

## EDITORIAL OPEN

## The efficacy and safety of inhaled corticosteroids: are we ignoring the potential advantages of ciclesonide?

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Inhaled corticosteroids (ICS) are still the cornerstone of treatment in asthma<sup>1</sup> and many physicians also prescribe ICS in chronic obstructive pulmonary disease.<sup>2,3</sup> The introduction of ICS in the late 1970s caused a large shift in clinical practice regarding the treatment of asthma. Before that time asthma was considered a disease predominantly caused by proliferation of the smooth muscle cell and the only treatment for this was considered to be bronchodilators. When the efficacy of ICS was finally proven by several studies,<sup>4</sup> and ICS were introduced in primary care, the number of deaths, hospital admissions and night-time calls for emergency primary care services almost disappeared. Furthermore, in many parts of the world ICS replaced the use of regular oral corticosteroids, and with that the incidence of severe side effects secondary to oral steroid treatment diminished.

Since then, asthma therapy guidelines have not changed much. The majority of asthma patients are still treated with a combination of ICS and short- or long-acting bronchodilators. Asthma guidelines recommend measuring asthma control and stepping-up of therapy when patients are uncontrolled on current therapy and stepping-down when patients are controlled.<sup>1</sup> However, guidelines may be too conservative in that they ignore some evidence from both mechanistic and clinical studies regarding the development of inhaled steroids and their effects.<sup>1</sup>

The most important factor when deciding on ICS treatment is the balance between benefit and harm of the ICS concerned. Research over the past decades has shown that not all patients are equal. Although some patients might experience very few side effects from even high-dose ICS, research from Foster *et al.*<sup>5</sup> showed that, when asked, most patients mentioned several side effects from their 'brown inhaler' such as hoarseness, thin skin and disturbed vision. Measurements of patient populations by Foster *et al.*<sup>5</sup> with the Inhaled Corticosteroids Questionnaire showed that 45% of patients reported moderate to severe side effects due to ICS when given in a dose  $\geq 800$  mcg budesonide daily.

A study from Chambers *et al.*<sup>6</sup> showed that 38% of patients who started ICS treatment subsequently stopped because of perceived side effects. However, side effects are seldom mentioned by patients, both in randomised controlled trials and in primary care consultations. Possibly this is because patients hesitate to talk about side effects because they are ashamed of discontinuing ICS treatment following such mild (but for them troublesome) side effects. Discontinuation of ICS is the most common cause of lack of asthma control in mild to moderate asthma. Therefore, the scientific community and interested clinicians should not only follow clinical guidelines but should also be alert to new drug developments.

Ciclesonide is an example of a new ICS that aims to reduce side effects by activation only in the lung, with low oral and

high pulmonary deposition, high first pass effect in the liver and high protein binding in the bloodstream.<sup>1</sup> Indeed, several clinical studies have confirmed these well documented claims. Agertoft and Pedersen<sup>7</sup> showed that ciclesonide compared to fluticasone showed no lower leg growth rate reduction whereas fluticasone did. Derom *et al.*<sup>8</sup> showed that cortisol suppression of ciclesonide was comparable with placebo whereas fluticasone led to marked cortisol suppression when both treatments were given at dosages leading to the same level of reduction in bronchial hyperresponsiveness. Finally, our group has shown that patients also perceive a difference in the number and intensity of side effects between ciclesonide and fluticasone in comparable dosages.<sup>9</sup>

The linked paper by Kuo-Chin *et al.*<sup>10</sup> is an open-label, 12-week study comparing the effect of ciclesonide (320 mcg) once daily with budesonide (400 mcg) twice daily in a group of 150 patients with mild to moderate well-controlled asthma (asthma control test score  $\geq 21$ ) who were stepped down from combination ICS/long-acting  $\beta_2$ -agonist therapy. Drug adherence was higher in the ciclesonide group (76.0% vs. 58.7%,  $P=0.03$ ) than in the budesonide group. The results of this study show that patients had favourable results from ciclesonide (once daily) over budesonide (twice daily) and, more importantly, fewer patients discontinued treatment with ciclesonide than with budesonide. More patients in the ciclesonide group (70%) were at least satisfied with their treatment, compared with 47.5% of patients in the budesonide group ( $P=0.02$ ). Unfortunately, the authors have not reported why the patients preferred ciclesonide. Was this because of the efficacy of the drug, the way of administering (once daily), or was the patient perceiving less side effects? One can only guess. However, the importance of the study lies in the withdrawal rate due to loss of asthma control—5% in the ciclesonide group versus 27% in the budesonide group. This might be related to the far worse adherence in the latter group, different ICS particle sizes, or differences in drug deposition.

Therefore, clinicians should be aware that the dosage, inhaler type and the administration frequency, but also the choice of ICS drug, is important in modern clinical practice. The difference might not lie in efficacy per microgram but mostly in patients' appreciation of the balance between efficacy and potentially troublesome side effects. Ciclesonide therefore has several advantages over the other ICS, resulting in high efficacy and low adverse events—both local and systemic—which are relevant for clinical practice.

## COMPETING INTERESTS

Tvd Molen received research funding from Nycomed, previous manufacturer of ciclesonide for the development of the Inhaled Corticosteroid Questionnaire. JWHK declares no competing interest.

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