Cystatin SN Upregulation in Patients with Seasonal Allergic Rhinitis

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Abstract

Seasonal allergic rhinitis (SAR) to the Japanese cedar, Cryptomeria japonica (JC) pollen is an IgE-mediated type I allergy affecting nasal mucosa. However, the molecular events underlying its development remain unclear. We sought to identify SAR-associated altered gene expression in nasal epithelial cells during natural exposure to JC pollen. We recruited study participants in 2009 and 2010 and collected nasal epithelial cells between February and April, which is the period of natural pollen dispersal. Fifteen patients with SAR-JC and 13 control subjects were enrolled in 2009, and 17 SAR-JC patients, 13 sensitized asymptomatic subjects (Sensitized), and 15 control subjects were enrolled in 2010. Total RNA was extracted from nasal epithelial cells and 8 SAR-JC patients and 6 control subjects in 2009 were subjected to microarray analysis with the Illumina HumanRef-8 Expression BeadChip platform. Allergen-stimulated histamine release was examined in the peripheral blood basophils isolated from patients with SAR. We identified 32 genes with significantly altered expression during allergen exposure. One of these, CST1 encodes the cysteine protease inhibitor, cystatin SN. CST1 expression in nasal epithelial cells was significantly upregulated in both the 2009 and 2010 SAR-JC groups compared with the control groups. Immunohistochemical staining confirmed the increased expression of CST1 in the nasal epithelial cells of SAR patients. Addition of exogenous CST1 to basophils inhibited JC allergen-stimulated histamine release in vitro. We propose that CST1 may contribute to inactivation of protease allergens and help re-establish homeostasis of the nasal membranes.

Introduction

Allergic diseases such as asthma and allergic rhinitis are major causes of morbidity in developed countries, and are increasing in frequency. In Japan, the prevalence of rhinitis was 16% in 1992 and 21% in 2002 [1]. Seasonal allergic rhinitis (SAR) due to the Japanese cedar (JC) Cryptomeria japonica pollen is an IgE-mediated type I allergy that affects the nasal mucosa and causes sneezing, itchiness, and nasal obstruction. It is one of the most common causes of allergy in Japan and is thus a major public health issue [2]. We have recently reported that the prevalence of allergic rhinitis in Japanese adults is 44.2%, and the most common allergen is the JC pollen [3]. Although all patients with SAR are positive for allergen-specific IgE, it is well known that the presence of specific IgE alone is insufficient to develop SAR. Therefore, there is a gap in our understanding of the events that link sensitization to the development of the disease. Two different basic theories on the pathogenesis of allergy have been suggested. One is that allergy is caused by unbalanced and overactive immunological responses against allergens, mostly driven by infiltration of type 2 helper T lymphocytes (Th2), mast cells, and eosinophils into the site of allergic diseases. The other more recent hypothesis is dysregulation of the epithelial barrier, so called “epithelial barrier dysfunction hypothesis,” that is, dysfunction and disruption of the epithelial barrier has been suggested for the primary defects in allergic diseases. Disruption of the skin barrier by filaggrin null mutations has been extensively reported to be associated with the development of atopic dermatitis and other allergic diseases [4,5]. Recently, Chen et al. reported that Pen c 13, a major allergen secreted by Penicillium citrinum, degraded occludin, zonula occludens (ZO)-1, and E-cadherin by nasal Pen c 13 challenge in BALB/c mice, showing increased airway hyperresponsiveness, inflammatory cell infiltration, and mucus overproduction in the lung, as well as increased total IgE and Pen c 13-specific IgE [6]. These data support the notion that dysfunction of epithelial barriers may lead to allergic diseases.

There have been few investigations at the molecular level of the characteristics that distinguish SAR patients from subjects who are allergen-specific IgE-positive, but remain asymptomatic. Such investigations will be essential to identify the molecular and cellular mechanisms by which allergic diseases develop, and they could lead to the discovery of novel preventive therapeutics.
One approach to understanding the molecular basis of allergic inflammation is to identify the differences in the gene expression during natural allergen exposure of patients with varying disease status. Microarray gene expression profiling permits thousands of genes to be analyzed simultaneously. Recently, we used microarray analysis to show the upregulation of interleukin-17 receptor B (IL17RB) expression in CD4+ T cells of SAR patients during natural allergen exposure [7]. Our study identified fewer genes that were modulated by allergen than those that had been observed in other microarray studies in which CD4+ cells were stimulated in vitro, suggesting that changes in the gene expression during natural exposure to allergen in vivo may be subdued.

However, it may be more informative to investigate SAR-related gene expression in nasal mucosa rather than in peripheral blood cells, because nasal mucosa is the site of the allergic inflammatory response. Indeed, a microarray analysis of nasal epithelial samples from patients with allergic patients revealed gene expression signatures that were strongly associated with exacerbated or uncontrolled asthma [8,9]. A recent study using nasal epithelial cells revealed a Th2-driven epithelial phenotype in pediatric patients with house dust mite allergy [8]. Although the entire nasal mucosa and underlying tissues are involved in the development of SAR, we may gain valuable insight into the molecular mechanisms of allergen sensitization and SAR development by comparing gene expression profiles in nasal epithelial cells from healthy subjects, patients with SAR, and sensitized subjects who are positive for allergen-specific IgE, but have not yet developed SAR.

In the present study, we performed a microarray analysis of nasal epithelial cells from SAR patients and Control subjects, and analyzed a candidate gene for RT-PCR using samples of Controls, SAR patients, as well as Sensitized subjects to identify gene expression signatures that reflect the status of the SAR patient during natural allergen exposure. We observed that the expression of the cysteine protease inhibitor cystatin SN (CST1) was upregulated specifically in SAR-JC patients during natural allergen exposure. Our results also showed CST1 may inhibit JC allergen-stimulated histamine release in vitro.

Methods

Ethics Statement

All participants provided written informed consent, and the study was approved by the ethics committees of the University of Tsukuba and the University of Fukui, Japan.

Subjects

Study participants were selected from 1575 hospital workers and university students, who participated in an epidemiological survey on allergic rhinitis between 2003 and 2007. All participants were of Japanese origin and were residents of Fukui prefecture, Japan. The characteristics of the study population have been described in detail elsewhere [3]. Total and allergen-specific IgE (specific for JC, Dermatophagoides [dust mites], Dactylis glomerata [orchard grass], Ambrosia artemisiifolia [ragweed], or the fungal allergens Candida albicans and Aspergillus spp.) were measured by the ImmunoCAP method (Pharmacia Diagnostics AB, Uppsala, Sweden).

We invited 14 of the 1575 survey subjects to participate in the microarray analysis study. Samples of nasal epithelial mucosa were collected by brushing the inferior turbinate, between February and April 2009—the period of natural exposure to JC pollen. Eight subjects had SAR due to JC pollen (SAR-JC group), which was diagnosed on the basis of a positive history of rhinitis between February and April and high levels of JC-specific serum IgE.

None of the SAR patients had detectable IgE specific for Dermatophagoides spp., Dactylis glomerata, A. artemisiifolia, C. albicans, or Aspergillus spp. and none of the subjects had been treated by oral corticosteroids during pollen season when we collected the samples.

They had also not taken intranasal and oral corticosteroids for at least 10 months before the collection of samples. All patients were classified as moderate to severe, according to the allergic rhinitis severity (ARIA) classification [10]. The Control group consisted of 6 subjects for whom the inclusion criteria were as follows: (1) no symptoms or history of allergic diseases, (2) no detectable IgE antibodies specific for the 6 common inhalant allergens, the same as the allergens mentioned above, and (3) total serum IgE levels below the general population mean. We also recruited an additional 7 SAR-JC patients and 7 Controls for real-time RT-PCR analysis.

In 2010, we enrolled an additional samples of 17 SAR-JC patients, 13 asymptomatic subjects with JC-specific IgE (Sensitized group), and 15 Control subjects into the study for RT-PCR validation of the microarray data. Subjects in the Sensitized group and SAR-JC group were negative for IgE specific for Dermatophagoides spp., Dactylis glomerata, A. artemisiifolia, C. albicans, or Aspergillus spp. The characteristics of the subjects are listed in Table 1.

Microarray analysis

Nasal epithelial cells were collected from the 14 subjects in the 2009 by brushing the inferior turbinate with a Cytosoft Cytology Brush (Medical Packaging Co, Camarillo, CA, USA). RNA was extracted using the RNeasy Mini Kit (Qiagen K.K., Tokyo, Japan) according to the manufacturer’s instructions. We assessed the quality of total RNA by performing agarose gel electrophoresis and used RNA showing 2 distinct ribosomal peaks corresponding to either 18S or 28S (28S>18S) for microarray and qRT-PCR experiments. Two samples (1 from a SAR patient and 1 from a Control subject) were excluded from the analysis because of the low quality of RNA. Therefore, a total of 12 samples were used for the subsequent microarray analysis.

Microarray assays were performed using the Illumina BeadArray single-color platform (Illumina, San Diego, CA). For the assay, cRNA was synthesized with an Illumina TotalPrep RNA Amplification Kit (Ambion, Austin, TX) according to the manufacturer’s instructions. In brief, aliquots of 500 ng of total RNA were reverse transcribed to first- and second-strand cDNA, which was purified by using spin columns, and then transcribed in vitro to yield biotin-labeled cRNA. A total of 750 ng biotin-labeled cRNA was hybridized to each Illumina HumanRef-8 BeadChip array (Illumina) at 55°C for 18 h. The hybridized BeadChip was washed and labeled with streptavidin-Cy3 (GE Healthcare, Waukesha, WI) and then scanned with the Illumina BeadStation 500 System (Illumina). The scanned image was analyzed by using the BeadStudio software (Illumina). In this system, 22,000 transcripts representing 8 whole-genome samples can be analyzed on a single BeadChip. We included at least 1 technical replicate (i.e., the same cRNA sample) for each BeadChip. We used cRNA from the same sample as our technical replicates to determine whether the experiment was performed properly. The correlation coefficients for replicate RNAs were 0.995–0.996 (r²). Microarray data was deposited in the Gene Expression Omnibus (www.ncbi.nlm.nih.gov/geo) under references GSE43523.

Quantitative real-time RT-PCR (qRT-PCR)

RNA was isolated from nasal epithelial cells obtained from the 45 subjects in 2010, as described above. We performed qRT-PCR on RNA by using the TaqMan Universal Master Mix and an Assays-on-Demand Gene Expression Kit (Applied Biosystems,
Foster City, CA) according to the manufacturer’s instructions. The relative expression was quantified by the ΔΔCt method by using GAPDH as the endogenous control and was performed using SDS software 2.2.0 (Applied Biosystems).

**Immunohistochemical staining**

In order to validate the quality of nasal epithelial cells by brushing the inferior turbinate with a Cytosoft Cytology Brush (Medical Packaging Co), nasal epithelial cells from 3 SAR-JCP and 3 control subjects were fixed on glass slides and stained for cytokeratin AE1/AE3 (Dako Corp., Carpinteria, CA) and vimentin (Dako).

Specimens of the inferior turbinate were collected from 6 patient with severe perennial nasal allergy and SYR symptoms (high levels of JC- and dust mite-specific serum IgE) and 5 Control subjects without nasal allergy. Characteristics of subjects for immunohistochemical staining is shown in Table S1. The inferior turbinate specimens were fixed in neutral buffered formalin (10%v/v formalin in water, pH 7.4) and embedded in paraffin wax. Immunohistochemical staining was performed on-μm-thick sections using a CST1-specific antibody (Abanova Corporation, Taipei, Taiwan), as described previously [11].

**Isolation of basophils and histamine release assay**

Peripheral blood mononuclear cells (PBMCs) were isolated with Lymphoprep (Cosmo Bio, Tokyo, Japan) from heparinized whole blood drawn from 6 patients with SAR (3 males and 3 females, aged 24–36 years, high levels of JC-specific serum IgE). Basophils were separated from PBMCs by using a human basophil isolation kit (Miltenyi Biotec GmbH, Bergisch Gladbach, Germany) according to the manufacturer’s instructions. Basophils (2×10⁶) were resuspended in RPMI1640 medium (Nissui Pharmaceutical, Tokyo, Japan) supplemented with 100 U/mL penicillin (Sigma-Aldrich, St. Louis, MO), 100 μg/mL streptomycin (Sigma-Aldrich), and 10% fetal calf serum (Thermo Fisher Scientific, Waltham, MA), and aliquoted at 200 μL/well in 96-well microtiter plates. The plates were then incubated at 37°C in a humidified 5% CO₂ atmosphere for 30 min. The JC pollen allergen Cry j1 (10 μg/mL; Torii, Tokyo, Japan) was preincubated with recombinant CST1 (200 ng/mL) for 30 min at 37°C, then aliquots of this mixture or Cry j1 alone were added to the basophils and the plates were incubated for an additional 30 min. The concentration of histamine released into the supernatant was measured with a Histamine Enzyme Immunoassay kit (Bertin Pharma, Montigny le Bretonneux, France) according to the manufacturer’s instructions.

**Statistical methods**

Total and specific IgE levels were expressed as the geometric means with 95% confidence intervals (CI). Total and specific IgE data were log-transformed and statistical analyses of IgE levels and age were performed by Student’s t-test or ANOVA. Statistical analysis of the gender ratio was evaluated with Fisher’s exact test.

For the microarray analysis, background-corrected values for each probe on the BeadChip array were extracted using GenomeStudio software (Illumina). Detection P-values were computed from the background model characterized by the chance that the target sequence signal was distinguishable from the negative controls on the same chip. This is based on the average of negative control genes and is termed “the method of background normalization” by Illumina. We performed normexp-by-control background correction, quantile normalization, and log2 transformation on the raw data using the “bcp” command of limma [12]. The normalized values were analyzed by using the R-2.13.1 statistical computing software (R-project, http://www.r-project.org/). Statistical significance of the microarray data was calculated using Welch’s t test, and multiple tests were corrected by the Benjamini and Hochberg false discovery rate [13]. A heatmap of genes differentially expressed between SAR patients and Controls was generated in this study, we extracted the values for the gene expression in eosinophils from the data reported by Nakajima et al [14] (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?token=killihiybhvphohj&term=GeneChip/public.htm). The qRT-PCR results were analyzed by Mann-Whitney’s U test.

**Gene Set Enrichment Analysis (GSEA)**

Gene Set Enrichment Analysis (GSEA) was performed using GSEA v2.0.9 [15] to explore the enriched gene sets related to SAR. We performed GSEA using a dataset provided by the GSEA website (c2.v2symbols.gmt, http://www.broadinstitute.org/gsea/downloads.jsp), as well as with gene sets we manually curated. These sets comprised differentially expressed nasal epithelial genes from children with dust mite allergy [MITE_ALLERGY], genes

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**Table 1. Characteristics of subjects for expression analysis.**

|                          | SAR-JC | Sensitized | Control | P value |
|--------------------------|--------|------------|---------|---------|
| year                     | 2009   | 2010       |         |         |
| Number of subjects       | 17     | 13         | 15      |         |
| Male/Female              | 8/9    | 4/9        | 6/9     | 0.64    |
| Age (mean ± SD)          | 30.47±7.83 | 24.9±5.75 | 24.6±5.46 | 0.24    |
| Total IgE (mean, 95%CI)  | 76.52 (53.30–109.84) | 61.26 (42.54–88.21) | 26.00 (12.06–56.07) | 0.0084  |
| Specific IgE (JC) (mean, 95%CI) | 16.19 (8.26–31.76) | 2.97 (1.37–6.43) | <0.34 | <0.0001 |

N.A, not applicable.
differentially expressed (2-fold upregulated) during in vitro induction by IL-4 and IL-13 (IL-4_AND_IL_13), and those genes induced by IFNs (2-fold up) (IFN\(\alpha\), IFN\(\beta\), and IFN\(\gamma\)) reported by Giovannini-Chami et al. [8]. The manually curated genes are listed in Table S2. The histamine release test results were analyzed using Wilcoxon signed-rank test.

Results

Highly expressed genes in nasal mucosa of subjects in the SAR-JC group

The immunohistochemical experiment revealed that most cells were positive for cytokeratin, suggesting most of the brushed nasal cells were of epithelial origin (Figure S1). To analyze differences in gene expression between the 7 SAR-JC subjects and 5 Control subjects in 2009, we first selected the transcripts that were expressed in at least 1 of the 12 samples and had detection P values of <0.01 on the BeadChip array. This yielded 13,188 expressed transcripts. Transcripts were considered as upregulated or downregulated if they showed an average of greater than 2-fold increase or decrease in expression, and exhibited a statistically significant difference between the SAR and Control groups (q<0.05). We identified 32 transcripts that were upregulated/downregulated in the nasal mucosa of the SAR subjects by at least 2-fold (q<0.05) (Figure 1 and Table 2). Among them, several genes are known to be strongly expressed in eosinophils [14] (Charcot-Leyden crystal protein [CLC] and cadherin-like 26 [CDH26]), and GSEA analysis using the dataset of c2.v2ymbols.gmtrevealed that the microarray dataset in the present study using 7 SAR patients and 5 Controls were enriched for the Top 30 increased eosinophil-specific transcripts (nominal P = 0.006 and FDR q = 0.043), suggesting that their upregulation in the SAR patients may partially be because of the infiltration of eosinophils into the nasal mucosa. GSEA using the gene dataset reported by Giovannini-Chami [8] revealed that genes upregulated in children with mite allergy (MITE ALLERGY) and genes induced by IL-4...
Table 2. Genes significantly altered in nasal epithelial cells in SAR-JC patients.

| Gene | Array address ID | NM numbers | Fold change | q values** |
|------|-----------------|------------|-------------|------------|
| CST1* | 6370541 | NM_001898.2 | 64.6 | 0.031 |
| ITLN1* | 7610170 | NM_017625.2 | 16.9 | 0.045 |
| CLC | 2030142 | NM_001828.4 | 5.9 | 0.032 |
| CDH26 | 4590541 | NM_021810.3 | 5.2 | 0.031 |
| NOS2A* | 4760324 | NM_153292.1 | 4.9 | 0.033 |
| HDC | 6280326 | NM_002112.2 | 3.4 | 0.049 |
| HIST1H2BD2 | 290730 | NM_138720.1 | 3.1 | 0.016 |
| RPL14 | 2140753 | NM_001034966.1 | 2.7 | 0.033 |
| HIST1H2BD2 | 6200669 | NM_001828.4 | 2.5 | 0.033 |
| OSTC | 4610722 | NM_021227.2 | 2.5 | 0.032 |
| IL20RB* | 3400995 | NM_144717.2 | 2.4 | 0.031 |
| ATP2C2 | 2510669 | NM_0018861.2 | 2.3 | 0.031 |
| RAS2* | 4570670 | NM_006270.3 | 2.2 | 0.047 |
| CFD1 | 6200494 | NM_006324.2 | 2.2 | 0.033 |
| SDPR* | 6290114 | NM_004657.4 | 2.0 | 0.031 |
| BCL2L15 | 1710575 | NM_001019022.2 | 2.0 | 0.045 |
| CAMK2G | 1010475 | NM_001222.2 | 2.0 | 0.033 |
| ZNF750 | 3420176 | NM_024702.2 | 2.1 | 0.032 |
| SLC6A6 | 2810280 | NM_003043.3 | 2.2 | 0.033 |
| NR1D2 | 2640707 | XM_001130839.1 | 2.2 | 0.031 |
| MAP3K2 | 3800762 | XM_001128799.1 | 2.3 | 0.031 |
| MAF | 3610440 | NM_005360.3 | 2.3 | 0.033 |
| SLC25A36 | 3870242 | NM_018155.1 | 2.3 | 0.041 |
| GKS | 6180543 | NM_001039547.1 | 2.4 | 0.041 |
| SYT7* | 5050390 | NM_004200.2 | 2.4 | 0.031 |
| AKR1B1* | 4260139 | NM_001628.2 | 2.4 | 0.025 |
| ST3GALS5* | 5260403 | NM_00142437.1 | 2.4 | 0.031 |
| OTX1* | 1740438 | NM_014562.2 | 2.8 | 0.049 |
| RORA | 1110180 | NM_002943.3 | 2.9 | 0.044 |
| PDE5A | 1770333 | NM_001083.3 | 3.1 | 0.033 |
| ANKRD36* | 5810050 | NM_198555.3 | 4.1 | 0.044 |
| ANKRD36B | 3600100 | NM_025190.2 | 5.5 | 0.031 |

*Absent call in Eosinophils according to the data by Nakajima et al. **Welch’s t test, Benjamini-Hochberg FDR.

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Table 3. GSEA using dataset of the study by Giovannini-Chami [5].

| Gene | Enrichment Score (ES) | Nominal ES (NES) | P value | q value |
|------|-----------------------|------------------|---------|---------|
| IL-4_AND_IL-13 | 0.73 | 1.85 | 0.004 | 0.027 |
| MITE_ALLERGY | 0.87 | 1.75 | 0.004 | 0.025 |
| IFN α | 0.21 | 0.74 | 0.78 | 0.89 |
| IFN γ | 0.19 | 0.66 | 0.83 | 0.75 |

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and IL-13 (IL-4_AND_IL-13) were enriched in SAR patients in our dataset (Table 3). However, genes related to IFN responses were not enriched in our dataset.

We found that CST1 (cystatin SN) was highly expressed in the SAR-JC subjects, undetectable in the Control subjects, and not expressed in eosinophils (absent call judged by the GeneChip Analysis Suite 5.0 program according to the data by Nakajima et al. [14]; hence, we focused on CST1 for further analysis. To validate the results of the microarray analysis, we measured CST1 expression in the nasal mucosal samples from the 2009 subjects by qRT-PCR. As can be seen in Figure 2A, CST1 expression was significantly higher in the SAR-JC group than in the Control group (P = 2.4 x 10^-7). To confirm these data, we also performed CST1 qRT-PCR on nasal mucosal samples in 2010 (Figure 2B). Here, CST1 expression was significantly higher in the SAR-JC group than in the Sensitized group (P = 0.00087) and Control group (P = 0.0006), whereas there was no significant difference in expression between the Sensitized and Control groups (P = 0.13).

We next investigated the expression of CST1 protein by immunohistochemical staining of the inferior turbinate samples from 6 patients with perennial nasal allergy and SAR symptoms and 5 Control subjects without allergy. As shown in Figure 3, CST1 was expressed in nasal epithelial cells in the SAR patients, but not in the Control subjects. To examine the effect of CST1 on allergen-stimulated histamine release, basophils were isolated from the peripheral blood of 6 patients with SAR, and stimulated with the JC allergen Cry j1 both in the presence and absence of CST1. Figure 4 shows that Cry j1-stimulated histamine release by basophils was effectively inhibited in the presence of CST1 (P = 0.032).

Discussion

In the present study, we performed microarray analyses to identify SAR-related genes in nasal epithelial cells. We identified 36 genes that showed statistically significantly differed in SAR subjects compared with the Control group, and we confirmed by qRT-PCR that the expression of CST1 was upregulated in SAR-JC patient compared with the Control group in both 2009 and 2010. We also showed that addition of exogenous CST1 to basophils inhibited JC allergen-stimulated histamine release in vitro.

Cry j1 is a major antigen of Japanese cedar pollen [16]. It is a basic glycoprotein localized in the pollen cell wall, which has pectate lyase activity [17]. The epitope of Cry j1 is recognized by specific IgE of Cry j1 and released histamine from basophils by interacting antigen and antigen-specific IgE. Alteration of Cry j1 structure by heat denaturation leads to reduction of IgE-binding activities [18,19]. Therefore, histamine release from the basophils of patients induced by adding Cry j1 might be due to the degranulation via crosslinking of Cry j1-specific IgE already bound on the IgE receptor. Takai et al. reported that interaction of Der f 1, one of the major allergens of house dust mite with protease activity, and cystatin A, which belongs to cystatin superfamily, inhibited the binding of IgE in mite-allergic volunteers’ sera, suggesting that cystatin A might change the binding site between antigen and antigen-specific IgE [20]. Thus, cystatin SN might affect the binding site between antigen and antigen-specific IgE, leading to the suppression of histamine release from basophils in the present study.

Cysteine proteases are widely expressed proteolytic enzymes that play roles in inflammatory tissue destruction as well as tissue remodeling [21]. The proteolytic activity of these enzymes is controlled by a family of inhibitors known as the cystatin superfamily [22], and cystatins inhibit cysteine proteases by forming tight but reversible complexes with their target enzymes.
Members of the type 2 cystatin subfamily are widely distributed [23], and include CST1, CST2, CST3, CST4, CST5, CSTP1, and CSTP2, which share >70% sequence similarity [24]. The CST1 is expressed in the submandibular gland, gall bladder, and uterus [24]. CST1 has been shown to bind tightly to the cysteine protease, papain, which is a potent allergen, and to inhibit the cysteine protease activity of papain [25].

Airborne allergens, such as those derived from pollen, cockroaches, fungi, and house dust mites, are proteases with the capacity to induce an inflammatory response. Previous studies have shown that the proteolytic activity of these allergens may facilitate sensitization, possibly by disrupting the balance between protease and antiprotease activities. This theory is supported by the observation that the protease allergens reduce the antiprotease activity of the mucosa [26,27,28,29]. Der p 1 and Der f 1, the major allergens derived from house dust mites, are cysteine proteases that degrade α1-antitrypsin inhibitor in the airway [28,29] and inactivate lung surfactant proteins A and D [26]. Cystatin A, a member of the cysteine protease inhibitor family, has been shown to suppress induction of Der f 1-specific IgE by modifying the interaction between Der f 1 and cystatin A, which may modulate the allergic inflammation and sensitization processes [20]. Thus, the cystatin family may play protective roles against allergens with protease activity.

Measurement of allergen-specific IgE in serum is widely used to diagnose allergic diseases such as SAR and asthma; however, subjects with allergen-specific IgE do not necessarily develop the allergic disease. Although many reports have described the pathophysiologic features of allergic inflammation, biomarkers or genes that predict the onset of allergic rhinitis are yet to be reported. In this study, we observed that CST1 was not expressed in cells from non-sensitized subjects, and that CST1 expression was significantly lower in the IgE-positive Sensitized subjects than in the SAR-JC subjects. These data suggest that CST1 may be a candidate marker gene to predict allergic rhinitis. We detected that 2 subjects in the Sensitized group showed particularly high CST1 expression in the nasal mucosa, and we speculate that these subjects may have been close to developing SAR symptoms.

A recent study using nasal epithelial cells from children with dust mite allergy showed that CST1 was the most highly upregulated gene and that CST1 was also strongly induced by stimulation with IL-4 and IL-13 [8]. Therefore, CST1 is likely to be upregulated in response to protease-containing allergens and the Th2 environment in the nasal mucosa. Because GSEA revealed that the genes upregulated in nasal brushing samples from mite allergic patients and the genes induced by IL-4 and IL-13 were significantly enriched in our dataset (Table 3), our data...
support the notion that pathways underlying allergic rhinitis are common regardless of the allergen.

The balance between protease and antiprotease activities might thus be a key to distinguish between the sensitization stage and allergic response, and we suggest that cystatins contributes re-establishing homeostasis of the nasal mucosa.

**Supporting Information**

**Figure S1 Immunohistochemical staining of cytokeratin and vimentin in brushed nasal epithelial cells.** Representative immunostaining of cytokeratin expression (A and B) and vimentin expression (C and D). Magnification: ×200. (PPTX)

**Table S1 Characteristics of subjects for immunohistochemical staining.** (XLSX)

**Table S2 Manually curated gene sets used for GSEA.** (XLSX)

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**Author Contributions**

Conceived and designed the experiments: Y. Imoto TA EN SF. Performed the experiments: Y. Imoto TT YM YH M. Ono TY Y. Ito. Analyzed the data: Y. Imoto TT EN. Contributed reagents/materials/analysis tools: Y. Imoto TY Y. Ito M. Okano SF. Wrote the paper: Y. Imoto TT TA EN SF.

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