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Suzuki, Yasuhiro, Shuji Kitahara, Takuya Suematsu, Masanobu Oshima, and Yasufumi Sato. 2017. "Requisite role of vasohibin2 in spontaneous gastric cancer formation and accumulation of cancer-associated fibroblasts." Cancer Science 108 (12): 2342-2351. doi:10.1111/cas.13411. http://dx.doi.org/10.1111/cas.13411.

Published Version
doi:10.1111/cas.13411

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Requisite role of vasohibin-2 in spontaneous gastric cancer formation and accumulation of cancer-associated fibroblasts

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Key words
Cancer-associated fibroblasts, epiregulin, gastric cancer, interleukin-11, vasohibin-2

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Funding Information
Japan Society for the Promotion of Science (Grant Number: 15K06821, 24501309).

Received May 1, 2017; Revised August 30, 2017; Accepted September 18, 2017

Cancer Sci 108 (2017) 2342–2351

doi: 10.1111/cas.13411

The vasohibin (VASH) family consists of two genes, VASH1 and VASH2. VASH1 is mainly expressed in vascular endothelial cells and suppresses angiogenesis in an autocrine manner, whereas VASH2 is mainly expressed in cancer cells and exhibits pro-angiogenic activity. Employing adenomatous polyposis coli gene mutant mice, we recently reported on the role of Vash2 in the spontaneous formation of intestinal tumors. In this study, we used K19-Wnt1/C2mE (Gan) mice and examined the role of Vash2 in spontaneous gastric cancer formation. Gan mice spontaneously develop gastric tumors by activation of Wnt and prostaglandin E2 signaling pathways in gastric mucosa after 30 weeks of age. Expression of Vash2 mRNA was significantly increased in gastric tumor tissues compared with normal stomach tissues. When Gan mice were crossed with the Vash2-deficient (Vash2lacZlacZ) strain, gastric cancer formation was significantly suppressed in Vash2lacZlacZ Gan mice. Normal composition of gastric mucosa was partially maintained in Vash2lacZlacZ Gan mice. Knockout of Vash2 caused minimal reduction of tumor angiogenesis but a significant decrease in cancer-associated fibroblasts (CAF) in tumor stroma. DNA microarray analysis and real-time RT-PCR showed that mRNA levels of epiregulin (Ereg) and interleukin-11 (Il11) were significantly downregulated in gastric tumors of Vash2lacZlacZ Gan mice. Furthermore, conditioned medium of gastric cancer cells stimulated migration of and α-smooth muscle actin expression in fibroblasts, whereas conditioned medium of VASH2 knockdown cells attenuated these effects in vitro. These results suggest that VASH2 plays an important role in gastric tumor progression via the accumulation of CAF accompanying upregulation of EREG and IL-11 expression.

Gastric cancer is the third leading cause of cancer-related death in Japan. Risk factors for gastric cancer include Helicobacter pylori infection, host genetic susceptibility and other environmental factors, and multiple genetic mutations, epigenetic alterations and aberrant molecular signaling pathways are involved in its development. (1)

The treatment of gastric cancer has evolved in the past three years. Molecular characterization of gastric cancer has provided clues for treatment development, and the introduction of agents targeting human epidermal growth factor (EGF) receptor 2 and vascular endothelial growth factor (VEGF) family/VEGF receptor has brought this disease into the era of molecular and personalized medicine. (2)

We searched for and isolated novel angiogenesis regulators that we have designated as vasohibin-1 (VASH1) and vasohibin-2 (VASH2). VASH1 is an endothelium-derived angiogenesis inhibitor, (3) whereas VASH2 is a homologue of VASH1 that acts as an angiogenesis stimulator. (4,5) These two regulators are devoid of the classical secretory signal sequence but are efficiently secreted when they form a complex with small vasohibin-binding protein. (6,7) Of note, VASH2 is produced by cancer cells such as ovarian carcinoma, hepatocellular carcinoma and intestinal adenocarcinoma, and promotes tumor growth by stimulating tumor angiogenesis. (8–10) Using xenograft models of human ovarian cancer, we recently reported that targeting of VASH2 by exogenous administration of siRNA or a neutralizing antibody specific for VASH2 inhibits tumor growth by attenuating tumor angiogenesis. (11,12) Furthermore, when an experimental model for spontaneous adenomatous polyposis, the adenomatous polyposis coli multiple intestinal neoplasia (ApcMin/+ ) mouse, is crossed with the Vash2lacZlacZ mouse, the number of intestinal tumors significantly decreases in association with normalization of tumor vessels. (10) The expression of VASH2 has also been documented in human gastric cancer cell lines, (13) but the significance of VASH2 in gastric cancer development is not well defined.

Spontaneous carcinogenic mouse models are useful for gaining insight into the pathogenesis and molecular mechanisms of cancers. The K19-Wnt1/C2mE (Gan) for Gastric neoplasia mouse was recently established by the transgenic expression of...
Wnt1, cyclooxygenase-2 and microsomal prostaglandin E synthase-1 under a Keratin 19 promoter in gastric epithelial cells, recapitulating human gastric tumors not only in terms of molecular mechanism but also tumor pathology.\(^{(14)}\) Here we used Gan mice, crossed with Vash\(_2\)lacZlacZ mice, and examined the significance of VASH2 in gastric tumor growth and the tumor stromal microenvironment.

Materials and Methods

Mouse models. As noted, Gan mice highly express Wnt1, cyclooxygenase-2 and microsomal prostaglandin E synthase-1 under Keratin 19 promoter activity in gastric epithelial cells and generate gastric tumors after 30 weeks of birth, as previously reported.\(^{(14)}\) Vash\(_2\)lacZlacZ mice were maintained as described earlier,\(^{(4)}\) and Gan mice were mated with Vash\(_2\)lacZlacZ mice to generate Vash\(_2\)lacZlacZ/Gan mice. For the tumor development analyses, mice were euthanized and examined at 30 weeks of age. All animal experiments were carried out according to the protocol approved by the Committee on Animal Experimentation of Tohoku University, Japan.

Histological and immunohistochemical analyses. Gan mice at age 30 weeks were anesthetized and transcardially perfused with PBS followed by 4% paraformaldehyde. Stomachs were extracted and incubated overnight, dehydrated in graded ethanol and xylene, and embedded in paraffin wax. Vertical sections (5 μm) of the entire gastric mucosa were prepared for conventional H&E staining or immunohistochemical staining. For immunohistochemical staining, tissue sections were autoclaved in citrate buffer (pH 6.0) for 5 min for antigen retrieval prior to incubation with primary antibodies. Antibodies against the H\(^+\)K\(^+\)-ATPase (MBL, Nagoya, Japan), F4/80 (Serotec, Oxford, UK), Keratin 19 (Nobori Biomedical, Nagoya, Japan) and SMA; Sigma, St. Louis, MO, USA) were used as the primary antibodies. Staining signals were visualized using HistoGene Simple Stain MAX PO (Nichirei, Tokyo, Japan) followed by counterstaining with hematoxylin. Microphotographs were obtained using an MC120 HD camera attached to a Leica DM 2000 LED microscope (Leica Microsystems K.K., Tokyo, Japan). For fluorescence immunostaining, antibodies conjugated with Alexa Fluor 488 or Alexa Fluor 555 (Thermo Fisher Scientific, Waltham, MA, USA) were used as the secondary antibodies. Nuclei were then counterstained using ToPro-3 (Thermo Fisher Scientific). Fluorescence images were captured using a Fluoview FV1000 confocal microscope system (Olympus, Tokyo, Japan).

Measurement of mucosal thickness. The mucosal thickness of gastric tumors was measured from micrographs of H&E-stained vertical sections of entire gastric tumors using LAS Software (Leica Microsystems K.K.). Relative mucosal thickness of samples from Vash\(_2\)lacZlacZ/Gan mice was calculated in comparison with the mean of the control Gan mice.

In situ hybridization. In situ hybridization was performed with the ISH Reagent Kit (Genostaff, Tokyo, Japan) according to the manufacturer’s instructions. Tissue sections were de-paraffinized with G-Nox and rehydrated through an ethanol series and PBS. The sections were fixed with 10% NBF (10% formalin in PBS) for 15 min at room temperature (RT) and washed in PBS; treated with 4 μg/mL Proteinase K (Wako Pure Chemical, Osaka, Japan) in PBS for 10 min at 37°C and washed in PBS; re-fixed with 10% NBF for 15 min at RT and washed in PBS; placed in 0.2 N HCl for 10 min at RT and washed in PBS; and placed within a Coplin jar containing 1× G-Wash (Genostaff), equal to 1× SSC. Hybridization was performed with probes corresponding to the nucleotide positions 1321–2001 of mouse Vash2 mRNA (GenBank accession number NM_144879.2) at concentrations of 300 ng/mL in G-Hybo-L (Genostaff) for 16 h at 60°C. After hybridization, the sections were washed in 1× G-Wash for 10 min at 60°C and 50% formamide in 1× G-Wash for 10 min at 60°C.

Sections then were washed twice in 1× G-Wash for 10 min at 60°C, twice in 0.1× G-Wash for 10 min at 60°C, and twice in TBST (0.1% Tween20 in TBS) at RT. After treatment with 1× G-Block (Genostaff) for 15 min at RT, the sections were incubated with anti-digoxigenin AP conjugate (Roche Diagnostics, Mannheim, Germany) diluted 1:2000 with ×50G-Block (Genostaff) in TBST for 1 h at RT. The sections were washed twice in TBST and then incubated in 100 mM NaCl, 50 mM MgCl\(_2\), 0.1% Tween20, 100 mM Tris-HCl, pH 9.5. Coloring reactions were performed with NBT/BCIP solution (Sigma-Aldrich, St. Louis, MO, USA) overnight and then washed in PBS. The sections were counterstained with Kernechtrot stain solution (Muto Pure Chemicals, Tokyo, Japan) and mounted with G-Mount.

RT-PCR and quantitative real-time RT-PCR. Total RNA was extracted from mouse tissues and cultured cells using QIAzol Lysis Reagent (QIAGEN, Hilden, Germany) and purified using the RNeasy Mini Kit (QIAGEN). First-strand cDNA was synthesized by reverse transcriptase using RevertAse Ace (TOYOBO, Osaka, Japan). PCR was performed using sets of primers specific for the target genes described below. Thermal cycler conditions consisted of an initial denaturation step at 95°C for 2 min, followed by 40 cycles of 10 s at 95°C, 10 s at 56°C and 30 s at 72°C. Relative mRNA levels of target genes were normalized to beta-2-microglobulin (B2m) mRNA level. The specific primer pairs for target genes are summarized in Table S1. Specific primer pairs for detection of CD44 splicing variant (CD44v) targeted exons 5 and 16 of CD44 mRNA.\(^{(15)}\)

Microarray analysis. Total RNA was extracted from gastric tumor tissues as described above. The quality of total RNA was confirmed by an Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA). The cRNA was synthesized and labeled with Cy3 using the Low Input Quick-Amp Labeling Kit (Agilent Technologies). The Cy3-labeled cRNA samples were subjected to hybridization to SurePrint G3 Mouse GE Microarray chips (Agilent Technologies) using a Gene Expression Hybridization Kit. Fluorescence signals in the chips were detected by an Agilent microarray scanner (Agilent Technologies) and quantified using Feature Extraction software (Agilent Technologies).

Cell cultures. FU97 cells (a human stomach cancer cell line) and SF-TY cells (a human skin fibroblast cell) were obtained from the Japanese Collection of Research Bioresources Cell Bank (Osaka, Japan). FU97 cells were cultured in DMEM (Wako Pure Chemical) supplemented with 10% heat-inactivated FBS (SAFC Biosciences, Lenexa, KS, USA) and 10 μg/mL of recombinant human insulin (Wako Pure Chemical). SF-TY cells were maintained in DMEM supplemented with 10% FBS and non-essential amino acids.
Knockdown of VASH2 by shRNA. FU97 cells were transfected with the VASH2 shRNA expression vector or its control mock vector using FuGENE HD Transfection Reagent (Promega, Madison, WI, USA) and were selected in culture medium containing 10 μg/mL puromycin (Wako Pure Chemical), as reported previously. (8) Fibroblast proliferation and migration. FU97 cells and VASH2 knockout (shVASH2) clones were plated in 60-mm dishes at 5 × 10^3 cells and cultured overnight in the culture medium. The following day, the medium was replaced by DMEM containing 0.5% FBS. The conditioned medium was collected 48 h later and filtered through a MILLEX-GP PES 0.22-μm filter (Millipore, Bedford, MA, USA). Cell proliferation was measured using a BrdU Cell Proliferation ELISA Kit (Abcam). Cells were plated in a 96-well plate at 5 × 10^3 cells per well and starved in DMEM containing 0.5% FBS for 16 h. Cells were then treated with conditioned media (CM) from FU97 cells or shVASH2 clones and labeled with BrdU for 24 h. Incubated BrdU was detected according to the manufacturer’s instructions. Migratory activity of fibroblasts was measured by modified Boyden chamber assay. (6) SF-TY cells that migrated across the membrane were fixed with methanol, or shVASH2 clones. After incubation for 4 h, SF-TY cells that migrated across the membrane were fixed with methanol, stained with DAPI (Sigma-Aldrich), and counted in nine fields per insert in a blinded manner.

Statistical analysis. Data are expressed as mean ± SD and were analyzed using unpaired Student’s t-tests. A value of P < 0.05 was considered to denote statistical significance.

Results

Increased expression of VASH2 in gastric tumor cells in Gan mice. Previous reports demonstrated that VASH2 is highly expressed in various types of human tumor tissues as well as in established cancer cell lines. (8,9,13,16–20) Shen et al. (13) reported that both human gastric cancer cell lines and experimentally prepared tumor-associated macrophages express VASH2 mRNA in an in vitro culture system. However, the roles of VASH2 in spontaneous gastric tumor growth are not fully understood. Gan mice spontaneously generate gastric tumors by simultaneous signal activation of both Wnt and prostaglandin E2 (PGE2) in gastric epithelia under keratin-14 promoter activity. (21–27) We first examined the expression level of Vash2 mRNA in gastric tumors spontaneously developed in Gan mice at 30 weeks of age. As shown in Figure 1(a), RT-PCR analysis showed enhanced expression of CD44 splicing variants (CD44v) and Vash2 in gastric tumors of Gan mice, whereas normal stomach tissues expressed low levels of Vash2 and no CD44v, as previously reported. (15,28) According to CD44v mRNA expression, increased expression of Vash2 mRNA was detected in Gan mice gastric tumors (Fig. 1a). Real-time RT-PCR analysis showed that the expression level of Vash2 mRNA in Gan mouse gastric tumors was approximately three times higher than in normal stomachs of wild-type mice (Fig. 1b).

We then performed in situ hybridization for mouse Vash2 transcript in tissue sections of Gan mouse gastric tumors at 30 weeks of age. Signals for Vash2 mRNA transcripts visualized by antisense probes were detected in gastric tumors of Gan mice but not in normal stomach (Fig. 1c). As shown in Figure 1(d), the signals were mainly localized in dysplastic epithelial cells (black arrowheads), whereas they were hardly observed in other stromal cells (white arrowheads). These results indicate that dysplastic epithelial cells express Vash2 mRNA in Gan mice.

Suppression of gastric tumor growth in Vash2-deficient Gan mice. To investigate the role of VASH2 in spontaneous gastric tumor development, we bred Gan mice with Vash2-deficient mice (Vash2LacZ/LacZ) to obtain Vash2-deficient Gan mice (Vash2LacZ/LacZ Gan) and examined the phenotypic differences...
pressed spontaneous gastric cancer development. These results indicate that the lack of the Vash2 gene suppression of gastric tumor growth by knockout of the Vash2 gene. (a) Representative macroscopic images of wild-type normal stomach and Gan and Vash2LacZ/LacZ Gan mouse gastric tumors at 30 weeks of age. (b) Representative images of H&E staining of Gan mouse (top) and Vash2LacZ/LacZ Gan mouse (bottom) gastric tumors at 30 weeks of age. Scale bar: 2 mm. (c) Relative gastric mucosal thickness of Vash2LacZ/LacZ Gan mouse gastric tumors (n = 11) to the mean level of Gan mouse tumors (n = 12) at 30 weeks of age. Each circle indicates the relative tumor thickness of individual mice. **P < 0.01 versus Gan mouse tumors.

in gastric tumors developed in those mice. As shown in Figure 2(a), development of gastric tumors was observed in Gan mice at 30 weeks of age, as reported previously. Of note, gastric tumor growth was obviously suppressed in Vash2LacZ/LacZ Gan mice compared with that in Gan mice (Fig. 2a,b). The mean of histologic tumor thickness was significantly reduced by knockout of Vash2 (Fig. 2b,c).

Immunohistochemical staining for Ki-67 antigen in paraffin sections of those gastric tumor tissues showed that Ki-67-positive proliferating cells were frequently observed in the entire gastric dysplastic epithelia of Gan mice tumors, whereas most proliferating cells were limited to the proliferating zone at the gland neck area in gastric tumors of Vash2LacZ/LacZ Gan mice (Fig. 3a,c). To confirm the distribution of gastric parietal cells in gastric tumors, we performed immunostaining with antibody for the proton pump (H+, K+-ATPase), used as a marker of parietal cells. As shown in the center panel of Figure 3(b), a small number of gastric parietal cells was observed in Gan mouse gastric tumors (Fig. 3d), indicating an architectural disorder of gastric tumor tissues, as reported previously. However, a numerous parietal cell population was still maintained in gastric tumors of Vash2LacZ/LacZ Gan mice (Fig. 3b,d). These results indicate that the lack of the Vash2 gene suppressed spontaneous gastric cancer development.

Effect of Vash2 on stromal activation in gastric cancer. The tumor stromal microenvironment (e.g. tumor angiogenesis, expansion of cancer associated fibroblasts (CAF) and infiltration of immune cells) plays crucial roles in tumor development and progression. Therefore, we examined whether Vash2 deficiency might affect the stromal microenvironment in gastric tumors. Immunofluorescence staining of tissue sections of Gan mouse gastric tumors with anti-CD31 antibody showed that a large number of capillary blood vessels are generated in these tumors (Fig. 4a, left panel). In comparison with Gan mice, slightly fewer capillaries were observed in gastric tumors of Vash2LacZ/LacZ Gan mice (Fig. 4a, right panel), but overall this difference was not statistically significant (Fig. 4b). Tumor blood vessels are typically fragile and leaky because of their immature structure accompanied by poor pericyte coverage. However, Vash2 deficiency did not affect pericyte coverage in the Gan mouse model (data not shown). Inflammatory responses via infiltrated monocytes/macrophages also have a critical effect on spontaneous gastric tumor growth. Immunohistochemical staining for the F4/80 antigen, a marker specific for mouse macrophages, revealed numerous macrophages infiltrating into gastric tumor stroma in Vash2LacZ/LacZ Gan and Gan mouse tumors (Fig. 4c,d), suggesting that Vash2 disruption might not affect macrophage infiltration in this mouse model.

Next, we evaluated the expansion of CAF in gastric tumor stroma by immunostaining for αSMA and VIM. Numerous VIM-positive and/or αSMA-positive CAF were detected in entire Gan mouse tumor tissues, whereas CAF were sparsely observed in Vash2LacZ/LacZ Gan tumors (Fig. 5a,b). Immunofluorescence staining for another CAF marker, Tenascin C (TNC), also showed that Vash2 deficiency significantly reduced the stromal expansion of TNC-positive CAF (Fig. 5c,d) compared with Gan mice. These results suggested that VASH2 produced by tumor cells might accelerate the expansion of CAF into tumor stroma in addition to its pro-angiogenic activity.

Downregulation of expression of Epiregulin and interleukin-11 by Vash2 depletion in gastrointestinal tumors. To identify genes with expression changes in gastric tumor tissues by knockout of the Vash2 gene, we comprehensively compared mRNA expression profiles between Gan mouse gastric tumors and Vash2LacZ/LacZ Gan mouse gastric tumors by a DNA...
mRNA levels of genes 3(a) and 4(a), respectively. A significant reduction in angiogenesis caused by Vash2 with the suppression of cancer cell proliferation and tumor growth through production of IL-6 family and EGF. Functional classification by Gene Ontology terms showed that upregulated genes were enriched for circadian rhythmic processes corresponding to Ontology terms. Downregulated genes were classified to many functional categories, including regulation of gene expression, biosynthesis, and defense response. Downregulation of stem cell/cancer stem cell markers (CD44 and CD133) and vascular endothelial cell markers (Edn1 and Tie2) correlated with the suppression of cancer cell proliferation and tumor angiogenesis caused by Vash2 disruption, as shown in Figures 3(a) and 4(a), respectively. A significant reduction in mRNA levels of CD44 and CD133 was confirmed by real-time RT-PCR analysis (Fig. 6b), whereas Vash2 deficiency did not significantly change mRNA levels of vascular endothelial cell markers and typical angiogenic factors, such as Vegfa and fibroblast growth factor 2 (Fig. 5b) (Fig. 6c,d). Here, we focused on interleukin-11 (IL-11, a member of the IL-6 family) and Epiregulin (Ereg, a member of the EGF family), listed in Table S3 as downregulated genes, because previous reports suggested that CAF might promote gastrointestinal tumor growth through production of IL-6 family and EGF. By real-time RT-PCR analysis we confirmed that the mRNA expression levels of Ereg, Il11 and Il6 were significantly reduced in Vash2LacZ/LacZ Gan mouse tumors compared to Gan mouse tumors (Fig. 6e).

Using ApcMin+ mice, we recently reported the role of VASH2 in the spontaneous formation of intestinal tumors. The Vash2-deficient ApcMin+ mice (Vash2LacZ/LacZ ApcMin+ ApcMin+) showed a reduced number of polyps alongside the attenuation of tumor angiogenesis as compared to ApcMin+ mice. We next carried out a DNA microarray analysis and identified 17 upregulated genes and 19 downregulated genes in intestinal polyps of Vash2LacZ/LacZ ApcMin+ ApcMin+ mice compared with ApcMin+ mice (Tables S4 and S5). Comparative analysis of mRNA expression profiles obtained from Gan mouse gastric tumors and ApcMin+ intestinal polyps revealed that three genes (Ereg, Il11 and Tmem190) were commonly downregulated in gastric tumors and intestinal polyps of Vash2-deficient mice (Fig. 5a). There was no overlap of upregulated genes (data not shown). Indeed, Ereg and Il11 mRNA was significantly downregulated in polyps of Vash2LacZ/LacZ ApcMin+ ApcMin+ mice compared to ApcMin+ mice (Fig. 5b), suggesting that VASH2 promotes expression of EREG and IL-11 in gastrointestinal tumorigenesis. Suppression of fibroblast activation by knockdown of VASH2 in gastric cancer cells in vitro. We finally examined whether VASH2 expressed in human gastric cancer cells affects fibroblast expansion in vitro. The expression of VASH2 mRNA in human gastric cancer cell line FU97 was significantly higher.
Fig. 4. Effect of Vash2 depletion on tumor angiogenesis and macrophage infiltration. (a) Representative images of immunohistochemical staining for CD31 (green) of Gan mouse (left) and Vash2LacZ/LacZ Gan mouse (right) gastric tumors at 30 weeks of age. Nuclei were stained with TOPRO-3 (blue). Scale bars: 200 μm. (b) Quantification of the percentage of CD31-positive vascular vessel area to total tumor area (n = 8 fields analyzed from three independent animals). (c) Representative images of immunohistochemical staining for F4/80 of Gan mouse (left) and Vash2LacZ/LacZ Gan mouse (right) gastric tumors at 30 weeks of age. Counterstaining of nuclei was performed with hematoxylin. Scale bars: 200 μm. (d) Quantification of the percentage of F4/80 positive area to total tumor area (n = 12 fields analyzed from three independent animals). All quantitative data show mean ± SD. *P < 0.05 versus Gan mouse tumors. N.S., not significant.

Fig. 5. Suppression of cancer-associated fibroblast expansion in gastric tumor stroma by Vash2 knockout. (a) Representative images of a double-fluorescent immunostaining for αSMA and VIM of Gan and Vash2LacZ/LacZ Gan mouse gastric tumors at 30 weeks of age. Nuclei were stained with TOPRO-3 (blue). Scale bars: 200 μm. Inset, high magnification. (b) Quantification of the percentages of αSMA-positive and VIM-positive areas to total gastric tumor area (n = 6 fields analyzed from three independent animals). (c) Representative images of a fluorescent immunostaining for TNC of Gan and Vash2LacZ/LacZ Gan mouse gastric tumors at 30 weeks of age. Nuclei were stained with TOPRO-3 (blue). Scale bars: 200 μm. (d) Quantification of the percentage of TNC-positive area to total gastric tumor area (n = 6 fields analyzed from three independent animals). All quantitative data show mean ± SD. *P < 0.05 versus Gan mouse tumors. N.S., not significant.
than that in human fibroblast SF-TY cells (Fig. 7a). We isolated VASH2 knockdown cell lines (shVASH2-1 and -2) using shRNA (Fig. 7b) and performed a loss-of-function experiment in vitro. CM of parent FU97 cells and mock control cells stimulated migration of and ACTA2 (αSMA) expression in SF-TY cells (Fig. 7c,d). Importantly, CM of shVASH2 cells significantly attenuated the stimulatory effects. There was no significant effect on proliferation of SF-TY cells (Fig. 7c).

These results suggested that VASH2 produced by cancer cells influences CAF expansion and gastrointestinal tumor growth through upregulation of EREG and IL-11 gene expression, in addition to its proangiogenic activity (Fig. 8).

Discussion

Previous studies have reported that VASH2 produced from cancer cells promotes tumor growth by stimulating tumor angiogenesis.\(^{8,9}\) VASH2 is secreted from cancer cells and acts on neighboring vascular endothelial cells as a paracrine angiogenesis stimulator. Most studies on the role of VASH2 in tumor growth have used allograft and xenograft models of established cancer cell lines. In this work, we employed Gan mice as a spontaneous gastric tumor developmental model and confirmed that knockout of Vash2 in Gan mice attenuates gastric tumor growth with a slight reduction in tumor angiogenesis. In addition to this proangiogenic activity, we found a novel role for VASH2 in the tumor stromal microenvironment, namely CAF. The increase in CAF is one of the characteristic features of gastric cancer in Gan mice.\(^{19}\) Vash2 depletion dramatically reduced the expansion of CAF in gastric tumor stroma along with downregulation of Ereg, Il6 and Il11 expression. We also confirmed that the expression of circadian genes (Dbp, Hlf, Per3 and Tef) and of cancer stem cell markers (CD44 and CD133) is upregulated and downregulated, respectively, in Vash2\(^{lacZ/lacZ}\) Gan mouse tumors. These changes might reflect suppression of tumor progression by knockout of Vash2 because previous reports have shown downregulation of circadian genes in various tumors.\(^{33–35}\)

Using the Apc\(^{Min/+}\) mouse model, we recently demonstrated that Vash2 depletion causes a reduced number of polyps alongside attenuation of tumor angiogenesis. Here, we also confirmed the reduced expression of Ereg and Il11 in intestinal polyps of Vash2-deficient Apc\(^{Min/+}\) mice as well as gastric tumors of Vash2-deficient Gan mice. EREG and IL-11 play a crucial role in gastrointestinal tumor growth, progression and chemoprevention through crosstalk between EGFR and gp130 signaling.\(^{24,32,36–39}\) Enhanced expression of the Ereg gene has also been found in gastric cancer cells.\(^{40}\) Oshima et al. report...
PGE2 pathway-dependent Ereg expression by gastric epithelial cells and macrophages in Gan mouse gastric tumors. Of importance, other recent reports have suggested that CAF as well as cancer cells are a major supplier of Ereg and IL-11 in tumor stroma. In this study, we showed that VASH2 disruption influences CAF expansion but not a population of macrophages infiltrating into the gastric tumor stroma. Therefore, it is possible that VASH2 stimulates the expression of EREG and IL-11 in CAF and cancer cells.

Cancer associated fibroblasts in Gan mice produce VEGFA and promote tumor angiogenesis; however, our data showed that Vash2 depletion did not affect Vegfa mRNA level despite reduced CAF expansion. We performed double immunostaining for αSMA and VIM and observed three types of CAF subpopulations (αSMA-positive, VIM-positive and double-positive cells), suggesting that CAF are a heterogeneous population in the tumor microenvironment. Ohlund et al. recently identified distinct populations of inflammatory and myofibroblastic CAF in pancreatic cancer. The inflammatory CAF had low αSMA expression and concomitantly produced inflammatory factors, such as IL-6, IL-11 and leukemia inhibitory factor. Therefore, it is possible that VASH2 may specifically affect certain subtypes of CAF that express higher levels of IL-11 and/or EREG than other subtypes.

Fig. 7. Knockdown of VASH2 in gastric cancer cells suppresses fibroblast activation in vitro. (a) VASH2 mRNA level in FU97 cells relative to that in SF-TY cells was determined by real-time RT-PCR analysis. The VASH2 mRNA expression levels were normalized to β-actin mRNA level. *P < 0.05 versus SF-TY cell. (b) VASH2 mRNA level in VASH2 knockdown clones and control mock transfectant relative to that in parental FU97 cells was determined by real-time RT-PCR analysis. *P < 0.05 versus parental FU97 cell. (c) Effect of conditioned media (CM) of gastric cancer cells on fibroblast migration and proliferation. SF-TY cells were treated with CM of indicated gastric cancer cells. Migration and proliferation of SF-TY cells were analyzed by a modified Boyden chamber assay and BrdU incorporation assay, respectively. *P < 0.05 versus mock CM. (d) Effect of CM of gastric cancer cells on ACTA2 mRNA expression in fibroblasts. The ACTA2 mRNA levels in SF-TY cells treated with CM of indicated gastric cancer cells for 48 h were examined by real-time RT-PCR. *P < 0.05 versus mock CM. All quantitative data show mean ± SD. All experiments were performed at least twice.

Fig. 8. Roles of VASH2 in gastric tumor growth. Schematic illustration of the roles of VASH2 in gastric tumor growth. VASH2 is produced from cancer cells and acts on neighboring EC as a proangiogenic factor in a paracrine manner. In contrast, VASH2 influences cancer cell proliferation and stromal activation, such as CAF expansion, by upregulating the expression of EREG, IL-11 and IL-6.
The authors have no conflict of interest to declare.

Acknowledgments

We wish to thank Dr. Fujinoya for excellent technical assistance. This study was supported by JSPS KAKENHI Grant Numbers 24501309 and 15K06821.

Disclosure Statement

Cancer Cell | December 2017 | vol. 108 | no. 12 | 2350

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42 Öhlund D, Handly-Santana A, Biffi G et al. Distinct populations of inflammatory fibroblasts and myofibroblasts in pancreatic cancer. J Exp Med 2017; 214: 579–96.
43 Kalluri R. The biology and function of fibroblasts in cancer. Nat Rev Cancer 2016; 16: 582–98.
44 Li H, Courtois ET, Sengupta D et al. Reference component analysis of single-cell transcriptomes elucidates cellular heterogeneity in human colorectal tumors. Nat Genet 2017; 49: 708–18.
45 Norita R, Suzuki Y, Furutani Y et al. Vasohibin-2 is required for epithelial–mesenchymal transition of ovarian cancer cells by modulating TGF-β signaling. Cancer Sci 2017; 108: 419–26.

Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

Fig. S1. Vash2 disruption commonly downregulates Ereg and Il11 genes in mouse gastric tumors and intestinal polyps.
Table S1. Primers used for RT-PCR and real-time RT-PCR analysis.
Table S2. Genes upregulated in gastric tumors of Vash2-deficient Gan mice.
Table S3. Genes downregulated in gastric tumors of Vash2-deficient Gan mice.
Table S4. Genes upregulated in intestinal polyps of Vash2-deficient Apc<sup>Min/+</sup> mice.
Table S5. Genes downregulated in intestinal polyps of Vash2-deficient Apc<sup>Min/+</sup> mice.