Hedgehog signaling regulates FOXA2 in esophageal embryogenesis and Barrett’s metaplasia

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Hedgehog (Hh) signaling is an important signaling pathway in vertebrates, involved in embryonic development, tissue regeneration, and tumorigenesis. In the context of the esophagus, Hh signaling is reactivated in Barrett’s esophagus, a condition in which the squamous epithelium is replaced by columnar epithelium. This metaplasia is thought to predispose to esophageal adenocarcinoma, the most rapidly increasing cancer in the United States.

The objective of this study was to identify Hh target genes involved in Barrett’s metaplasia. Using microarray analysis, we found that the transcription factor FOXA2 is more highly expressed in murine embryonic esophagus compared with postnatal esophagus. Conditional activation of Shh in mouse esophageal epithelium induced FOXA2, while FOXA2 expression was reduced in Shh knockout embryos, establishing FOXA2 as an esophageal Hh target gene. Evaluation of patient samples revealed FOXA2 expression in Barrett’s metaplasia, dysplasia, and adenocarcinoma but not in esophageal squamous epithelium or squamous cell carcinoma. In esophageal squamous cell lines, Hh signaling upregulated FOXA2, which induced expression of MUC2, an intestinal mucin found in Barrett’s esophagus, and the MUC2-processing protein AGR2. Together, these data indicate that Hh signaling induces expression of genes that determine an intestinal phenotype in esophageal squamous epithelial cells and may contribute to the development of Barrett’s metaplasia.
shown that reactivation of epithelial Hh signaling in the mouse esophagus might result in columnar metaplasia (19), but to establish a potential role for Hh signaling in the pathogenesis of human Barrett’s esophagus, it is important to demonstrate that this signaling can produce cells with intestinal features.

The objective of our current study was to identify additional Hh pathway target genes that participate in the phenotypic specification of epithelial cells during normal esophageal development and to determine whether those genes might contribute to the formation of Barrett’s specialized intestinal metaplasia. By microarray analysis of the developing mouse esophagus, we found that Foxa2 (also known as Hnf3b) is more highly expressed in the embryonic esophagus than in the postnatal esophagus. Foxa2 is known to be a Sonic hedgehog (SHH) target gene in the central nervous system and a required foregut endodermal transcription factor (27–29). We demonstrate that FOXA2 is a target of Hh signaling in the embryonic mouse esophageal epithelium as well as in human Barrett’s epithelial cells and esophageal squamous epithelial cells derived from patients with Barrett’s esophagus. Moreover, we found that overexpression of FOXA2 in human esophageal squamous epithelial cells induces expression of MUC2 and AGR2 (an endoplasmic reticulum protein involved in MUC2 processing) (25, 30–33). These studies suggest that the Hh target gene FOXA2 is involved in the pathogenesis of the specialized intestinal type of Barrett’s metaplasia.

Results

FOXA2 is expressed in columnar epithelial cells of the embryonic esophagus and the adult intestine. During normal development, the mouse embryonic esophagus initially is lined by a columnar epithelium that expresses cytokeratin 8 (CK8). Later, this differentiates into a stratified squamous epithelium that expresses CK14 phenotype through decreased expression of squamous cell transcription factors, such as p63 (10, 11) or SOX2 (12, 13), or through increased expression of columnar cell transcription factors, such as CDX1 (14, 15), CDX2 (16, 17), SOX9 (18, 19), or MATH1 (20). Indeed, decreased expression of squamous transcription factors and increased expression of columnar transcription factors have been reported in biopsy specimens of metaplastic epithelia from patients with Barrett’s esophagus (19, 21–23).

Early in embryonic life, when the esophagus is lined by columnar epithelium, esophageal Hh signaling is active. Later, as the embryonic columnar lining differentiates into stratified squamous epithelium, Hh signaling is extinguished. In earlier studies, we showed that the surgical induction of gastroesophageal reflux in the mouse could reactivate esophageal Hh signaling (19). We also demonstrated that the Hh signaling pathway is reactivated in Barrett’s metaplasia, and we found evidence of epithelial-mesenchymal Hh signaling (with expression of Hh ligands by epithelial cells and BMP4 expression by mesenchymal cells) in Barrett’s tissue specimens but not in specimens of squamous-lined esophagus (19). We further showed that BMP4 could stimulate esophageal squamous cells to produce the transcription factor SOX9, which induces the squamous cells to express columnar cytokeratins (19).

Several types of columnar epithelia have been described in Barrett’s esophagus, including an atrophic gastric fundic-type epithelium, a junctional-type epithelium (also called cardiac epithelium), and an intestinal-type epithelium (also called specialized intestinal metaplasia) that has goblet cells and expresses the intestinal-type mucin MUC2 (24, 25). Only specialized intestinal metaplasia clearly predisposes to adenocarcinoma, which is why American gastroenterology societies require the demonstration of specialized intestinal metaplasia in esophageal biopsy specimens as a diagnostic criterion for Barrett’s esophagus (26). We have shown that reactivation of epithelial Hh signaling in the mouse esophagus might result in columnar metaplasia (19), but to establish a potential role for Hh signaling in the pathogenesis of human Barrett’s esophagus, it is important to demonstrate that this signaling can produce cells with intestinal features.

The objective of our current study was to identify additional Hh pathway target genes that participate in the phenotypic specification of epithelial cells during normal esophageal development and to determine whether those genes might contribute to the formation of Barrett’s specialized intestinal metaplasia. By microarray analysis of the developing mouse esophagus, we found that Foxa2 (also known as Hnf3b) is more highly expressed in the embryonic esophagus than in the postnatal esophagus. Foxa2 is known to be a Sonic hedgehog (SHH) target gene in the central nervous system and a required foregut endodermal transcription factor (27–29). We demonstrate that FOXA2 is a target of Hh signaling in the embryonic mouse esophageal epithelium as well as in human Barrett’s epithelial cells and esophageal squamous epithelial cells derived from patients with Barrett’s esophagus. Moreover, we found that overexpression of FOXA2 in human esophageal squamous epithelial cells induces expression of MUC2 and AGR2 (an endoplasmic reticulum protein involved in MUC2 processing) (25, 30–33). These studies suggest that the Hh target gene FOXA2 is involved in the pathogenesis of the specialized intestinal type of Barrett’s metaplasia.

Results

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(Figure 1A and ref. 34). Realizing that this pattern of epithelial replacement is the opposite of what occurs during the formation of columnar metaplasia in Barrett’s esophagus (Figure 1A), we began our studies by isolating whole C57BL/6 mouse esophagi from embryonic day 15.5 (E15.5) embryos and 2-day-old (P2) pups. These time points were chosen because the columnar-lined esophagus showed the earliest signs of squamous differentiation, with initial expression of CK14 in basal cells between E15.5 and E16.5, and because in the P2 esophagus the cytokeratin expression profile was markedly more squamous (Figure 1B).

Total RNA from separately pooled E15.5 and P2 esophagi was reverse transcribed and hybridized to an Agilent Whole Mouse Genome 44k microarray (Figure 2A). Overall, we found a pattern of differentially regulated genes that was consistent with high Hh activity in the columnar esophageal epithelium at E15.5 and with low Hh activity in the stratified squamous epithelium at P2 (Table 1). For example, we found that Hh pathway components such as Ptc, Hh interacting protein (Hhip), and Gli2 were more highly expressed in the E15.5 esophagi, while the P2 esophagi exhibited higher expression of squamous cytokeratins (Ck4, Ck6, Ck10, Ck13, and Ck14). We also found that Sox9, a gene we implicated previously in the development of Barrett’s esophagus (19), was more highly expressed in the E15.5 esophagi. Although we identified a number of candidate columnar genes, we chose to focus on the transcription factor Foxa2, which was expressed at levels 2.14 times higher in the E15.5 esophagi than in the P2 esophagi. We chose Foxa2 because it is a putative SHH target gene in the developing neural tube, it is required for the proper differentiation of endoderm, and it can induce expression of the goblet cell mucin MUC2 (which is found in Barrett’s specialized intestinal metaplasia) in intestinal epithelium (35, 36). Quantitative real-time PCR (qRT-PCR) was used to validate the microarray results for expression of Hh pathway genes and Foxa2 (Figure 2B).

To confirm a previous report of FOXA2 expression in the E15.5 mouse esophagus (37), we next sought to determine FOXA2’s temporal expression pattern during mouse embryonic esophageal development. We found that nuclear FOXA2 was expressed as early as E11.5 in mouse esophageal epithelium (data not shown) and continued to be expressed in a subset of esophageal epithelial cells between E14.5 and E17.5 (Figure 2C and data not shown for E17.5), correlating with esophageal expression of cytoplasmic

![Figure 2](https://example.com/figure2.png)

**Figure 2. Microarray analysis identifies Foxa2.** (A) MvA plot of E15.5 esophagi versus P2 esophagi microarray experiment. (B) Validation of microarray results with qRT-PCR for Hh pathway/target genes and Foxa2. **P < 0.01 as compared with E15.5. (C) FOXA2 immunohistochemistry of E14.5 to E16.5 esophagus. Scale bars: 50 μm.

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Note that positive fold change values indicate higher expression in E15.5, while negative fold change values indicate higher expression in P2.
CK8 (Figure 1B), but disappeared by P2 (Figure 2C) when the esophageal epithelium was distinctively squamous (Figure 1B). Neither FOXA2 nor CK8 was expressed in adult mouse esophageal epithelium, but both were expressed in the adult mouse small intestine (Supplemental Figure 1, A–D; supplemental material available online with this article; doi:10.1172/JCI66603DS1). We next performed FOXA2 immunohistochemistry on adult human gut tissues and found FOXA2 expression in columnar epithelia of the stomach, small intestine, and colon but not in squamous esophageal epithelia (Supplemental Figure 1, E–H). These findings suggest that FOXA2 expression plays a role in columnar differentiation in both mouse and human gastrointestinal tract epithelia.

FOXA2 is induced by Hh signaling in mouse esophageal epithelial cells. Since proper development of the mouse esophagus requires Hh signaling (38, 39) and since Foxa2 is a Hh target gene during neural tube formation (27), we postulated that FOXA2 expression in the mouse esophageal epithelium might be dependent on SHH. We previously developed an in vivo transplant culture system, which recapitulates the 3D structure of the mouse esophagus, with stratified squamous epithelium overlying fibroblasts (19). Cultures made with esophageal epithelia from conditionally activated Shh transgenic (ShhTg) mice expressed epithelial Shh and exhibited columnar phenotypic features, including nuclei oriented perpendicular to overlying stratified squamous epithelium (19). These columnar features were not seen in similar cultures made from wild-type esophageal epithelia. FOXA2 immunohistochemistry showed that in vivo transplant cultures made from conditionally activated ShhTg epithelial cells expressed nuclear FOXA2 (Figure 3B), while cultures made from wild-type esophageal epithelial cells did not (Figure 3A).

**Figure 3.** **Foxa2 is an esophageal Hh target gene.** FOXA2 expression is upregulated in ShhTg mouse esophageal epithelium. FOXA2 immunohistochemistry in 3D in vivo transplant cultures made with (A) wild-type or (B) ShhTg esophageal epithelial cells. FOXA2 expression is downregulated in Shh−/− embryos. FOXA2 immunohistochemistry in E14.5 (C) esophagus and trachea and (E) distal foregut and E16.5 (D) proximal foregut and (F) distal foregut. (G and H) Higher magnification images of sections shown in E and F, respectively. Gastrointestinal (G) and respiratory (R) epithelium, cartilage (C), and cystic lung (L) are indicated. Scale bars: 50 μm.

FOXA2 induction by Hh signaling is independent of Sox9. Since FOXA2 is regulated by Sox9 in the pancreas (40) and our ShhTg in vivo transplant cultures are induced to express epithelial SOX9, we next sought to determine whether FOXA2 expression in mouse esophageal epithelium was regulated by SOX9. Since global knockout of Sox9 (Sox9−/−) in mice leads to embryonic lethality at E11.5 (41), we generated mice with esophageal-specific deletion of Sox9 by crossing mice containing Cre recombinase knocked into the endogenous Shh locus (Shh-Cre−/−) mice (42) with mice homozygous for a conditional floxed allele of Sox9 (Sox9fl/fl) mice (ref. 43 and Supplemental Figure 2A). These Shh-Cre−/−Sox9fl/fl mice died perinatally. FOXA2 immunohistochemistry revealed similar FOXA2 expression in the esophageal epithelia of E17.5 Shh-Cre−/−Sox9fl/fl embryos and wild-type littermate embryos (Supplemental Figure 2B). This establishes that SOX9 is not required for FOXA2 expression in esophageal epithelium.

Differential regulation of FOXA2 by Hh signaling occurs in respiratory and gastrointestinal epithelia of the developing foregut. Shh knockout (Shh−/−) embryos were then generated as previously described (44) or by crossing Shh-Cre−/− mice to Shh-Cre−/− mice. In contrast to Shh-Cre−/−Sox9fl/fl embryos, which express FOXA2 in the absence of SOX9, esophageal FOXA2 expression was lost in Shh−/− embryos (Figure 3, C–H). Shh−/− embryos are known to develop foregut abnormalities in which the trachea and esophagus do not separate properly, and the combined trachea/esophagus structure ends distally in both lungs and stomach (38). We found that, in E14.5 Shh−/− embryos, the epithelium lining the trachea expressed FOXA2, but the epithelium lining the esophagus did not (Figure 3C). More distally, we found a distinct boundary in FOXA2 expression between respiratory and gastrointestinal epithelia.

The respiratory epithelium lining a cystic lung expressed FOXA2, while stratified squamous epithelium of the esophagus and forestomach did not (Figure 3, E and G). Further analysis of the E14.5 Shh−/− gastrointestinal epithelium revealed spotty expression of squamous CK14 in basal cells of the esophagus and forestomach (Figure 4, C and D) 1 day earlier than in wild-type gastrointestinal epithelium (Figure 1B and Figure 4, A and B). This suggests that
esophageal development to human disease, we next examined FOXA2 expression in human esophageal tissue specimens. We performed FOXA2 immunohistochemistry on tissue microarrays of esophagectomy specimens from 37 patients who had Barrett’s esophagus with or without associated esophageal adenocarcinoma. Among the 37 esophagectomy specimens, we obtained samples of nondysplastic Barrett’s metaplasia from 15 patients, Barrett’s metaplasia with low-grade dysplasia from 7 patients, Barrett’s metaplasia with high-grade dysplasia from 14 patients, and esophageal adenocarcinoma from 27 patients. Nuclear FOXA2 expression was not present in any normal esophageal squamous epithelium but was seen in 80% of patients with nondysplastic Barrett’s metaplasia, 86% of patients with low-grade dysplasia, 93% of patients with high-grade dysplasia, and 78% of patients with esophageal adenocarcinoma (Figure 5, A–C, and E). We then performed qRT-PCR for FOXA2 on frozen tissues from 6 esophagectomy cases. Consistent with our immunohistochemical results, we found considerable expression of FOXA2 in the Barrett’s dysplasia of all 6 cases, but the accompanying squamous epithelium exhibited no FOXA2 expression in all but 1 case, which showed minimal expression (Figure 5D). We next performed FOXA2 immunohistochemistry on a tissue microarray containing 11 cases of esophageal squamous cell carcinoma and did not find FOXA2 expression in any patient (Figure 5F). Taken

Shh loss leads to premature squamous epithelial differentiation of the dorsal foregut. At E16.5, the ventral portion of the Shh−/− combined foregut structure lined by respiratory epithelium continued to express nuclear FOXA2, while the dorsal gastrointestinal epithelium did not (Figure 3, D, F, and H). These findings are consistent with previous reports that loss of Shh expression in the lungs does not affect FOXA2 expression by respiratory epithelium (38, 45). Moreover, the differential regulation of FOXA2 by Hh signaling in the ventral and dorsal foregut suggests that there is an early developmental commitment to either a respiratory or a gastrointestinal epithelial cell fate.

**FOXA2 is expressed in Barrett’s esophagus and esophageal adenocarcinoma.** To explore the relevance of our findings in mouse esophageal development to human disease, we next examined FOXA2 expression in human esophageal tissue specimens. We performed FOXA2 immunohistochemistry on tissue microarrays of esophagectomy specimens from 37 patients who had Barrett’s esophagus with or without associated esophageal adenocarcinoma. Among the 37 esophagectomy specimens, we obtained samples of nondysplastic Barrett’s metaplasia from 15 patients, Barrett’s metaplasia with low-grade dysplasia from 7 patients, Barrett’s metaplasia with high-grade dysplasia from 14 patients, and esophageal adenocarcinoma from 27 patients. Nuclear FOXA2 expression was not present in any normal esophageal squamous epithelium but was seen in 80% of patients with nondysplastic Barrett’s metaplasia, 86% of patients with low-grade dysplasia, 93% of patients with high-grade dysplasia, and 78% of patients with esophageal adenocarcinoma (Figure 5, A–C, and E). We then performed qRT-PCR for FOXA2 on frozen tissues from 6 esophagectomy cases. Consistent with our immunohistochemical results, we found considerable expression of FOXA2 in the Barrett’s dysplasia of all 6 cases, but the accompanying squamous epithelium exhibited no FOXA2 expression in all but 1 case, which showed minimal expression (Figure 5D). We next performed FOXA2 immunohistochemistry on a tissue microarray containing 11 cases of esophageal squamous cell carcinoma and did not find FOXA2 expression in any patient (Figure 5F). Taken

**Figure 4. Squamous CK14 is prematurely expressed in Shh−/− embryos.** CK14 immunohistochemistry in E14.5 esophagus and forestomach from (A and B) wild-type littermate controls and (C and D) Shh−/− embryos. Esophagus (E) is indicated. Scale bars: 50 μm.

**Figure 5. FOXA2 expression in benign and malignant human esophageal tissue.** FOXA2 immunohistochemistry of (A) normal esophageal squamous epithelium, (B) nondysplastic Barrett’s metaplasia, and (C) dysplastic Barrett’s metaplasia. (D) FOXA2 qRT-PCR of normal esophageal squamous epithelium and Barrett’s dysplasia from esophagectomy specimens. FOXA2 immunohistochemistry of esophageal (E) adenocarcinoma and (F) squamous cell carcinoma. Scale bars: 50 μm.
together, these data suggest that FOXA2 is expressed minimally if at all in normal esophageal squamous epithelium and in esophageal squamous cell carcinomas, but FOXA2 is expressed in Barrett’s metaplasia, dysplasia, and cancer.

We then characterized FOXA2 expression by qRT-PCR and Western blot in 6 telomerase-immortalized human esophageal epithelial cell lines developed from endoscopic biopsies taken from patients who had GERD with and without Barrett’s esophagus (refs. 16, 46, 47, and Figure 6, A and B). In squamous epithelial cell lines derived from patients who had GERD without Barrett’s esophagus (NES-G2T, NES-G4T), minimal FOXA2 transcript was seen and no protein was detected by Western blot. In contrast, the squamous epithelial cells derived from patients who had GERD with Barrett’s esophagus (NES-B3T, NES-B10T) showed slightly increased FOXA2 transcript levels, with some expression of FOXA2 protein. Compared with the squamous cell lines, Barrett’s epithelial cell lines (BAR-T, BAR-10T) showed increased levels of both FOXA2 transcript and protein. We next performed qRT-PCR for FOXA2 on 3 esophageal squamous cell carcinoma cell lines (KYSE70, KYSE180, KYSE510) and 3 esophageal adenocarcinoma cell lines (OE33, FLO-1, JHESoAd1) and found that FOXA2 was substantially higher in the adenocarcinoma lines (Figure 6C).

Figure 6. FOXA2 expression in benign and malignant human esophageal cell lines. (A) FOXA2 qRT-PCR and (B) FOXA2 Western blot of telomerase-immortalized esophageal cell lines. NS, not significant. ***P < 0.001 as compared with NES-G2T. (C) FOXA2 qRT-PCR of esophageal squamous cell carcinoma and adenocarcinoma cell lines. FOXA2 immunofluorescence in (D) BAR-T cells incubated with IgG control (red). (E) BAR-T cells incubated with FOXA2 (red) primary antibody, and (F) OE33 cells incubated with FOXA2 primary antibody (positive control). Scale bars: 50 μm.

FOXA2 is regulated by Hh signaling in Barrett’s epithelial cells and in esophageal squamous cells from patients who have GERD with Barrett’s esophagus. Given our findings of SHH regulation of FOXA2 in the esophageal epithelia of embryonic mice, we sought to determine whether FOXA2 is downstream of Hh signaling in adult human esophageal epithelial cells. We characterized Hh pathway status by performing qRT-PCR in NES-B3T, NES-B10T, BAR-T, and BAR-10T cells for Hh pathway components SHH, PTCH, GLI1, GLI2, and GLI3 (Figure 7A). We selected only the NES-B3T and NES-B10T esophageal squamous cell lines for use, because, unlike NES-G2T and NES-G4T cells that did not express FOXA2 protein, NES-B3T and NES-B10T expressed low baseline levels of FOXA2 protein (Figure 6B). All 4 cell lines expressed similar levels of PTCH. Compared with NES-B3T cells, the LES-B10T, BAR-T, and BAR-10T cells expressed increased GLI1 and GLI2 transcripts. Expression of SHH was highest and expression of GLI3 was lowest in BAR-10T cells, suggesting that BAR-10T cells have the highest active Hh signaling among our 4 esophageal cell lines. To determine whether our telomerase-immortalized cells could secrete active SHH, we cocultured the 4 benign esophageal epithelial cell lines with Shh-Light2 cells. Shh-Light2 cells are 3T3 cells stably transfected with a firefly luciferase reporter gene driven by 8 tandem GLI1 binding sites and a renilla luciferase reporter gene driven by the TK promoter (48). Dual-luciferase assay after 48 hours revealed that active SHH was secreted the most by the BAR-10T Barrett’s cell line (Figure 7B). We next sought to determine whether FOXA2 expression...
was dependent on Hh signaling in BAR-10T cells. Following GLI1 siRNA-mediated knockdown, we found that FOXA2 expression decreased (Figure 7C). This suggests that in Barrett’s epithelial cells, FOXA2 is dependent on Hh signaling.

To determine whether FOXA2 expression can be activated by Hh signaling in esophageal squamous epithelial cells from patients who have GERD with Barrett’s esophagus, we used recombinant SHH and plasmids containing either mouse Shh or Gli1. We electroporated NES-B3T and NES-B10T cells with pSRα-FLAG-Gli1 and analyzed them by qRT-PCR 48 hours later. We found that GLI1, as expected, induced expression of Hh pathway targets PTCH and GLI1 itself (Figure 8, A and C). In both squamous cell lines, GLI1 overexpression also led to FOXA2 induction (Figure 8, A and C). Electroporation of NES-B3T cells with pRK5-Shh (Figure 8B) or treatment of NES-B10T cells with recombinant SHH (Figure 8D) led to increased PTCH, GLI1, and FOXA2 expression by qRT-PCR, suggesting that esophageal squamous epithelial cells from patients who have GERD with Barrett’s esophagus can respond to Hh signaling.

To further characterize how GLI1 mediates activation of FOXA2, we analyzed the promoter region of FOXA2 for putative GLI binding sites (GLI-Bs), but none were identified. In 1997, Sasaki and colleagues described a minimal enhancer 3′ of the mouse Foxa2 coding sequence, which contains an essential GLI-B with the sequence 5′-GAACACCCA-3′ (27). This GLI-B was found 6,577-bp downstream of the stop codon. We analyzed the human FOXA2 gene and found this same GLI-B 6,584-bp from the stop codon (Figure 8E). We next obtained luciferase reporter constructs driven by a basal δ-crystalline promoter and 8 tandem copies of either the GLI-B or a mutated GLI-B (mutGLI-B) with the sequence 5′-GAAGTGGGA-3′. We electroporated these constructs, along with mouse GLI1, into NES-B10T cells (in which we had shown that exogenous GLI1 upregulates FOXA2 expression). Luciferase activity increased significantly when GLI1 was coexpressed with GLI-B but not when GLI1 was coexpressed with mutGLI-B (Figure 8E). This demonstrates that GLI1 binds to this FOXA2 enhancer and that FOXA2 is a direct target of the Hh-GLI signaling cascade.

FOXA2 regulates AGR2 and MUC2 in Barrett’s epithelial cells and in esophageal squamous cells from patients who have GERD with Barrett’s esophagus. We next sought to identify FOXA2 target genes that are relevant to Barrett’s esophagus. Barrett’s specialized intestinal metaplasia is characterized by prominent goblet cells and other cells that express MUC2, a protein disulfide isomerase that is localized to the endoplasmic reticulum and required for proper MUC2 protein processing (49). Agrp−/− mice lack normal goblet cells and have decreased intestinal mucus (50). Agrp−/− mice lack normal goblet cells and have decreased intestinal mucus (50). AGR2 promoter and upstream sequence contain 4 FOXA2 consensus binding sites [VA(A/T)TRTT(G/T)]
RYTY, where V = A, C, G; R = purine; and Y = pyrimidine] (51). Recognizing that Barrett’s epithelial cells express more FOXA2 than squamous cells from patients with Barrett’s esophagus, we sought to delineate FOXA2’s function through siRNA-mediated knockdown of FOXA2 in Barrett’s cells and through overexpression of FOXA2 in squamous cells. pIRES-FOXA2 was electroporated into NES-B3T and NES-B10T cells, and FOXA2 expression was confirmed by immunofluorescence, which demonstrated strong nuclear staining (data not shown). We then examined the effect of FOXA2 overexpression in NES-B3T and NES-B10T cells by Western blot and found that this led to increased expression of AGR2 protein (Figure 9A). siRNA-mediated knockdown of FOXA2 in BAR-10T and BAR-T cells led to decreased expression of AGR2 protein (Figure 9B; BAR-T data not shown). These findings suggest that AGR2 is a direct target gene of FOXA2. Since AGR2 is required for proper processing of mature MUC2 protein, we examined the effect of FOXA2 overexpression on MUC2 expression by performing immunofluorescence for MUC2 on NES-B3T and NES-B10T cells electroporated with FOXA2. We found that MUC2 protein expression increased in both squamous cell lines electroporated with FOXA2 (Figure 9C; NES-B10T data not shown). FOXA2 can regulate MUC2 expression either directly (via transcriptional activation) or indirectly (through AGR2 induction). Therefore, we performed qRT-PCR for AGR2 and RT-PCR for MUC2 in NES-B3T and NES-B10T cells that were electroporated with FOXA2 (Figure 9D) or in BAR-T cells, in which FOXA2 expression was knocked down with siRNA (Figure 9E). We found that AGR2 transcript increased in NES-B3T and NES-B10T cells and decreased in BAR-T cells; similarly, MUC2 transcript increased in NES-B3T and NES-B10T cells and decreased in BAR-T cells (Figure 9, D and E). This suggests that MUC2 is transcriptionally regulated by FOXA2 in esophageal squamous and Barrett’s epithelial cells. FOXA2-mediated transcriptional regulation of AGR2 would also affect MUC2 protein levels, because AGR2 is required for MUC2

Figure 8. Hh signaling induces FOXA2 in telomerase-immortalized squamous esophageal cells. FOXA2 qRT-PCR following GLI1 electroporation in (A) NES-B3T and (C) NES-B10T cells or following (B) electroporation of SHH in NES-B3T cells or (D) treatment with recombinant SHH in NES-B10T cells. (E) NES-B10T cells were electroporated with a firefly luciferase reporter gene driven by 8 tandem copies of the GLI-BS found within a 3′ enhancer of the Foxa2 gene or a mutGLI-BS with sequence 5′-GAACTGCGA-3′ and either empty vector or GLI1. Data were normalized using TK-Renilla luciferase activity. Schematic of the genomic sequence of mouse Foxa2 and human FOXA2 with the GLI-BS 3′ to the coding sequence (CDS). Stop codon is shown in red. *P < 0.05. **P < 0.01 as compared with vector or vehicle control.
The origin of Barrett’s metaplasia is not known. One hypothesis holds that Barrett’s metaplasia develops from a unique population of progenitor cells that are capable of differentiating into both squamous and columnar epithelial cell types. Our studies in embryonic mice demonstrate the early commitment of foregut epithelium to either a respiratory or gastrointestinal cell fate. Using Shh–/– embryos, we have shown that Hh regulation of FOXA2 differs between the gastrointestinal epithelium lining the esophagus and stomach and the respiratory epithelium lining the lungs. Esophageal and foregut progenitor cells share a common foregut origin. These findings suggest that molecular mechanisms in respiratory epithelium cannot necessarily be extrapolated to those in the esophagus and underscore the importance of studies focused specifically on esophageal epithelium.

Discussion

Using a developmental approach, we identified the transcription factor FOXA2 as being expressed by embryonic mouse esophageal columnar epithelium. Our studies performed with conditionally activated ShhTg mouse esophageal epithelium and Shh–/– mouse embryos established that Foxa2 is an esophageal Hh target gene. In humans, FOXA2 was expressed in Barrett’s metaplasia, dysplasia, and adenocarcinoma tissue specimens; in nonneoplastic Barrett’s epithelial cell lines; and in esophageal squamous cell lines derived from patients with Barrett’s metaplasia. In the squamous cell lines, FOXA2 was upregulated by Hh signaling, causing expression of MUC2, an intestinal mucin found in Barrett’s specialized intestinal metaplasia, and AGR2, an endoplasmic reticulum protein involved in MUC2 processing. This suggests that reactivation of Hh signaling in esophageal squamous epithelial cells can induce genes that determine an intestinal phenotype and may contribute to the development of the specialized intestinal type of Barrett’s metaplasia.

Our studies in embryonic mice demonstrate the early commitment of foregut epithelium to either a respiratory or gastrointestinal cell fate. Using Shh–/– embryos, we have shown that Hh regulation of FOXA2 differs between the gastrointestinal epithelium lining the esophagus and stomach and the respiratory epithelium lining the lungs. Esophageal and foregut progenitor cells share a common foregut origin. These findings suggest that molecular mechanisms in respiratory epithelium cannot necessarily be extrapolated to those in the esophagus and underscore the importance of studies focused specifically on esophageal epithelium.

The origin of Barrett’s metaplasia is not known. One hypothesis holds that Barrett’s metaplasia develops from a unique pop-
that contention, we found differences between esophageal squamous cell lines from patients who have GERD with and without Barrett's esophagus in how acid activates ERK1/2, in baseline expression of an inhibitory form of MEK1, and in how acid and bile salts increase expression and activation of CDX2 (16, 56, 57). In the present study we have found expression of FOXA2, a key developmental transcription factor involved in specifying an intestinal-type columnar-lined esophagus, in esophageal squamous epithelial cells from patients with Barrett's esophagus but not in esophageal squamous epithelial cells from patients who have GERD without Barrett's esophagus. Such differences might underlie the development of Barrett's metaplasia in some individuals with GERD. We have shown that FOXA2 is a direct Hh target gene in human esophageal squamous and Barrett's epithelium. In squamous cells, FOXA2 can induce expression of MUC2, an intestinal mucin found in specialized intestinal metaplasia, and AGR2, an endoplasmic reticulum protein required for proper processing of MUC2. In earlier studies, we showed that the transcription factor SOX9, found in columnar mouse embryonic esophageal epithelium and in human Barrett’s metaplasia, is an indirect Hh target (indirect because SOX9 expression is upregulated by stromal BMP4 signaling induced by epithelial Hh signaling). Unlike SOX9, FOXA2 is a direct Hh target gene that regulates the expression of MUC2, a protein involved in the intestinal alterations characteristic of specialized intestinal metaplasia (25). Based on these findings, we now propose a molecular model of Barrett’s metaplasia in which Hh signaling, through direct and indirect pathways, induces

![Figure 10. Proposed role of FOXA2 in a molecular model of transcommitment causing Barrett’s metaplasia.](http://www.jci.org)

**A** The components of gastroesophageal reflux, acid and bile acids, cause secretion of SHH by esophageal squamous cells. **B** SHH signals to stroma to induce expression of BMP4 by esophageal fibroblasts, which then signal back to the squamous epithelial cells to induce SOX9 expression (epithelial-mesenchymal crosstalk). SHH can also induce expression of FOXA2 in esophageal squamous epithelial cells. **C** Activation of SOX9 and FOXA2 leads to expression of genes involved in columnar and intestinal goblet cell differentiation, such as CK8 and MUC2.
separate transcription factors relevant to the columnar intestinal metaplasia of Barrett’s esophagus (Figure 10).

Though our studies demonstrate that FOXA2 upregulates MUC2 transcript and protein expression in esophageal squamous epithelial cells from patients who have GERD with Barrett’s esophagus, we did not observe a switch to a full goblet cell phenotype. To our knowledge, direct nongoblet-to-goblet transition has not been demonstrated in esophageal epithelium in vitro. In vivo, treatment with γ-secretase inhibitors and resultant loss of Notch signaling in a surgical rat model of reflux esophagitis and Barrett’s metaplasia leads to esophageal goblet cell hyperplasia within the reflux-induced metaplasia (58). During mouse esophageal development, hypomorphic SOX2 or complete loss of expression of either p63 or Noggin causes the appearance of goblet-like cells within the epithelium (53, 59, 60). It is possible that FOXA2 overexpression combined with knockdown of SOX2 or p63, inhibition of Notch signaling, or activation of BMP signaling would lead to a full goblet cell phenotype in our esophageal squamous cell lines.

MUC2 regulation within Barrett’s metaplasia appears to be complex. CDX2 has been accepted as a major regulator of MUC2 expression, and CDX2 has been shown to be upregulated by acidic pH and bile acids in esophageal squamous epithelial cells (16). Another study suggests that Notch inhibition with γ-secretase inhibitors leads to loss of repression of ATOH1, which activates CDX2, which in turn leads to MUC2 expression (61). FOXA2 can also transcriptionally activate CDX2, but this appears to be repressed in normal squamous esophageal epithelium by the coexpression of the transcriptional repressor RXF1, which is lost in Barrett’s metaplasia (52). We now demonstrate in esophageal squamous epithelial cells that FOXA2 regulates MUC2 mRNA in a CDX2-independent fashion. In addition, we have shown that FOXA2 upregulates AGR2, a protein that has been associated with Barrett’s metaplasia and is required for MUC2 protein processing. Thus, the combinatorial effect of CDX2 and FOXA2 may act to increase expression of MUC2 in Barrett’s metaplasia.

In summary, using a developmental approach, we have identified Foxa2 as a Hh esophageal target gene. FOXA2 can be directly activated by Hh signaling in human esophageal squamous epithelium, and FOXA2 is expressed in Barrett’s metaplasia, in which we previously demonstrated active Hh signaling. FOXA2 can induce expression of MUC2 and AGR2, two genes that are important for acquisition of the intestinal cell phenotype characteristic of Barrett’s specialized intestinal metaplasia. These data suggest that FOXA2 regulates genes involved in the pathogenesis of specialized intestinal metaplasia, the type of columnar metaplasia associated with cancer risk in Barrett’s esophagus. Moreover, our findings provide a molecular rationale for future studies using Hh pathway inhibitors to prevent the development and neoplastic progression of Barrett’s metaplasia in patients who have GERD.

Methods

Mice. Timed-pregnant C57BL/6 female mice were sacrificed, and whole embryos were harvested between E11.5 and E17.5. The day the presence of a vaginal plug was detected was designated as E0.5. Gastrointestinal organs were harvested from sacrificed P1 and P2 pups and adult C57BL/6 mice. Shh−/− mice were generated by replacing the second exon and surrounding introns with a PGK-neo selection cassette, as previously described (44), and were maintained in CD1 mice. Alternatively, Shh−/− mice were generated by crossing Shh-Cre+/- mice (JAX strain 005622) (42). Shh−/− embryos were harvested from timed-pregnant females at E14.5 and E16.5, and genotypes were confirmed. Esophageal-specific deletion of Sox9 was obtained by crossing male Shh-Cre-/+ mice (42) with female Sox9fl/fl mice (JAX strain 013106) (43). N1 male Shh-Cre+/- Soxxfl/fl mice were then crossed with female Sox9fl/fl mice to generate the desired genotype. To confirm esophageal epithelium-specific expression of Shh-Cre, we crossed Shh-Cre+/- mice with Gr(ROSA)-LacZ mice (JAX strain 003309) (62) and stained tissue sections with X-gal to determine β-galactosidase activity. Wild-type control embryos were littermates of experimental animals.

Microarray. Microarray analysis was performed as previously described (63). Total esophageal RNA from E15.5 and P2 C57BL/6 mice was isolated with Trizol (Invitrogen) and then further purified using RNeasy (Qiagen). RNA was quantitated with a NanoDrop ND-100, and the quality was determined using an Agilent 2100 Bioanalyzer. Sample amplification and labeling procedures were carried out according to the manufacturer’s protocol using the Low RNA Input Fluorescent Linear Amplification Kit (Agilent) and 0.3 μg total RNA as starting material. The Agilent Whole Mouse Genome microarray (G4122A), with 41,174 unique transcripts, was used. Following hybridization according to the Agilent microarray processing protocol, microarrays were scanned using an Agilent G2505B Scanner controlled by Agilent Scan Control 7.0 software. Data were extracted with Agilent Feature Extraction 9.1 software.

For analysis, the base 2 log ratios of the red (Cy5) and green (Cy3) signal intensities, \( M = \log_2 R / G \) (where \( M \) stands for signal intensity, \( R \) stands for Cy5 intensity, and \( G \) stands for Cy3 intensity), were calculated and used in the downstream analysis. Lowess normalization was performed on \( G = \log_2 R / G \) with the K package SMA (http://www.stat.berkeley.edu/~terry/zarray/Software/smacode.html) to remove systematic obscuring variation, such as the fluorescence dye-labeling bias effect, which arises from the microarray technology rather than from the differences of the biological conditions. The log posterior odds of the differential gene expression in mouse E15.5 esophagi as compared with P2 esophagi were computed with the stat.Newton function of SMA (64, 65) (http://www.stat.berkeley.edu/~terry/zarray/Software/smacode.html) according to a hierarchical gamma-gamma model. The criteria of the log odds > 1, which means the posterior odds favoring change were used to produce the differentially expressed gene list. The differential gene expression was visualized with an MvA plot, the intensity log-ratio \( M = \log_2 R / G \) vs. the mean log-intensity \( A = 1/2 \times (\log_2 R + \log_2 G) \) plot. All computation was performed using the statistical computing software R (http://www.r-project.org). The microarray data have been deposited to the NCBI GEO database, with accession no. GSE56528.

In vivo transplant culture system. Esophageal epithelial cells were isolated from wild-type and conditional ShhTg mice as previously described (19). The epithelial cells were infected with adenoviral-cre to activate the conditional transgene. Following this, the epithelial cells along with C57BL/6 esophageal fibroblasts were placed into a denuded rat trachea. The ends of the trachea were ligated, and this culture system was placed subcutaneously into a NOD/SCID mouse for 4 weeks.

Clinical samples. Tissue microarrays containing esophageal squamous epithelium, Barrett’s esophagus, dysplastic Barrett’s esophagus, esophageal adenocarcinoma, stomach, and fibrous tissue from...
37 patients were obtained from the Johns Hopkins Tissue Microarray facility. An additional tissue microarray containing normal esophagus and esophageal squamous carcinoma from 11 patients was purchased from US Biomax (ES242). These tissue microarrays were submitted for immunohistochemistry as described below. Six frozen esophagectomy cases containing normal squamous epithelium and dysplastic Barrett’s were obtained from the Johns Hopkins Tissue Bank. Tissue was cut from each case, and the diagnosis was confirmed by a gastrointestinal pathologist. RNA was isolated from the tissue as described below.

**Immunohistochemistry and immunofluorescence.** Formalin-fixed paraffin-embedded sections were submitted to epitope retrieval with heated citrate buffer and quenching of endogenous peroxidase. Following incubation with primary antibody for 1 hour, immunohistochemistry was completed using the Vectastain ABC Kit (VectorLabs) and DAB reagent (Sigma-Aldrich). Primary antibodies used for immunohistochemistry include FOXA2 (Lifespan) at 1:2,000, CK8 (DSHB) tochemistry was completed using the Vectastain ABC Kit (VectorLabs) and DAB reagent (Sigma-Aldrich). Primary antibodies used for immunohistochemistry include FOXA2 (Lifespan) at 1:2,000, CK8 (DSHB) tochemistry was completed using the Vectastain ABC Kit (VectorLabs) and DAB reagent (Sigma-Aldrich). Primary antibodies used for immunohistochemistry include FOXA2 (Lifespan) at 1:2,000, CK8 (DSHB) tochemistry was completed using the Vectastain ABC Kit (VectorLabs) and DAB reagent (Sigma-Aldrich). Primary antibodies used for Western blotting include FOXA2 (Santa Cruz) at 1:50 and MUC2 (Novocastra) at 1:100.

**Cell lines.** Human esophageal epithelial cells were isolated from endoscopic biopsies of squamous and Barrett’s epithelium (defined as the presence of specialized intestinal metaplasia) and telomerase immortalized as previously described (46). Squamous cell lines from patients with GERD with (NES-B3T, NES-B10T) and without Barrett’s esophagus (NES-G2T, NES-G4T) and metaplastic Barrett’s esophageus cell lines (BAR-T, BAR-10T) were used in experiments. These benign cell lines were maintained, without a fibroblast feeder layer, on dishes precoated with 0.01 mg/ml fibronectin (Sigma-Aldrich), 0.03 mg/ml type I collagen (BD Biosciences), and 0.01 mg/ml bovine serum albumin (Sigma-Aldrich) in keratinocyte medium for 1 hour and then washed twice with PBS. Esophageal squamous carcinoma cell lines (KYSE70, KYSE180, KYSE510) were provided by John Harmon (Johns Hopkins University, Baltimore, Maryland, USA). Esophageal adenocarcinoma cell lines were purchased from ATGC (OE33) or provided by James Eshleman, (Johns Hopkins University) (JHEsoAd1) and David Beer (University of Michigan, Ann Arbor, Michigan, USA) (FLO-1). Esophageal epithelial cells were electroplanted using Nucleofector 4 (Axamca) or transfected using Lipofectamine LTX (Invitro-gen) with the indicated plasmids. Reactions were performed in 6-well plates with a total 6 μg DNA for electroporations and 3 μg DNA for transfections. FOXA2 was PCR amplified from a full-length cDNA (Open Biosystems) and cloned into pIREs.

**Hh signaling assays.** Untreated telomerase-immortalized epithelial cells were cocultured with Shh-Light2 cells (48) for 48 hours. Luciferase activity was then assayed by the Dual-Luciferase Reporter System (Promega). Esophageal squamous epithelial cells from patients with Barrett’s metaplasia were treated with 1 μg/ml rSHH (R&D Systems) or vehicle for 48 hours in serum-free medium and then RNA harvested for analysis.

**Enhancer assay.** Reporter constructs 8x3’GLI-BS-delta51-LucII (GLI-BS) and 8xm3’GLI-BS-delta51-LucI (mutGLI-BS) (27), which had been previously deposited by Hiroshi Sasaki (Kumamoto University), were obtained from the RIKEN BioResource Center, Tsukuba, Japan. NES-B10T cells were electroporated with TK-Renilla luciferase, either GLI-BS or mutGLI-BS reporter constructs, and either pcDNA3.1 (vector) or pSh-FLAG-Gli. Luciferase activity was then assayed by the Dual-Luciferase Reporter System (Promega) 48 hours later.

**siRNA and Western blotting.** siRNA-mediated knockdown was performed as previously described (19) using individual FOXA2 or SMARTpool GLI1 siRNAs (Dharmacon). Protein lysates were harvested at 48 hours and electrophoresed on a Bis-Tris gel (Invitrogen). Following transfer to nitrocellulose and blocking with 5% milk in TBST, primary antibody was applied overnight. The following day, membranes were incubated with secondary antibody for 1 hour, followed by protein detection with chemoluminescence. Primary antibodies used for Western blotting include FOXA2 (Santa Cruz) at 1:100, AGR2 (Sigma-Alldrich) at 1:500, and GAPDH (Millipore) at 1:5,000.

**RT-PCR and qRT-PCR.** Total RNA was isolated with Trizol (Invitrogen), and 1 μg RNA was reverse transcribed with Quantitect (Qiagen). cDNA was then used as a template to perform qualitative PCR on a Perkin Elmer GeneAmp 9700 and qRT-PCR with an Applied Biosystems StepOnePlus machine. ΔΔct method was used to determine relative transcript levels on qRT-PCR. Ct values were normalized to β-actin. Primer sequences are provided in Supplemental Table 1.

**Statistics.** Quantitative data are expressed as the mean ± SEM. Data with multiple comparisons were initially tested for normality. ANOVA followed by the Tukey honest significant difference post-hoc test was used to analyze normally distributed data, and the Kruskal-Wallis test was used to analyze nonnormally distributed data. Comparison of 2 means was analyzed using an unpaired 2-tailed Student’s t test. These tests for statistical significance were performed with GraphPad Prism version 4.00 for Windows, GraphPad Software (http://www.graphpad.com). A P value of less than 0.05 was considered significant.

**Study approval.** All work with animals was approved by the Institutional Animal Care and Use Committees of Johns Hopkins University and the VA North Texas Health Care System and the Peter MacCallum Cancer Centre Animal Experimentation Ethics Committee. Research with deidentified human clinical specimens was granted an exemption from the requirement for informed consent and approved by the Johns Hopkins University Institutional Review Board.

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