Genome-wide association studies of asthma have identified genetic variants in the *IL1RL1* gene, but the molecular mechanisms conferring risk are unknown. *IL1RL1* encodes the ST2 receptor (ST2L) for IL-33 and an inhibitory decoy receptor (sST2). IL-33 promotes type 2 inflammation, which is present in some but not all asthmatics. We find that two single nucleotide polymorphisms (SNPs) in *IL1RL1* — rs1420101 and rs11685480 — are strongly associated with plasma sST2 levels, though neither is an expression quantitative trait locus (eQTL) in whole blood. Rather, rs1420101 and rs11685480 mark eQTLs in airway epithelial cells and distal lung parenchyma, respectively. We find that the genetically determined plasma sST2 reservoir, derived from the lung, neutralizes IL-33 activity, and these eQTL SNPs additively increase the risk of airway type 2 inflammation among asthmatics. These risk variants define a population of asthmatics at risk of IL-33–driven type 2 inflammation.
**IL1RL1 asthma risk variants regulate airway type 2 inflammation**

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**Introduction**

Advances in molecular phenotyping reveal heterogeneity within the asthmatic population (1–3). The most consistent endotype to emerge is characterized by excessive airway type 2 inflammation (type 2-high asthma) (2–6). IL-33 signaling through the ST2 receptor (encoded by the **IL1RL1** gene) is an important driver of type 2 immune responses (7, 8). Genetic variation in both the **IL33** and **IL1RL1** genes is strongly associated with asthma in genome-wide association study (GWAS) meta-analyses of European and North American populations (9, 10). The molecular mechanism and the influence of these risk variants on airway type 2 inflammation is unknown.

Two major transcript variants are produced from the **IL1RL1** gene (11). Transcript variant 1 is a long transcript that encodes the membrane-bound ST2 receptor (ST2L) for IL-33, while transcript variant 2 is short and encodes a soluble decoy receptor (sST2). ST2 transcripts can be expressed from two spatially distinct promoters. The distal promoter can drive either **ST2L** or **sST2** expression, while the proximal promoter directs expression of **sST2** only. Two studies indicate that single nucleotide polymorphisms (SNPs), some overlapping with the asthma GWAS variants, are strongly associated with **sST2** plasma levels, though neither is an expression quantitative trait locus (eQTL) in whole blood. Rather, rs1420101 and rs11685480 mark eQTLs in airway epithelial cells and distal lung parenchyma, respectively. We find that the genetically determined plasma **sST2** reservoir, derived from the lung, neutralizes IL-33 activity, and these eQTL SNPs additively increase the risk of airway type 2 inflammation among asthmatics. These risk variants define a population of asthmatics at risk of IL-33-driven type 2 inflammation.

Genome-wide association studies of asthma have identified genetic variants in the **IL1RL1** gene, but the molecular mechanisms conferring risk are unknown. **IL1RL1** encodes the ST2 receptor (ST2L) for IL-33 and an inhibitory decoy receptor (sST2). IL-33 promotes type 2 inflammation, which is present in some but not all asthmatics. We find that two single nucleotide polymorphisms (SNPs) in **IL1RL1**—rs1420101 and rs11685480—are strongly associated with plasma **sST2** levels, though neither is an expression quantitative trait locus (eQTL) in whole blood. Rather, rs1420101 and rs11685480 mark eQTLs in airway epithelial cells and distal lung parenchyma, respectively. We find that the genetically determined plasma **sST2** reservoir, derived from the lung, neutralizes IL-33 activity, and these eQTL SNPs additively increase the risk of airway type 2 inflammation among asthmatics. These risk variants define a population of asthmatics at risk of IL-33-driven type 2 inflammation.
To answer these questions, we explored the role of IL1RL1 genetics in determining sST2 plasma levels, blood and lung sST2 transcript expression, and airway type 2 inflammation in a large cohort of asthmatics and controls (Figure 1 and Supplemental Table 1; supplemental material available online with this article; doi:10.1172/jci.insight.87871DS1). IL1RL1 genetic variant expression effects were studied in different lung cell types using a large dataset of cultured primary human airway epithelial cells and a publicly available expression database of distal lung parenchymal tissue (Figure 1). Finally, we evaluated the relationship between IL1RL1 eQTL variants that mark distinct lung cell–specific expression and risk for the type 2-high asthma endotype (Figure 1).

**Results**

**IL1RL1 asthma risk variants are associated with sST2 plasma protein levels but not whole blood sST2 gene expression levels.** To examine genetic influences on plasma sST2 protein levels, we selected 5 candidate protein quantitative trait loci (pQTL) SNPs at the IL1RL1 locus (Figure 2A and Supplemental Figure 1A). Based on 1,000 genomes data in the CEU population (Utah residents with ancestry from northern and western Europe), these SNPs tag 96 variants distributed across the locus. Minor allele frequencies and linkage disequilibrium maps for our study population can be found in Supplemental Figure 1, A and B. These variants are eQTL candidates based on reported genetic associations with asthma, allergic diseases, eosinophilia, sST2 plasma protein levels, and in vitro sST2 expression (9, 10, 12, 13, 15–18) (Supplemental Figure 1A). Among 90 self-identified white non-Hispanic asthmatic and healthy control subjects, we found strong associations between 3 of the SNPs, and liquid chromatography mass spectroscopy (LC-MS) determined sST2 plasma protein levels (Figure 2B). These 3 associations were also observed in the full mixed-race cohort of 171 asthmatic and healthy control subjects (Figure 2B). The direction of effect we observed for these variants (Supplemental Figure 2, A and B) is consistent with prior reports from a pediatric Dutch cohort and from the Framingham Offspring Cohort, which also found that rs1420101 and rs11685480 were among the most significant associations with circulating sST2 levels (12, 13). We hypothesized that these variants mediate their effect on protein levels through regulation of gene expression. To test this, we measured sST2 transcript expression levels in whole blood RNA from 63 asthmatics and 23 healthy control subjects (Figure 2B). None of the SNPs tested were associated with sST2 transcript or sST2 protein levels (Figure 2B, and Supplemental Figure 3, B and C).

rs11685480 and rs1420101 mark cell type–specific lung eQTL. The strong genetic association of IL1RL1 variants with sST2 plasma protein but not blood gene expression measurements suggests that other tissues may be a source of circulating sST2 protein. Given the importance of the airway epithelium in asthma, we examined sST2 transcript expression and protein secretion in cultured primary airway epithelial cells from 141 donors. These airway epithelial cells were harvested from cadaver tracheas and cultured at air-liquid interface to produce mucociliary epithelium that was examined in the unstimulated condition and following stimulation with the type 2 cytokine IL-13. We found the A allele of the rs1420101 SNP, which was associated with lower plasma sST2 protein levels, was also associated with lower sST2 gene expression and sST2 protein secretion both in the unstimulated and the IL-13–stimulated culture conditions (Figure 3, A and B, and Supplemental Figure 2C). In contrast, the other blood pQTLs, rs11685480 and rs1921622, were not associated with sST2 transcript or sST2 protein levels (Figure 3, A and B, and Supplemental Figure 2C).
We thus considered the possibility that the rs11685480 SNP influences gene expression in another lung cell type. We tested the association between the blood pQTL SNPs and sST2 expression in distal lung tissue using data from 237 white non-Hispanic donors generated by the Genotype-Tissue Expression Project (GTEx) consortium. Notably, whole lung specimens are largely composed of alveolar epithelial and microvascular endothelial cells, with few airway epithelial cells. In contrast to airway epithelial cells, we found that both rs11685480 and rs1420101 SNPs were strongly associated with distal lung expression of sST2 (Figure 3C). Again, the same allele of both of these SNPs was associated with lower sST2 gene expression (Supplemental Figure 2, E and F).

The differences in eQTL status between the rs1420101 and rs11685480 SNPs by lung cell type suggest these SNPs mark independent, cell type–specific molecular effects. Given that sST2 can be expressed from either or both the distal and proximal promoters (Figure 3D) (18), differential promoter usage by cell type could explain this effect. Supporting this, the rs11685480 SNP is located in the distal IL1RL1 promoter and has previously been shown to regulate distal promoter sST2 transcript expression (18). We investigated promoter usage in airway epithelial cells, alveolar epithelial, and lung microvascular endothelial cells using quantitative PCR (qPCR) assays to amplify exon 1a or exon 1b of ST2 transcripts (Figure 3D). Exon 1a is exclusively found in transcripts derived from the distal IL1RL1 promoter, while exon 1b is found in transcripts derived from the proximal promoter. We find that airway epithelial cells and lung microvascular

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**Figure 2. Single nucleotide polymorphisms (SNPs) in the IL1RL1 gene influence plasma protein sST2 levels but not whole blood cell sST2 expression.** (A) Schematic showing the IL1RL1 gene transcript structure highlighting 5 candidate expression quantitative trait loci (eQTL) SNPs. Blue exons are specific to the ST2L transcript, green exons are specific to the sST2 transcript, and red exons are shared by both transcripts. The distal promoter can produce both the long ST2L and the short sST2 transcript, while the proximal can produce only the short sST2 transcript. (B) sST2 plasma protein levels (red) as measured by liquid chromatography-mass spectroscopy (LC-MS) are associated with 3 SNPs in the IL1RL1 gene: rs11685480, rs1420101, and rs1921622 in white non-Hispanic subjects only (WNH) and among the entire cohort (All). The tested allele in all cases is associated with lower-circulating sST2 levels. None of the tested SNPs was an eQTL for sST2 expression in whole blood RNA (blue). SNP regression coefficient and the corresponding 95% CIs are plotted with P values listed below. Additive linear regression models were used. Nonitalicized P values indicate the WNH, while the italicized P values shown for the entire cohort (All).
endothelial cells largely use the proximal promoter, while alveolar epithelial cells use the proximal and distal promoters equally (Figure 3E). Our results identifying rs11685480 as an eQTL for distal lung tissue but not airway epithelial cells are consistent with rs11685480 effects on distal promoter–driven expression in alveolar but not airway epithelial cells. Likewise, our results identifying rs1420101 as an eQTL in both the airway epithelium and distal lung tissue suggest that the expression effect is mediated through the proximal promoter that is used in both cell types.

Plasma sST2 levels are correlated with whole blood EC50 for IL-33. To determine the contribution of plasma sST2 levels to IL-33 signaling, we examined IL-33–stimulated PBMCs and whole blood cells from 4 healthy subjects. This paired design allowed us to examine the effect of plasma proteins on IL-33 signaling. We find the half maximal effective concentration (EC50) for IL-33 across 4 donors showed little variability in PBMCs, while the EC50 in whole blood is markedly higher and considerably more variable, suggesting the presence of an IL-33 inhibitor in whole blood (Figure 4, A and B). Soluble ST2 directly inhibits IL-33 signaling in both mouse models and in vitro cultures (19). Consistent with this, we find that plasma sST2 levels are positively correlated with whole blood cell EC50 to IL-33 (Figure 4C), suggesting that soluble ST2 in circulating plasma is a functional inhibitor of IL-33 signaling in whole blood.

Airway epithelial sST2 expression is associated with type 2 inflammation and is directly regulated by IL-13. In order to understand the significance of a genetically determined deficiency of sST2 in promoting airway

Figure 3. Differences in airway epithelial cell and distal lung tissue expression genetics is driven by differential promoter usage. In cultured airway epithelial cells, rs1420101 (blue) — but not rs11685480 (red) or rs1921622 (black) — is associated with sST2 gene expression (n = 127) (A) and sST2 secreted protein levels (n = 141) (B) in both the unstimulated and IL-13–stimulated conditions. White non-Hispanic subjects are indicated with a circle, while the entire cohort is indicated with a square. SNP regression coefficient and the corresponding 95% CIs are plotted. Additive linear regression models were used. *P < 0.05, **P < 0.01. (C) In contrast, both rs1420101 (blue) and rs11685480 (red) are associated with sST2 gene expression in distal lung parenchymal tissue in GTEx cadaveric donors (n = 237). SNP regression coefficient and the corresponding 95% CIs are plotted with P value listed below. Additive linear regression models were used. (D) Schema showing that sST2 transcripts can be derived from either the IL1RL1 distal or proximal promoters. sST2 transcripts derived from the distal promoter will contain exon 1a, while those derived from the proximal promoter will contain exon 1b. Primers were designed to specifically amplify exon 1a and exon 1b, as indicated by black arrowheads. (E) Cultured human airway epithelial cells (n = 6) and cultured human lung endothelial cells (n = 3) use the proximal promoter, while human alveolar epithelial cells (n = 7) use both promoters. *P < 0.05, ***P < 0.001 using 2-tailed paired t test.
inflammation in asthma, we examined sST2 expression in bronchial brush RNA from 22 healthy and 83 asthmatic subjects. In previous studies, the degree of airway type 2 inflammation was measured in these asthmatics using a gene expression signature consistent with IL-13 activation of the epithelium (2, 20). A single metric derived from the qPCR-based expression of POSTN, CLCA1, and SERPINB2 in airway epithelial cells was used to classify asthmatics as type 2-high or -low. We find that sST2 gene expression is increased in asthmatics compared with healthy controls (Supplemental Figure 3D); however, this effect is present only in the subset with evidence of airway type 2 inflammation (type 2-high asthma) (Figure 5A). These findings suggest that sST2 expression is regulated by type 2 cytokines in airway epithelial cells.

To test this directly, we examined sST2 gene expression and protein secretion in cultured human airway epithelial cells from 141 donors in unstimulated and IL-13–stimulated conditions (21). IL-13 markedly increased sST2 gene expression and secreted sST2 protein in these cells (Figure 5, B and C). We did not find significant associations between IL1RL1 SNPs and in vivo bronchial brush sST2 gene expression (Supplemental Table 3), which may be due to confounding mediated by differences in type 2 inflammation across the subjects. This finding contrasts with the eQTL and pQTL effects of rs1420101 revealed in the unstimulated and defined exposure to IL-13 in the in vitro airway epithelial cultures (Figure 3, A and B). Thus, there appears to be dual regulation of airway epithelial sST2 gene expression in vivo. The genetic effect of rs1420101 acts through the proximal IL1RL1 promoter to lower sST2 gene expression. At the same time, the type 2 cytokine IL-13 dramatically upregulates sST2 expression by airway epithelial cells. Both in vivo and in vitro, sST2 acts as a negative regulator of type 2 inflammation via direct inhibition of IL-33 signaling (19, 22–25). Our results suggest that this negative regulation is most prominent in asthmatics with...
excessive type 2 inflammation and lead us to predict that IL1RL1 lung eQTL SNPs, which are associated with decreased sST2 expression, will increase the risk of type 2-high asthma endotype.

rs11685480 and rs1420101 eQTL variants are associated with the type 2-high asthma endotype. To determine if IL1RL1 eQTL/pQTL SNPs are associated with airway type 2 inflammation in asthma, we examined 150 asthmatics with airway type 2 endotyping data (Supplemental Table 2). Type 2 endotype (high vs. low) was determined in our prior studies by the presence of airway gene expression signatures consistent with type 2 inflammation in either bronchial brush or sputum cell RNA (1, 3). Type 2-high asthmatics demonstrate markedly increased blood eosinophil counts, serum IgE measures, and a fraction of exhaled nitric oxide (FeNO), as well as lower prebronchodilator forced expiratory volume in 1 second (FEV1) percent predicted compared with type 2-low asthmatics (Supplemental Table 2). In our genetic analysis, we find that both eQTL/pQTL SNPs, rs11685480 and rs1420101, are associated with increased risk of the type 2-high asthma endotype (odds ratio [OR] 2.32, \( P = 0.002 \); OR 1.74, \( P = 0.032 \), respectively) (Figure 6, A and B). We observed this association despite the fact that inhaled steroid use can reduce airway type 2 inflammation (20), an effect that could result in subject misclassification and a bias toward a null result. Consistent with the function of sST2 as an inhibitor of IL-33 and type 2 inflammation, the allele for both SNPs that is associated with lower sST2 gene and sST2 protein expression is associated with increased risk of the type 2-high endotype. Based on our promoter usage data suggesting an independent effect for the rs11685480 and rs1420101 SNPs on sST2 expression, we considered if carrying risk alleles from both SNPs would be associated with further decreases in sST2 plasma levels. We found a strong negative relationship between the number of rs11685480/rs1420101 risk alleles carried and circulating sST2 plasma protein levels (Figure 6C). Consistent with our model, the risk estimate for type 2-high asthma in the combined 3–4 allele–carrying group versus the 0–2 allele–carrying group was higher than the risk estimate for either SNP alone (OR 2.85, \( P = 0.02 \)) (Figure 6C). These results provide strong evidence that reduced sST2 expression is a risk factor for the type 2-high endotype of asthma.
Discussion

In the last decade, both clinical and translational research has revealed heterogeneity in the pathological mechanisms underlying asthma. The most clinically important disease endotype described to date is characterized by excessive airway type 2 inflammation and affects about half of asthmatics (2, 26). Using genetically modified mice, many of the molecular mechanisms of type 2 inflammation have been discovered (27); however, little is known regarding the genetic predisposition to this endotype in human asthma. In this study, we demonstrate that asthma risk variants in the IL1RL1 gene specifically increase the risk of type 2-high disease. Moreover, we show that these variants confer risk by reducing plasma and airway levels of sST2, an inhibitor of IL-33 signaling and type 2 inflammation. Together, our findings define a population of asthmatics that is at risk for IL-33-driven type 2 inflammation.

To date, 7 candidate gene studies and 4 GWAS studies — including European and North American GWAS meta-analysis studies involving 26,000 and 17,000 subjects, respectively — have implicated genetic variation at the IL1RL1/IL18R1 locus in the development of asthma (9, 10, 12, 15–17, 28–32). Studies showing a relationship between several IL1RL1 GWAS variants and plasma sST2 protein levels suggest that these variants function as eQTLs (12, 13, 18). Most notably, a GWAS for sST2 plasma levels in the Framingham Offspring Cohort found strong evidence for genetic regulation of circulating protein (13).

The most strongly associated SNP in this study was rs950880, which is in complete linkage disequilibrium with rs1420101 and within the haplotype we implicated as a blood pQTL. The rs1420101 SNP was also associated with plasma sST2 levels in a large pediatric Dutch cohort (PIAMA) (12). We find that rs1420101 is an eQTL in airway epithelium and distal lung parenchyma, tissues that use the proximal IL1RL1 promoter. These results are consistent with the effect of the rs1420101 SNP being mediated through this promoter. Analyzing CEU data from the 1,000 genomes project, we find 12 other variants in strong linkage disequilibrium (r² > 0.8) with the rs1420101 variant. While rs1420101 and rs950880 are located in the first and fifth introns, respectively, 3 variants within this block are located in the proximal promoter. Molecular genetic studies of these variants in airway epithelial cells will be needed to determine if the eQTL effect of the rs1420101 block is mediated through the proximal promoter and to further narrow the causal variant within this SNP block.

One of the SNPs most strongly associated with soluble ST2 levels in the above referenced Framingham cohort, rs953934, is a tag for the rs11685480 (CEU r² = 0.98) SNP that we have implicated as a blood pQTL and eQTL driving expression through the distal IL1RL1 promoter in the lung parenchyma, specifically within alveolar epithelial cells (13). Moreover, the rs11685424 SNP, which is in complete LD with rs11685480 SNP in the CEU population, is strongly associated with sST2 expression in parenchymal lung tissue measured by microarray (14), consistent with the distal lung tissue expression data presented here. Notably, rs11685480 resides within a distal promoter haplotype that is associated with atopic dermatitis and modulates ST2 expression in a human mast cell line (18). Our observation that both the distal and proximal promoters are used by alveolar epithelial cells presents the possibility that ST2L is also expressed and affected by this eQTL. However, we note that ST2L is weakly expressed in the GTEx lung tissue data versus sST2 (median RPKM expression levels sST2 = 56.4 vs. ST2L = 0.83). Taken together, these data strongly suggest that rs11685480 alone or in combination with other SNPs alters sST2 expression through the distal promoter in alveolar epithelial cells and thereby contributes to circulating sST2 levels.

Our experimental findings support the presence of lung cell type-specific eQTLs tagged by SNPs rs1420101 and rs11685480. We did find a significant association between the rs1921622 SNP and circulating levels of sST2 protein, though the association was less significant than for rs1420101 or rs11685480. Moreover, we found no association with this SNP and airway epithelial cell sST2 gene expression and only a weak association with distal lung tissue sST2 expression. We hypothesize that the rs1921622 SNP effect may be explained by linkage disequilibrium with the rs1420101 SNP or that this SNP may function as an eQTL in another untested cell type. Further molecular studies will be needed to delineate this mechanism. While our results support that the identified circulating sST2 pQTLs are the result of these SNPs functioning as eQTLs in lung cells but not blood cells, we acknowledge that eQTL effects of these SNPs in specific blood cell types could be missed by our mixed blood cell type analysis. Additionally, these SNPs could mediate posttranslational changes to sST2 protein and thus contribute to the observed pQTL.

Our data suggest that the rs11685480 and rs1420101 SNPs or variation in LD with these SNPs mark independent eQTL loci that contribute to asthma pathogenesis by reducing circulating and lung sST2 levels to promote type 2 airway inflammation. This is consistent with several well-powered genetic epidemiology
studies of asthma and allergic phenotypes (12, 15). Specifically, the A allele of rs1420101 SNP is strongly associated with increased risk of asthma in a comparison of 44,890 controls versus 7,996 European asthmatics (15). Stratification of asthmatics in that cohort indicates that the association is strongest among atopic asthmatics, consistent with our finding that the rs1420101 A allele is associated with the type 2-high asthma endotype. The rs1420101 variant is also one of the 5 genome-wide significant hits from a GWAS of blood eosinophil levels conducted in over 20,000 European and 5,000 Asian patients (15). In that study, the A allele of rs1420101 is associated with higher levels of both eosinophils and IgE. Although the rs11685480 SNP has not been screened in GWAS studies, the allele of this SNP that is associated with lower sST2 levels is also associated with type 2-high disease, matching the rs1420101 findings. Moreover, carrying multiple risk alleles for rs1420101 and rs1685480 is associated in further reductions in plasma sST2 levels and an even higher risk of type-2 high asthma, suggesting that these 2 loci function independently.

Our genetic mechanism is strongly supported by mouse models demonstrating that IL-33 promotes type 2 inflammation and is inhibited by sST2 (19). We find that this inflammatory pathway is altered in the airway of asthmatics with excessive type 2 inflammation. In these subjects, sST2 is markedly upregulated, an effect that is mediated by the type 2 cytokine IL-13. Genetic variation that reduces local airway sST2 production and disrupts a finely tuned balance between IL-33–driven type 2 cytokine production and the IL-13–induced negative feedback loop may be the primary driver of the risk we observe. However, leakage of plasma proteins into the airways during inflammation may allow a reservoir of biologically active sST2 to restrain type 2 inflammation. In this way, the genetically determined level of plasma sST2 protein, derived from the lung, would influence risk of type 2 inflammation in asthma.

Our translational study of in vivo and in vitro lung tissue genetics allows us to develop a model linking the strong IL1RL1 genetic epidemiology to basic immunology and to demonstrate a key role for IL-33 in asthmatic type 2 inflammation (Supplemental Figure 4) (8, 33–35). This multicellular model suggests that local airway sST2 levels, as well as circulating plasma sST2 levels, contribute to neutralization of IL-33 in the tissue. A genetically determined deficiency in sST2 in the airway (driven by the proximal promoter), as well as a deficiency in sST2 in the plasma (driven though both the proximal and distal promoters), act to increase IL-33 signaling and promote airway type 2 inflammation. These genetic variants define an asthmatic population that is at increased risk for IL-33–driven type 2 inflammation and that may benefit from targeted blockade of the IL-33/ST2 signaling axis.

Methods

UCSF Airway Tissue Bank (UCSF ATB)

We studied human subjects who had biospecimens stored in the UCSF ATB. We included subjects who had available DNA, as well as any of the following: plasma, whole blood RNA, bronchial brush RNA, or induced sputum cell RNA (Figure 1 and Supplemental Table 1). These samples were collected in research studies at UCSF from 2000–2012 and stored in the UCSF ATB. Subject characterization and biospecimen collections were performed according to standardized and uniform protocols. These biospecimens have been used in other reported studies from our group (1–3, 20, 36–39). Asthma was defined as physician diagnosis and confirmed by measures of airway hyperresponsiveness or reversible airway obstruction. Healthy control subjects were nonsmokers with no history of lung disease and no evidence of airway hyperresponsiveness.

Measurement of sST2 protein

sST2 was measured by ELISA (Presage ST2 Assay, Critical Diagnostics) per the manufacturer’s instructions and by immunocapture LC-MS (IC LC-MS). The IC LC-MS was performed by first incubating plasma overnight with biotinylated anti-ST2 (BAF523, R&D Systems), followed by bead capture, acid elution, and tryptic digestion. Peptides unique to ST2 were quantitated against a peptide control and standard curve with ST2-Fc. Plasma measurements of sST2 made with the Critical Diagnostics assay correlated strongly with the measurements made by IC LC-MS (Supplemental Figure 3A).

Cultured human airway epithelial cells

Airway epithelial cells were harvested from the tracheas of 141 organ donors obtained via the California Transplant Donor Network (21). DNA was extracted from cells following harvest using DNeasy Blood and Tissue Kit (Qiagen). Harvested cells were also expanded for one passage only in basal epithelial growth conditions.
media (Lonza) and then plated to 12-mm diameter transwells. Cells were grown for 21–23 days in 2% Ultroser G (Pall Corporation) in DMEM/F12 media and then stimulated with media alone or IL-13 (10 ng/ml) (R&D Systems) every 24 hours for 2 days. RNA was extracted using RNeasy kits (Qiagen). Basal conditioned media was collected and analyzed by ELISA for sST2 protein (sST2 Duoset, R&D Systems).

Lung microvascular endothelial cell culture
Human lung microvascular endothelial cells (HMVEC-L) were purchased from Lonza and grown according to the distributor’s instructions in EGM-2MV Bullet Kit Medium (Lonza) for one passage. Cells were harvested at confluence, and RNA was extracted as described above.

Lung alveolar epithelial cells
Alveolar type II epithelial cells were isolated from deidentified organ donors whose lungs were not suitable for transplantation and donated for medical research through the National Disease Research Interchange and the International Institute for the Advancement of Medicine. The type II cells were dissociated from the lung by treatment with elastase, and they were purified by discontinuous density gradient centrifugation and positively selected by binding to magnetic beads through CD326 (EP-CAM) (Miltenyi Biotec) (40, 41).

Measurement of gene expression
Whole blood cell (PAXGENE) RNA and bronchial brush RNA expression levels of sST2 were measured by qPCR. Briefly, 20 ng of RNA was reverse transcribed using the SuperScript VILO cDNA Synthesis Kit and Master Mix (Invitrogen). Bronchial brush cDNA (4 ng) was PCR amplified in multiplex for housekeeping genes PPIA, RPL13A, and EEF1A1 and sST2; 4 ng of PAXGENE cDNA was PCR amplified for housekeeping genes GAPDH, PPIA, and RPL37A and sST2. Nested taqman primers and probes were then used in a standard 40-cycle taqman reaction using the preamplified PCR product and Taqman Universal Master Mix (Invitrogen). Primer and probe sequences can be found in Supplemental Table 4. The gene expression levels of sST2 in cultured airway epithelial cells (control and IL-13 stimulated) were determined from whole transcriptome sequencing. Specifically, Kapa Biosystems mRNA-seq library kits (catalog KK8421) were used to generate sequencing libraries with barcodes and adapters compatible with the Ion Torrent Proton sequencing instrument (Thermo Fisher Scientific). Sequencing was conducted on an Ion Torrent Proton sequencer using P1 chips. Sequencing reads for each sample from different sequencing runs were consolidated and downsampled to the same number of reads within each control and IL-13–treated pair, with a total number of reads within 8 million to 12 million reads per sample. Downsampled reads from each sample were pseudoligned and quantified using Kallisto (version 0.42.3) (42) and the human transcriptome sequence index generated from the iGenomes transcript annotation file (downloaded from Illumina’s FTP site on 01/15/2013). Estimated transcript counts from the Kallisto abundance files were rounded to the nearest integer and merged to form a count table for all 254 paired samples. Variance stabilizing transformation (VST) (DESseq2) was performed with an experimental design labeling samples based on IL-13 and control stimulation (blind=FALSE) (43). The NM_003856 transcript was used for expression of sST2. For the distal lung eQTL analysis, we used data generated by the GTEx consortium downloaded from database of Genotypes and Phenotypes (dbGap; http://www.ncbi.nlm.nih.gov/gap) (dbGap Study Accession: phs000424.v6.p1). We used raw counts generated by FluxCapacitor for the 320 GTEx_MidPoint_RNaseq samples (phe0000006.v2) from lung tissue, marked as best suited for eQTL analysis. We then selected 242 individuals with reported white race and available GTEx_MidPoint_Imputation genotypes (phg000520.v2). We then performed PCA analysis and identified 5 subjects who did not cluster with the remaining white-race individuals and those were removed from subsequent analyses, resulting in a total of 237 individuals with complete genotype and expression data. We performed variance stabilizing transformation on GTEx transcript expression. Transcript quantitation for GTEx was performed with the ensembl GTF; therefore, sST2 expression for this cohort was represented by the ENST00000311734.2 transcript. The ENST00000311734.2 transcript is equivalent to the RefSeq NM_003856 transcript, differing only in the 5′ end of the first exon.

qPCR for IL1RL1 exon 1a and 1b was performed on primary cultured airway epithelial cell RNA, lung microvascular endothelial cell RNA, and alveolar epithelial cell RNA. RNA was reverse transcribed as described above. Taqman primers and probes were then used in a standard 40-cycle taqman reaction using 2 μg of unamplified cDNA and Taqman Universal Master Mix (Invitrogen) to measure exon 1a, exon 1b,
and housekeeping genes PPIA, RPL13A, and EEF1A1. Primer and probe sequences are indicated in Supplemental Table 4.

**Blood Cell Cultures**

Human heparinized whole blood or mononuclear cells (PBMCs) were primed with IL-12 for 2 hours followed by the addition of IL-33 Cys-Ser variant (var) for 15–20 hours. IL-33 var was expressed and purified from *E. coli* by expression of a variant of IL-33, amino acids 112–270, with all 4 cysteines replaced by serine. The activity of IL-33 var is more stable in tissue culture than WT IL-33 (44). IFNγ was measured in the supernatant by mesoscale discovery immunoassays. EC50 of IL-33 was calculated following log transformation of the IFNγ levels. The type 1–stimulating conditions (IL-12 and IL-33) were used in these whole blood and PBMC assays rather than type 2–stimulating conditions (IL-33 and TSLP) because we find very few circulating type 2 cytokine–producing cells in the blood of healthy volunteers, resulting in very little type 2 cytokine response.

**Airway type 2 classification**

Data for the classification of asthmatics as type 2-high or -low was available from our prior studies (3, 20). In 76 subjects, the classification was made using a 3-gene expression signature of IL-13–responsive epithelial cell genes from bronchial brush RNA (1). The 3 genes are POSTN, SERPINB2, and CLCA1, and their individual expression values can be combined into a single 3-gene mean metric to determine the presence of type 2 inflammation in the airway epithelium, as previously described (20). In 84 subjects, the classification was made using gene expression for type 2 cytokines in induced sputum cells (3). The three type 2 cytokine genes are IL4, IL5, and IL13, and their individual expression values are combined into a similar 3-gene metric to determine the presence of type 2 inflammation in the airway lumen, as previously described (20). Ten subjects had measures of both bronchial brush and induced sputum type 2 inflammation. Subjects were classified as type 2-high if either the bronchial brush or the induced sputum cell standardized measurement exceeded 2 standard deviations of the mean value for healthy controls (3, 20).

**Genotyping**

UCSF ATB samples were genotyped for IL1RL1 SNPs rs1041973, rs1420101, rs10192157, and rs1921622 by Taqman genotyping assays. We used LNA Primetime probes and primers (IDT DNA) to genotype the rs11685480 SNP. Taqman and LNA reactions were performed with 5–10 ng of DNA according to manufacturer’s recommendations using either an ABI 7900HT or Quantstudio 6 Flex real-time thermocyclers (Invitrogen). Genotypes of the rs1420101 and rs1921622 SNPs for the donor cultured airway epithelial cells were extracted from Illumina Infinium Genome-wide genotyping array data. The rs11685480 SNP was genotyped in donor cultured airway epithelial cells with the assay and method described above. Genotype data for the rs11685480, rs1420101, and rs1921622 IL1RL1 SNPs in the GTEx lung donors was obtained from National Human Genome Research Institute (NHGRI) GTEx data release phs000424.v6.p1.

**Statistics**

*UCSF ATB genetic analyses.* Protein and gene expression data was transformed where appropriate to more closely approximate a normal distribution for genetic analysis. Specifically, normalized blood paxgene expression data was log2 transformed; normalized bronchial brush expression data was log2 transformed and then square root transformed. Blood sST2 protein levels were natural log (ln) transformed. Genetic association between IL1RL1 SNPs and gene and protein expression levels was tested by linear regression models coding the SNPs in an additive fashion (e.g., 0 = GG, 1 = AG, 2 = AA). UCSF ATB analyses of only white non-Hispanic subjects included asthma status, inhaled steroid use, age, and sex as covariates. The analysis of all subjects included the formerly stated covariates with the addition of 5 dichotomous variables for race/ethnicity based on self-identification, namely (i) white non-Hispanic, (ii) black non-Hispanic, (iii) Asian non-Hispanic and Native Hawaiian and Other Pacific Islanders, (iv) Hispanic any race, (v) and other, which included subjects reporting as mixed race non-Hispanic, white with no ethnicity, or non-Hispanic with American Indian/Alaska native race.

*Cultured human airway epithelial cells and GTEx lung genetic analyses.* VST normalized gene expression values, from airway epithelial cell cultures, were used to perform probabilistic estimation of expression residuals (PEER) normalization (45) and to generate PEER residuals corrected for plink (version 1.07)
determined sex, genome-wide genetic data determined genetic ancestry principle components (pc1, pc2, and pc3), and 64 hidden PEER factors (corresponding to ~25% of sample size). Similarly, GTEx lung VST-normalized gene expression data was used to perform PEER normalization and to generate PEER residuals corrected for age, sex, genome-wide genetic data–determined genetic ancestry principle components (pc1 and pc2), and 59 hidden PEER factors (corresponding to ~25% of sample size) for the sST2 transcript. Both the GTEx and cultured airway epithelial cell sST2 gene expression PEER residuals were analyzed for association with IL1RL1 SNPs by linear regression analysis using an additive genetic model. The airway epithelial bank IL-13–stimulated sST2 peer residuals were ln transformed for analysis. The untransformed control and square root–transformed IL-13–stimulated sST2 protein levels were also tested for association with IL1RL1 SNPs by linear regression analysis using an additive genetic model including sex and genetic principle components as covariates. Logistic regression was used to test the association between the rs1420101 and rs11685480 genotype and type 2-high vs. type 2-low status, including age, sex, steroid use, and racial/ethnic group (as defined above) as covariates. Between-group differences were assessed using a 2-tailed paired t test (cultured epithelial cell protein and gene expression unstimulated vs. IL-13–stimulated; distal vs. proximal promoter gene expression; EC50 IL-33 in whole blood vs. PBMC); 2-tailed unpaired t test (plasma sST2 protein, sST2 gene expression, or sST2 bronchial brush gene expression in healthy vs. asthma); ordinary 1-way ANOVA with Tukey’s multiple correction test (sST2 bronchial brush gene expression in healthy vs. type 2-high vs. type 2-low asthmatics; or ordinary 1-way ANOVA with post-test for linear trend (plasma sST2 protein in subjects carrying 0–5 risk alleles for rs1420101 or rs11685480). Linear regression was used to assess the relationship between whole blood EC50 for IL-33 and plasma sST2 protein LC-MS, as well as the relationship between plasma sST2 measured with LC-MS and critical diagnostics ELISA. A P value less than 0.05 was considered statistically significant. All statistical analyses were either performed in STATA/SE v11.0 or Graphpad Prism v6.0d.

Study approval
Research protocols were approved by the IRB at UCSF, and all subjects provided written informed consent for the study in which they originally participated. They also provided written informed consent for their biospecimens to be placed in the UCSF ATB for studies, in addition to the original protocol.

Author Contributions
EDG, MAS, and JVF conceived of the study. EDG, JP, MELS, RTS, KHN, JVF, and MAS designed experiments. RJM, JWL, and MAM provided reagents and technical support. EDG, JP, CLR, MELS, LZS, JLE, HJM, RTS, and MCP conducted experiments. EDG, AWA, LR, KHN, and MAS analyzed data. EDG, KHN, JVF, and MAS interpreted and synthesized the data. EDG generated figures. EDG and MAS wrote manuscript.

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