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Polymorphisms in the Muscarinic Receptor 1 Gene Confer Susceptibility to Asthma in Japanese Subjects

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Rationale: The human cholinergic receptor muscarinic-1 (CHRM1) is widely distributed in the lungs. In patients with asthma, CHRM1 may be involved in airway constriction, airway epithelial cell proliferation, and airway inflammation. The *CHRM1* gene is located on chromosome 11q13, which is one of the candidate loci for asthma and atopy.

Objectives: To determine the role of the *CHRM1* gene polymorphisms in asthma.

Methods: We studied nine single-nucleotide polymorphisms (–18379G > A, –9697C > T, –6965T > C, –4953A > G, +267A > C, +1353C > T, +3970C > G, +5418C > G, and +5455G > T) in a case-control study using 326 patients with asthma and 333 healthy control subjects. We also examined functional consequences of the –9697C > T and –4953A > G polymorphisms at the regulatory region using an mRNA reporter assay.

Measurements and Main Results: Two common single-nucleotide polymorphisms (–9697C > T and –4953A > G) were associated with asthma. The odds ratio for the TT homozygotes at the –9697C > T polymorphism was 0.29 compared with the CC homozygotes (95% confidence interval, 0.12–0.73; $p = 0.008$), and the odds ratio for the GG homozygotes at the –4953A > G polymorphism was 1.86 compared with the AA homozygotes (95% confidence interval, 1.04–3.34; $p = 0.038$). Haplotype analysis showed that the –9697T/–6965T/–4953A haplotype was associated with a lower risk of asthma ($p = 0.00055$) and the –9697C/–6965T/–4953G haplotype was associated with an increased risk of asthma ($p = 0.020$). The –9697T/–4953A haplotype was also associated with lower luciferase activity *in vitro* compared with the –9697C/–4953G haplotype.

Conclusions: This study, together with an *in vitro* functional study, suggests that the *CHRM1* gene is an important susceptibility locus for asthma on chromosome 11q13.

Keywords: case-control studies; IgE; muscarinic cholinergic receptor-1; single-nucleotide polymorphism

The cholinergic nerves are the dominant neural bronchoconstrictor pathway in humans (1). They release acetylcholine onto muscarinic receptors causing cholinergic bronchoconstriction (2), mucous hypersecretion, and edema in the airways. Increases in cholinergic nerve activity and cholinergic hypersensitivity are associated with asthma, and patients with asthma are hypersensitive to the bronchoconstricting actions of muscarinic agonists (3). The human cholinergic receptor muscarinic 1 (CHRM1; Online Mendelian Inheritance of Man database no. 118510) is widely localized in the human lung, including the alveolar walls,

AT A GLANCE COMMENTARY

Scientific Knowledge on the Subject

Genetic studies repeatedly have linked asthma and asthma-related phenotypes to chromosome 11q13, on which several biological candidate genes are located.

What This Study Adds to the Field

Gene coding the human cholinergic receptor muscarinic-1 (CHRM1) is an important susceptibility locus for asthma at chromosome 11q13.

bronchial epithelial cells, parasympathetic ganglia, neuromuscular junction, and submucosal glands (4). Studies using pirenzepine, a muscarinic antagonist selective for M1 receptors, have shown that M1 muscarinic receptors are involved in vagally induced bronchoconstriction (5–7). M1 receptor-deficient mice showed increased bronchoconstriction in response to 10^{-8} M muscarine in peripheral airways (8), suggesting the existence of an M1 receptor-dependent pathway counteracting cholinergic bronchoconstriction. M1 receptors also play a role in mast cell function (9), epithelial cell proliferation in the trachea (10), release of neutrophil and monocyte chemotactic activity from epithelial cells (11), acetylcholine-induced relaxation of the human pulmonary veins (12), and regulation of water and electrolyte secretion on submucosal glands (13). Taken together, CHRM1 is critically involved in the pathophysiology of asthma.

The gene encoding CHRM1 exists on chromosome 11q13, which has been linked to asthma and asthma-related phenotypes in several genomewide searches (14–17). Given the important role of muscarinic cholinergic mechanisms in asthma, the *CHRM1* gene is biologically an excellent candidate for asthma susceptibility in the region of chromosome 11q13. Thus, in the current study, we examined whether genetic variations in the *CHRM1* gene are associated with asthma. To gain insight into the possible molecular basis of the disease association, we also examined functional consequences of single-nucleotide polymorphisms (SNPs) at the regulatory region of the *CHRM1* gene.

METHODS

See online supplement for additional details.

Study Subjects

A total of 659 unrelated Japanese adults were enrolled in the study (Table 1). Asthma was defined on the basis of recurrent episodes of at least two of three symptoms (cough, wheeze, and dyspnea) that are associated with demonstrable reversible airflow limitation (15% variability in FEV₁ or in peak expiratory flow rate either spontaneously or with an inhaled, short-acting β_2 -agonist), or increased airway responsiveness to methacholine, or both, as described elsewhere (18).

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TABLE 1. BASIC CHARACTERISTICS OF 659 JAPANESE SUBJECTS

	Healthy Control Subjects (n = 333)	Subjects with Asthma (n = 326)	p Value*
Age, yr, median (range)	41 (18–72)	45 (16–79)	< 0.0001
Sex, n (male/female)	208/125	148/178	< 0.0001
Smoking, n (never/ex-/current)	226/12/95	190/69/67	< 0.0001
Atopy, n (%)	170 (51)	237 (73)	< 0.0001
Total serum IgE, log IU/ml, mean (SD)	1.84 (0.627)	2.40 (0.622)	< 0.0001
FEV ₁ , % predicted, mean (SD)	—	69.2 (13.5)	
% Reversibility in FEV ₁ , median (range)	—	16.9 (0–211)	

* One-way analysis of variance or χ^2 test was used where appropriate.

Identification of Polymorphisms

Genomic DNA from Japanese subjects was genotyped for +267A > C (rs2067477) and +1353C > T (rs2067480), because an association of these polymorphisms at the muscarinic M1 receptor gene with cognitive function in schizophrenic patients has been reported (19). We selected an additional seven SNPs for genotyping based on the frequency and location of SNPs and the linkage disequilibrium (LD) structure in and around the *CHRM1* gene. We initially obtained genotyping data of 26 HapMap SNPs (spanning 31.6 kb around the gene) from 45 unrelated Japanese subjects at the International HapMap Project (available online at <http://hapmap.org/>). To select tagSNPs in this region, we used the multimarker predictor method implemented in the Tagger program (20). Tag set was generated (using a threshold r^2 of 0.8) using 14 common SNPs with a minor allele frequency of more than 0.05 in the Japanese population.

As the +267A > C and +1353C > T polymorphisms were in a complete LD, we genotyped a total of eight SNPs (–18379G > A [rs1938677], –9697C > T [rs2075748], –6965T > C [rs542269], –4953A > G [rs1942499], +1353C > T [rs2067480], 3970C > G [rs4963323], 5418C > G [rs11601597], and 5455G > T [rs11605665]) for all individuals (n = 659). These SNPs were typed using the assay that combines kinetic (real-time quantitative) polymerase chain reaction (PCR) with allele-specific amplification, as described elsewhere (18). The PCR products were detected using the ABI 7700 Sequence Detection System with the dsDNA-specific fluorescent dye SYBR Green I (Applied Biosystems, Foster City, CA). The –4953A > G polymorphism was typed using TaqMan assay (Applied Biosystems).

Statistical Analysis

The association of the *CHRM1* gene polymorphism was measured by odds ratio (OR) with 95% confidence intervals (CI) as estimates of relative risk for the development of asthma. We used the Hardy-Weinberg equilibrium (HWE) program (21) to compare observed numbers of genotypes with the numbers of genotypes expected under HWE. For haplotype analyses, we used the Haplo.score program, which adjusts for covariates and calculates simulation p values for each haplotype (22).

Luciferase Reporter Gene Assay

We constructed two promoter reporter plasmids by placing two haplotypes (–9697C/–4953G and –9697T/–4953A) into the pGL3-Basic vector. Human neuroblastoma IMR32 cells (1×10^6) were transiently transfected with 9.5 μ g of one of the two constructs and 0.5 μ g of the pRL-TK vector, an internal control for transfection efficiency. After 24 h, we measured luciferase activity using the Dual-Luciferase Reporter Assay System (Promega, Tokyo, Japan). Results were expressed as means \pm SEM and were compared by paired *t* test.

Electrophoretic Mobility Shift Assay

Transcription factor (nuclear factor [NF]- κ B or upstream stimulating factor [USF]-1)-DNA binding activity was analyzed using the electrophoretic mobility shift assay (EMSA) kit (Panomics, Redwood, CA), according to the manufacturer's instructions.

RESULTS

Characteristics of the 333 healthy control subjects and 326 subjects with asthma are shown in Table 1. The median age of

subjects with asthma was significantly higher than in healthy control subjects ($p < 0.0001$). There were significantly more females in the asthma group than in the control group ($p < 0.0001$). Subjects with asthma were more likely to be atopic and had higher levels of total serum IgE than did healthy control subjects (χ^2 test or analysis of variance, $p < 0.0001$). More than 50% of the control subjects were atopic, which is consistent with

TABLE 2. COMPARISONS OF ALLELE AND GENOTYPE FREQUENCIES OF EIGHT *CHRM1* SINGLE-NUCLEOTIDE POLYMORPHISMS BETWEEN PATIENTS WITH ASTHMA AND CONTROL SUBJECTS

SNP	Allele/Genotype	HC n (%)	BA n (%)	p Value*
–18379 (rs1938677)	G	361 (54.5)	335 (48.1)	0.33
	A	301 (45.5)	311 (51.9)	
	GG	107 (32.3)	92 (28.5)	0.56
	GA	146 (44.1)	151 (46.7)	
	AA	78 (23.6)	80 (24.8)	
–9697 (rs2075748)	C	508 (76.3)	533 (81.7)	0.015
	T	158 (23.7)	119 (18.3)	
	CC	195 (58.6)	216 (66.3)	0.039
	CT	118 (35.4)	101 (31.0)	
	TT	20 (6.0)	9 (2.7)	
–6965 (rs542269)	T	491 (73.9)	472 (73.1)	0.72
	C	173 (26.1)	174 (26.9)	
	TT	184 (55.4)	175 (54.2)	0.94
	TC	123 (37.1)	122 (37.8)	
	CC	25 (7.5)	26 (8.0)	
–4953 (rs1942499)	A	477 (71.6)	434 (66.6)	0.047
	G	189 (28.4)	218 (33.4)	
	AA	174 (52.3)	147 (45.1)	0.15
	AG	129 (38.7)	140 (42.9)	
	GG	30 (9.0)	39 (12.0)	
+1353 (rs2067480)	C	615 (92.3)	608 (93.3)	0.52
	T	51 (7.7)	44 (6.7)	
	CC	284 (85.3)	287 (88.0)	0.19
	CT	47 (14.1)	34 (10.5)	
	TT	2 (0.6)	5 (1.5)	
+3970 (rs4963323)	C	533 (80.3)	524 (80.9)	0.78
	G	131 (19.7)	124 (19.1)	
	CC	216 (65.1)	211 (65.1)	0.75
	CG	101 (30.4)	102 (31.5)	
	GG	15 (4.5)	11 (3.4)	
+5418 (rs11601597)	C	414 (62.3)	385 (59.6)	0.31
	G	250 (37.7)	261 (40.4)	
	CC	132 (39.7)	113 (35.0)	0.44
	CG	150 (45.2)	159 (49.2)	
	GG	50 (15.1)	51 (15.8)	
+5455 (rs11605665)	G	538 (81.5)	496 (77.5)	0.074
	T	122 (18.5)	144 (22.5)	
	GG	218 (66.1)	192 (60.0)	0.19
	GT	102 (30.9)	112 (35.0)	
	TT	10 (3.0)	16 (5.0)	

Definition of abbreviations: BA = bronchial asthma; HC = healthy controls.

* χ^2 Test.

recent findings that the prevalence of atopy (as indicated by specific IgE against common inhaled allergens) among Japanese subjects is increasing (23, 24). Prebronchodilator baseline FEV₁ at an initial visit to our hospital was examined for 293 subjects with asthma (89.9%), and improvement of FEV₁ after bronchodilator therapy (400 µg salbutamol) or after a course of standard asthma medications (inhaled corticosteroids, long-acting β₂-agonists, theophylline, or leukotriene modulators) was recorded for 214 (65.6%) subjects with asthma (Table 1).

All eight of the SNPs investigated were within the HWE in the control group ($p > 0.05$). The overall success rate for genotyping was 99.6%. Of the eight SNPs, two common SNPs ($-9697C > T$ [rs2075748] and $-4953A > G$ [rs1942499]) in the regulatory region of the *CHRM1* gene had a significant association with asthma (Table 2). Both of these SNPs were significantly associated with asthma when the analysis was adjusted for age, sex, smoking status, and atopic status (Table 3). The OR for the TT homozygotes of the $-9697C > T$ polymorphism was 0.29 compared with the CC homozygotes (95% CI, 0.12–0.73; $p = 0.008$), and the OR for the GG homozygotes of the $-4953A > G$ polymorphism was 1.86 compared with the AA homozygotes (95% CI, 1.04–3.34; $p = 0.038$).

We analyzed data from the eight SNPs with the Haploview program (25) and identified two haplotype blocks (Figure 1) in our case-control population. Haplotype block I comprised three SNPs in the regulatory region ($-9697C > T$ [rs2075748], $-6965T > C$ [rs542269], $-4953A > G$ [rs1942499]), and haplotype block II comprised three SNPs ($+1353C > T$ [rs2067480], $+3970C >$

G [rs4963323], $+5418C > G$ [rs11601597]) in the coding exon and the 3'-UTR. Haplotype analyses were performed in both blocks I and II. The frequency of *CHRM1* haplotypes is shown in Table 4. In block I, the $-9697T/-6965T/-4953A$ haplotype was associated with a significantly lower risk of asthma ($p = 0.00055$) and the $-9697C/-6965T/-4953G$ haplotype was associated with a significantly increased risk of asthma ($p = 0.020$). Inspection of specific haplotypes revealed that this association is most likely due to $-9697C > T$ and $-4953A > G$, because the same allele for $-6965T > C$ is part of both risk and protective haplotypes. In contrast, none of the haplotypes in block II was associated with asthma.

In the case-only study, associations between asthma-related phenotypes, such as total serum IgE levels and atopy, and the polymorphisms of *CHRM1* were also investigated. We could not find any significant association between the genotypes of the eight SNPs and total serum IgE levels or atopy (see Tables E1 and E2 in the online supplement).

The transcriptional activity of the *CHRM1* SNPs at the regulatory region was compared between the $-9697C/-4953G$ haplotype and the $-9697T/-4953A$ haplotype transiently transfected into human neuroblastoma IMR32 cells. Luciferase activity in cell extracts was assessed 24 h after transfection, and was expressed as fold increase in the activity of the *CHRM1* reporter constructs compared with the pRL-TK vector. Figure 2 shows that the reporter plasmid carrying the $-9697T/-4953A$ promoter displayed 37% lower transcriptional activity compared with the plasmid carrying the $-9697C/-4953G$ promoter ($p = 0.019$).

TABLE 3. GENETIC IMPACT ON ASTHMA OF EIGHT SINGLE-NUCLEOTIDE POLYMORPHISMS IN AND AROUND THE *CHRM1* GENE

SNP	Genotype	OR (95% CI)	
		Adjustments (–)	Adjustments (+)*
$-18379G > A$ (rs1938677)	GG	1 (Reference)	1 (Reference)
	GA	1.20 (0.84–1.72)	1.14 (0.77–1.71)
	AA	1.19 (0.79–1.81)	1.33 (0.84–2.12)
$-9697C > T$ (rs2075748)	CC	1 (Reference)	1 (Reference)
	CT	0.77 (0.56–1.07)	0.73 (0.51–1.06)
	TT	0.41 (0.18–0.91) [†]	0.29 (0.12–0.73) [‡]
$-6965T > C$ (rs542269)	TT	1 (Reference)	1 (Reference)
	TC	1.04 (0.75–1.44)	1.00 (0.70–1.44)
	CC	1.09 (0.61–1.97)	1.04 (0.54–2.02)
$-4953A > G$ (rs1942499)	AA	1 (Reference)	1 (Reference)
	AG	1.28 (0.93–1.78)	1.38 (0.96–1.98)
	GG	1.54 (0.91–2.60)	1.86 (1.04–3.34) [†]
$+1353C > T$ (rs2067480)	CC	1 (Reference)	1 (Reference)
	CT	0.72 (0.45–1.15)	0.61 (0.37–1.02)
	TT	2.47 (0.48–12.9)	1.96 (0.31–12.5)
$+3970C > G$ (rs4963323)	CC	1 (Reference)	1 (Reference)
	CG	1.03 (0.74–1.44)	1.12 (0.78–1.62)
	GG	0.75 (0.34–1.67)	0.78 (0.33–1.86)
$+5418C > G$ (rs11601597)	CC	1 (Reference)	1 (Reference)
	CG	1.24 (0.88–1.73)	1.29 (0.89–1.88)
	GG	1.19 (0.75–1.89)	1.16 (0.70–1.93)
$+5455G > T$ (rs11605665)	GG	1 (Reference)	1 (Reference)
	GT	1.25 (0.90–1.74)	1.28 (0.89–1.85)
	TT	1.82 (0.81–4.1)	1.73 (0.73–4.09)

Definition of abbreviations: CI = confidence interval; OR = odds ratio.

* Adjustment for matching factors and potential confounding factors, including sex, age, smoking status, and atopic status, was performed by unconditional logistic regression analysis.

[†] $p < 0.05$.

[‡] $p < 0.01$.

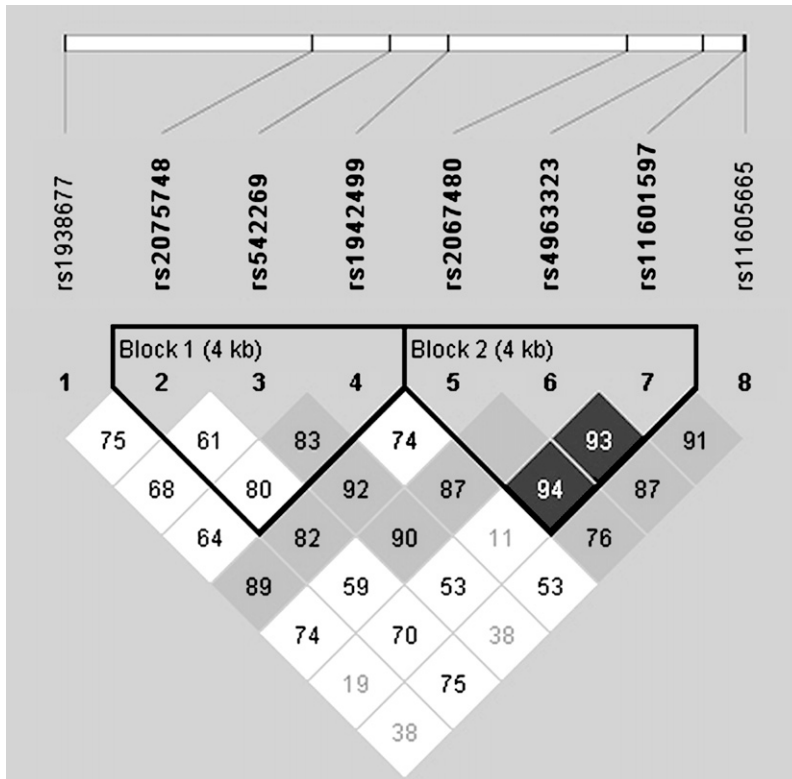


Figure 1. Locations and linkage disequilibrium (LD) map structure of single-nucleotide polymorphisms (SNPs) in and around the *CHRM1* Haploview plot shows pairwise LD (D' values) for 8 SNPs based on genotypes of 659 individuals of the case-control study. The eight SNPs include $-18379G > A$ [rs1938677], $-9697C > T$ [rs2075748], $-6965T > C$ [rs542269], $-4953A > G$ [rs1942499], $+1353C > T$ [rs2067480], $+3970C > G$ [rs4963323], $+5418C > G$ [rs11601597], and $+5455G > T$ [rs11605665]. LD blocks are framed in *black* and were classified according to the "solid spine" option (25). Each square plots the level of LD (D' values) between a pair of SNPs.

This difference in transcriptional activity was consistent in eight independent experiments.

EMSA failed to show a robust difference in binding affinity of NF- κ B to the $-9697T$ or the $-9697C$ allele, or in binding affinity of USF-1 to the $-4953A$ or the $-4953G$ allele (data not shown).

DISCUSSION

Given a high *a priori* biological plausibility for asthma, we tested the hypothesis that the allelic variants at the regulatory region

of the *CHRM1* confer susceptibility to asthma by conducting a case-control study in a relatively large population of unrelated Japanese subjects. In accordance with our primary hypothesis, we found that the presence of the $-9697CC$ genotype, $-4953GG$ genotype, or the $-9697C/-4953G$ haplotype at the regulatory region was significantly associated with a diagnosis of asthma. Our genetic association study had several strengths: first, muscarinic receptors, including M1, have been biologically implicated in the pathogenesis of asthma; second, the gene encoding the *CHRM1* is located on chromosome 11q13, a genomic region

TABLE 4. ESTIMATED HAPLOTYPE FREQUENCIES OF THE *CHRM1* GENE POLYMORPHISMS

		Haplotype Frequency		Haplotype-specific Score	p Value* (Empirical)
		Control (n = 333)	Asthma (n = 326)		
Block I					
Haplotype	$-9697/-6965/-4953$				
1	T T A	0.207	0.158	-3.03	0.00055
2	C C A	0.239	0.234	-0.23	0.827
3	C T A	0.250	0.254	0.057	0.959
4	C T G	0.268	0.303	2.15	0.020
Block II					
Haplotype	$+1353/+3970/+5418$				
1	T C C	0.371	0.391	0.120	0.131
2	C C C	0.353	0.349	0.808	0.814
3	C G C	0.192	0.179	0.926	0.918
4	C C G	0.371	0.391	0.454	0.481

Haplotype frequencies were estimated using the Haplo.Stats program. In Block I (regulatory region of the gene), haplotype analyses showed that the $-9697T/-6965T/-4953A$ haplotype was associated with a significantly lower risk of asthma ($p = 0.00055$) and the $-9697C/-6965T/-4953G$ haplotype was associated with a significantly increased risk of asthma ($p = 0.020$). In contrast, in Block II, none of haplotypes showed a significant association with asthma. Note that haplotype-specific scores give effect estimates; negative haplotype-specific scores are associated with a protective effect, and positive haplotype-specific scores are associated with an increased risk. Haplotypes with frequencies less than 0.05 were excluded from the analyses.

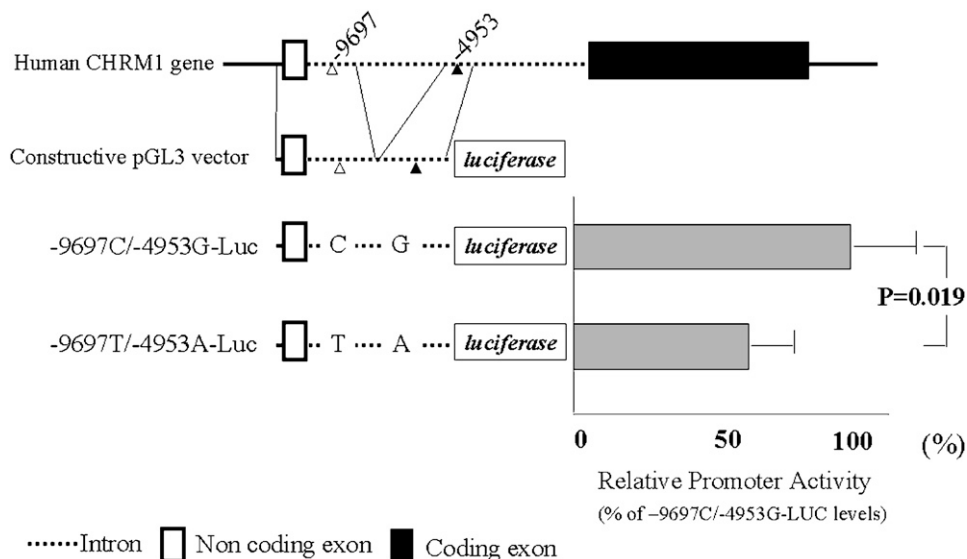


Figure 2. Transcription activity analysis of the promoter region of *CHRM1* IMR32 cells were transiently transfected with 9.5 μ g of *CHRM1* luciferase reporter constructs (schematically depicted to the left) plus 0.5 μ g of pRL-TK vector. Luciferase activities were normalized against the internal control Renilla values. The data represent means \pm SD for the entire dataset (four independent experiments, each in duplicate), and are expressed as a percentage of the $-9697C/-4953G$ LUC activity. The difference between these constructs was significant at $p = 0.019$ (paired *t* test; $n = 8$).

linked to the diagnosis of asthma and atopy in several genome-wide scans (14–17); third, *in vitro* functional analyses have shown that the haplotype at the regulatory region has an effect on a basal promoter function in IMR32 cells, with the $-9697T/-4953A$ haplotype associated with 37% decreased promoter activity compared with the $-9697C/-4953G$ haplotype. Accordingly, our data suggest that the $-9697T/-4953A$ haplotype may influence the affinity of a particular nuclear protein to the regulatory region of the *CHRM1* gene, resulting in altered transcriptional activity and ultimately leading to a higher or lower risk of asthma.

Although the exact mechanisms underlying the involvement of the *CHRM1* gene in the pathogenesis of asthma remain to be identified, several reports indicate that the cholinergic pathway has an important role in the pathogenesis of asthma, in particular in the regulation of bronchoconstriction, airway inflammation, and airway remodeling. An M1 receptor-dependent pathway counteracts cholinergic bronchoconstriction, possibly via the release of a relaxing agent (8); both respiratory epithelia and sympathetic nerve terminals within bronchial smooth muscle are equipped with M1 receptors (7, 26) and releasable bronchodilating agents, such as nitric oxide and prostaglandin E_2 (27). Studies with the M1 receptor-preferring antagonist, pirenzepine, have also suggested the existence of pulmonary M1 receptors modulating airway diameter (28). Furthermore, Jones and colleagues (29) demonstrated that stable expression in RBL-2H3 mast cells of the M1 muscarinic acetylcholine receptor leads to carbachol-stimulated mast cell degranulation. An animal model of asthma showed that anticholinergic agents protect against allergen-induced airway remodeling (30). Together with these *in vivo* and *in vitro* findings, our findings support the contention that *CHRM1* plays an important role in the pathogenesis of asthma. Our findings may be of considerable relevance to asthma treatment, providing an important basis for identification of individuals for whom the cholinergic pathway could be targeted.

Sequence analysis indicated that the T allele at the $-9697C > T$ polymorphism creates a potential NF- κ B binding site and that the A allele at the $-4953A > G$ polymorphism creates a potential USF-1 transcription factor binding site by reference to the MatInspector or TFSEARCH database (31). We, however, failed to see any difference in binding intensities of these nuclear factors to the $-9697C > T$ or $-4953A > G$ polymorphism.

Therefore, we cannot exclude the possibility that these SNPs might not be causative in nature, but are in LD with a true susceptibility allele in the regulatory region of the *CHRM1* gene. Population stratification may influence the observed associations (32). However, our population is racially homogeneous, as all subjects recruited in the study were from the Japanese population, which is considered monoracial; thus, our subjects had a relatively low risk of population stratification effects. Furthermore, we recruited all participants in the current study from a single institute to minimize the chance of mixing populations with inherently diverse allele frequencies of a susceptibility gene. In addition, all SNPs were in HWE in a set of unrelated healthy subjects. Therefore, we believe that the usual problems associated with population stratification may be of limited importance in the present study. Nevertheless, we acknowledge that population stratification may have influenced the present findings, and that the findings of the current study are preliminary and do not, by themselves, conclusively confirm an etiologic relationship. A more comprehensive approach that examines the functional consequences of the *CHRM1* promoter polymorphisms and identifies the possible promoter-dependent mechanism for an association between *CHRM1* and asthma is required.

In conclusion, given the important role of muscarinic cholinergic mechanisms in pulmonary disease, this case-control study, together with an *in vitro* functional analysis, suggests that the *CHRM1* gene is an important susceptibility locus for asthma at chromosome 11q13. The $-9697T/-4953A$ haplotype at the regulatory region of the gene may contribute to the development of asthma by altering the human lung muscarinic receptor system in ways that could account for the increased *in vivo* lung cholinergic hyperresponsiveness found in patients with asthma.

Conflict of Interest Statement: None of the authors has a financial relationship with a commercial entity that has an interest in the subject of this manuscript.

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