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Schlafen 4–expressing myeloid-derived suppressor cells are induced during murine gastric metaplasia

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Chronic Helicobacter pylori infection triggers neoplastic transformation of the gastric mucosa in a small subset of patients, but the risk factors that induce progression to gastric metaplasia have not been identified. Prior to cancer development, the oxyntic gastric glands atrophy and are replaced by metaplastic cells in response to chronic gastritis. Previously, we identified schlafen 4 (Slfn4) as a GLI1 target gene and myeloid differentiation factor that correlates with spasmolytic polypeptide-expressing metaplasia (SPEM) in mice. Here, we tested the hypothesis that migration of SLFN4-expressing cells from the bone marrow to peripheral organs predicts neoplastic changes in the gastric microenvironment. Lineage tracing in Helicobacter-infected Slfn4 reporter mice revealed that SLFN4+ cells migrated to the stomach, where they exhibited myeloid-derived suppressor cell (MDSC) markers and acquired the ability to inhibit T cell proliferation. SLFN4+ MDSCs were not observed in infected GLI1-deficient mice. Overexpression of sonic hedgehog ligand (SHH) in infected WT mice accelerated the appearance of SLFN4+ MDSCs in the gastric corpus. Similarly, in the stomachs of H. pylori–infected patients, the human SLFN4 ortholog SLFN12L colocalized to cells that expressed MDSC surface markers CD15+CD33+HLA-DR+. Together, these results indicate that SLFN4 marks a GLI1-dependent population of MDSCs that predict a shift in the gastric mucosa to a metaplastic phenotype.

Introduction

Helicobacter pylori infection induces chronic inflammation in the stomach, which eventually leads to atrophy of the acid-producing glands, metaplasia, dysplasia, and then gastric cancer in some infected patients (1–3). The timeline for gastric cancer development occurs over several decades, raising the likelihood that Helicobacter infection is only one of several factors contributing to transformation of the normal gastric epithelium. Indeed, results from The Cancer Genome Atlas (TCGA) for gastric cancer reported only sporadic molecular signatures of Helicobacter in primary tumors (4). Therefore, infection with Helicobacter alone is not sufficient to predict who among the small subset of infected patients will develop gastric cancer. In most instances, the extent of the inflammatory response also contributes to cancer susceptibility (5, 6). Nevertheless, at least 50% of infected individuals develop chronic gastritis without symptoms or further progression of the mucosa to metaplasia and dysplasia (7). Both intestinal metaplasia and spasmolytic polypeptide-expressing metaplasia (SPEM) are histologic lesions strongly associated with neoplastic transformation (8–11). Identifying markers predictive of preneoplasia would allow clinicians to risk-stratify the subset of individuals at greater risk for progression to gastric cancer and who subsequently require more frequent monitoring, regardless of Helicobacter prevalence (12).

Hedgehog (HH) ligands expressed in the gastrointestinal epithelium activate GLI transcription factors in stromal and immune cells (13). Zavros and coworkers showed that the sonic hedgehog (SHH) ligand secreted from gastric parietal cells within 2 days after the infection is required for myeloid cell recruitment to the infected mouse stomach (14). After 2 months of Helicobacter infection, SHH expression in parietal cells gradually diminishes in response to chronic gastritis, despite adjacent mucous cells retaining ligand expression (15). Moreover, GLI1+ myeloid cells are recruited to the stomach within the initial months of the infection (16). By 6 months, most of the parietal and zymogenic chief cells have atrophied and are replaced by SPEM. However, deletion of one or both Gli1 alleles prevents Helicobacter-induced SPEM, indicating that targets of the HH pathway are required for SPEM development (16). Since HH signaling in the stomach is paracrine (17), blocking HH-dependent crosstalk between infiltrating immune cells and the gastric epithelium is sufficient to prevent histologic transition from chronic gastritis to metaplasia.

A myeloid cell differentiation factor and GLI1 target gene (18) called schlafen 4 (Slfn4) was identified in a microarray analysis of gastric tissue from Helicobacter-infected WT versus Gli1−− mice (16). SLFN4+ myeloid cells express IL-1β and TNF-α (16), cytokines associated with myeloid-derived suppressor cell (MDSC) regulation (6, 19). MDSCs are a heterogeneous myeloid cell population that develops under conditions of infection, tissue injury, autoimmune disease, and cancer (20). Their ability to suppress T cell function dampens the immune response and creates a microenvironment favoring neoplastic transformation (21).

Here we tested the hypothesis that Slfn4 expression marks a GLI1-dependent population of myeloid cells with phenotypic characteristics of MDSCs. We found that SLFN4+ cells in the mouse stomach coincided with SPEM in the setting of chronic gastritis before the development of dysplasia. Since type I interferon induction of Slfn4 gene expression required Gli1, we concluded that HH signaling synergizes with regulatory cytokines to create a permissive environment for gastric metaplasia, a harbinger of possible neoplastic transformation.
two cell-specific markers confirmed the presence of SPEM in the infected mice with WT marrow (Supplemental Figure 1; supplemental material available online with this article; doi:10.1172/JCI82529DS1). WT mice developed metaplastic changes only if infected, and the SPEM correlated with an increase in CD11b+ and SLFN4+ cells (Figure 1B). However, the absence of SPEM in the Gli1–/– mice correlated with reduced numbers of SLFN4+, but not a significant reduction of CD11b+ cells (Figure 1B), suggesting a shift in the composition of the myeloid population rather than a defect in total myeloid cell recruitment.

Results

SPEM development requires Hedgehog signaling. Helicobacter-induced SPEM in WT mice was observed at 6 months after infection. Infected Gli1fl/fl;Hsd:1SH mice (referred to here as Gli1+ mice) did not develop SPEM as previously reported, which implicates a significant role for canonical HH signaling in the emergence of this preneoplastic lesion (Figure 1A and ref. 16). In the stomach, stromal cells, specifically myofibroblasts and immune cells, express GLI1 (16, 17, 22). To determine whether the metaplastic change was due to bone marrow-derived cells (BMDCs), we transferred marrow from Slfn4-eGFP-CreER2 Rosa26-LSL-ttdTomato (Slfn4-ttT; see below) mice (that are Gli1+/−) into irradiated WT (WT w/bone marrow transplant [BMT]) or Gli1–/– mice (Gli1–/– w/BMT) prior to infection with Helicobacter for 6 months. Infected Gli1−/− w/BMT mice that received WT marrow developed SPEM, demonstrating that BMDCs were sufficient to transfer gastric susceptibility to Helicobacter-induced metaplasia (Figure 1A). SPEM is defined by the combined expression of chief cell (e.g., gastric intrinsic factor) and mucous neck cell (e.g., TFF2 and GSII-lectin) markers that represent reprogramming of the mature gastric chief cells in the setting of parietal cell atrophy (23–25). Overlap between these

Figure 1. Bone marrow–derived cells are sufficient to induce SPEM with H. felis. (A) Representative H&E images (n = 6) for each group as indicated with and without bone marrow transplant (BMT) (from Slfn4-CreER2 Rosa26-LSL-ttdTomato [Slfn4-ttT] mice) with and without H. felis infection (Hf). UI, uninfected. Scale bars: 50 μm. (B) Flow analysis of SLFN4-ttT+ or CD11b+ cells versus side scatter (SSC). The percentage of SLFN4+ or CD11b+ cells per mouse is shown in the representative histograms and plotted in the adjacent scatter graphs for n = 6 mice per group over 3 experiments. Horizontal lines represent the median and interquartile range. Significance was determined using Kruskal-Wallis ANOVA with Dunn’s test of multiple comparisons. *P < 0.05.
infection, there was a rapid increase in the blood levels of SHH, which peaked within days after the infection and then returned to baseline levels after 1 month (Figure 2A). The result was consistent with a prior report indicating that H. pylori infection increases circulating levels of SHH, which originates from gastric parietal cells (14). H. felis–infected pCMV-Shh mice exhibited greater inflammation in the gastric corpus at 2 months, and subsequently SPEM was observed at 4 months, which was 2 months earlier than observed in WT littermates (Figure 2B). The amount of polymorphonuclear (PMN) leukocytes infiltrating the mucosa and the extent of SPEM, as quantified by morphometry, were greater in the SHH-overexpressing mice (Figure 2, C and D). SPEM development in pCMV-Shh mice was also documented by de novo expression of clusterin, as well as coexpression of GSII lectin and GIF at the oxyntic gland base, with atrophy of H’-K’-ATPase–positive parietal cells (Figure 2E). Real-time quantitative PCR (RT-qPCR) analysis for H’-K’-ATPase α subunit (Atp4a), Gif, mucous neck cell and SPEM cell marker trefoil factor 2 (Tff2), and clusterin mRNAs were used to quantify the extent of the metaplastic epithelial response (Supplemental Figure 3). Unlike the corpus, the antrum exhibited no phenotypic changes (Supplemental Figure 4).

SHH accelerates the appearance of SLFN4. Since ectopic expression of SHH accelerated SPEM development, we queried whether the appearance of SLFN4+ cells also occurred earlier. A Slfn4-eGFP-CreERT2 BAC transgene was generated to trace SLFN4+ cells from the bone marrow by breeding this transgenic line to Rosa26-LSL-tdTomato mice to produce the transgenic line Slfn4-eGFP-CreERT2 Rosa26-LSL-tdTomato (referred to here as Slfn4-tdT mice), which served as bone marrow donors (Figure 3A). Bone marrow cells from the Slfn4-tdT mice were injected into 8-week-old Slfn4−/− mice, and 6 weeks later, they were injected with H. felis (8). Bone marrow–derived cells from transgenic mice were detected by eGFP fluorescence at 4 months postinfection (Figure 3B). Histologic scoring of gastric PMN infiltration and SPEM for n = 8–10 mice per time point over 3 experiments. Kruskal-Wallis ANOVA with Dunn’s test of multiple comparisons was performed. *P < 0.05; horizontal lines represent the median and interquartile range. (E) Images of 4-month-infected gastric corpus stained with GSII lectin (green), clusterin (CLU, green), GIF (red), and H’-K’-ATPase (HK, white) in WT and pCMV-Shh mice. Scale bars: 50 μm. n = 5 mice per group for B and E.
dependent increase in SLFN4+ cells that correlated with the duration of Helicobacter infection (Figure 3E and Supplemental Figure 5B).

Slfn4 mRNA and SLFN4 protein were detectable in total cell lysates from the corpus, and expression peaked at 4 and 6 months after H. felis infection with the appearance of SPEM in the pCMV-Shh and WT mice, respectively (Figure 3F and Figure 2B). Using gastric lysates, a Western blot analysis of the tdTomato (tdT) protein (red fluorescent protein) matched the pattern of endogenous SLFN4 expression, except in mice without BMT (untreated [UT]), which did not express tdT. Over the subsequent months after infection, both endogenous SLFN4 protein and the tdT protein were at low levels until 4 months after the infection, when Slfn4 expression peaked in the pCMV-Shh mice (Figure 3G). Epifluorescence revealed the presence of SLFN4+ cells in the lamina propria of the infected pCMV-Shh corpus 4 months after infection (Figure 3H).

Figure 3. SLFN4+ cells track to the stomach after H. felis infection. (A) Schematic diagram of Slfn4-tdT (Slfn4-CreERT2 Rosa26-LSL-tdTomato) mice generated. (B) Protocol to lineage trace SLFN4-expressing cells from the bone marrow to peripheral organs. WT versus pCMV-Shh chimeric mice reconstituted with Slfn4-tdT bone marrow were treated with Tx 2 weeks prior to euthanization. Shown is the percentage of SLFN4+ cells in the (C) bone marrow, (D) spleen, and (E) corpus by flow cytometry, plotted as the median and interquartile range for n = 8–10 mice per group over 3 experiments. Kruskal-Wallis ANOVA with Dunn’s test of multiple comparisons was performed for C–E; *P < 0.05. The corpus was analyzed for (F) Slfn4 mRNA by RT-qPCR and (G) tdT, SLFN4, and GAPDH protein by Western blot for n = 8–10 mice. One-way ANOVA followed by Tukey’s multiple comparisons test on log-transformed RT-qPCR values was performed for F; **P < 0.01. UT, untreated. (H) Representative images from n = 5 pCMV-Shh mouse corpora infected for 4 months showing SLFN4-tdT red+ cells with E-cadherin (green) and DAPI. Left: original magnification, ×200. White box region enlarged in the middle (tdT and DAPI) and right (DAPI only) panels (magnification, ×1,000). Asterisks indicate granulocytic nuclear morphology. (I) Giemsa stain of SLFN4-tdT+ cells sorted from n = 3 infected pCMV-Shh mouse stomachs (original magnification, ×600) showing granulocytic and monocytic cells enumerated per 100 cells. Black box indicates enlarged region (original magnification, ×1,000).

old irradiated WT and pCMV-Shh mice. After a 6-week engraftment period, the two groups were infected with H. felis, then pulsed with tamoxifen (Tx, 0.1 mg/g body weight/d) 2 weeks before euthanization at the indicated time points (Figure 3B). The percentage of SLFN4+ cells from the bone marrow, spleen, liver, intestine, and stomach corpus was analyzed. When we compared the uninfected pCMV-Shh mice with WT mice, there was a notable increase in the percentage of SLFN4+ cells in the bone marrow and spleen, with a pronounced trend toward significance (Supplemental Figure 5A), but no differences were observed in the corpus (stomach), liver, or intestine in the absence of the Helicobacter infection (Supplemental Figure 5, A, C, and D). After infection with H. felis, the number of SLFN4+ expressing cells remained similar in the bone marrow and spleen of the pCMV-Shh mice compared with WT mice (Figure 3, C and D, and Supplemental Figure 5B). Moreover, there was no significant increase in the number of SLFN4+ cells in the liver and intestine with infection over 6 months (Supplemental Figure 5, C and D). However, in the gastric corpus, there was a time-dependent increase in SLFN4+ cells that correlated with the duration of Helicobacter infection (Figure 3E and Supplemental Figure 5B). Slfn4 mRNA and SLFN4 protein were detectable in total cell lysates from the corpus, and expression peaked at 4 and 6 months after H. felis infection with the appearance of SPEM in the pCMV-Shh and WT mice, respectively (Figure 3F and Figure 2B). Using gastric lysates, a Western blot analysis of the tdTomato (tdT) protein (red fluorescent protein) matched the pattern of endogenous SLFN4 expression, except in mice without BMT (untreated [UT]), which did not express tdT. Over the subsequent months after infection, both endogenous SLFN4 protein and the tdT protein were at low levels until 4 months after the infection, when Slfn4 expression peaked in the pCMV-Shh mice (Figure 3G). Epifluorescence revealed the presence of SLFN4+ cells in the lamina propria of the infected pCMV-Shh corpus 4 months after infection (Figure 3H). A magnified view of the tdT+ cells in the lamina propria showed the presence of granulocytic nuclei in the SLFN4-tdT+ cells. To further identify the nuclear morphology of the SLFN4+ cells from
SLFN4 marks MDSCs in the gastric mucosa. Ectopic expression of SHH increased the number of cells that were CD11b+Gr-1+SLFN4+ 4 months after infection compared with WT mice that acquired the same cell population at 6 months (Supplemental Figure 6). Mouse MDSC markers are defined as CD11b+Gr-1+ and subdivided into monocytic (Ly6G–Ly6Chi) and granulocytic (Ly6G+Ly6Clo) subtypes. Analysis for Gr-1 subunits confirmed that the SLFN4+ cells from bone marrow, spleen, and stomach were Ly6G+Ly6Clo, consistent with a granulocytic MDSC phenotype (Gr-MDSCs) (Figure 4A). Next, we queried whether the gastric SLFN4+ population acquired the ability to suppress T cells. MDSCs suppress T cells in part by depleting l-arginine from the microenvironment. l-Arginine depletion by MDSCs suppresses CD3-ζ chain biosynthesis, which subsequently blocks T cell proliferation during the G_{1} phase of the cell cycle, ultimately reducing T cell cytokine production (20). MDSCs express high levels of inducible NOS (iNOS, or NOS2) and arginase I (ARG1), which consume l-arginine in the microenvironment to produce nitric oxide and ROS (26). We found that the SLFN4+ cells were positive for iNOS in the corpus but not in the bone marrow or spleen, suggesting that acquisition of the MDSC phenotype occurred in the stomach (Figure 4A). Gated SLFN4+iNOS+ cells from the corpus were further identified as ARG1+ cells. Indeed, further analysis of these cells in the stomach demonstrated colocalization of SLFN4-tdT with iNOS and ARG1 (Figure 4B). A CFSE-based T cell suppression assay was performed to detect the suppressive function of SLFN4+ cells. When the SLFN4+ cells from the stomach were sorted and cocultured with activated T cells, we observed a 52% suppression of T cell activation with the SLFN4+ cells isolated from the stomach, but only minimal suppression when the SLFN4+ cells were isolated from the bone marrow or spleen (Figure 4C). Tregs (CD4+CD25+) from spleen were used as the positive control. Kruskal-Wallis ANOVA with Dunn’s test of multiple comparisons was performed. *P < 0.05.

**Figure 4. SLFN4+ cells from gastric corpus suppress T cell proliferation.** (A) SLFN4+ cells from bone marrow, spleen, and corpus were analyzed for Ly6G, Ly6C, and iNOS expression. Gated SLFN4+‘iNOS’ cells from the corpus were further analyzed for arginase I (ARG1). Median percentage (interquartile range) for each gated subpopulation shown in red for n = 4 mice per tissue. (B) Representative images from n = 4 mouse stomachs of SLFN4-tdT+ cells (red) colocalized with ARG1 (green) and iNOS (green). Scale bars: 20 μm. Asterisks indicate colocalization. (C) SLFN4+ cells were sorted from the indicated tissues of 3 pooled 4-month-infected infected pCMV-Shh mice and then cocultured with naive splenic T cells prestained with the CFSE dye. T cell proliferation was initiated with anti-CD3/CD28 microbeads cultured with or without SLFN4+ cells for 72 hours, and CFSE+ T cells were quantified by flow cytometry. The median percentage of proliferating T cells is shown in the representative histograms (nos. 1–6) plotted for n = 5 experiments in the scatter graph. Tregs (CD4+CD25+) from spleen were used as the positive control. Kruskal-Wallis ANOVA with Dunn’s test of multiple comparisons was performed. *P < 0.05.
a three-step process requiring a pool of circulating SLFN4+ cells (which was expanded by SHH overexpression), accumulation of these cells in the corpus in response to the Helicobacter infection, and then acquisition of the ability to suppress T cell proliferation.

**Type I IFNs regulate SLFN4.** Van Zuylen et al. previously showed that type I interferons induce Slfn4 mRNA expression in bone marrow–derived macrophages through a MYD88-independent mechanism (18). Moreover IFN-α strongly induces all human schlafen family members (27). Since induction of Slfn4 expression correlated with the acquisition of MDSC function coincident with Tukey’s multiple comparisons test on log-transformed values was performed. *P < 0.05, ***P < 0.001. Immunofluorescence images for G IFN-J and (H) cleaved caspase-3 with H+-K+-ATPase (HK-ATPase) in the gastric corpus of 2-month H. felis–infected pCMV-Shh mice. Inset: Image of uninfected pCMV-Shh gastric corpus stained for cleaved caspase-3. Asterisks and arrows indicate nonapoptotic and apoptotic cells, respectively. G, left, original magnification, ×200; G, right, and H, magnification, ×400); (I) Images for IFN-α with E-cadherin in 4-month-infected pCMV-Shh mice. I, left: original magnification, ×200; I, right: magnification, ×600. White boxes indicate enlarged regions. Images in G–I are representative of n = 3 mouse stomachs.

**Figure 5. Analysis of DAMP signaling components in H. felis–infected corpus.** Total RNA was extracted from H. felis–infected WT and pCMV-Shh corpus lysates and analyzed by RT-qPCR for (A) Trlr9, (B) Myd88, (C) Irfb1, (D) Irf7, and (E) Irfn mRNA at the indicated time points, and then expressed as fold change relative to uninfected WT mice (UI). (F) Serum IFN-α levels determined by ELISA. Shown are the median and interquartile range for n = 8–10 mice per group over 3 experiments. One-way ANOVA followed by Tukey’s multiple comparisons test on log-transformed values was performed. *P < 0.05, ***P < 0.001. Immunofluorescence images for G IFN-J and (H) cleaved caspase-3 with H+-K+-ATPase (HK-ATPase) in the gastric corpus of 2-month H. felis–infected pCMV-Shh mice. Inset: Image of uninfected pCMV-Shh gastric corpus stained for cleaved caspase-3. Asterisks and arrows indicate nonapoptotic and apoptotic cells, respectively. G, left, original magnification, ×200; G, right, and H, magnification, ×400); (I) Images for IFN-α with E-cadherin in 4-month-infected pCMV-Shh mice. I, left: original magnification, ×200; I, right: magnification, ×600. White boxes indicate enlarged regions. Images in G–I are representative of n = 3 mouse stomachs.
Coexpression of H-‘K’-ATPase and cleaved caspase-3 suggested that the dying parietal cells were the likely source of DAMP signals (Figure 5H). IFN-α was expressed on both lamina propria cells and E-cadherin –CD8α+ cells, which are likely cytotoxic T or dendritic cells (31, 32). Since peak gene expression correlated with SPEM transcription (18) (Figure 6D). The SLFN4+ cells responded directly to IFN-α, and that both required GLI1 (Figure 6B). To directly confirm that Slfn4 induction by either IFN-α or IFN-β revealed that IFN-α was a more potent inducer of Slfn4 gene expression than IFN-β and that both required GLI1 (Figure 6B). To directly confirm that the SLFN4+ cells responded directly to IFN-α, we generated primary myeloid cells from the Slfn4-tdT and Slfn4-eGFP-CreER T2 mouse lines and then treated both populations of primary myeloid cultures with IFN-α. No epifluorescence was observed in the absence of Tx/with IFN-α or with Tx/without IFN-α (Figure 6C). However, after Tx and IFN-α treatment, epifluorescent SLFN4+ cells were observed in primary myeloid cells isolated from the Slfn4-tdT but not the Slfn4-tdT Glit1+/– mouse lines (Figure 6C).

The Slfn4 promoter consists of a Glit binding site at −3323 (16) and several STAT/ISRE elements at −239 near the start site of transcription (18) (Figure 6D). Slfn4 promoter activity required GLI1; however, elevated SHH levels did not appear sufficient to boost Slfn4 expression in the corpus (Figure 3F), whereas peak induction required IFN-α, suggesting synergy between the HH pathway and an inflammatory signal. To test this concept in the context of a Helicobacter infection, we treated peritoneal myeloid cells from WT and Glit1+/– mice with recombinant SHH (rSHH), H. felis, or both rSHH and H. felis. Although rSHH and H. felis treatment was...
that IFN-α strongly induced Slfn4 expression in only a subset of CD11b^+GR-1^+ cells. Gli1-dependent iNos and Arg1 mRNA expression were highest in the SLFN4^+ compared with SLFN4^− cells after IFN-α treatment (Supplemental Figure 9, A and B). Moreover, the ability to suppress T cell function was greatest only in the CD11b^+Gr-1^+SLFN4^+ cells compared with the SLFN4^− population (Supplemental Figure 9C).

SLFNs associated with Gr-MDSCs and intestinal metaplasia in human stomach. Since the mouse data suggested that SLFNs might function as potential biomarkers for gastric cancer precursor lesions such as intestinal metaplasia in human subjects, we examined whether SLFNs are expressed in H. pylori–infected individuals and related to MDSCs. Human MDSC markers are defined as CD33^+ HLA-DR^−/lo, and subdivided into monocytic (CD14^+) and granulocytic (CD15^+) subtypes. Since SLFN4 does not exist in the human genome, we analyzed SLFN12L as the human ortholog to the mouse Slfn4 gene based on sequence homology (18) and SLFN5 as the predominant type I IFN–regulated human SLFN gene in human tissues (27, 36). Normal gastric mucosa without H. pylori infection was neg-
ative for both SLFN12L- and SLFN5-expressing immune cells, but SLFN$^+$ cells were observed in \textit{H. pylori}-infected gastric mucosa with intestinal metaplasia (Figure 7A). These immune cells were also positive for the Gr-MDSC marker CD15 in and each patient were near metaplastic glands in at least 2 fields. Further analysis showed that SLFN12L$^-$ cells are CD15$^-$CD33$^+$HLA-DR$^+$, consistent with the Gr-MDSC designation (Figure 7, B and C). Therefore, human SLFN12L expression increases in \textit{H. pylori}-infected individuals with intestinal metaplasia and marks a Gr-MDSC subpopulation.

Discussion

Myeloid cells home from the bone marrow to peripheral organs to fight invading organisms. Infection by \textit{Helicobacter} induces acute and then chronic gastritis, initiating the development of gastric metaplasia in some individuals and cancer in about 1% of those infected (I). Yet it is not well understood what component of the chronic inflammatory milieu provides the impetus to propel the mucosa toward cellular atrophy and metaplasia, ultimately presaging the development of cancer. Petersen et al. recently reported that M2 macrophages promote the advancement of SPEM in the presence of inflammation induced by treatment of mice with DMP-777 and L635 (37). However, with chemical induction, oxyntic gland metaplasia develops very rapidly relative to bacterial pathogenesis and may not necessarily recapitulate the events occurring over a prolonged time course as observed in humans. Here we report that canonical HH signaling plays an essential role in the transition from chronic inflammation to gastric metaplasia (SPEM). Subtyping a population of immune cells responsible for the metaplasia, we found that the GLI1 target gene \textit{Slfn4} marks a subset of CD11b$^+$Gr-1$^+$Ly6G$^+$ Ly6G$^{lo}$ cells, which acquire the ability to suppress T cell proliferation within the infected stomach. Examining the infected gastric mucosa in 2-month intervals over 6 months revealed that induction of mucosal IFN-$\alpha$ correlated with both SPEM development and the emergence of SLFN4$^+$ cells. Recombinant SHH, live \textit{Helicobacter}, or IFN-$\alpha$ treatment of peritoneal-derived myeloid cell cultures induced \textit{Slfn4} mRNA, providing an ex vivo approach to study the process of MDSC development. The data presented here support a novel pathway for metaplastic induction, where HH signaling synergizes with regulatory cytokines to create a permissive environment for neoplastic transformation of the gastric mucosa.

The early phase of gastric cancer pathogenesis, especially with intestinal metaplasia, tends to express high levels of SHH ligand (38). However, the effect of SHH on gastric tumor initiation and growth has not been well documented. \textit{Acute Helicobacter infection} in mice increased SHH blood levels (within 2 days), in agreement with a report by Schumacher et al. that secreted SHH diffuses from the stomach through the circulation and is interpreted by the immune system as a help signal to direct circulating immune cells to the infected stomach (14). Once specific myeloid cells express GLI1 in response to SHH ligand, they become primed to respond to accumulating DAMPs and TLR9 signal transduction, which ultimately stimulates the release of IFN-$\alpha$ into the microenvironment, perhaps with the initial intent to quench the inflammatory response initiated by \textit{Helicobacter} (32).

Since oxyntic glands atrophy during chronic \textit{Helicobacter} infection (39), stressed parietal cells would typically be the optimal candidates to release DAMP signals prior to undergoing apoptosis (30). We found that parietal cells express IFN-$\beta$, a proximal cytokine mediator of DAMP signaling that peaked 2 months after the \textit{H. felis} infection. The time course for mucosal \textit{Ifna} and \textit{Slfn4} mRNA expression was consistent with activation of this regulatory pathway, suggesting that induction of DAMPs and \textit{Tlr9} expression occurs 2 months after infection. We speculate that the lag period between chronic inflammation and metaplasia permits the accumulation of a sufficient amount of DAMPs from dying parietal cells that gradually increase \textit{Tlr9} and type I interferon expression. Indeed, initiation of DAMP signaling coincided with expression of cleaved caspase-3 by parietal cells, marking their entry into an apoptotic phase during this 2-month lag period prior to complete atrophy and their replacement by metaplastic mucous cells. Similarly, a recent study reported that \textit{H. pylori} infection in mice gradually induces \textit{Tlr9} expression in the gastric mucosa over 6 months, enhances neutrophil recruitment, and dampens the proinflammatory cytokine response (28).

This cascade of events leads to induction of IFN-$\alpha$ coincident with the emergence of SLFN4-expressing MDSCs and SPEM. Thus, GLI1-dependent \textit{Slfn4} expression within myeloid cell subpopulations appears to be a reliable marker of the phenotypic transition, since it requires both the epithelial HH ligand and immune-modulatory cytokine IFN-$\alpha$ for peak expression. Type I interferons have not been frequently found to play a role in the proinflammatory response observed in a \textit{Helicobacter} infection (40). Since myeloid cells normally encounter \textit{Helicobacter}-mediated events only in the infected stomach, peak induction of \textit{Slfn4} appears to occur only when tissue IFN-$\alpha$ levels rise above a certain threshold, and could account for the time lag observed prior to the acquisition of the SLFN4$^+$ MDSC phenotype. This transition period represents an important interventional window, since it is well accepted that once metaplasia segue into dysplasia, medical intervention is unlikely to reverse the histologic progression to frank cancer due to the genetic heterogeneity of many epithelial cancers.

Schlafens in immune cells were recently summarized as being a heterogeneous group of genes that potently suppress cell growth, are strongly induced by type I interferons, and primarily exhibit antiviral and antineoplastic effects (35, 41, 42). Yet little is understood regarding epithelial remodeling by infiltrating SLFN-expressing cells. Although we primarily used this GLI1 gene target to probe the role of HH signaling in neoplasia, the current understanding of the function of schlafen family members suggests that they exhibit potent regulatory effects depending on the cell type (43, 44). Indeed, the analysis of natural mutations in both mouse and human schlafen loci suggests that they normally depress the cell cycle (45, 46). Specifically, the \textit{elektra} phenotype in mice represents a missense mutation in \textit{Slfn2} that allows T cells to remain in a semiactivated state, suggesting a role in mediating T cell quiescence (46, 47). We report here that \textit{Slfn4} expression marks a GLI1-dependent population of myeloid cells with the ability to suppress T cell function and phenotypic characteristics of granulocytic MDSCs, consistent with one study showing that gastric cancer patients have elevated levels of MDSCs with a granulocyte-like phenotype (48). Moreover, we showed here that the human homologs SLFN12L and SLFN5 correlated with \textit{H. pylori}...
infection and intestinal metaplasia, and coincided with Gr-MDSC formation after H. pylori infection in mice (53). The difference between that report and our study might be related to the time points they studied (49 days after infection) compared with the extended time line reported here (180 days).

We conclude that Sfn4 is perhaps one of several HH-dependent immune genes expressed at low levels in circulating myeloid cells that becomes primed to respond to a threshold level of specific immune mediators after an acute increase in SHH ligand. In this way, Sfn4-MDSCs would remain quiescent until they engage regulatory signals (HH ligand and type I IFNs) at the appropriate time to dampen the proinflammatory response in the stomach. Presumably, sustained immune dysregulation creates a microenvironment capable of supporting cancer development.

**Methods**

**Transgenic mice.** Generation of the pCMV-Shh transgenic mouse line was described previously, and it was genotyped using the following primer pairs: forward 5′-CTGGCTAACTAGAGAACC; reverse 5′-CTTTTGGATTTCTAGTAGACC (16). Glil-nLacZ reporter mice were a gift from Alex Joyner (Memorial Sloan Kettering Cancer Center, New York, New York, USA). The mice contain a knock-in of the cDNA for the Glil gene, which disrupts expression, creating a null allele. The mice were maintained as heterozygous (Glil+nLacZ) and homozygous (Glil−/−; Glil−/−) mice.

**Generation of the Slfn4-CreER T2 Rosa26-tdTomato reporter mice.** To lineage trace the SLFN4+ cells, Sfn4−eGFP-CreER T2 Rosa26-tdTomato transgenic mice were bred to the Rosa26-LSL-tdT-Tomato reporter mice to generate Sfn4−CreER T2 Rosa26-tdT-Tomato (Sfn4−tdT) mice. The Sfn4−tdT mice and Glil−/− mice were bred to generate Sfn4−CreER T2 Rosa26-tdT-Tomato Glil−/− (Sfn4−tdT Glil−/−) mice. All mice were cohoused under the same specific pathogen–free conditions and fasted overnight prior to use, with free access to water. All mice were maintained on a C57BL/6 genetic background.

**BMT.** The recipient mice (WT, pCMV-Shh, Glil−/−, or Glil−/−) were lethally irradiated using 2 doses of 5 Gy with a 1-hour interval between doses. Bone marrow cells (2 × 10^6) collected from hind leg long bones of 8- to 12-week-old donor mice (Sfn4−tdT mice) were injected into the tail vein of the irradiated recipient mice. After a 6-week engraftment period, chimeric mice were infected with H. felis to determine BMT efficiency. Mice were fasted overnight before euthanasia by CO2 inhalation. The number of CFU per 100 μl was determined by dilution and plating of bone marrow cells on selective plates. The optical density of H. felis was measured, and the final concentration was adjusted to 10^8 organisms per ml of saline. Mice were gavaged 3 times over 5 days with 10^8 H. felis CFU per 100 μl. Control mice were gavaged with 100 μl saline alone.

**Tissue preparation.** Mice were fasted overnight before euthanization. The squamous forestomach was removed prior to bisecting the stomach along the greater and lesser curvatures and then sectioning each half stomach into longitudinal strips. Tissue from the lesser
and greater curvatures was fixed overnight in 4% paraformaldehyde and transferred to 70% ethanol for subsequent paraffin embedding. The remaining corpus tissue was minced for total RNA and protein extraction as well as tissue dissociation in EDTA for flow cytometry.

**Flow cytometry.** Stomach or small intestine was minced, incubated in 20 ml Ca²⁺/Mg²⁺-free HEPES-buffered saline solution (CMF-HBSS) supplemented with 5% FBS (Thermo Fisher Scientific) and 2 mM EDTA (S1201, Lonza) with orbital shaking at 250 rpm for 20 minutes at 37°C. The non-dissociated tissue was removed by passing the 20 ml cell suspension through a 100-μm cell strainer by gravity (Fisher Scientific). The tissue collected in the cell strainer was further digested a second time in 20 ml EDTA media (CMF-HBSS/5% FBS with 2 mM EDTA) at 250 rpm for 20 minutes at 37°C. After a second round of EDTA chelation, the cell suspension was passed through the cell strainer a third time, minced rapidly, and then transferred to 20 ml of collagenase solution (1.5 mg/ml type VIII collagenase dissolved in prewarmed CMF-HBSS/5% FBS with 40 μg/ml DNase I) for digestion at 200 rpm for 10–20 minutes at 37°C. After passing through the 100-μm cell strainer and centrifuging at 400 g for 5 minutes at 4°C, the cell pellet was resuspended in PBS containing 5% BSA. To collect bone marrow cells, the marrow from the femurs was directly flushed with PBS, and red blood cells were lysed in ACK lysis buffer (10-548E, Lonza). Cells were collected at 400 g and resuspended in 200 μl PBS containing 5% BSA (Roche). To collect spleenocytes and hepatocytes, the spleen and liver were triturated with a 1-ml syringe and flushed through a 70-μm cell strainer into a 50-ml conical tube. The strainer was rinsed twice with HBSS to recover all cells. The cells were collected at 300 g for 10 minutes, and the pellet was resuspended in ACK lysis buffer with gentle swirling for 2 minutes to lyse red blood cells. The lysis was stopped by adding 12 ml of HBSS, centrifuging at 300 g for 10 minutes, and then resuspending the cells in PBS containing 5% BSA. Cells (10⁴ in a 100-μl total volume) were blocked with purified rat anti-mouse CD16/CD32 Fc block (BD Biosciences) at 4°C for 5 minutes, prior to incubation with 0.25 μg CD11b-FTTC (BioLegend), Gr-1–APC (BioLegend), Ly6G–APCCy7 (BioLegend), Ly6C–Pacific Blue (BioLegend), CD8α–APC (BioLegend), PDCA-1–PE-Cy7 (eBioscience), rabbit anti-E-cadherin (sc-7870, Santa Cruz Biotechnology Inc.), NOS2–FITC (Santa Cruz Biotechnology Inc.), rabbit anti-arginase I (sc-20150, Santa Cruz Biotechnology Inc.), IFN-α–FITC (Interferon Source). Prior to staining for intracellular proteins, cells were fixed and permeabilized with the Fix and Perm Kit (Thermo Fisher). Prior to initiating the multicolor analysis, we ran the fluorescence minus one (FMO) control to define the spectral overlap. A single-stained control for each fluorochrome was made for compensation for each experiment. A separate set of cells was stained with isotype control antibodies for the negative control. Isotype control antibodies used were PE isotype control (BioLegend, 400608), Pacific Blue isotype control (BioLegend, 400627), PE-Cy7 isotype control (BioLegend, 400618), FITC isotype control (eBioscience, 11-4614-80), APC isotype control (eBioscience, 11-4321-41), APC-Cy7 (BioLegend, 400623). Cell viability was determined with propidium iodide (Abcam, ab14083). Fluorescent-conjugated secondary antibodies goat anti-rabbit IgG APC conjugate (A-10931) and goat anti-rabbit IgG Alexa Fluor 555 conjugate (A27017) were purchased from Life Technologies. The cells were analyzed using an iCyt Synergy SY3200 flow sorter (Sony Biotechnology Inc.).

**Histological analysis.** Five-micrometer sections were prepared, deparaffinized, and then rehydrated. Each slide contained sections of well-oriented gastric glandular mucosa, which extended along the greater curvature of the stomach. Slides were stained with H&E to evaluate the degree of PMN accumulation and metaplasia in the corpus. A veterinary pathologist blinded to the experimental conditions used a semiquantitative scale from 0 to 3 on ×200 histologic fields to score gastric lesions. The absence of inflammatory aggregates, atrophy, or metaplasia was given a score of 0. For PMNs, more than 3 occasional clusters (3 or more contiguous cells) was scored 1, widespread PMN aggregates at the base of glands were scored 2, and aggregates throughout the full mucosal thickness were scored 3. For mucus cell metaplasia (loss of parietal cells with replacement by mucous-type cells), the presence of “foamy” epithelial cells in 1–2 glands was scored 1, foamy cells in most fields but less than the entire field was scored 2, and the entire glandular epithelium replaced by foamy mucus cells was scored 3. The weighted score = \((\text{number of fields with a score} = 1) + (2 \times \text{number of fields with a score} = 2) + (3 \times \text{number of fields with a score} = 3) / (\text{total number of fields} \times 3)\). For each slide, around 12–20 microscopic fields were scored.

Specific protein expression was identified by immunostaining. Antigen retrieval was performed using 10 mM citric acid buffer (pH 6). Slides were washed in 0.01% Triton X-100 in PBS twice, incubated with the serum of the animal in which the secondary antibody was raised for 1 hour. For immunofluorescence staining, slides were incubated with the following antibodies overnight: GSII-FTTC (1:1,000, FL-1211, Vector Laboratories), arginase I (1:100, sc-20150, Santa Cruz Biotechnology Inc.), IFN-β (1:100, AP30401PU-N, Acris Antibodies), IFN-α–FITC (1:200, 22100-3, Interferon Source), H–K–ATPase–β (1:1,000, D032-3, Medical and Biological Laboratories), caspase-3 (1:400, 9661, Cell Signaling Technology), clusterin (1:2,000, sc-6420, Santa Cruz Biotechnology Inc.), E-cadherin (1:1,000, sc-7870, Santa Cruz Biotechnology Inc.), CD15 (1:7,500, 13-0159-1631, ebioscience), SFLN12L (1:100, NB01-91060, Novus). Alexa Fluor–conjugated secondary antibodies (Molecular Probes, Invitrogen) were used to detect primary antibodies at a dilution of 1:500 and then mounted with ProLong Gold Antifade reagent with DAPI (Life Technologies). Species-matched secondary antibodies without primary antibodies were used as the negative controls. Immunofluorescence was visualized using a Nikon Eclipse E800 microscope or Nikon A-1 confocal microscope with NIS acquisition software. For light microscopy, the Vectastain ABC kit (Vector Laboratories) and DAB substrate kit (ab64238, Abcam) were used. Primary antibodies used for DAB staining were SFLN12L (1:100, NB01-91060, Novus), CD15 (1:7,500, 13-0159-1631, ebioscience), HLA-DR (1:50, 340688, BD), CD33 (1:300, 303414, BioLegend), SLFN5 (1:500, ab121537, Abcam). Secondary antibodies used were goat anti-rabbit (1:1,000, 656140, Invitrogen) and donkey anti-mouse (1:500, BA9200, Vector Laboratories).

**T cell suppression assay.** T cells from the mouse spleen were isolated using the EasySep Mouse T Cell Isolation Kit (StemCell Technologies) in which magnetically labeled cells to be excluded are retained by the magnetic field, while unlabeled T cells pass through and were collected for CFSE labeling using the CellTrace CFSE Cell Proliferation Kit (C34554; Molecular Probes). Purified T cells were cultured in RPMI supplemented with 10% FBS, 2 mM l-glutamine, 1 mM sodium pyruvate, 100 mM nonessential amino acids, 5 mM HEPES free acid, 10 ml of 0.05 μM l-mercaptoethanol, and 100 U/ml penicillin/streptomycin with 10% heat-inactivated FBS. To stimulate proliferation, T cells were cultured with anti-CD3/CD28–coated sulfate latex beads in the medium. Suppression of T cell proliferation was assayed after the
addition of SLFN4+ (tdT, red) cells for 3 days at a T cell/SLFN4+ cell ratio of 10:1. SLFN4+ cells were sorted from H. felis-infected mouse stomach, bone marrow, or spleen. To ensure that the EDTA-collagenase isolation procedures used to isolate the gastric SLFN4+ (tdT, red) cells did not differentially induce the gastric cell suppressor function relative to the SLFN4+ (tdT, red) cells, bone marrow cells and splenocytes were prepared in parallel with the gastric tissue. In brief, in order to isolate cells from the bone marrow and spleens to be used in the T cell suppression assay, the bone marrow was flushed from the long bones with the HBSS/FBS EDTA media (as described above), and spleens were triturated and passed the cell strainer into the EDTA media, with shaking for 20 minutes. The chelation step was repeated. The bone marrow and spleen cells were then incubated in the HBSS/FBS collagenase/DNase solution for 20 minutes at 37°C with agitation as described above in detail for the miniced gastric tissue. The CFSE content in T cells was analyzed by flow cytometry in which low CFSE labeling indicated more proliferation and high-intensity CFSE indicated less proliferation. T lymphocytes were cultured with (CD4+CD25+) splenic Tregs at the same 10:1 ratio were used as the positive control for T cell suppressor activity.

Thioglycollate-elicited peritoneal cells. Mice were inoculated intraperitoneally with 1 ml of 3% thioglycollate 2 days prior to isolation of peritoneal myeloid cells. The harvested peritoneal cells were plated onto 6-well plates and allowed to adhere overnight to the plastic. The nonadherent cells were removed by aspiration. Flow cytometry was used to determine that most of the adherent peritoneal cells were positive for CD11b and therefore were designated primary myeloid cells. These cells were treated with 800 U/ml IFN-γ, 800 U/ml IFN-α, 200 ng/ml IFN-β, 200 ng/ml recombinant mouse SHH (rmSHH, R&D Systems), or 107 CFU of H. felis per well (~10 CFU:1 myeloid cell) for 12 hours prior to analysis. Primary myeloid cells prepared from the Sfn4-tdT mice were cultured in DMEM media with 10% FBS at 37°C and treated with 100 nM Tx (dissolved in DMSO) for 24 hours to induce Cre recombinase activity ex vivo.

Real-time quantitative PCR. Total RNA from the corpus was extracted from freshly collected tissue in TRIzol (Invitrogen). The number of mice per group and replicate experiments are indicated in the figure legends.

Statistics. For all ELISA and qPCR experiments, statistical analysis for significance was performed on the log-transformed values using 1-way ANOVA with Tukey’s post hoc test for multiple comparisons. Cell subpopulation computed as percentages and the weighted scores used in the morphometric analysis were analyzed using Kruskal-Wallis 1-way ANOVA with Dunn’s test for multiple comparisons (GraphPad Prism). All data were expressed as the median with the interquartile range. P values less than 0.05 were considered statistically significant. The number of mice per group and replicate experiments are indicated in the figure legends.

Study approval. Animal use was approved by the University of Michigan Institutional Animal Care and Use Committee. The human tissue samples used in this study were deidentified archived specimens that do not require IRB regulation, as reviewed by the University of Michigan Medical School Institutional Review Board (IRBMED; ID: HUM00108090).

Author contributions. LD designed and conducted the experiments, analyzed the results, and drafted and edited the manuscript. MHH, KAE, QL, and ROR acquired data. AP performed immunohistochemical analysis. JLM designed the experiments, analyzed results, and wrote the manuscript.

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42. Derbinski J, Schulte A, Kyeswski B, Klein L. Pillars


