Variant LTC₄ synthase allele modifies cysteinyi leukotriene synthesis in eosinophils and predicts clinical response to zafirlukast

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Clinical trials with leukotriene synthesis inhibitors and cysteinyi leukotriene (cys-LT) receptor antagonists have shown that cys-LTs (LTC₄, LTD₄, and LTE₄) are the predominant mediators of bronchoconstriction in response to diverse asthma triggers including allergen, PAF, sulphur dioxide, and non-steroidal anti-inflammatory drugs (NSAIDs). In addition, inhalation of a single dose of a cys-LT causes persistent eosinophilia in the bronchial mucosa and sputum of asthmatic patients, and LT modifier drugs significantly reduce airway and blood eosinophilia in clinical asthma.

We have shown that, in the asthmatic airway, eosinophils represent the majority of cells that express the terminal enzyme in the cys-LT pathway, LTC₄ synthase, and hence have the capacity to generate cys-LT upon stimulation. Inflammation and bronchoconstriction in the asthmatic airway may be sustained by a vicious cycle of cys-LT synthesis and eosinophil recruitment (fig 1). Anomalies in the genetic and cytokine factors that regulate cys-LT pathway enzymes in eosinophils may thus be central to the pathophysiology of asthma.

Cys-LTs are synthesised from membrane derived arachidonic acid. During stimulus-specific cell activation, arachidonic acid released by phospholipases including cytosolic phospholipase A, (PLA₂) is translocated to the 5-lipoxygenase activating protein (FLAP) and converted in two steps to leukotriene (LT)₄, by 5-lipoxygenase (5-LO). LT₄ is converted to LTD₄ by cells expressing LT₄ hydrolase, and/or to LTE₄ by cells expressing LTC₄ synthase, which conjugates LT₄ to reduced glutathione. After carrier mediated cellular export of LTC₄, the sequential cleavage of glutamate and glycine residues provides the extracellular receptor active metabolites LTD₄ and LTE₄, respectively. Cys-LTs act at specific cys-LT receptors on target cells including bronchial smooth muscle and inflammatory leucocytes.

The cys-LT pathway in eosinophils may be modulated by the cytokine microenvironment in the bone marrow, circulation, and airway. Maturation of eosinophils in vitro by the eosinophilopoietic cytokines interleukin (IL)-5 and IL-3 is accompanied by sequential expression of PLA₂, 5-LO, FLAP, and LTC₄ synthase, associated with acquisition of a cys-LT synthesising phenotype. In asthmatics, enhanced capacity of blood eosinophils to generate LTC₄, may be caused by cytokine induction of 5-LO pathway enzyme expression, including FLAP. Induction of 5-LO pathway enzyme expression by the cytokine microenvironment may also be observable within the bronchial mucosa of mild asthmatics naturally exposed to a seasonal aeroallergen.

The capacity for cys-LT synthesis of eosinophils may also be influenced by promoter polymorphism in the genes encoding cys-LT pathway enzymes. Promoter polymorphism in 5-LO is linked to modified gene transcription and a diminished clinical response to an LT synthase inhibitor in a small proportion (∼5%) of asthmatics, but this cannot account for the large subgroup of poor responders to LT modifier drug therapy.

In bronchial biopsy specimens of patients with aspirin intolerant asthma (AIA) who have a constitutively high baseline production of cys-LTs and generally good clinical responses to leukotriene modifier drugs, the numbers of cells expressing LTC₄ synthase are four times more prevalent than in biopsy specimens from aspirin tolerant asthmatics (ATA) and 19 times more prevalent than in normal bronchial biopsy specimens. The increased prevalence of LTC₄ synthase in biopsy specimens from patients with AIA is partly the result of an

Figure 1  Schematic diagram illustrating that genetic and cytokine factors modulate the vicious cycle of cysteinyi leukotriene synthesis and eosinophil recruitment in the asthmatic airway.
increased proportion of eosinophils expressing the enzyme (50% versus 21% in patients with ATA) and correlates exclusively with raised cys-LTs levels in bronchoalveolar lavage (BAL) fluid and with bronchial hyperresponsiveness to inhaled lysine-aspirin. A single nucleotide polymorphism in the LTC₄ synthase gene promoter (A₁₄C) has been reported in 76% of patients with AIA compared with 42–44% in those with ATA and normal subjects. The variant (C) allele is associated with higher levels of LTC₄ synthase mRNA in blood eosinophils, suggesting that the variant allele increases expression of LTC₄ synthase and hence the capacity of these cells to synthesise cys-LT.

Since the variant LTC₄ synthase genotype is found in a significant proportion of patients with ATA and normal subjects, we hypothesised, firstly, that eosinophils from subjects with the variant genotype may have a predisposition to enhanced synthesis of cys-LTs when the 5-LO pathway is stimulated. Secondly, we hypothesised that the variant LTC₄ synthase genotype may be more prevalent in severe persistent asthmatics, irrespective of aspirin sensitivity, than in normal subjects. Thirdly, we hypothesised that severe asthmatics with the variant genotype may represent a subgroup of patients in whom increased cys-LT synthesis is particularly important in impairing lung function, and hence that these patients may be synonymous with a phenotype of “responders” to anti-LT therapy.

Methods
Normal subjects (n = 31) were adult volunteers with no history of chronic respiratory disease who were taking no medication. Four normal subjects were atopic as judged by skin prick tests. The 23 asthmatic patients (6M, 17F; aged 25–76 years) had a clinical history of chronic, severe asthma uncontrolled by their medication. Three asthmatic patients reported a clinical history of aspirin sensitivity. All 23 asthmatic subjects were receiving inhaled corticosteroids, and 10 were also receiving oral steroids, 14 were receiving a long acting β₂ agonist, four were receiving theophylline, and one cyclosporin. At pre-study assessment, forced expiratory volume in one second (FEV₁) varied between 0.24 and 2.5 litres and peak expiratory flow (PEF) between 106 and 408 l/min. During the study asthmatic subjects abstained from all medication except inhaled corticosteroids and inhaled β₂ agonists as required, and received oral zafirlukast (20 mg bd) for two weeks when lung function assessments were repeated. Although an open study of zafirlukast, patients and investigators were blinded to patient genotype.

Eosinophils were purified from the blood of normal subjects and their capacity for LTC₄ synthesis assessed as previously described. Briefly, mononuclear cells and platelets were removed by Percoll discontinuous density gradient centrifugation, erythrocytes by hypotonic lysis, and neutrophils by negative immunomagnetic selection (MAC system) using anti-CD16 magnetic beads. Resulting eosinophils (>98% pure, >96% viable) were cultured (10⁷ cells/0.5 ml) in HEPES buffered RPMI1640 medium at 37°C for 15 minutes in the presence and absence of calcium ionophore A23187 (1 µM) and in the presence of the cyclo-oxygenase inhibitor indomethacin (10 µM). Following extraction of intracellular LTs by methanol treatment, total cys-LT synthesis was quantified by Biotrak enzyme immunoassay (EIA; Amersham, UK). Urinary LTE₄ levels were assessed as a marker of whole body cys-LT synthesis in normal subjects by EIA and corrected for creatinine.

Genomic DNA was isolated from whole blood using the QIAamp DNA Blood Mini Kit (Qiagen). The region flanking the A₁₄C polymorphism was amplified using PCR (forward primer 5’ CTC CAT TCT GAA GCC AAA G 3’; reverse primer: 5’ AGA CCG CCT CAC CAC TT 3’). Alleles were genotyped using MspI restriction digestion of the 296 bp PCR product; an endogenous MspI site provided an internal control for the assay.

The study was approved by the Southampton & SouthWest Hampshire Joint Research ethics committee and all subjects gave informed written consent to participate.

Results
The variant LTC₄ synthase genotypes (C/C and C/A) were observed in 13 out of 23 severe asthmatic patients (56%) compared with only 10 of 31 normal subjects (32%). The frequency of the variant allele (q) was thus 0.27 in the asthmatic patients and 0.19 in the normal subjects. Despite the small size of the subject...
results presented here provide preliminary evidence that, irrespective of aspirin sensitivity, the variant genotypes confer an increased capacity for cys-LT synthesis in eosinophils, the predominant cell type expressing LTC₄ synthase in the asthmatic airway, and that in severe asthmatic patients, the variant genotypes may be markers of a leukotriene dependent phenotype of asthma that responds well to treatment with a leukotriene receptor antagonist.

Differences in LTC₄ synthase expression in eosinophils and other leukocytes may only be rate limiting for LTC₄ synthesis when the 5-LO pathway is highly stimulated. Thus, in eosinophils from normal subjects, there was no difference in LTC₄ synthesis in vitro between the variant and wild type genotypes in the absence of ionophore stimulation, and low levels of leukocyte activation in vivo may also explain the lack of difference in urinary LTE₄ levels (fig 2B). However, stimulation of normal eosinophils with calcium ionophore revealed significantly increased LTC₄ synthesis in subjects with the variant LTC₄ synthase genotype (fig 2A). The difference became statistically significant in the presence of indomethacin, which prevents shunting of arachidonate along the cyclo-oxygenase (COX) pathway. This may explain why in most patients with AIA, who have a high prevalence of the variant LTC₄ synthase allele, exposure to aspirin and other COX inhibitors consistently provokes an acute surge of cys-LT synthesis leading to acute bronchoconstriction.

The increased prevalence of the variant LTC₄ synthase genotype in the patients with severe asthma (56%) compared with the normal subjects (32%), and the concomitant increase in the capacity for LTC₄ production in eosinophils when stimulated, suggests that a genetic propensity for enhanced cys-LT production in vivo may be a predisposing factor for severe asthma irrespective of aspirin sensitivity.

The study of two weeks treatment with oral zafirlukast in patients with severe chronic asthma was an open study involving a small number of patients (n = 23). Withdrawal of oral corticosteroids and other asthma medication before the study probably accounts for the deterioration in lung function observed in patients with the wild type LTC₄ synthase genotype. Against this background, patients with the variant genotypes showed a tendency to improvement in two measures of lung function (FEV₁ and FVC; fig 3). Although not reaching statistical significance (p = 0.1), the results suggest that enhanced cys-LT synthesis contributes disproportionately to the pathophysiology of asthma in the 56% of patients with the variant LTC₄ synthase genotype, and that these patients may represent the target group of good responders to anti-leukotriene therapy. Large scale clinical studies are planned to investigate this further.

**Figure 3** Percentage changes in lung function compared with pre-study values in severe asthmatics with variant (C/C and C/A) and wild type (A/A) LTC₄ synthase genotypes following oral therapy with the leukotriene receptor antagonist zafirlukast (Accolate; 20 mg bd) for two weeks.

In normal subjects spontaneous production of LTC₄ from unstimulated blood eosinophils was not significantly different between the two genotype subgroups. However, when stimulated with calcium ionophore A23187 (1 µM) in the presence of indomethacin (10 µM) to block shunting of arachidonate substrate along the prostanoid pathway, blood eosinophils from patients with variant genotypes (C/C and C/A) produced approximately three times more LTC₄ (7.8 (2.9) ng/10⁶ cells, n = 13) than eosinophils from patients with the wild type genotype (A/A, 2.7 (0.4) ng/10⁶ cells, n = 10; p = 0.04, Mann-Whitney test; fig 2A). There was no significant difference between normal subjects with the variant genotypes and those with the wild type genotype in urinary levels of LTE₄ (145 (21) pmol/mmol versus 128 (8) pmol/mmol creatinine, respectively; p > 0.2; fig 2B).

In asthmatic subjects with variant LTC₄ synthase genotypes, oral administration of zafirlukast (20 mg bd) for two weeks increased FEV₁ by 9 (12)% and forced vital capacity (FVC) by 15 (18)% compared with pre-study values (n = 13), with decreases of –12 (18)% and –18 (15)% respectively, in patients with the wild type genotype (n = 10; both p = 0.1; fig 3). Home PEF values did not differ between the groups.

**Discussion**

The capacity of leukocytes to generate cys-LTs in response to allergic and non-allergic triggers may be modulated by the cytokine microenvironment in the asthmatic airway and by genetic factors (fig 1). The single nucleotide polymorphism (A₄₄₄C) described in the LTC₄ synthase gene promoter has been postulated to enhance gene transcription and enhance cellular capacity for cys-LT synthesis.¹¹ The variant LTC₄ synthase genotypes are reported to be highly prevalent (76%) in patients with aspirin intolerant asthma¹⁶ who have constitutively increased cys-LT synthesis and good clinical responses to anti-leukotriene therapy, and are also common (42–44%) in normal subjects and patients with aspirin tolerant asthma. The results presented here provide preliminary evidence that elevated LTC₄ synthase genotypes may be important in the evaluation of disease severity, and that these patients may represent the target group of good responders to anti-leukotriene therapy. Large scale clinical studies are planned to investigate this further.

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