NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Health Technology Evaluation

Depemokimab for treating severe eosinophilic asthma in people 12 years and over

Final scope

Draft remit/evaluation objective

To appraise the clinical and cost effectiveness of depemokimab within its marketing authorisation for severe eosinophilic asthma in people aged 12 years and over who are inadequately controlled on medium- to high-dose inhaled corticosteroids (ICS) plus another asthma controller.

Background

Asthma is a chronic inflammatory disease associated with variable airflow obstruction and airway hyperresponsiveness. It is characterised by exacerbations where people may experience breathlessness, chest tightness, wheezing, sputum production and cough. Eosinophils, which are a type of white blood cell, are thought to play a major role in airway inflammation in asthma. Severe eosinophilic asthma is a type of asthma that is associated with high levels of eosinophils and recurrent exacerbations. Eosinophilic nasal polyps may also be present.

People with severe asthma often have a severely impaired quality of life which can lead to fatigue, absence from school or work and psychological problems including stress, anxiety and depression. Exacerbations may result in hospitalisation and can be life-threatening. In the UK, up to 7.2 million people have asthma, with around 200,000 people having severe asthma.¹ In England in 2023, there were 1,195 recorded deaths from asthma.²

NICE guideline <u>NG245</u> and guidelines from the <u>Global Initiative for Asthma</u> (GINA) recommend a stepwise approach for treating asthma. Control is maintained by stepping up treatment as necessary and stepping down when control is good. The recommendations are summarised as follows:

- Offer a low-dose inhaled corticosteroid (ICS)/formoterol combination inhaler
 to be taken as needed for symptom relief to people with newly diagnosed
 asthma. If control is inadequate on an as-needed inhaler, or if the
 presentation is more severe at the point of diagnosis, people will then be
 offered daily maintenance treatment with MART (maintenance and reliever
 therapy) in addition to using it as a reliever therapy when needed.
- If control is inadequate on moderate-dose MART, but neither Fractional Exhaled Nitric Oxide nor eosinophil count is raised, people may take either a leukotriene receptor antagonist (LTRA) or a long-acting muscarinic receptor antagonist (LAMA), or both, in addition to moderate-dose MART.

Severe asthma is associated with a high number of exacerbations treated with high dose oral corticosteroids. Some patients take daily oral steroids as maintenance therapy where symptoms are not well controlled despite other treatments. Both daily

maintenance oral steroids and repeated short courses of high dose steroids for exacerbations can cause long term side effects.

People may be offered the following biological treatments if they meet the eligibility criteria:

NICE <u>TA880</u> recommends tezepelumab as an add-on maintenance treatment for severe asthma when treatment with high-dose inhaled corticosteroids plus another maintenance treatment has not worked well enough. It is recommended only if people:

- have had 3 or more exacerbations in the previous year, or
- are having maintenance oral corticosteroids.

NICE <u>TA751</u> recommends dupilumab as an add-on maintenance therapy for treating severe asthma with type 2 inflammation that is inadequately controlled, despite maintenance therapy with high-dose inhaled corticosteroids and another maintenance treatment, only if:

- the person has a blood eosinophil count of 150 cells per microlitre or more and fractional exhaled nitric oxide of 25 parts per billion or more, and has had at least 4 or more exacerbations in the previous 12 months
- the person is not eligible for mepolizumab, reslizumab or benralizumab, or has asthma that has not responded adequately to these biological therapies

NICE <u>TA565</u> and NICE <u>TA671</u> recommend benralizumab and mepolizumab in adults as add-ons for treating severe refractory eosinophilic asthma, only if:

- the blood eosinophil count has been recorded as 300 cells per microlitre or more with 4 or more exacerbations needing systemic corticosteroids in the previous 12 months, or the person has had continuous oral corticosteroids of at least the equivalent of prednisolone 5 mg per day over the previous 6 months, or
- the blood eosinophil count has been recorded as 400 cells per microlitre or more with 3 or more exacerbations needing systemic corticosteroids in the past 12 months.

NICE <u>TA479</u> recommends reslizumab as an add-on for treating severe eosinophilic asthma that is inadequately controlled in adults despite maintenance therapy with high-dose ICS plus another drug, only if the blood eosinophil count has been recorded as 400 cells per microlitre or more with 3 or more severe asthma exacerbations needing systemic corticosteroids in the past 12 months.

NICE <u>TA278</u> recommends omalizumab for treating severe persistent confirmed allergic IgE-mediated asthma as an add-on to optimised standard therapy in people aged 6 and older who need continuous or frequent treatment with oral corticosteroids (4 or more courses in the previous year). Optimised standard therapy is defined as a full trial of and, if tolerated, documented compliance with high-dose ICS, LABAs, LTRAs, theophyllines, oral corticosteroids, and smoking cessation if clinically appropriate.

The technology

Depemokimab (brand name unknown, GlaxoSmithKlein UK Ltd) does not currently have a marketing authorisation in the UK for treating severe eosinophilic asthma. It is

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being studied in phase 3 clinical trials as add add-on treatment for people 12 years and older with severe uncontrolled eosinophilic asthma. It is compared with placebo, mepolizumab and benralizumab.

Intervention(s)	Depemokimab
Population(s)	People 12 years and over with severe eosinophilic asthma which is inadequately controlled on medium- to high-dose inhaled corticosteroids (ICS) plus another asthma controller
Subgroups	If the evidence allows, the following subgroups will be considered: • baseline eosinophil levels • baseline fractional exhaled nitric oxide levels
	people who take maintenance oral corticosteroid treatment
	people who require frequent oral corticosteroid treatment
	Number of exacerbations in previous year
Comparators	For people for whom biologics are indicated or suitable according to NICE guidance, in addition to standard therapy:
	 Tezepelumab Reslizumab Benralizumab Mepolizumab Dupilumab Omalizumab For people for whom currently available biologics are not indicated or suitable: Optimised standard therapy without biologics

Outcomes	The outcome measures to be considered include:
	asthma control
	 incidence of exacerbations, including those which require unscheduled contact with healthcare professionals or hospitalisation
	use of oral corticosteroids
	 patient and clinician evaluation of response
	 lung function (objective measures such as forced expiratory volume, FEV₁; peak expiratory flow, PEF).
	immunogenicity
	mortality
	time to discontinuation
	adverse effects of treatment
	health-related quality of life.
Economic analysis	The reference case stipulates that the cost effectiveness of treatments should be expressed in terms of incremental cost per quality-adjusted life year.
	The reference case stipulates that the time horizon for estimating clinical and cost effectiveness should be sufficiently long to reflect any differences in costs or outcomes between the technologies being compared.
	Costs will be considered from an NHS and Personal Social Services perspective.
	The availability of any commercial arrangements for the intervention, comparator and subsequent treatment technologies will be taken into account.
Other considerations	The availability and cost of biosimilar and generic products should be taken into account.
	Guidance will only be issued in accordance with the marketing authorisation. Where the wording of the therapeutic indication does not include specific treatment combinations, guidance will be issued only in the context of the evidence that has underpinned the marketing authorisation granted by the regulator.
Related NICE recommendations	Related technology appraisals:
	Tezepelumab for treating severe asthma (2023) NICE technology appraisal guidance 880
	Dupilumab for treating severe asthma with type 2 inflammation (2021) NICE technology appraisal guidance 751

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Mepolizumab for treating severe eosinophilic asthma (2021) NICE technology appraisal guidance 671

Benralizumab for treating severe eosinophilic asthma (2019) NICE technology appraisal guidance 565

Reslizumab for treating severe eosinophilic asthma (2017) NICE technology appraisal guidance 479

Omalizumab for treating severe persistent allergic asthma (2013) NICE technology appraisal guidance 278

Related technology appraisals in development:

Depemokimab for treating chronic rhinosinusitis with nasal polyps. NICE technology appraisal guidance (ID6449) Publication date to be confirmed.

Related NICE guidelines:

Asthma: diagnosis, monitoring and chronic asthma management (BTS, NICE, SIGN) (2024) NICE guideline 245

Asthma pathway (BTS, NICE, SIGN) (2024) NICE guideline 244

Related interventional procedures:

Smart Peak Flow for monitoring asthma (2022) NICE MedTech innovation briefing 282

OxyMask for delivering oxygen therapy (2018) NICE MedTech innovation briefing 160

<u>Smartinhaler for asthma</u> (2017) NICE MedTech innovation briefing 90

Alair bronchial thermoplasty system for adults with severe difficult to control asthma (2016) NICE MedTech innovation briefing 71

Related quality standards:

Asthma (2013) NICE quality standard 25

References

https://www.ons.gov.uk/peoplepopulationandcommunity/birthsdeathsandmarriages/d eaths/bulletins/deathsregistrationsummarytables/2023, Accessed: 04/02/2025

¹ What is Asthma? (2024) Asthma + Lung UK, Available: https://www.asthmaandlung.org.uk/conditions/asthma/what-asthma, Accessed: May 2025

² Deaths registered in England and Wales: 2023, (2024) Office for National Statistics. Available: