity of MTA3, its physiologic role in whole mammary gland development remains unknown.

The Wnt family of secreted proteins, which bind to specific cell surface receptors, influences mammary gland development and tumorigenesis. Wnt1 and Wnt3 were the first Wnt family members implicated in mouse mammary carcinomas (Nusse and Varmus 1992). Wnt4 is maximally expressed in the mammary epithelium during early to mid pregnancy (Bradbury et al. 1995). Mutant Wnt4-null mammary glands in early pregnancy showed a specific defect in progesterone-induced ductal side branching, implicating Wnt4 as a downstream effector of progesterone signaling (Brisken et al. 2000). In an attempt to achieve a greater clarity of MTA3’s biological functions in a physiological setting, here we investigated the effect of targeted overexpression of MTA3 in mouse mammary gland. We found that Wnt4 is a direct target of MTA3 and that phenotypic changes in the MTA3 transgenic mammary glands remarkably resemble those from the Wnt4-null mice.

Results and Discussion

MTA3 deregulation in mouse mammary epithelium leads to a defective ductal branching

To understand the influence of MTA3 on mammary gland development and morphogenesis, we generated a transgenic mice expressing human T7-MTA3 cDNA under the control of the mouse mammary tumor virus long terminal repeat [Supplementary Fig. S1A]. Three transgenic founder mouse lines with variable copy number of MTA3 transgene were generated [Supplementary Fig. S1B,C]. Transgene T7-MTA3 was expressed in a variety of organs with the highest expression in the mammary gland [Supplementary Fig. S1D]. Immunofluorescence of a mammary gland with an anti-T7 Ab further revealed a mosaic nature of MTA3 transgene expression in the nucleus of epithelial cells [Supplementary Fig. S1E]. Expression of T7-MTA3 in mammary glands was further confirmed by immunoprecipitation followed by Western blot analysis using an anti-T7 mAb [Supplementary Fig. S1F].

To determine the contribution of MTA3 to mammary gland development, we analyzed the status of ductal branching in the virgin glands from wild-type and MTA3-TG female mice. Glands were isolated and whole mounted, and compared at the developmental stage (4 wk) and mature stage (10–18 wk). At 4 wk of age, termi-
nal end buds were present and appeared normal in the transgenic mice (data not shown). At maturity, the ducts had reached nearly to the end of the fat pads; however, significantly less secondary and tertiary ductal branches were observed in the MTA3-TG mouse glands. These phenotypic changes were observed in mammary glands in three founder lines (Fig. 1A). The number of ductal side branches in wild-type branches (107.8 ± 13.7, n = 12) was higher than the number in transgenic branches (48.8 ± 7.1, n = 11) (Fig. 1B).

Ductal branch defect is due to impaired epithelial cell proliferation

To determine whether the defect in mammary gland development was the result of reduced cell proliferation or increased cell death, BrdU incorporation and TdT-mediated dUTP-X nick-end labeling (TUNEL) assays were performed. Wild-type and MTA3 virgin females (16-wk-old) were administered BrdU for its incorporation into DNA (Fig. 1C). The proliferation rate of the MTA3-TG mammary epithelium (4.8% ± 2.7, n = 6) was approximately half that of the wild-type mammary epithelium (10.2% ± 2.2, n = 5) (Fig. 1D). There were no significant differences in the apoptosis rates, measured by TUNEL assay, between wild-type and MTA3 mammary glands (data not shown). These results suggested that MTA3 status might be a determinant of mammary epithelium proliferation. Interestingly, there was no effect of MTA3-TG on the levels of Snail (a direct target of MTA3 in cancer cell lines) or E-cadherin in the mammary gland (Supplementary Fig. S2A,B). The noticed lack of any significant effect of MTA3-TG on the level of Snail expression is not in line with a previous report showing Snail repression in cell lines (Fujita et al. 2003). These differences may be possibly due to the lack of a normal epithelial cell line in the previous study or MTA3 represses Snail expression only in cancer cell line or both.

Defective Wnt4 pathway in mammary gland epithelial cells of MTA3-TG mice

Since progesterone receptors (PRs) and Wnt signaling are the major regulators of ductal branching in mammary glands (Robinson et al. 2000) and because PR and Wnt knockout females exhibit a defect in mammary gland development (Lydon et al. 1995) similar to that in the MTA3-TG females, we next determined whether deregulation of MTA3 affects PR or Wnt expression and function in mammary epithelium. We first examined PR expression in the wild-type and MTA3-TG virgin mammary glands. IHC analysis revealed that wild-type cells were 34.7% ± 12.7 positive for PR versus 31.2% ± 14.2 in the MTA3-TG cells. Similarly, levels of ERα-positive cells (47.2% ± 9.2 wild-type cells and 46.3% ± 8.3 TG cells were ER positive) showed no variation between the wild-type and TG virgin mammary glands, suggesting that the observed phenotypic changes in the MTA3-TG mice are not due to the change in the ER or PR status (Supplementary Fig. S3A,B).

Next we examined the levels of β-catenin, a key component of the canonical Wnt pathway. We found that β-catenin levels were much lower in the MTA3-TG mammary glands than in the wild-type glands (Fig. 2A), suggesting a potential role of MTA3 in repressing Wnt pathway. Accordingly, the levels of cyclin D1, one of the Wnt targets, were reduced to 14.6% ± 8.6, n = 5 in the MTA3-TG as compared with 35.4% ± 7.3 cyclin D1-positive cells in the wild-type mammary epithelial cells (Fig. 2B). Interestingly, we also discovered a considerable decrease in the in situ levels of Wnt4 as well as of WNT4 protein in the MTA3-TG mammary gland when compared with the wild-type mammary gland (Fig. 2C,D).
Secretion of Wnt4 ligand is suppressed by MTA3

To elucidate the role of MTA3 in the Wnt pathway in mammary gland epithelium, we next generated stable pooled clones of T7-MTA3 in the HC11 murine mammary epithelial cell line (Fig. 3A), which has been extensively used as a valid model to study the proliferative and differentiating effects of Wnt genes (Humphreys and Rosen 1997; Civenni et al. 2003). To assess the effect of MTA3-TG on the WNT4 pathway, we compared the levels of Wnt4 mRNA in the HC11/MTA3 cells with the levels in the control pcDNA clones, and found a drastic reduction in the level of WNT4 mRNA in the HC11/MTA3 cells [Fig. 3B]. The observed decreased WNT4 expression in the HC11/MTA3 cells was accompanied by reduced Wnt4 secretion in the conditioned media as compared with the control pcDNA cells (Fig. 3C). These findings suggested a negative regulatory role of MTA3 on Wnt4 expression and secretion in mammary epithelial cells.

MTA3 suppresses the canonical Wnt pathway

The drastic down-regulation of Wnt4 secretion in HC11/MTA3 cells is expected to inhibit the canonical Wnt pathway. Accordingly, we found a significant reduction in the level of β-catenin in the HC11/MTA3 clones as compared with the levels in the HC11/pcDNA clones (Fig. 3D). We also found lower levels of phosphorylated Ser9-GSK3β in the HC11/MTA3 clones than in the HC11/pcDNA clones [Fig. 3D] showing more active GSK3β, which leads to greater phosphorylation and degradation of β-catenin in HC11/MTA3 clones. To validate the repression of the canonical Wnt pathway by MTA3, we show that MTA3 suppresses the β-catenin/LEF signaling using a TOP-Flash-luc activity [Fig. 3E]. We also found a significant inhibition in the levels of WISP2 mRNA, a Wnt target gene, in the HC11/MTA3 clones [Fig. 3F].

MTA3 is a physiologic negative regulator of Wnt4 expression

To further strengthen the noticed causative correlation between the levels of MTA3 and repression of the Wnt4 pathway, we next silenced the endogenous MTA3 expression in the HC11/pcDNA cells by an MTA3-specific small interfering RNA (siRNA) and assessed the status of Wnt4 pathway. We found that a conditional reduction in the amount of MTA3 leads to an enhancement in the levels of Wnt4 and its target WISP2 mRNA (Fig. 3G,H). We next explored the potential involvement of the MTA3/NuRD complex in the repression of Wnt4 expression by MTA3. Treatment of the HC11/MTA3 cells with a general HDAC inhibitor TSA resulted in an increased Wnt4 expression [Fig. 3I] as well as accumulation of the secreted Wnt4 [Fig. 3J] in the HC11/MTA3 cells as compared with its levels in the control cells. These findings suggested that MTA3 represses Wnt4 expression in an HDAC-dependent manner in mammary epithelial cells, and that MTA3 status may be an upstream determinant of the Wnt4 expression, and consequently, functions in mammary epithelial cells.

MTA3/NuRD complex associates with two putative regions of the Wnt4 regulatory chromatin

The negative regulatory chromatin region of the mouse Wnt4 gene has been mapped up to 7 kb upstream of the start site (Devagan et al. 2005). To examine the physical interaction of the MTA3/NuRD complex with the Wnt4 chromatin, we carried out a promoter walk for the 10-kb region upstream of the Wnt4 gene using chromatin immunoprecipitation (ChIP) analysis. Initially, ChIP ly-

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**Figure 3.** MTA3 status affects Wnt4 expression and signaling. (A) Expression of T7-MTA3 protein in the indicated pooled HC11 clones. (B) Northern blot analysis of the Wnt4 mRNA. (C) Western blot showing Wnt4 protein in the culture media. (D) Western blot analysis of β-catenin and Ser9 phospho-GSK3β. (E) Top-Flash-Luc activity in the HC11 clones. (RU) Relative light unit. (F) RT-PCR analysis of the WISP2 mRNA. (G,H) Effect of MTA3-siRNA on the levels of Wnt4 and WISP2 mRNAs. (I,J) Effect of TSA on the levels of Wnt4 mRNA by RT-PCR. (I) Effect of TSA on secreted Wnt4 protein.
MTA3, HDAC2, and Mi2 binding to the 1-kb and 1.5-kb regions of the Wnt4 gene. ChIP assay showing differential recruitment of the MTA3-binding regions of the Wnt4 chromatin. (C) A double ChIP assay, first with T7 Ab followed by the HDAC2 Ab for the two regions of Wnt4 chromatin. (E) Status of the acetylated histone H3 in the MTA3–NuRD complex-bound Wnt4 chromatin with the region 6 (−2681 bp to −3234 bp) and 15 (−7688 bp to −8243 bp) regions of the Wnt4 chromatin in the HC11/MTA3 cells as compared with the corresponding Wnt4 regions in the HC11/pcDNA cells. Further, we performed a double ChIP assay in the HC11/MTA3 clones; the initial ChIP analysis was done using the anti-HDAC2 antibody. Results showed the simultaneous association of MTA3 and HDAC2 with the region 6 (−2681 bp to −3234 bp) and regions encompassing 15 and 16 (−7688 bp to −8801 bp) but not regions 7 and 17 (Fig. 4D), confirming that the MTA3/NuRD complex is associated with these two specific regions of the Wnt4 chromatin. Regions 3 and 4 of the Wnt4 chromatin, where no association of MTA3 was seen in ChIP assays were used as negative controls. To explore the potential decreased acetylation status of the histones associated with the MTA3-interacting Wnt4 regions, we performed ChIP using acetyl-Histone3 (H3)-specific antibody, and found a considerable decrease in levels of the acetylated H3 associated with the 1-kb (−2681 bp to −3793 bp) and with the 1.5-kb (−7688 bp to −9352 bp) regions of the Wnt4 chromatin. Further, HDAC1 and HDAC2 are integral components of the MTA3/NuRD complex and MTA3 represses WNT4 expression in an HDAC-dependent manner (Fig. 3I, J), we next tested whether HDAC2 is also recruited to the MTA3-interacting Wnt4 chromatin. ChIP analysis with an HDAC2-specific antibody showed that HDAC2 was present on the same 1.0-kb (−2681 bp to −3793 bp) and 1.5-kb (−7688 bp to −9352 bp) regions of the Wnt4 chromatin as MTA3, raising the possibility of recruitment of the MTA3/NuRD complex to the above regions of the Wnt4 chromatin (Fig. 4B). ChIP in HC11/MTA3 cells for Mi2, the largest subunit of the NuRD complex, showed the recruitment of Mi2 to both the 1-kb and the 1.5-kb regions of the Wnt4 chromatin (Fig. 4B, lower panel). In the 1-kb region, a greater association of Mi2 was observed with the region 6 (−2681 bp to −3234 bp), whereas a stronger association of Mi2 was seen to the regions 16 and 17 (−8802 bp to −9352 bp) within the 1.5-kb region. We next performed ChIP analysis of lysates from HC11/pcDNA and HC11/T7-MTA3 cells, and found a clear increased recruitment of T7-MTA3 to regions 6 (−2681 bp to −3234 bp) and 15 (−7688 bp to −8243 bp) of Wnt4 chromatin in HC11/T7-MTA3 cells over the albeit basal levels in HC11/pcDNA cells (Fig. 4C).

MTA3 suppresses mammary gland ductal side branching during early pregnancy

Wnt4 is essential for mammary gland development and side branching in early pregnancy [Brisken et al. 2000]. We examined the effect of MTA3-TG upon the ductal branching in mammary glands from pregnant day 2 female mice. Similar to the virgin mammary gland, we found reduced branching and BrDU incorporation in the early pregnancy [Supplementary Fig. S5], suggesting a direct role of MTA3 in repressing Wnt4 transcription.

Figure 4. MTA3/NuRD complex targets Wnt4 regulatory chromatin. (A) Schematic diagram of 10-kb regulatory chromatin of mouse Wnt4 gene. (B) ChIP analysis showed T7-MTA3, HDAC2, and Mi2 binding to the 1-kb and 1.5-kb regions of the Wnt4 chromatin. Regions 3 and 4 were the negative control. (C) ChIP assay showing differential recruitment of T7-MTA3 to regions 6 and 15 of Wnt4 chromatin in HC11/MTA3 cells (D) A double ChIP assay, first with T7 Ab followed by the HDAC2 Ab for the two regions of Wnt4 chromatin. (E) Status of the acetylated histone H3 in the MTA3–NuRD complex-bound Wnt4 chromatin in the HC11/pcDNA and HC11/MTA3 by ChIP analysis [left] and the relative acetyl-H3 levels at MTA3-binding regions of the Wnt4 chromatin [right].
pcDNA and HC11/MTA3 cells, MTA3 was found to be a potent repressor of progesterone-stimulated activity of PR (Supplementary Fig. S6C). Consistent with a potential suppression of the PR action, we found an inability of progesterone to stimulate Wnt4 expression in the HC11/MTA3 cells as compared with HC11/pcDNA cells [Supplementary Fig. S6D]. Next, we tested the ability of progesterone to modulate the level of MTA3, an inhibitor of Wnt4 gene expression [this study]. However, we found no effect of progesterone upon the level of MTA3 mRNA in the HC11 cells [Supplementary Fig. S6E]. Together, these results suggested that the noticed proliferation defect and hypobranching in the MTA3-TG mammary epithelium might be causally linked with the repressed Wnt4 pathway.

**Mammary gland differentiation and morphogenesis is hindered by MTA3 overexpression**

Wnt signaling is also required for differentiation processes during organogenesis [Moon et al. 2002]. Therefore, we investigated whether the inhibition of Wnt4 pathway in HC11/MTA3 cells would affect their ability to undergo differentiation. One of the physiological effects of the differentiation-stimulating signals in mammary epithelial cells is to accumulate neutral lipid droplets, which can be easily detected by oil red-O staining. We compared the extent of differentiation in the HC11/MTA3 clones with the extent in the control HC11/pcDNA clones in response to the lactogenic hormones dexamethasone, insulin, and prolactin [DIP]. We found that HC11/pcDNA cells underwent complete differentiation, whereas cellular differentiation in HC11/MTA3 clones was considerably suppressed [Fig. 5A].

Whole-mount analysis of the lactating mammary gland from the MTA3-TG mice revealed a profound suppression of mammary gland morphogenesis as compared with the wild-type mice [Fig. 5B]. Histologic study of the mammary gland on lactation day 2 by H&E staining showed a lower density of alveolar lobules in the MTA3-TG mice than in the wild-type mice, another indicator of a decreased differentiation [Fig. 5B]. The inhibitory effect of the MTA3-TG upon differentiation was further validated by a distinct reduction of levels of the phosphorylated Tyr694-STAT5a in the MTA3-TG mice compared with the wild-type mice [Fig. 5B]. Since phosphorylated STAT5a has been closely linked to mammary gland differentiation, whereas cellular differentiation in HC11/MTA3 clones was considerably suppressed [Fig. 5A].

The expression of MTA3 was examined its effect on lactation. To evaluate whether MTA3 could affect the level of MTA3, an inhibitor of Wnt4 gene expression [this study]. However, we found no effect of progesterone upon the level of MTA3 mRNA in the HC11 cells [Supplementary Fig. S6E]. Together, these results suggested that the noticed proliferation defect and hypobranching in the MTA3-TG mammary epithelium might be causally linked with the repressed Wnt4 pathway.

Next we examined the status of Wnt4 and its negative regulator MTA3 during mammary gland development in the wild-type mice. Western blot analysis showed an inverse correlation of the levels of Wnt4 and MTA3 protein in the virgin and pregnant mammary glands, and to some extent, in the lactating mammary glands [Fig. 5C]. The suppression of the mammary gland morphogenesis in the MTA3-TG mice indicated the possible hindrance of Wnt4 pathway. Moreover, in the lactating mammary glands (Fig. 5C). The expression of Wnt4 and its negative regulator MTA3 during mammary gland development is hindered by MTA3 overexpression.

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Since MTA3 transgene resulted in reduced secretion of Wnt4, we explored the effect of Wnt4 expression on β-ca-
sein gene in the HC11 cells. Transient overexpression of Wnt4 resulted in noticeable change in the β-casein luc activity [Supplementary Fig. S7A], whereas Wnt4 knockdown with Wnt4-specific siRNA showed a modest reduction in the β-casein luc activity upon DIP treatment [Supplementary Fig. S7B]. To further decipher the effect of Wnt4 on the differentiation of mammary cells, Wnt4 expression was silenced in the HC11 cells using Wnt4 siRNA and cells were stimulated with DIP for 48 h. Oil red-O staining showed a noticeable decrease in the accumulation of neutral oil droplets in cells with Wnt4 knockdown in comparison with the control cells (Supplementary Fig. S7C). The effect was not as pronounced as in the HC11/MTA3 clones [Fig. 5A], where it might be due to long-term depletion of Wnt4.

In addition to the β-casein gene, we also observed reduced steady-state mRNA levels of another marker of lactogenesis, whey acidic protein1 (WAP1) in mammary glands of four independent MTA3-TG mice in comparison with the levels of WAP1 mRNA in the wild-type mice [Fig. 5C]. The repressive effect of MTA3 overexpression on the β-casein gene and WAP1 suggested a potential lactation defect in the MTA3-TG mice. This was supported by the presence of milk in the belly of 6-d-old pup born to the wild-type female mice, whereas there was no or albeit milk in the belly of age-matched pups from the MTA3-TG female [Supplementary Fig. S8]. The body weight of the pups in two MTA3-TG founder lines, 5 and 10 d after birth, was consistently lower than was the weight of wild-type pups of the same age (Fig. 5H), presumably because of the reduced milk production in the MTA3-TG mice. These findings supported the notion that a reduction in the level of Wnt4 may be closely associated with a reduced differentiation, which along with reduction in milk proteins may contribute to the observed lactation defect in the MTA3-TG mice.

A principle finding of this study is an inhibitory effect of MTA3 transgene on the ductal side branching in mammary glands due to the suppression of Wnt4 signaling resulting in reduced cell proliferation and differentiation. MTA3 was a direct repressor of Wnt4 transcription, and this repression could not be rescued by progesterone signaling in HC11 mammary epithelial cells. It is of interest to note that the effects of MTA3 overexpression in the mammary gland development were opposite of those seen in the MTA1-TG mice. Overexpression of MTA1-TG resulted in an excessive branching and precocious development of the mammary gland due to a higher degree of proliferation and differentiation [Bagheri-Yarmand et al. 2004]. These findings also support the notion of physiologic differences between the MTA family members, and lack of functional redundancy between MTA3 and MTA1. In summary, we identified Wnt4 as a novel target of MTA3, making it the first direct corepressor of Wnt4 transcription in a physiologic setting.

Materials and methods

Generation of transgenic mice and genotyping

An MMTV-human MTA3-TG construct was created by subcloning T7-tagged MTA3 cDNA into MMTV–SV40–8AsK vector [Bagheri-Yarmand et al. 2004], and summarized in the Supplemental Material.

Whole-mount analysis and histology

The whole-mount analysis, and histology was performed as previously described [Bagheri-Yarmand et al. 2004], and summarized in the Supplemental Material.

In situ hybridization, RT–PCR, and ChIP

Digoxigenin-labeled in vitro transcripts were generated from plasmid containing mouse Wnt4 fragments [Briskin et al. 2000]. RT–PCR and ChIP were performed as summarized in the Supplemental Material.

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