CRTH2 antagonists in asthma: current perspectives

Abstract: Chemoattractant receptor-homologous molecule expressed on T_{H2} cells (CRTH2) binds to prostaglandin D_{2}. CRTH2 is expressed on various cell types including eosinophils, mast cells, and basophils. CRTH2 and prostaglandin D_{2} are involved in allergic inflammation and eosinophil activation. Orally administered CRTH2 antagonists are in clinical development for the treatment of asthma. The biology and clinical trial data indicate that CRTH2 antagonists should be targeted toward eosinophilic asthma. This article reviews the clinical evidence for CRTH2 involvement in asthma pathophysiology and clinical trials of CRTH2 antagonists in asthma. CRTH2 antagonists could provide a practical alternative to biological treatments for patients with severe asthma. Future perspectives for this class of drug are considered, including the selection of the subgroup of patients most likely to show a meaningful treatment response.

Keywords: CRTH2, clinical trial, eosinophilic asthma, prostaglandin D_{2}

Introduction

Asthma is characterized by variable airflow obstruction, bronchial hyperreactivity, and airway inflammation. The presence of allergy is common in asthma patients and can cause bronchoconstriction and promote chronic airway inflammation. However, many asthma patients have no evidence of allergy. The heterogeneous nature of asthma means that a “one size fits all” approach to pharmacotherapy is unlikely to be successful. The development of novel asthma treatments requires an individualized approach, where medicines are targeted toward subgroups of patients with distinct characteristics who are most likely to benefit.

A clinical phenotyping approach has been advocated in order to identify patient subgroups with clinical characteristics that are associated with a treatment response or prognosis. Endotyping is the identification of a patient subgroup defined by the presence of a biological mechanism. Recently, biological treatments for asthma have been developed that use biomarkers to identify patients with specific mechanisms (i.e., endotypes). The use of both clinical phenotype information and biomarkers to select patients for novel anti-inflammatory treatments aligns to the precision medicine strategy that takes an individualized approach to pharmacotherapy to optimize the benefit versus risk ratio.

Our understanding of the complex nature of inflammation in asthma has evolved far beyond the simple dichotomy of allergic versus nonallergic asthma. The subset of lymphocytes called T-helper 2 cells were thought to drive allergic inflammation, leading to the term “T_{H2} inflammation” being associated with asthma. We recognized that many
cytokines involved in allergic inflammation are also released from other cell types, including the recently identified innate lymphoid cells (ILCs). This has led to a change in terminology to the more general “T2” inflammation. Furthermore, T2 inflammation, such as eosinophilic inflammation, can exist in the absence of allergy. To add further complexity, non-T2 inflammation can also contribute to asthma pathophysiology, such as through the IL-17 cytokine family, which is associated with neutrophilic inflammation.

The most commonly used pharmacological treatments for asthma are inhaled beta-agonists and inhaled corticosteroids (ICS), which provide bronchodilator and anti-inflammatory effects, respectively. Combination inhalers containing ICS plus a long-acting beta-agonist (ICS/LABA) have shown greater clinical efficacy than ICS alone, providing a treatment option that is widely used for many asthma patients. However, there is an unmet medical need, as many patients remain poorly controlled while taking ICS/LABA combinations. The use of long-acting muscarinic antagonists provides an additional bronchodilator option for these patients. Monoclonal antibodies targeting eosinophilic inflammation and T2 inflammation have been developed for asthma, but there is a need for additional novel anti-inflammatory treatments.

Chemoattractant receptor-homologous molecule expressed on T_{H}2 cells (CRTH2) is a G-protein coupled receptor that binds to the ligand prostaglandin D_{2} (PGD_{2}). There is evidence from in vitro studies, as well as animal and human investigations, that CRTH2 is involved in allergic and eosinophilic inflammation. A number of orally administered CRTH2 antagonists have been developed for the treatment of asthma in recent years. This article reviews the evidence for the involvement of the CRTH2 pathway in asthma and the results of clinical trials of CRTH2 antagonists in asthma patients. We also consider future perspectives for this class of drug, including considerations of which asthma subgroup is most likely to show a clinically meaningful treatment response, and whether biomarkers can be used to identify these patients.

PGD_{2}–CRTH2 biology

Arachidonic acid metabolism by cyclooxygenase enzymes and, subsequently, prostaglandin synthases leads to the production of prostaglandin. PGH_{1} is converted to PGD_{2} by PGD_{2} synthase in various cell types including mast cells and leukocytes. Mast cells are an important source of PGD_{2} in tissues, with lower levels produced by T_{H}2 lymphocytes, dendritic cells, and eosinophils. PGD_{2} undergoes rapid metabolism, with a short half-life of ~30 min in the circulation. The main products of PGD_{2} metabolism are D_{1}PGJ_{2} and 9α1βPGF_{2α}, which also have agonist effects at PGD_{2} receptors.

The biological effects of PGD_{2} are mediated by three G-protein-coupled receptors: CRTH2 (which is also called the D prostanoid receptor 2 [DP_{2} receptor]), DP_{1}, and T prostanoid (TP) receptors. The interaction of PGD_{2} and DP_{1} increases smooth muscle relaxation, vasodilation, vascular permeability, and epithelial CCL22 production, all of which may assist in the recruitment of leukocytes to the sites of inflammation, as well as in the inhibition of T_{H}1 development and function. CRTH2 activation increases intracellular calcium levels and reduces intracellular cyclic adenosine monophosphate levels, and activates various signaling pathways including phospholipase C, phosphatidylinositol 3-kinase, and p38 mitogen-activated kinase. CRTH2 is expressed by T_{H}2 cells, eosinophils, basophils, epithelial cells, and innate lymphoid type 2 cells (ILC2). Mast cells also express CRTH2, but only internally, and treatment with PGD_{2} does not induce CRTH2-dependent changes in Ca^{2+}, suggesting the receptor has a different function to that in other inflammatory cells.

The PGD_{2}–CRTH2 interaction is strongly implicated in allergic inflammation (summarized in Figure 1). The T_{H}2 cells orchestrate the allergic inflammatory response by releasing mediators such as IL-4, IL-5, and IL-13 that promote the recruitment and activation of inflammatory cells. Also, T_{H}2 cells demonstrate increased CRTH2 expression compared with other lymphocyte subtypes. Furthermore, T_{H}2 cells migrate toward PGD_{2} in vitro, and this chemotaxis is blocked by CRTH2 antagonism. PGD_{2}–CRTH2 signaling is also involved in the recruitment and activation of eosinophils and basophils. There is recent evidence that ILC2 cells play a role in airway inflammation in asthma by secreting T2 cytokines. ILC2 cells also show CRTH2-dependent migration after exposure to PGD_{2}. Animal studies have also shown a role for CRTH2 in allergic lung inflammation, with PGD_{2} treatment causing significantly increased eosinophilic lung inflammation in ovalbumin-challenged mice.

Evidence for increased PGD_{2}–CRTH2 activity in asthma

It is not clear if PGD_{2} levels in bronchoalveolar lavage (BAL) are increased in patients with mild asthma compared to healthy subjects as reports are contradictory. However, there is evidence that PGD_{2} levels increase in more
severe disease, as BAL PGD$_2$ levels are increased in severe asthma patients compared with mild and moderate asthma patients.$^{15,44}$

Gene and protein expressions of hematopoietic prostaglandin synthase (HPGDS; responsible for PGD$_2$ formation) are elevated in the bronchial epithelium of patients with moderate and severe asthma, compared with healthy subjects.$^{15}$ Furthermore, the source of HPGDS appeared to be mast cells embedded in the epithelium, as HPGDS levels correlated with mast cell tryptase. Increased PGD$_2$ levels are observed in severe asthma patients taking high-dose ICS despite the known effect of ICS in terms of reducing the number of PGD$_2$-producing mast cells in bronchial tissue.$^{44}$

Levels of PGD$_2$ are rapidly released following allergen challenge, with a 150-fold increase in BAL levels within 9 min of exposure.$^{42,45}$ Levels probably return to normal within a short period of time as no increase in PGD$_2$ was seen in sputum samples collected 24 h post allergen exposure.$^{46}$ Ex vivo studies using human lung tissue have demonstrated that allergen induction of PGD$_2$ is dependent on mast cell activation.$^{47}$

Mutarithas et al investigated the expression of CRTH2 by T lymphocytes in the blood and BAL fluid of asthma patients and healthy controls using flow cytometry.$^{48}$ The proportion of T-cells expressing CRTH2 was low in both blood and BAL.

The numbers of CRTH2$^+$ blood lymphocytes were similar in healthy and asthma subjects, but there was a significant increase in the number of BAL CRTH2$^+$ T-cells in patients with asthma (2.3% versus 0.3%, p<0.05). In both blood and BAL, T$_\text{H2}$ cells producing either IL-4 or IL-13 showed greater expression of CRTH2 than IFN$\gamma$ producing T$_\text{H1}$ cells. Based on the results of this relatively small study (n=11 asthma patients and n=7 healthy subjects), the authors proposed that the low expression levels indicated a limited role of CRTH2 in the control of lymphocyte activity in asthma and that CRTH2 antagonism may not diminish T-cell recruitment to the asthma lung.

Fajt et al reported a larger bronchoscopy study involving mild asthma (steroid naïve; n=11), moderate asthma (using ICS; n=22), and severe asthma patients (n=46), along with a healthy control group (n=33).$^{15}$ In this study, the highest CRTH2 expression levels in BAL cells were observed in severe asthma patients, measured through gene expression and immunocytochemical studies. Interestingly, there were also associations between PGD$_2$, HPGDS, and CRTH2 expression levels and worse clinical outcomes, including exacerbations and asthma control. Stinson et al also reported increased numbers of CRTH2$^+$ inflammatory cells in severe asthma patients compared with healthy subjects, this time in the submucosa of bronchial biopsies.$^{33}$

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**Figure 1** Effects of PGD$_2$–CRTH2 signaling in asthma.

Notes: PGD$_2$ is predominantly released from mast cells following allergen stimulation; other cells such as eosinophils and T$_\text{H2}$ cells may also contribute to PGD$_2$ levels. Interaction of PGD$_2$ with CRTH2 stimulates the recruitment of T2-associated cells to the airways and release of associated cytokines, as well as eosinophil and basophil degranulation and epithelial metaplasia.

Abbreviations: CRTH2, chemoattractant receptor-homologous molecule on T helper type 2 cells; ILC2, innate lymphoid type 2 cells; IL, interleukin; PGD$_2$, prostaglandin D$_2$; T$_\text{H2}$, T-helper type 2 cell.
Bronchial epithelial cells are known to express CRTH2, and in vitro studies using primary human cells have shown that activation of CRTH2 induces epithelial cell migration, mucin production, and metaplasia. Stinson et al showed that the number of CRTH2+ epithelial cells decreased with asthma severity and that this reduction in CRTH2 expression was specifically related to areas of the epithelium that were undergoing metaplasia, which was increased in biopsies from more severe patients. It is yet unclear if the increase in metaplasia in severe asthma is induced by PGD2 or if such an activation results in a reduction in CRTH2 expression. PGD2 activation in other cell types, such as TH2 cells, is known to downregulate CRTH2 expression.

Overall, these studies in asthma have shown a pattern of increased PGD2 and CRTH2 expressions in more severe asthma. CRTH2 expression has also been shown to be elevated in other allergic diseases. Blood eosinophils from atopic kerato-conjunctivitis patients express higher levels of CRTH2 than those from healthy subjects, and these cells showed enhanced migration toward PGD2 and its stable metabolite 13,14-dihydro-15-keto-PGD2, which could explain the increased number of eosinophils in atopic tissue.

The effects of CRTH2 antagonists in asthma

Small-molecule antagonists of the CRTH2 receptor designed for oral administration have been synthesized, with some showing potency in preclinical studies to warrant evaluation in asthma clinical trials. Clinical trials of CRTH2 antagonists on asthma clinical end points are summarized in Table 1. The first CRTH2 antagonist to be evaluated in patients with asthma was OC000459. A 4-week, placebo-controlled parallel group study was conducted in 132 patients who were not using ICS. There was an improvement in FEV1 with OC000459 (200 mg administered twice daily) compared to placebo in the “full analysis population” (7.1% versus 4.3%, respectively). This difference was not statistically significant, but was significant in the “per protocol population” which included only patients who completed the study with valid spirometry measurements (9.8% [210 mL] versus 1.8% [30 mL] improvement; \( p = 0.037 \)). Significant improvements in quality of life and nighttime symptoms were observed with OC000459 compared with placebo in populations in both analyses. Induced sputum was conducted in a subset (n=27), and it showed a significant reduction in eosinophil counts with active treatment, but not placebo. This study provided the first clinical evidence that CRTH2 antagonism could benefit patients with asthma, targeting eosinophilic inflammation.

OC000459 was subsequently assessed at lower doses (25 mg once daily, 200 mg once daily, and 100 mg twice daily) in a placebo-controlled study over 12 weeks in asthma patients not using ICS (n=512 randomized). There was a similar increase in FEV1 in each OC000459 treatment group, with the pooled active treatment dose groups showing a 95 mL difference compared to placebo at week 12 (\( p = 0.024 \)). There were also improvements in the asthma control questionnaire (ACQ) and asthma quality of life questionnaire (AQLQ) scores with OC000459 compared with placebo. The effect of OC000459 on lung function appeared to be greater in patients with atopy and those with higher blood eosinophil counts (>250 cells/µL).

OC000459 has been studied using the inhaled allergen challenge model. This drug attenuated the late asthmatic response, with a less pronounced fall in FEV1 after inhaled allergen, compared with placebo in steroid-naïve patients with asthma. OC000459 also reduced the magnitude of sputum eosinophilia caused by allergen exposure. There was a significant period effect in this placebo-controlled crossover design, with no therapeutic effect on FEV1 observed in patients who received active treatment first. At first glance, one would assume that this was a carryover effect in a crossover study. However, the screening allergen challenge data showed that this group actually had no therapeutic response with OC000459 compared to the screening challenge, suggesting that the drug had no effect in this group rather than a carryover effect. These patients with no drug response had a significantly lower FEV1 at screening, suggesting a variation between individuals in drug response, which is dependent on clinical characteristics. Nevertheless, the CRTH2 antagonist setipiprant also inhibited the allergen-induced late asthmatic response, providing further confirmation of the effects of this drug class on allergic inflammation.

The effects of the CRTH2 antagonist BI671800 have been investigated in asthma patients not using ICS and in a population of asthma patients taking ICS. BI671800 (50, 200, and 400 µg), inhaled fluticasone propionate (220 µg), and placebo (all administered twice daily) were studied over 6 weeks using a parallel group design in asthma patients not using ICS (Trial 1), while a similar design was used in asthma patients taking ICS to evaluate BI671800 400 mg twice daily, montelukast 10 mg once daily, and placebo (Trial 2). In Trial 1, BI671800 caused FEV1 improvements of ~3%–4% compared with placebo (3.98% or 134 mL, \( p = 0.0078 \) at
that showed a significant response.\textsuperscript{56} asthma patients using ICS. No subgroups could be identified AMG853 had no effect on lung function, ACQ, AQLQ, or (using a threshold of 350 cells/mm\textsuperscript{3}). observed in patients with higher blood eosinophil counts 80 mL). In both studies, a greater effect of BI671800 was with montelukast was not statistically significant (2.37%, 142 mL, with placebo (3.87%, 142 mL, \( p \leq 0.005 \)), while the change with montelukast was not statistically significant (2.37%, 80 mL). In both studies, a greater effect of BI671800 was observed in patients with higher blood eosinophil counts (using a threshold of 350 cells/mm\textsuperscript{3}).

The CRTH2 antagonist AZD1981 was also investigated in one trial involving asthma patients where previous ICS treatment was withdrawn at randomization (Study 1), and in another trial in asthma patients where ICS treatment was continued (Study 2).\textsuperscript{18} Study 1 was a parallel group study comparing AZD1981 1000 mg twice daily with placebo, with low-dose ICS being withdrawn on the day of randomization. Study 2 was also a placebo-controlled, parallel group study, but patients continued medium- to high-dose ICS, with any low-acting beta-agonist use stopped. The active treatments were randomized. There were 13 fevipiprant-treatment arms, and studied for 12 weeks. The primary end point was trough FEV\textsubscript{1} at 12 weeks; fevipiprant caused a model-averaged FEV\textsubscript{1} change of 112 mL compared with placebo \( (p=0.0014) \) with fevipiprant (geometric means at baseline and 12 weeks were 5.4\% and 1.1\%, respectively) compared with placebo (geometric means at baseline and 12 weeks were 4.6\% and 3.9\%, respectively). There was also a significant reduction in bronchial submucosal eosinophil counts. This study was not sufficiently powered to properly assess lung function and symptoms, but it confirmed that the principal mechanism of action of CRTH2 antagonists is through blocking eosinophil activation and chemotaxis.

A recent publication has described a dose-ranging, parallel-group Phase II study of fevipiprant in patients with uncontrolled, allergic asthma \((n=1058\) randomized).\textsuperscript{59} Asthma patients taking ICS with evidence of allergy (by skin prick or Immunoglobulin E test) were recruited and treated with inhaled budesonide 200 µg twice daily during the run-in and treatment periods. Patients with an ACQ score \( \geq 1.5 \) at the end of the run-in period, indicating uncontrolled symptoms, were randomized. There were 13 fevipiprant-treatment arms, a placebo arm, and a montelukast arm. Fevipiprant doses from 1 to 450 mg were used, with once daily and twice daily regimes, and studied for 12 weeks. The primary end point was trough FEV\textsubscript{1}, at 12 weeks; fevipiprant caused a model-averaged FEV\textsubscript{1} change of 112 mL compared with placebo \( (p=0.0035) \), while for montelukast, 134 mL was observed \( (p=0.0033) \). The total fevipiprant daily dose of 150 mg caused the greatest FEV\textsubscript{1} change, with 179 and 164 mL observed after 75 mg twice daily \( (p=0.0059) \) and 150 mg once daily.

### Table 1: Clinical trials of CRTH2 antagonists in asthma

<table>
<thead>
<tr>
<th>Study</th>
<th>Drug</th>
<th>ICS use</th>
<th>Number of patients randomized</th>
<th>Duration(weeks)</th>
<th>Primary outcome (mean; 95%CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Barnes et al\textsuperscript{16}</td>
<td>OC00459</td>
<td>No</td>
<td>132</td>
<td>4</td>
<td>No change in FEV\textsubscript{1} (2.44%; (-4.42, 9.31))*</td>
</tr>
<tr>
<td>Pettipher et al\textsuperscript{17}</td>
<td>OC00459</td>
<td>No</td>
<td>512</td>
<td>12</td>
<td>Improved FEV\textsubscript{1} (95 mL)</td>
</tr>
<tr>
<td>Kuna et al\textsuperscript{18}</td>
<td>AZD1981</td>
<td>No</td>
<td>113</td>
<td>4</td>
<td>No difference in PEFR</td>
</tr>
<tr>
<td>Kuna et al\textsuperscript{18}</td>
<td>AZD1981</td>
<td>Yes</td>
<td>368</td>
<td>4</td>
<td>No difference in PEFR</td>
</tr>
<tr>
<td>Hall et al\textsuperscript{19}</td>
<td>BI671800</td>
<td>No</td>
<td>389</td>
<td>6</td>
<td>Improved FEV\textsubscript{1} (134 mL)</td>
</tr>
<tr>
<td>Hall et al\textsuperscript{19}</td>
<td>BI671800</td>
<td>Yes</td>
<td>243</td>
<td>6</td>
<td>Improved FEV\textsubscript{1} (142 mL)</td>
</tr>
<tr>
<td>Busse et al\textsuperscript{20}</td>
<td>AMG853</td>
<td>Yes</td>
<td>397</td>
<td>13</td>
<td>No difference in ACQ</td>
</tr>
<tr>
<td>Gonen et al\textsuperscript{21}</td>
<td>QAW039</td>
<td>Yes</td>
<td>61</td>
<td>12</td>
<td>Reduced sputum eosinophils (ratio 3.5: 1, 7, 7.0)**</td>
</tr>
<tr>
<td>Bateman et al\textsuperscript{22}</td>
<td>QAW039</td>
<td>Yes</td>
<td>1058</td>
<td>12</td>
<td>Improved FEV\textsubscript{1} (112 mL: 4, 175)**</td>
</tr>
</tbody>
</table>

**Notes:** Only placebo-controlled trials of \( \geq 4 \) weeks duration have been included. Primary outcome shows the difference versus placebo; only statistically significant changes are shown. 95\% CI are shown if presented in the publication. *A secondary analysis in the per protocol population was significant; mean difference 7.66\% (0.49, 14.82). **Ratio of geometric mean sputum eosinophil changes for fevipiprant versus placebo. ***Model-averaged FEV\textsubscript{1} for different fevipiprant doses.

**Abbreviations:** ACQ, asthma control questionnaire; CRTH2, chemoattractant receptor-homologous molecule on T helper type 2 cells; FEV\textsubscript{1}, forced expiratory volume in one second; ICS, inhaled corticosteroids; PEFR, peak expiratory flow rate.

400 mg), while a greater improvement was observed with fluticasone propionate (8.62\% or 293 mL, \( p<0.0001 \)). In Trial 2, BI671800 significantly improved FEV\textsubscript{1} compared with placebo (3.87\%, 142 mL, \( p=0.005 \)), while the change with montelukast was not statistically significant (2.37\%, 80 mL). In both studies, a greater effect of BI671800 was observed in patients with higher blood eosinophil counts (using a threshold of 350 cells/mm\textsuperscript{3}).
doses ($p=0.0075$), respectively. No changes in symptoms for
fevipiprant compared to placebo were observed, but there was
a numerical reduction in exacerbations with both fevipiprant
and montelukast treatments compared with placebo. This
study provides information on the optimal doses to be used
in Phase III studies of fevipiprant.

In general, CRTH2 antagonists have been well tolerated
in these clinical trials. The adverse events profile in these
studies has been reported to be similar to placebo.

**The effects of CRTH2 antagonists on nasal allergic inflammation**

CRTH2 antagonism has been demonstrated to attenuate
nasal allergic inflammation. A two-way, placebo-controlled
cross-over study in patients with allergic rhinitis (n=36)
using OC000459 (200 mg twice daily) administered for 8
days exposed subjects to grass pollen for 6 h in an environ-
mental challenge chamber on days 2 and 8 of each treatment
period.59 The primary end point was the total nasal symptom
score (TNSS) averaged over 6 h of allergen challenge on day
8; although there was a significant reduction in TNSS with
OC000459 ($p=0.035$), there was also evidence of a significant
carryover effect between treatment periods.

Krug et al performed a randomized, double-blind,
placebo-controlled, two-way (partial) cross-over study in
patients with allergic rhinitis.60 Patients (n=146) were ran-
donized to receive placebo and one of the following active
treatments; BI671800 twice daily (50, 200, or 400 mg bid),
montelukast 10 mg once daily, or fluticasone propionate
200 µg nasal spray once daily. Patients received treatment for
2 weeks, with a grass pollen challenge in an environmental
challenge chamber performed at the end of each treatment
period. The primary efficacy end point was the TNSS area
under the curve from 0 to 6 hours (AUC$_{0-6h}$) during the aller-
gen challenge. Additional end points included measurement
of nasal secretion weight, nasal flow (rhinomanometry),
nasal secretion inflammatory mediators, and nasal secretion
inflammatory cells. BI671800 200 mg caused a statistically
significant difference in adjusted mean TNSS AUC$_{0-4h}$ values
versus placebo (−17%; $p=0.0026$). BI671800 50 and 400 mg
causd numerically lower TNSS AUC$_{0-4h}$ values compared
with placebo, but these did not reach statistical significance.
Statistically significant differences were observed in the
montelukast and the fluticasone propionate (−15%; $p=0.0115$
and −33%; $p<0.0001$, respectively). BI671800 significantly
reduced nasal eosinophil counts at all the doses studied. In
general, fluticasone propionate had a greater effect than
BI671800 on secondary end point measurements.

Nasal challenge studies are technically demanding, and
the carryover effect observed in the study with OC000459
highlights the practical issues that may be encountered.
Nevertheless, these nasal challenge studies show that CRTH2
antagonism inhibits allergic nasal inflammation, although the
effect is lower in magnitude compared to ICS.

**Future perspectives**

Clinical studies of CRTH2 antagonists have shown a degree
of efficacy in patients with mild to moderate asthma.16,17,53,55
However, the unmet medical need in these patients is low,
with ICS or ICS/LABA combinations providing effective
treatment. Furthermore, CRTH2 antagonists are unlikely
to replace ICS treatment, as the clinical benefits of the for-
mer appear to be lower than the latter in mild to moderate
asthma.55 There is a better way of using CRTH2 antagonists
as an additional anti-inflammatory treatment in patients
already using ICS, and early phase clinical trial evidence
has demonstrated efficacy of this approach.55

There is an upregulation of the PGD$_{2}$–CRTH2 signal-
ing axis in severe asthma compared with mild to moderate
asthma patients and healthy controls.15 Also, there is an unmet
medical need for additional anti-inflammatory treatments in
patients with severe asthma, and the future development of
CRTH2 antagonists should be focused on this patient subset.
Fevipiprant caused a reduction in airway eosinophil numbers
in patients with severe asthma,57 while other CRTH2 anti-
gonists have shown increased clinical efficacy in patients
with higher blood eosinophil counts.59,60 Overall, these findings
suggest that CRTH2 antagonists would be better targeted
toward severe asthma patients with eosinophilic inflamma-
tion. This strategy is supported by the well-described inhibi-
tory effect of CRTH2 antagonists on eosinophil activation
and chemotaxis.52,61,62

Blood eosinophils have been used as a biomarker of
eosinophilic airway inflammation, although the correlation
between these measurements is often modest, leading to false-
positive and false-negative results.63 Nevertheless, blood
eosinophil counts remain a practical method for analyzing
eosinophils in clinical practice, in contrast to sputum analysis
or bronchoscopic sampling. The cut-off point that should be
used to identify eosinophilic asthma is controversial, with
150, 300, and 400 cells/µL being used in various clinical trials
of novel drugs.64,66 If one assumes that the effects of CRTH2
antagonists follow a (blood eosinophil) concentration–
clinical response curve, then the drug effect sizes will show
incremental increases at higher blood eosinophil concentra-
tions, rather than being an “all or nothing” phenomenon.
Exacerbations are the key clinical end points to be assessed in severe asthma long-term Phase III studies. The effect size of CRTH2 antagonists on exacerbations is unknown, but is likely to be greater at higher blood eosinophil concentrations.

Biological treatments targeting eosinophilic inflammation have been successfully developed and are now being used in clinical practice. There are also other biological treatments targeting T2 cytokines, such as IL-4 and IL-13, that are in the late-stage clinical development. The optimum patient selection for these different classes of biological treatments remains under debate, as individuals with T2 inflammation could be potential candidates for anti-eosinophil treatments (such as mepolizumab or benralizumab) or anti-IL4/IL-13 treatment with dupilumab. There are no head-to-head studies to compare such treatments, so clinicians may need to make decisions based on criteria such as higher blood eosinophil counts, which could favor anti-IL5 treatment, and the presence of atopic dermatitis, which may favor treatment with dupilumab. Although these biological treatments have shown good efficacy in severe asthma, there are practical issues to be addressed concerning the regular administration of these systematically administered drugs at intervals such as fortnightly and the high costs of biological treatments. Furthermore, there are some concerns with the long-term side effects of such treatments. Orally administered CRTH2 antagonists may offer an alternative to these biologics based on the good safety profile observed in clinical trials performed to date and the practically easier method of administration. The possibility for CRTH2 antagonists to carve a space in the marketplace based on practicality and safety is dependent on the clinical benefits on exacerbations to be proved in Phase III studies.

Conclusion
CRTH2 antagonists provide a potentially convenient oral treatment for severe asthma. The results of long-term studies in severe asthma focusing on exacerbations will dictate whether these drugs have sufficient efficacy to be approved by drug regulatory authorities for the treatment of asthma. The biology and clinical data indicate that CRTH2 antagonists should be targeted toward eosinophilic asthma. CRTH2 antagonists could provide a practical alternative to biological treatments for this subgroup of patients.

Disclosure
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