

Treatment strategies for allergy and asthma

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Abstract | Allergic diseases have reached epidemic proportions worldwide. An understanding of the cellular and soluble mediators that are involved in allergic inflammatory responses not only helps in understanding the mechanisms of current treatments, but is also important for the identification of new targets that are amenable to both small-molecule and biological interventions. There is now considerable optimism with regards to tackling the allergy epidemic in light of improvements in systemic and mucosal allergen-specific immunotherapy, the identification of key cytokines and their receptors that drive T-helper-2-cell polarization, a clearer understanding of the pathways of leukocyte recruitment and the signalling pathways that are involved in cell activation and mediator secretion, and new approaches to vaccine development.

T helper 2 cells

(T_H2 cells). A T-helper-cell subset that has an important role in humoral immunity and in allergic responses. T_H2 cells produce cytokines that regulate IgE synthesis (IL-4), eosinophil proliferation (IL-5), mast-cell proliferation (IL-9) and airway hyper-responsiveness (IL-13). A T_H2 -cell pattern of cytokine expression is observed in allergic inflammation and in parasitic infections, conditions that are both associated with IgE production and eosinophilia.

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Allergic diseases are those that are mediated by the expansion of the T helper 2 cell (T_H2 -cell) subset of T cells, together with isotype switching of B cells to generate IgE antibodies specific for common environmental allergens¹. Although almost half of the urban population worldwide is atopic (genetically predisposed to produce IgE antibodies in serum) and most allergy sufferers are atopic, it is possible to develop allergies in the absence of atopy — common examples of this are wasp and bee allergies. Allergic reactions are symptomatic responses to a normally innocuous environmental antigen. Allergic diseases are highly patient specific and include asthma, rhinoconjunctivitis, sinusitis, food allergy, atopic dermatitis, angioedema and urticaria, anaphylaxis, and insect and drug allergy, all of which can occur either alone or in combination. Allergic diseases have reached epidemic proportions worldwide and their incidence is continuing to increase in association with the Western lifestyle, making it imperative that we continue to improve our understanding of the mechanisms of allergic disease². Allergies can affect all age groups and can appear at any time, but it is the marked increase of allergies in children and young adults that is of particular concern³. It is probable that environmental exposure to potential allergens early in life or during pregnancy has a pivotal role in the development of allergy and its expression in different organs (BOX 1).

Since the first description of 'reagin' in 1922 and its characterization as IgE in 1966 (REF. 4), considerable advances have been made in understanding the immunological pathways that lead to allergy, the chemical

mediators involved and the innate and adaptive immune responses that regulate its expression. Progress has also occurred in understanding the structural and intrinsic biology of environmental agents that determine their allergenicity, and the role of other environmental factors, such as pollutants and microorganisms, in augmenting or inhibiting allergen sensitization⁵. As most allergic disorders express themselves clinically at epidermal or mucosal surfaces, a breakdown of the physical barrier that is normally provided by these surfaces and altered innate immunity is also recognized to be of great importance in allergic reactions. The study of interactions between many susceptibility genes and the environment is revealing new pathophysiological mechanisms and creating unique opportunities for the prevention and treatment of allergy.

The time has arrived to take a fresh look at the new treatment strategies that are now appearing on the horizon. This Review provides an overview of possible innovative therapeutics, focusing on some exciting new immunological and molecular targets and on vaccine approaches for allergy and asthma.

The allergic cascade

The allergic inflammatory response is characterized by selection of the T_H2 -cell pathway, which is initiated by the uptake of allergens by professional antigen-presenting cells (APCs) that present selected peptides on MHC class II molecules to naive T cells, thereby directing them in favour of a T_H2 -cell phenotype in which the transcription factor GATA3 (GATA-binding

Box 1 | Early-life influences in the development of allergy and asthma

Environmental exposure to allergens that occurs early in life or during pregnancy is thought to initiate the allergic response to increase T helper 2 (T_H2)-cell and decrease T_H1-cell and regulatory T-cell subpopulations.

Strong allergy-protective factors have been observed in farming environments. This protective effect can be explained by dietary habits, lifestyle and environmental exposures, and it is associated with an increased production of the T_H1-cell cytokines tumour-necrosis factor and interferon- γ by cord-blood mononuclear cells. Nutritional factors that affect the development of allergic diseases early in life include breastfeeding, dietary fatty acids, antioxidants, and foods that affect the gut microflora. In many industrialized societies, the decreased consumption of ω 3-fatty acids, which have been replaced by an increased intake of ω 6-fatty acids, favours a T_H2-cell phenotype during ontogeny and development.

An imbalance between reactive oxygen and antioxidants is another factor that contributes to the chronic inflammatory process in asthma. Deficiency in dietary antioxidants and vitamins in the Western diet, with decreased consumption of fresh fruit and vegetables, correlates with changes in the prevalence of allergies and asthma.

The mammalian gastrointestinal tract harbours a complex ecosystem consisting of bacteria that are in homeostasis with the host immune system, and a disturbance of this homeostasis might have a role in the development of allergy. Exposure to microbial products skews the T-cell balance towards a T_H1-cell response by interacting with Toll-like receptors (TLRs) expressed by dendritic cells. Environmental indicators of microbial exposure — such as the presence of endotoxin, muramic acid, β (1,3)-glucans and extracellular polysaccharides from *Aspergillus* spp. and *Penicillium* spp. — are inversely related to the incidence of allergic diseases. Although there have been attempts to harness protection against allergy by using microbial products such as *Mycobacterium vaccae* to stimulate a T_H1-cell response, a more promising approach is the application of specific TLR ligands such as CpG oligonucleotides (which bind TLR9) and chitin (which binds TLR2 and TLR4).

protein 3) mediates cytokine secretion⁶ (FIG. 1). This is in contrast to the T_H1-cell phenotype that is dominant in autoimmune diseases, in which T-bet controls cytokine secretion — for example, the secretion of interferon- γ (IFN γ)⁷. The crucial role played by dendritic cells (DCs) acting as professional APCs in this sensitization process is reviewed by Hamida Hammad and Bart Lambrecht in this issue⁸. B cells are also important for allergen capture and processing, especially in the presence of small amounts of allergen⁹.

In the presence of co-stimulation, T cells coordinately upregulate expression of a cluster of genes encoded on human chromosome 5q31–33 that include the genes encoding interleukin-3 (IL-3), IL-4, IL-5, IL-9, IL-13 and granulocyte/macrophage colony-stimulating factor (GM-CSF)¹⁰. These cytokines are involved in the class-switching of B cells to IgE synthesis (IL-4 and IL-13), the recruitment of mast cells (IL-4, IL-9 and IL-13) and the maturation of eosinophils (IL-3, IL-5 and GM-CSF) and basophils (IL-3 and IL-4), which are the main mediator-secreting effector cells of the allergic response. The imbalance between T_H2-cell and T_H1-cell (that is, IFN γ -producing) responses has formed the basis for our understanding of allergic immune responses for more than two decades¹¹. More recently, regulatory T cells (T_{Reg} cells) have been discovered as another pivotal subset of CD4⁺ T cells with implications for allergic disease. These cells are characterized by expression of the transcription factor FOXP3 (Forkhead box P3) and the IL-2 receptor (CD25)¹², but in contrast to activated effector T cells, they express low levels of CD127 (REF. 13). Extensive studies in mouse models

strongly implicate T_{Reg} cells in the suppression of allergic responses¹⁴, and there is emerging evidence that T_{Reg} cells also control T_H2-cell responses in humans through the inhibitory cytokines IL-10 and transforming growth factor- β (TGF β), with atopy resulting from an imbalance between T_H2 cells and T_{Reg} cells¹⁵.

Another newly identified type of CD4⁺ T cell has been named the T_H17 cell on the basis of secretion of IL-17A and IL-17F, which are associated with neutrophilic inflammation¹⁶. The transcription factor ROR γ t (retinoic-acid-receptor-related orphan receptor- γ t) identifies T_H17 cells and is selectively activated by IL-1 β and IL-6 (REF. 17), with IL-23 being responsible for the proliferation of these cells. IL-17A is overexpressed in asthmatic airways in association with neutrophil influx¹⁸ and it induces production of the neutrophil chemoattractant IL-8 (CXCL8) by human airway smooth muscle cells¹⁹.

Once an individual is sensitized to a particular allergen, subsequent encounters with that allergen cause crosslinking of IgE bound to the high-affinity IgE receptor (Fc ϵ RI) to stimulate the release of granule-associated and newly generated mediators that are responsible for the early allergic response (which occurs within 1–30 minutes), together with the release of cytokines and chemokines that recruit macrophages, eosinophils and basophils that comprise the late response (within 6–72 hours)²⁰. Autacoid mediators such as histamine and the cysteinyl leukotrienes (CysLTs) increase endothelial expression of P-selectin and E-selectin to initiate leukocyte rolling, followed by the expression of intercellular adhesion molecule 1 (ICAM1) and vascular cell adhesion molecule 1 (VCAM1), which interact with integrin receptors to arrest the leukocyte and assist its passage into the perivascular space²¹. Chemokines, such as CC-chemokine ligand 2 (CCL2), CCL8, CCL7 and CCL13 (monocyte chemoattractant proteins 1–4), CCL3, CCL24 and CCL26 (eotaxins 1–3) and CCL5 (RANTES), direct and prime leukocytes for mediator secretion²². Although adhesion molecules and chemokines might seem to be attractive therapeutic targets at which to direct inhibitors, none has yet been translated into clinical use²³. The epithelium has a particularly important role in mucosal and skin allergy as the source of crucial cytokines and chemokines — such as CCL17 (TARC), CCL22 (MDC), IL-25 (IL-17E) and IL-33 — that promote T_H2-cell function, as well as thymic stromal lymphopoietin (TSLP), which interacts with DCs and mast cells to increase the T_H2-cell response^{24,25}.

In asthma and atopic dermatitis, the epithelium is more susceptible to oxidant stress and injury²⁶. In asthma, the presence of IL-13, as well as epidermal, neural, vascular and fibroblast growth factors, leads to mucous metaplasia and remodelling of the airway walls²⁷ (FIG. 2). However, it is still not known how these inflammatory and remodelling events relate to the increase in airway smooth muscle and its increased responsiveness in asthma. In the case of atopic dermatitis, T_H1-cell pathways are also important through the release of IFN γ , which induces keratinocyte apoptosis and the recruitment of further T cells into the lesion²⁸.

Atopy

This term (from the Greek *ατοπία*, meaning *placelessness*) refers to the susceptibility to develop exaggerated IgE responses to common environmental allergens, defined clinically by the presence of one or more positive skin-prick tests. Atopy represents a genetic predisposition towards allergic diseases.

Regulatory T cells

(T_{Reg} cells). These are specialized cells that act to suppress the function of other cells. In allergic inflammation, T_{Reg} cells can have an important role in limiting allergic responses by suppressing the function of T_H2 cells. The molecular mechanism by which T_{Reg} cells exert their activity is either through cell-to-cell contact with the cell being suppressed or through secretion of the immunosuppressive cytokines TGF β and IL-10.

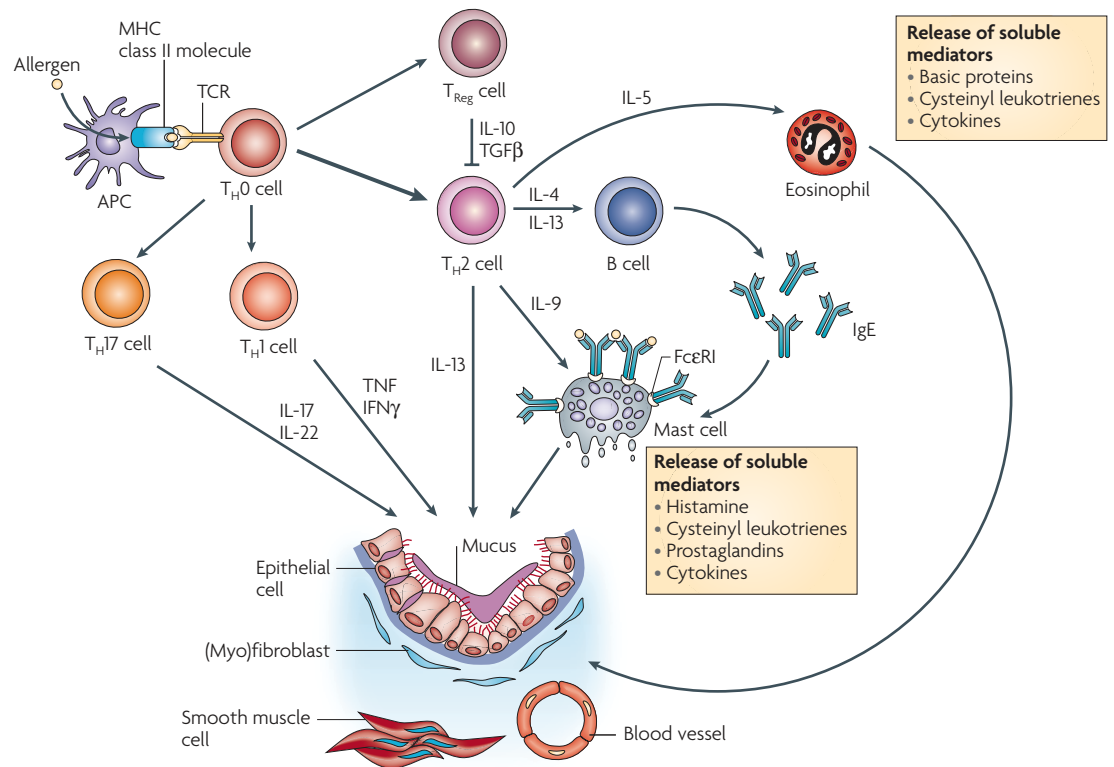


Figure 1 | Allergic mechanisms. In predisposed individuals, initial exposure(s) of professional antigen-presenting cells (APCs) to allergen leads mainly to the activation of allergen-specific T helper 2 (T_H2) cells and IgE synthesis, which is known as allergic sensitization. Subsequent exposures to allergen cause inflammatory-cell recruitment and activation and mediator release, which are responsible for early (acute) allergic responses and late allergic responses. In the early allergic response, within minutes of contact with allergen, IgE-sensitized mast cells degranulate, releasing both pre-formed and newly synthesized mediators in sensitized individuals. These include histamine, cysteinyl leukotrienes and cytokines, which promote vascular permeability, smooth-muscle contraction and mucus production. Chemokines released by mast cells and other cell types direct the recruitment of inflammatory cells that contribute to the late allergic response, which is characterized by an influx of eosinophils and T_H2 cells. Eosinophils release a large number of pro-inflammatory mediators, including cysteinyl leukotrienes and basic proteins (cationic proteins, eosinophil peroxidase, major basic protein and eosinophil-derived neurotoxin), and they might be an important source of pro-inflammatory cytokines such as interleukin-3 (IL-3), IL-5 and IL-13. There is now evidence that T_H1-cell responses might also be responsible for some of the pathogenic features in patients suffering from chronic forms of atopy, including epithelial apoptosis and smooth-muscle-cell activation. Regulatory T (T_{Reg}) cells are another important subset of CD4⁺ T cells with implications for the suppression of T_H2-cell responses in humans involving the inhibitory cytokines IL-10 and transforming growth factor-β (TGFβ). Another newly identified CD4⁺ T-cell subset, known as T_H17 cells on the basis of secretion of IL-17A and IL-17F, seems to be specifically associated with the neutrophilic inflammatory events that occur during disease exacerbation and in tissue remodelling. FcεRI, high-affinity receptor for IgE; IFNγ, interferon-γ; TCR, T-cell receptor; TNF, tumour-necrosis factor.

Allergen avoidance

In atopic individuals, allergen sensitization is fundamental to the development of any allergic disease. Therefore, avoidance of allergens before or after sensitization should be beneficial as primary or secondary prophylaxis. In the case of house dust mites, birth-cohort studies have shown that the level of allergen exposure early in life correlates with the extent of sensitization²⁹. For domestic pets, the situation is more complex, with early-life exposure decreasing rather than increasing allergen sensitization, possibly as a result of simultaneous exposure to inhibitory products from non-pathogenic microorganisms, whereas exposure later in childhood leads to sensitization. Trials looking at the effects of decreased exposure to house dust mites in early childhood have produced mixed

results, with most of the trials either showing no effect³⁰ or increased IgE sensitization³¹. This disappointing response to primary prophylaxis can be explained by the fact that extremely low allergen exposures can lead to sensitization and, as a result, anything less than complete allergen avoidance is unlikely to be successful. Greater success has been obtained by using multiple early-life interventions in addition to the avoidance of house dust mites and pets, such as breast-feeding with the mother on a low-allergen diet or giving extensively hydrolysed formula milk to babies³². A similar situation applies to peanut allergy, in which avoidance during pregnancy and early infancy can increase rather than protect against sensitization³³. Therefore, exposure to a high dose of peanut, rather than avoidance in infancy, might be a way to induce protective tolerance³⁴.

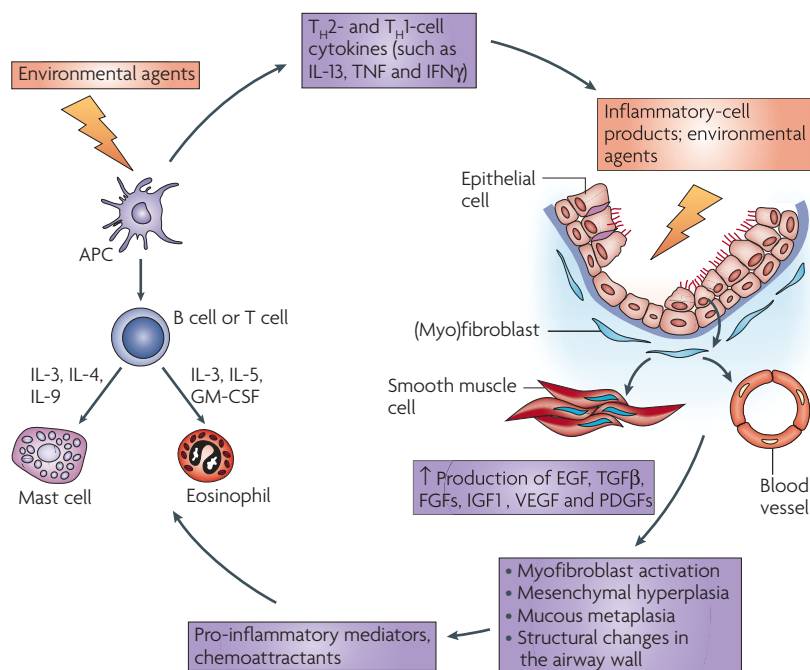


Figure 2 | Inflammatory and remodelling responses in asthma with activation of the epithelial-mesenchymal unit. The airways in asthma show characteristics of a chronic wound with evidence of ongoing epithelial injury and repair. As in any wound, responses to tissue injury create the necessary stimuli to recruit the underlying mesenchyme to participate in the repair process through the release of various growth factors and cytokines. Interleukin-13 (IL-13) is a key cytokine in this process, driving goblet-cell metaplasia and myofibroblast differentiation, and supporting immunoglobulin class-switching to IgE. Both in human asthma and in mouse models of allergen-induced remodelling, epidermal growth factors (EGFs), transforming growth factor- β (TGF β), fibroblast growth factor 2 (FGF2) and FGF7 (KGF), insulin-like growth factor 1 (IGF1), platelet-derived growth factor AA (PDGFAA) and PDGFBB, vascular endothelial growth factor (VEGF) and IL-5 are all implicated. The epithelium is more susceptible to oxidant stress and to injury by prematurely entering into apoptosis. Impaired wound repair leads to the increased production of growth factors by damaged epithelium, activated structural cells and infiltrating inflammatory cells. In asthma this leads to myofibroblast activation, mesenchymal hyperplasia, mucous metaplasia and the induction of structural changes throughout the airway wall. The ongoing eosinophil- and mast-cell-driven inflammatory responses are responsible for the maintenance and progression of tissue injury and repair. The activation of T helper 1 (T_H1)-cell pathways is also important for tissue inflammation and remodelling through the release of interferon- γ (IFN γ), which induces epithelial apoptosis and the recruitment of further T cells into the lesion) and tumour-necrosis factor (TNF, which elicits the proliferation and activation of myofibroblasts and fibroblasts and induces a hypercontractile phenotype in airway smooth muscle cells). APC, antigen-presenting cell; GM-CSF, granulocyte/macrophage colony-stimulating factor.

In children who are already sensitized, single or combination interventions to decrease exposure to both dietary and aeroallergens result in a meaningful and sustained improvement in the control of asthma and rhinitis. However, in adults, the data are far less convincing, probably because of the many allergenic and non-allergenic factors that contribute to ongoing disease. In the case of asthma and rhinitis, a double-blind, randomized, placebo-controlled study of allergen-impermeable bed covers involving 1,122 adults with asthma and 279 patients with allergic rhinitis failed to show any beneficial effects of reducing exposure to house dust mites^{35,36}.

Transactivation
A transcriptional mechanism by which gene transcription is induced resulting in the *de novo* synthesis of susceptible proteins.

Transrepression
A transcriptional mechanism by which gene transcription is prevented resulting in an overall repressive effect.

Established treatments

Inhaled corticosteroids and short- and long-acting β 2-adrenoceptor agonists (SABAs and LABAs) are now the mainstay of asthma treatment as advocated by disease-management guidelines. In the case of rhinitis, α -adrenoceptor agonists are used to relieve nasal congestion, and non-sedating H1-antihistamines and topical corticosteroids are well-established control therapies. So, for most allergic disorders, a combination of symptom-relieving and control therapies forms the basis of management guidelines. Asthma provides a particularly good example of how an understanding of the pathophysiology of the disease has underpinned the treatment approaches that are in use.

Corticosteroids. T_H2-cell-mediated inflammation in asthmatic airways is suppressed by corticosteroids through the inhibition of expression of cytokines, chemokines and adhesion molecules³⁷, whose encoding genes are regulated by transcription factors such as nuclear factor- κ B (NF- κ B) and activator protein 1 (AP1)³⁸. Free corticosteroids diffuse across the cell membrane, where they interact with cytoplasmic glucocorticoid receptors. This results in the activation of these receptors and their subsequent translocation to the nucleus, where the transcriptional activity of target genes is modulated by several different mechanisms, including gene transactivation and gene transrepression³⁹ (FIG. 3).

Whereas inhaled corticosteroids are highly effective at suppressing airway inflammation, they do not influence the natural history of the disease, even when treatment is started early in childhood^{40,41}. Inhaled corticosteroids are largely ineffective in virus-induced exacerbations⁴² and in those asthmatics who smoke⁴³.

β 2-adrenoceptor agonists. Inhaled SABAs such as salbutamol and turbutaline are the most effective bronchodilators currently available for the rapid relief of asthma symptoms. After binding of these agonists to the β 2-adrenoceptor, adenylate cyclase is stimulated by the signal-transducing G_s protein to increase production of cyclic adenosine 3'5'-monophosphate (cAMP), thereby activating protein kinase A. This mediates smooth-muscle relaxation through the phosphorylation of myosin light-chain kinase and by opening Ca²⁺-dependent K⁺ (KCa) channels, which relieves bronchoconstriction in asthma.

The two inhaled LABAs, formoterol and salmeterol, induce bronchodilation for at least 12 hours⁴⁴ and are used as a supplementary therapy for asthma that is not controlled by inhaled corticosteroids. It has also been proposed that LABAs increase the efficacy of inhaled corticosteroids⁴⁵, although this is controversial. Monotherapy with a LABA is not recommended as it could mask worsening inflammation, with potentially serious consequences⁴⁶. Increased patient adherence and convenience have led to the widespread use of combination inhalers containing inhaled corticosteroids and LABAs.

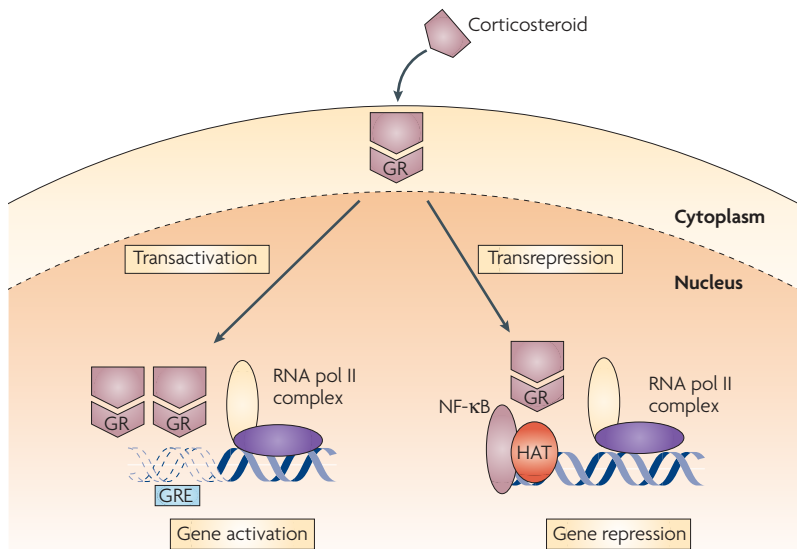


Figure 3 | Anti-inflammatory actions of corticosteroids. Corticosteroids enter the cell, bind to the glucocorticoid receptor (GR) in the cytoplasm and translocate to the nucleus, where the transcription of target genes is initiated. Many genes contain glucocorticosteroid response elements (GREs) in their promoters. Through transactivation, binding of the activated glucocorticoid receptor homodimer to a GRE in the promoter region of steroid-sensitive genes leads to the transcription of genes encoding anti-inflammatory mediators such as annexin-1 (lipocortin-1), secretory leukoprotease inhibitor (SLPI), interleukin-10 (IL-10) and the inhibitor of nuclear factor-κB (IκBα). Through transrepression, the glucocorticoid receptor-corticosteroid complex interacts with large co-activator molecules with intrinsic histone acetyltransferase (HAT) activity (such as cyclic AMP response element binding protein, CBP), which are activated by pro-inflammatory transcription factors (such as NF-κB and AP1), thus switching off expression of the inflammatory genes that are activated by these transcription factors. RNA pol II, RNA polymerase II.

Mediator antagonists and synthesis inhibitors. H1-antihistamines such as chlorpheniramine were the first specific agents used to treat allergic reactions. Although the early products were effective at controlling the symptoms of allergy, their sedative and anti-cholinergic side-effects were problematic. These have been overcome by the development of a second generation of drugs — including cetirizine, levocetirizine, loratadine and desloratadine — that have a decreased capacity to cross the blood–brain barrier, greater efficacy and selectivity, and decreased cardiac toxicity⁴⁷. The CysLTs are some of the most potent contractile agonists of airway smooth muscle and they also have effects on microvessels, mucous glands, eosinophils and nerves by interacting with the CysLT receptor 1 (CysLTR1). During active asthma and rhinitis, increased levels of CysLTC₄, CysLTD₄ and CysLTE₄ have been detected in biological fluids. Neither the biosynthesis nor the actions of CysLTs are inhibited by corticosteroids⁴⁸. The currently available oral leukotriene modifiers are CysLTR1 antagonists (montelukast, zafirlukast and pranlukast), but despite numerous attempts, only one 5-lipoxygenase inhibitor, zileuton, has so far passed clinical trials⁴⁹. Although CysLTR antagonists can be used as a monotherapy in mild to moderate asthma (particularly in children), their main use is as a supplementary therapy to inhaled corticosteroids⁵⁰. Leukotriene inhibitors are also effective

Xanthine

A purine base found in most body tissues and fluids as a result of purine degradation. Theophylline is a methylated xanthine with activities as both a cyclic-AMP phosphodiesterase inhibitor and adenosine-receptor antagonist, which is commonly used for its effects as a mild stimulant and as a bronchodilator.

for the treatment of allergic rhinoconjunctivitis (which frequently coexists with asthma)⁵¹, but not for the treatment of atopic dermatitis⁵².

Phosphodiesterase inhibitors. Theophylline is a xanthine with activity as both a cAMP phosphodiesterase (PDE) inhibitor and an adenosine-receptor antagonist. Theophylline has been used to treat asthmatic bronchoconstriction, but the cardiac and central-nervous-system side-effects that occur at doses similar to those required to generate therapeutic effects (low therapeutic index) have led to a marked reduction in its use. Although it has been proposed that theophylline also has some anti-inflammatory effect, the evidence is rather weak. More effective PDE inhibition has been achieved by targeting the type-4 isoenzyme with non-xanthine drugs such as rofumulast⁵³.

Drugs for refractory disease. There are some patients with asthma whose symptoms are not adequately controlled by conventional treatments. Lack of adherence to treatment is an important cause, but otherwise refractory asthma sometimes responds to immunomodulators such as low-dose methotrexate, azathioprine or cyclosporine. However, randomized, controlled trials of these immunomodulators have given mixed results and side-effects are common⁵⁴. By contrast, calcineurin inhibitors that inhibit T-cell responses, such as oral cyclosporine A and locally applied tacrolimus and pimecrolimus, are effective treatments for atopic dermatitis that is refractory to corticosteroid treatment⁵⁵.

The failure of corticosteroids to decrease the level of expression of tumour-necrosis factor (TNF) and other T_H1-cell-associated cytokines in asthmatic airways might explain why corticosteroids have limited effects in more severe forms of the disease⁵⁶. Based on the increased expression of TNF in the airways⁵⁷ and in blood mononuclear cells⁵⁸ in severe asthma, two small studies have reported the efficacy of the soluble TNF-receptor fusion protein etanercept^{57,59}. Large multi-centre trials with etanercept and TNF-specific monoclonal antibodies are now in progress. Inhibitors of p38 mitogen-activated protein kinase and IκB kinase (IKK), such as SB 220025 (REF. 60) and TPCA-1 (REF. 61) respectively, are also attractive new therapeutic approaches for refractory asthma, as a result of their ability to inhibit the production of pro-inflammatory cytokines such as TNF and IL-1.

Allergen-specific immunotherapy

Allergen-specific immunotherapy (SIT) is an immunemodifying therapy that has been recommended for the treatment of allergic rhinitis, venom hypersensitivity, some drug allergies and mild bronchial asthma. SIT induces immunological tolerance and the induction of blocking IgG4 antibodies through repeated exposure to allergen(s). After experimental or natural exposure to allergens, SIT decreases the recruitment of mast cells, basophils and eosinophils in the skin, nose, eye and bronchial mucosa (FIG. 4). SIT produces an increase in the level of allergen-specific IgA and IgG4 antibodies, and a decrease in the level of allergen-specific IgE antibodies.

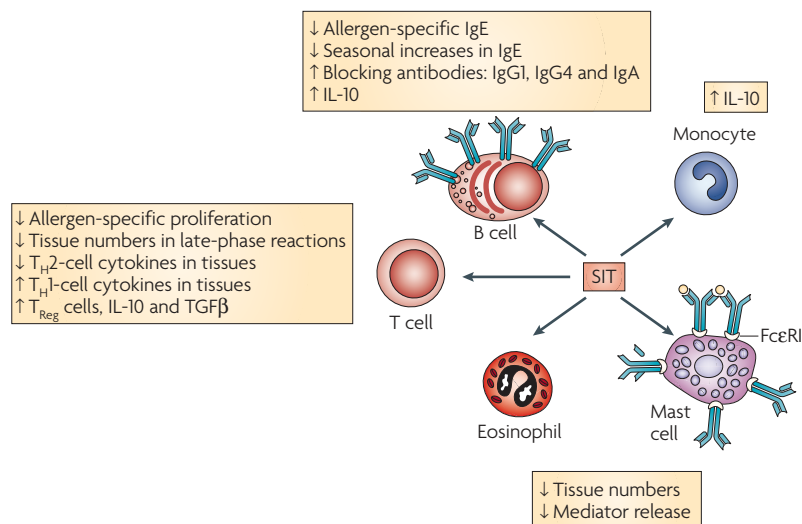


Figure 4 | Effects of allergen-specific immunotherapy (SIT) on immune parameters. SIT modifies cellular and humoral responses to allergen. The ratio of T helper 1 (T_H1)-cell cytokines to T_H2 -cell cytokines is increased after SIT, and functional regulatory T (T_{Reg}) cells are induced. The production of interleukin-10 (IL-10) by monocytes, macrophages, B cells and T cells is increased. The production of transforming growth factor- β (TGF β) is increased and, together with IL-10, TGF β might contribute to T_{Reg} -cell function and immunoglobulin class-switching to IgA, IgG1 and IgG4. These immunoglobulins compete with IgE for allergen binding, thereby decreasing the allergen capture and presentation that is facilitated by IgE in complex with the high-affinity receptor for IgE (Fc ϵ R1) or the low-affinity receptor for IgE (Fc ϵ R2, CD23). In addition, SIT decreases the number of mast cells and the ability of mast cells to release mediators. The recruitment of eosinophils and neutrophils to sites of allergen exposure is also decreased.

CpG oligonucleotide motifs

DNA oligonucleotide sequences that include a cytosine–guanosine sequence and certain flanking nucleotides. They have been found to induce innate immune responses through interaction with TLR9. When coupled to allergens, CpG DNA seems to increase immunological tolerance by shifting the balance of T-cell phenotypes from T_H2 to T_H1 cells. CpG motifs are also known as immunostimulatory oligodeoxynucleotides (ISS ODNs).

Airway hyper-responsiveness

An abnormally increased sensitivity of the airways to otherwise innocuous stimuli, resulting in increased responses to inhaled allergen and airway smooth-muscle spasmogens (for example, methacholine or histamine). In humans, this is generally defined by PC_{20} (the provocation concentration that causes a 20% decrease in forced expiratory volume in one second, FEV $_1$).

It also induces CD4 $^+$ CD25 $^+$ FOXP3 $^+$ T_{Reg} cells that produce high levels of IL-10 and/or TGF β , two cytokines that are known to attenuate allergen-specific T_H2 -cell responses. IL-10 suppresses mast-cell, eosinophil and T-cell responses⁶², and the pleiotropic functions of TGF β maintain a diverse and self-tolerant T-cell repertoire, including T_{Reg} cells⁶³.

Subcutaneous immunotherapy (SCIT) involves the regular subcutaneous injection of allergen extracts or recombinant allergens using incremental regimes, with the induction of tolerance taking from several days to several months depending on the regime used. The usual approach is a build-up phase (consisting of weekly injections) followed by a maintenance phase (consisting of monthly injections). Once tolerance is induced it can last for several years without further treatment⁶⁴. The limiting factor in SCIT is anaphylactic side-effects, which vary in incidence from 0.1–5% of individuals depending on severity⁶⁵. Improved efficacy with decreased side-effects is the aim of the new approaches to SCIT, including T-cell-reactive peptides⁶⁶, hypoallergenic recombinant allergens⁶⁷, chemically modified allergens (allergoids)⁶⁸, replacing adjuvants such as alum with those containing tyrosine or calcium phosphate, incorporation of immunomodulators such as monophosphoryl lipid A (MPL)⁶⁹, and embedding of allergens into nanoparticles⁷⁰. Attaching CpG oligonucleotide motifs to purified allergens seems to be a particularly promising approach to SCIT by increasing the efficacy and decreasing the side effects, as recently reported for the novel ragweed-allergen conjugate⁷¹.

The administration of allergens to the oral mucosa as a route for immunotherapy has only recently gained acceptance (sublingual immunotherapy, SLIT). Although much higher doses of allergen are required than are used for SCIT, the side-effect profile is impressively mild, making this therapy an attractive alternative and suitable for children^{72,73}. Several clinical trials and meta-analyses show that SLIT is effective for the treatment of pollenosis caused by grass, *Parietaria judaica*, olive, ragweed and birch, as well as rhinitis that is associated with house-dust-mite and cat allergies, but the benefit is less than that of topical corticosteroids and antihistamines⁷⁴.

The efficacy and safety of SCIT in patients with allergic asthma is controversial. In the 2003 Cochrane review of 75 trials covering a total of 3,188 patients with asthma, SCIT led to a significant reduction in asthma symptom scores, medication use and airway hyper-responsiveness, with evidence of a dose-related effect⁷⁵. Dose-response studies carried out for dog⁷⁶, cat⁷⁷, ragweed⁷⁸ and grass⁷⁹ allergies show consistent efficacy of SCIT for between 5 and 20 mcg of the major allergen.

SCIT and SLIT also decrease the development of sensitization to new allergens and decrease the risk of new asthma in both adults and children with rhinitis. A significantly decreased rate for the development of new allergen sensitizations has been shown in mono-sensitized patients who received SCIT compared with untreated, matched controls^{80–82}. Several studies have indicated that allergic rhinitis often precedes asthma and therefore that rhinitis might be an important risk factor for the development of asthma^{83,84}. A recent retrospective cohort study of 332 subjects with allergic rhinitis showed that 53.1% of subjects who were not treated with SCIT developed asthma, whereas 41.6% of subjects who received SCIT were diagnosed with asthma at the end of the observation period⁸⁴. The observed effect of SCIT in decreasing the incidence of new asthma cases is of clinical importance, with a significant reduction in the prevalence of physician-diagnosed asthma in adults with allergic rhinitis. These findings seem to be in agreement with the results of a ground-breaking, prospective, multicentre study (the PAT study) of 205 children with a clinical history of grass- or birch-induced rhinoconjunctivitis, who were randomized to receive SCIT for 3 years or to an open control group⁸⁵. There was a statistically significant decrease in risk for the development of asthma during the treatment period (odds ratio, 2.52 in favour of SCIT for the prevention of asthma). At the 5-year follow-up (2 years after discontinuation of SCIT), the preventive effect of SCIT was still evident. Of the 142 children without asthma before the start of the study, those that had received SCIT had significantly less asthma than those in the control group: 16/48 with asthma in the SCIT group compared with 24/29 in the untreated control group (odds ratio, 2.68 in favour of SCIT for the prevention of asthma)⁸⁶. In a randomized, placebo-controlled 3-year study of allergen immunotherapy in non-asthmatic, rhinitic adults monosensitized to *Parietaria* pollen, it was reported that 47% of patients in the placebo group developed asthma symptoms by the end of the study, compared with only

14% of the patients treated with SCIT⁸⁷. This significant reduction in the incidence of new asthma cases seems to be confirmed also in studies with SLIT. In a recent prospective, multicentre study of 113 non-asthmatic children with a clinical history of hay fever to grass who were randomized to receive SLIT for 3 years or to an open control group, there was a statistically significant reduction in risk for the development of asthma during the treatment period (odds ratio, 3.8 in favour of SLIT for the prevention of asthma)⁸⁸.

IgE as a therapeutic target

The sentinel role of IgE in increasing allergen uptake by DCs and activating mast cells and basophils for mediator release is reviewed in this issue by Hannah Gould and Brian Sutton⁸⁹. IgG antibodies specific for the C3 domain of IgE that block IgE binding to FcεRI (and FcεRII, CD23) were shown to inhibit allergen-induced inflammatory responses in mice and in humans⁹⁰. Omalizumab, a humanized IgE-specific, non-anaphylactic IgG1 has been developed for the treatment of severe allergic asthma⁹¹ (FIG. 5). Clinical trials in adults, adolescents and children with poorly controlled IgE-mediated asthma have shown that omalizumab administered subcutaneously 2–4 times per week (in proportion to the total level of IgE in the patient's serum and to the patient's body weight) improves symptom control and allows patients to be managed with lower doses of inhaled corticosteroids. Omalizumab has been well-tolerated in clinical trials lasting as long as 52 weeks. Although the level of circulating free IgE decreases rapidly after the first dose of omalizumab, up to 16 weeks of treatment is required before optimal clinical effects are seen. Omalizumab is also effective for the treatment of allergic rhinoconjunctivitis, but therapy has to begin long before the pollen season^{92,93}. There is interest in developing a peptide-based vaccine for active immunization to elicit long-term, protective, IgE-specific antibodies. In sensitized rats, autoantibodies specific for IgE can be

induced by IgE peptide-based vaccines⁹⁴. RP01, a novel immunotherapeutic that induces the production of IgE-specific antibodies, has just entered Phase II clinical trials for allergic asthma⁹⁵. Lumiliximab, an antibody specific for the low-affinity IgE receptor FcεRII, also decreases circulating IgE levels and has passed Phase 1 trials for mild to moderate allergic asthma⁹⁶.

Inhibitors of mast cells

The archetypal mast-cell-stabilizing drug sodium cromoglicate (SCG) was first introduced as a treatment for asthma in 1968 and was followed by nedocromil sodium in 1984 (REF. 97). After inhalation, both drugs inhibit the allergen-induced early- and late-phase responses in the upper and lower airways and conjunctiva, where mucosal mast cells are crucially involved in the allergic response⁹⁸. Nedocromil sodium and SCG inhibit the flux of chloride ions in mast cells, epithelial cells and neurons to increase their threshold for activation⁹⁹. Human mast cells also express a Ca²⁺-activated K⁺ channel — K(Ca)3.1 — that promotes mast-cell chemotaxis and increases IgE-dependent mast-cell activation, which indicates that the inhibition of K(Ca)3.1 with drugs such as clotrimazole and TRAM-34 (1-[(2-chlorophenyl) diphenylmethyl]-1H-pyrazole) might be a new approach to mast-cell inhibition^{100,101}.

Another newly identified cell-surface target on mast cells is CD63, a member of the tetraspanin family of proteins. CD63 interferes with integrin signalling, localization or trafficking and interacts with the α3, α4 and α6 chains of β1 integrins to modify adhesion to fibronectin and vitronectin¹⁰². A monoclonal antibody that blocks CD63 inhibits FcεRI-mediated activation of mast cells that are adherent to extracellular matrix proteins but not of nonadherent cells.

The SRC tyrosine kinases FYN and LYN are important modulators of the molecular events that are initiated by engagement of FcεRI¹⁰³. A novel strategy to inhibit mast-cell activation is to inactivate the SYK that propagates FcεRI signalling¹⁰⁴. A cell-based, high-throughput screen for small molecules that block IgE signalling identified a 2,4-diaminopyrimidine (R112) as a reversible mast-cell SYK inhibitor¹⁰⁵. In allergic rhinitis, intranasal administration of R112 followed by allergen challenge inhibited nasal obstruction and rhinorrhoea, together with the inhibition of prostaglandin D₂ production¹⁰⁶. In patients with seasonal rhinitis who were exposed for 2 days to a high pollen count, R112 was effective in reducing global symptoms of rhinitis with rapid onset¹⁰⁷.

The interaction of stem-cell factor (SCF) with its tyrosine-kinase receptor KIT is obligatory for mast-cell development, proliferation, survival, homing and adhesion, and for optimal IgE-induced mast-cell degranulation and cytokine production¹⁰⁸. Drug candidates that target SCF or KIT include SCF-specific antibodies, antisense oligonucleotides, KIT inhibitors and inhibitors of downstream signalling molecules¹⁰⁹. Imatinib mesylate (imatinib), nilotinib and dasatinib are tyrosine-kinase inhibitors that were developed to treat BCR-ABL-expressing leukaemia and gastrointestinal stromal tumours. By also inhibiting a mutant form

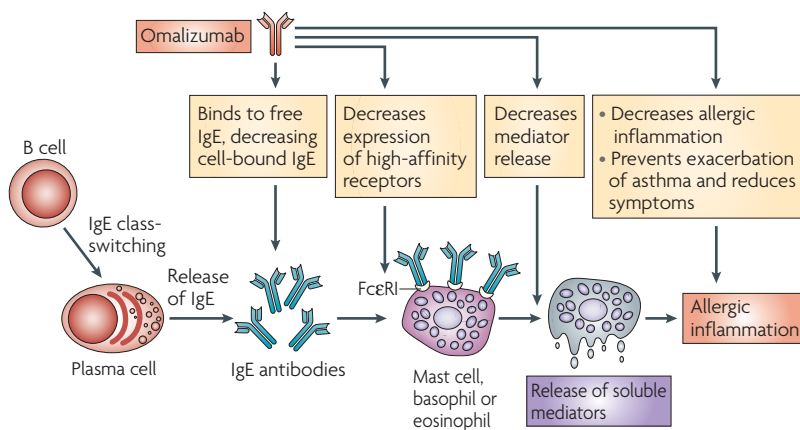


Figure 5 | The effects of targeting IgE on allergic responses. The monoclonal IgE-specific antibody omalizumab can decrease free IgE levels and decrease the amount of IgE binding to both high-affinity (FcεRI) and low-affinity (FcεRII) IgE receptors, resulting in an attenuation of allergic reactions. In addition to inhibiting the binding of IgE to mast cells, basophils and eosinophils, omalizumab downregulates the expression of FcεRI. This will also decrease the amplification of the inflammatory response mediated by T helper cells to prevent IgE-dependent allergen presentation.

of KIT, they induce the apoptosis of mast cells, which is of value in the treatment of some forms of systemic mastocytosis. This approach is also being evaluated for treating allergic disease¹¹⁰.

Modulating the expression of activating and inhibitory receptors is an important mechanism for regulating immune responses. Cells that are activated through ligation of receptors bearing immunoreceptor tyrosine-based activation motifs (ITAMs) can be negatively regulated by other receptors bearing immunoreceptor tyrosine-based inhibitory motifs (ITIMs)¹¹¹. Animals that are deficient in some of these ITIM-containing receptors — for example, animals that are deficient in *FcγRIIB*, gp49B1 (LILRB4) or paired immunoglobulin-like receptor B (PIRB) — have increased allergic responses¹¹². IgG can completely suppress IgE-mediated anaphylaxis by interacting with *FcγRIIB*¹¹³, which leads to activation of the SRC homology 2 (SH2)-domain-containing inositol polyphosphate 5'-phosphatase (SHIP) through recruitment of DOK (docking protein) and RasGAP to *FcεRI*¹¹⁴. Similar inhibitory mechanisms are invoked when gp49B1 on mast cells is activated by its integrin ligand, αβ3 (REF. 115). So, the immunoglobulin-like receptors and their intracellular signalling molecules provide important new therapeutic targets to inhibit mast cells, as well as T cells, involved in the allergic cascade. A human bifunctional *Fcγ-Fcε* fusion protein designed to crosslink *FcγRIIB* and *FcεRI* on human mast cells and basophils inhibits IgE-dependent degranulation and allergic reactions in mice transgenic for human *FcεRIα*¹¹⁶. A chimeric human–cat *Fcγ-Fel d 1* fusion protein inhibits allergic responses in mice sensitized to the major cat allergen *Fel d 1*, and this strategy therefore has potential as an enhanced form of immunotherapy¹¹⁷.

Cytokine-based immunotherapies

Because of the sentinel role that T_H2 cytokines have in orchestrating allergic inflammation, they and their receptors are key therapeutic targets (FIG. 6). With almost no exceptions, this approach has required the application of biological agents in the form of blocking monoclonal antibodies, fusion proteins and, most recently, inhibitors of the T_H2 -cell transcription factors STAT6 (signal transducer and activator of transcription 6) and GATA3.

IL-4. Both IL-4 and IL-13 have a crucial role in the immunoglobulin isotype switching of B cells to produce IgE, whereas IL-4 alone is crucial for maintaining the T_H2 -cell phenotype, which makes both cytokines attractive therapeutic targets. A large number of animal studies have shown that blocking production or inhibiting the effects of IL-4 has profound effects on the allergic phenotype. A soluble, recombinant, human IL-4 receptor (altrakincept) consists of the extracellular portion of human *IL-4Rα* and is non-immunogenic. A small proof-of-concept trial of nebulized inhaled altrakincept for 12 weeks in patients with mild to moderate asthma indicated efficacy by allowing withdrawal from treatment with inhaled corticosteroids without relapse¹¹⁸, and this result was subsequently confirmed in a larger trial¹¹⁹. However, a Phase III trial failed to confirm the efficacy

of altrakincept for the treatment of asthma. This trial does not invalidate IL-4 as a target for the treatment of allergy and asthma, as there were concerns over the bio-availability of altrakincept in this study. Further Phase II studies are in progress using humanized IL-4-specific and IL-4Rα-blocking antibodies such as pascolizumab (SB240,683)¹²⁰. Two vaccines against IL-4 have been tested in mice, one in which IL-4 is chemically coupled to limpet haemocyanin¹²¹ and the other in which a 14-amino-acid peptide from IL-4 is inserted into variant hepatitis B virus core antigen¹²². Both vaccines induced high antibody titres specific for mouse IL-4 and inhibited antigen-induced lung inflammation. However, using costimulation blockade in a mouse model of allergy to grass pollen¹²³, it was reported that the secondary IgE response is not T-cell dependent, thereby raising doubts over the usefulness of IL-4 blockade for treating established allergic disease.

IL-13. The numerous functions of IL-13 in regulating IgE production, eosinophilic inflammation, airway-smooth-muscle hyperplasia, the induction of goblet-cell hyperplasia with mucus production, and the recruitment of monocytes, macrophages and T cells into the airway spaces make it a key therapeutic target in allergy and asthma¹²⁴. IL-13 binds to a low-affinity *IL-13Rα1* subunit and a high-affinity complex comprised of *IL-13Rα1* and *IL-4Rα*. Binding to this high-affinity complex leads to the phosphorylation-dependent activation of Janus kinase 1 (JAK1), JAK2 and STAT6. *IL-4Rα* also stabilizes the binding of IL-13 to its receptor to augment IL-13-mediated responses. However, a non-signalling, high-affinity IL-13-binding chain, *IL-13Rα2*, strongly inhibits the activity of IL-13 (REF. 125). Selective blockade of IL-13 has been achieved in mice using a soluble form of *IL-13Rα2*, which competes for binding to IL-13 but not to IL-4, and this led to the reversal of airway hyper-responsiveness and mucus production in allergen-exposed sensitized mice¹²⁶. A soluble form of *IL-13Rα2* that binds IL-13 with 100-fold greater affinity than does *IL-13Rα1* is present in mouse but not human serum.

Antagonizing the effects of IL-13 could also be achieved by administering soluble IL-13 receptors or IL-13R-specific monoclonal antibodies. In cynomolgus monkeys sensitized to *Ascaris suum* and then challenged with antigen from this nematode, a mouse antibody specific for human IL-13 (mAb13.2) and the humanized counterpart (IMA-638) inhibited eosinophil and neutrophil influx into the lungs as assessed by bronchoalveolar lavage¹²⁷. Phase I trials of the IL-13-specific monoclonal antibody CAT-354 in 34 mildly asthmatic patients have been successfully completed and Phase II trials are in progress. Subcutaneous or inhaled pitrakinra, a mutant IL-4 protein that inhibits the binding of IL-4 and IL-13 to *IL-4Rα* complexes, has recently shown efficacy in the treatment of allergen-induced asthma¹²⁸. A novel, recombinant IL-13 peptide-based vaccine has also been shown to reduce allergic inflammatory responses in mice¹²⁹. As STAT6 is the common transcription factor for both IL-4 and

Immunoreceptor tyrosine-based activation motif (ITAM). Activating receptors often have ITAMs consisting of a consensus amino-acid sequence with paired tyrosines and leucines (YxxI/Lx_{6–12}YxxI/L). These are normally located in the cytoplasmic domains of ligand-binding transmembrane receptors (such as *FcεRI* and TCR) and they mediate interaction between the transmembrane receptor complex and protein tyrosine kinase activity, which is required to initiate early and late signalling events.

Immunoreceptor tyrosine-based inhibitory motif (ITIM). Inhibitory receptors often have one or more ITIMs (consensus, S/I/V/LxYxxI/V/L). Ligand engagement by inhibitory receptors (such as CTLA4 in T cells) results in ITIM phosphorylation and the recruitment of phosphotyrosine phosphatases, which leads to decreased tyrosine phosphorylation of activation-pathway effectors.

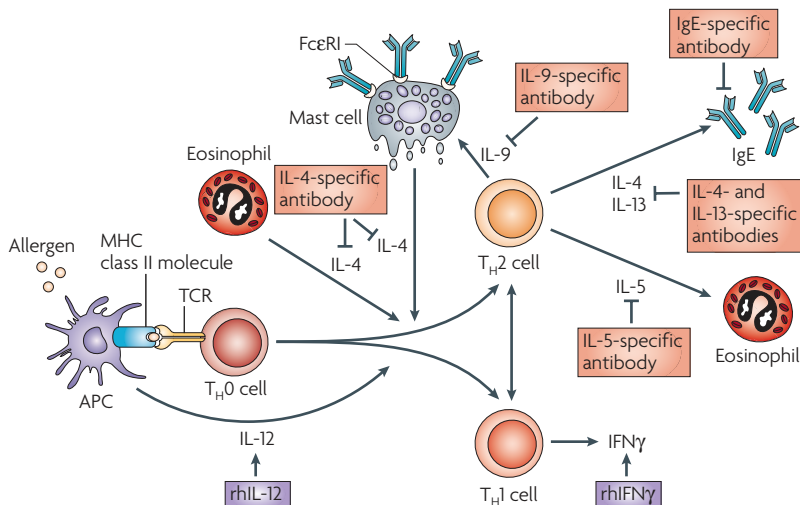


Figure 6 | Cytokine-based therapies in asthma. Allergic inflammation has been characterized as a mainly T helper 2 (T_H2)-cell disease; therefore, efforts to alter the T_H2 – T_H1 -cell balance in asthma have been aggressively pursued, either by inhibiting T_H2 -cell cytokines (in particular, interleukin-4 (IL-4), IL-13 and IL-5) or promoting T_H1 -cell responses (interferon- γ (IFN γ) and IL-12). Inhibition of the allergic component of atopic asthma can also be achieved using IgE-specific monoclonal antibodies. APC, antigen-presenting cell; Fc ϵ RI, high-affinity receptor for IgE; rh, recombinant human; TCR, T-cell receptor.

IL-13 signalling, it is also an attractive therapeutic target using a dominant-negative peptide¹³⁰ and anti-sense RNA-based approaches. Anti-sense and RNA-interference-based therapeutic strategies are being explored to target various upstream signalling molecules in asthma and allergy, including Fc ϵ RI α , cytokine receptors, adhesion molecules, ion channels, cytokines and related factors, intracellular signal-transduction molecules and transcription factors involved in T_H2 -cell differentiation and allergic inflammation¹³¹.

IL-5. Rodent and non-human primate studies have indicated an important role for IL-5 in various models of asthma. Inhaled IL-5 modulates the number of eosinophil progenitors in both the airways and bone marrow of asthmatic individuals and induces local eosinophilia in non-asthmatic individuals¹³². Two humanized, human-IL-5-specific monoclonal antibodies, Sch-55,700 and mepolizumab (SB-240,563), have been developed for the treatment of asthma. In a small, double-blind trial, mepolizumab produced a rapid dose-dependent reduction in the number of circulating and sputum eosinophils that persisted for 3 months but, surprisingly, this had no effect on either the late asthmatic response or on airway hyper-responsiveness¹³³. In a group of patients with severe persistent asthma, treatment with Sch-55,700 resulted in a decrease in the number of blood eosinophils, but over the course of 10 weeks it had no effect on any measures of asthma outcome¹³⁴, an observation that has recently been confirmed in a large trial with mepolizumab¹³⁵.

A further study using mepolizumab confirmed the persistent suppression of eosinophilia in blood, bone marrow and airway lavage, but in airway biopsies, there was only a 55% reduction in the number of tissue eosinophils¹³⁶. As a proportion of eosinophils in the airways

of patients with asthma lack IL-5R^{137,138}, it was suggested that this might explain the apparent lack of clinical efficacy of targeting IL-5 (REF. 139). IL-5 could have more subtle effects on asthmatic airways — for example, mepolizumab treatment decreases immunostaining for tenascin, lumican and procollagen III in the bronchial mucosal subepithelial basal lamina¹⁴⁰ and in allergen-challenged skin¹⁴¹. In addition, IL-5-specific treatment resulted in a parallel decrease in the number of airway eosinophils expressing mRNA for TGF β 1 and of TGF β 1 levels in bronchoalveolar-lavage fluid, which indicates a possible role for IL-5 in airway remodelling¹⁴². In contrast to asthma, mepolizumab is highly efficacious in the treatment of hypereosinophilic syndrome¹⁴³ and eosinophilic oesophagitis¹⁴⁴, but not atopic dermatitis¹⁴⁵. A therapeutic DNA-based vaccine against IL-5 is also being developed.

IL-9. Blocking the actions of IL-9 reduces allergen-induced airway inflammation and airway hyper-responsiveness in mouse models. Two Phase I dose-escalation studies of an IL-9-specific monoclonal antibody (MEDI-528) in healthy volunteers have been completed without problems¹⁴⁶. Phase II trials are in progress for treating symptomatic, moderate to severe, persistent asthma.

Interferons. Of the T_H1 -cell-associated cytokines, IFN γ is the most potent in suppressing T_H2 -cell-mediated allergic inflammation, and the exogenous administration of IFN γ suppresses allergic airway inflammation in animal models. IFN γ is also strongly induced during allergen-specific immunotherapy. However, studies of the subcutaneous administration of recombinant human IFN γ in asthma have been disappointing¹⁴⁷. In mice, increased production of IFN γ together with an established T_H2 -cell response results in increased inflammation, possibly by damaging the epithelial barrier¹⁴⁸. By contrast, two small trials have shown that systemic administration of IFN γ for 18 months is effective for the treatment of severe corticosteroid-refractory asthma, with one study showing reversal of the T_H2 -cell cytokine profile in blood mononuclear cells after treatment of patients with severe asthma^{149,150}.

The recent demonstration that epithelial cells from asthmatic individuals *in vitro* have an impaired protective IFN β response to infection with the common cold virus^{151,152} has prompted clinical trials of recombinant human IFN β by inhalation to prevent severe virus-induced asthma exacerbations¹⁵³.

IL-12. IL-12 sends a powerful signal to naive precursor T cells, directing their differentiation to T_H1 cells *in vitro* and *in vivo* and shifting the immune response towards cell-mediated immunity. In animal models, administration of IL-12 during sensitization suppresses allergen-induced T_H2 -cell responses in favour of T_H1 -cell development and inhibits airway hyper-responsiveness and airway eosinophilia after antigen challenge. In asthma, the production of IL-12 by whole blood cells and its expression in airway biopsies is impaired. So, IL-12 has the potential to modify allergic diseases. Injection of recombinant human IL-12 in patients with mild asthma decreased the number

of circulating blood eosinophils after allergen challenge (but not sputum eosinophilia, the late-phase response or airway hyper-responsiveness)¹⁵⁴ and this was accompanied by flu-like symptoms, abnormal liver-function tests and cardiac arrhythmias. This approach is also brought into question by the observation that in mice, DCs retrovirally overexpressing IL-12 induce strong T_H1-cell responses to inhaled antigen in the lung but fail to suppress T_H2-cell polarization after sensitization¹⁵⁵.

IL-10. IL-10 inhibits the expression of many pro-inflammatory cytokines and chemokines, as well as pro-inflammatory enzymes, and it is the main inhibitory cytokine produced by T_{Reg} cells in allergen immunotherapy. IL-10-deficient mice have increased airway inflammation after allergen challenge¹⁵⁶, and in outbred mice, this was inhibited by the intratracheal transfer of the gene encoding IL-10 (REF. 157). Administration of IL-10 to normal volunteers decreases the numbers of circulating CD4⁺ and CD8⁺ T cells, with suppression of mitogen-induced T-cell proliferation and endotoxin-driven TNF and IL-1β production¹⁵⁸. Recombinant human IL-10 has been developed and is currently being tested in rheumatoid arthritis, inflammatory bowel disease, psoriasis, organ transplantation and chronic hepatitis C, but its effect in asthma has yet to be studied.

Targeting barrier function

The increased access of allergens to immune cells in the skin could explain the convincing association between loss-of-function mutations of the barrier protein filaggrin and the increased occurrence of atopic dermatitis¹⁵⁹. In the case of asthma, impaired barrier function is a consequence of defective tight-junction assembly²⁵, which can be restored both in human cells *in vitro* and in a mouse model *in vivo*¹⁶⁰ by the topical application of epidermal growth factor or keratinocyte growth

factor. The restoration of barrier function by growth-factor administration has been highly successful in the treatment of inflammatory bowel disease¹⁶¹ and oral mucositis associated with cytotoxic therapy¹⁶².

Concluding remarks

Despite an enormous increase in our understanding of the immune mechanisms involved in allergic diseases, it is disappointing that this knowledge has not been translated into new treatments. One possible reason for this is our lack of understanding of disease chronicity and the environmental factors that contribute to disease in addition to allergen exposure. The recent success of omalizumab as the first biological agent for the treatment of allergy opens new opportunities for using biological agents to target specific cytokines and cell-surface proteins that are involved in the allergic cascade, such as CCR4 in T_H2-cell recruitment¹⁶³, and TSLP and IL-25, which link the epithelium with DC activation and the T_H2-cell response. Several new cytokine therapeutic targets are shown in TABLE 1.

The chemical engineering of monoclonal antibodies to increase their antibody-dependent cytotoxic potential seems to be a particularly promising approach when directed against cells expressing T_H2-cell markers such as IL-5R and CCR4 (REF. 164). Vaccine approaches to allergy prevention and treatment are developing rapidly. In addition to the use of recombinant allergens and peptide fragments to increase efficacy and decrease side-effects, the coupling of allergens to oligonucleotide stimulatory sequences such as CpG DNA and other ligands of pathogen-recognition receptors to tip the balance towards a T_H1-cell response, and the development of allergen-based DNA vaccines will revolutionize allergen immunotherapy. Vaccine strategies are also looking promising for generating endogenous blocking antibodies

Table 1 | Potential new cytokine therapeutic targets in allergy and asthma

Cytokine target	Cellular origin	Predicted effects	Nature of intervention	Refs
IL-15	Leukocytes, including phagocytes. Also, neurons and muscle	Increases T _H 2-cell, B-cell, NK-cell, macrophage and monocyte responses	Blocking antibody and soluble IL-15Rα	166
IL-17A	Subset of CD4 ⁺ T cells	Neutrophil influx	Blocking antibody	167
IL-17F	Subset of CD4 ⁺ T cells	Antagonizes the effects of IL-17A	Receptor agonist	168
IL-17E (IL-25)	Subset of CD4 ⁺ T cells	Increases the T _H 2-cell response and BHR	Blocking antibody	169
IL-33 (IL-1F11) (IL-1/TLR superfamily)	Epithelium	Increases the mast-cell response and induces the production of T _H 2-cell cytokines	Soluble receptor, ST2R	170
IL-31	T _H 2 cells	Increases the T _H 2-cell response, pruritis and dermatitis	None identified	171
IL-21 (homology with IL-2, IL-4 and IL-15)	CD4 ⁺ T cells	Augments CD4 ⁺ and CD8 ⁺ T cells, NK cells and B cells	None identified	172
TSLP (IL-7 superfamily)	Epithelium and mast cells	Augments co-stimulation by DCs to increase the T _H 2-cell response and activates mast cells	Blocking antibody	173
IL-18 (IL-1 superfamily)	Macrophages and activated T cells	Increases IFNγ production by T cells. Additional targets are macrophages, neutrophils, DCs and endothelial cells	IL-18	174

BHR, bronchial hyper-responsiveness; DC, dendritic cell; IFNγ, interferon-γ; IL, interleukin; NK cell, natural killer cell; T_H2 cell, T helper 2 cell; TLR, Toll-like receptor; TSLP, thymic stromal lymphopoietin

specific for targets that are known to be involved in the allergic phenotype. As tissue injury and aberrant repair are an important component of chronic allergic diseases such as asthma, attempts are now being made to develop new therapeutic approaches to target these aspects, an example of which is bronchial

thermoplasty, which removes airway smooth muscle in asthma¹⁶⁵. However, what is now needed is a clearer understanding of the origins of allergy and the factors responsible for the increasing incidence of allergic diseases with a view to developing preventive, as well as therapeutic, strategies.

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Competing financial interests

The authors declare competing financial interests: see [web version](#) for details.

DATABASES

Entrez Gene: <http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene>
 CD63 | CysLTR1 | Fc ϵ RI | Fc ϵ RII | Fc γ RIIB | GM-CSF | IFN γ | IL-3 | IL-4 | IL-4R α | IL-5 | IL-9 | IL-10 | IL-12 | IL-13 | IL-13R $\alpha 1$ | IL-13R $\alpha 2$ | IC(Ca)3.1 | KIT | SCF | TGF β | TNE

FURTHER INFORMATION

Stephen Holgate's homepage: <http://www.som.soton.ac.uk/research/iir/members/sth/default.htm>

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