

**MHRA**  
10 South Colonnade  
Canary Wharf  
London  
E14 4PU  
United Kingdom

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## **Decision Cover Letter**

### **Decision of the licensing authority to:**

grant a product specific waiver

MHRA-101654-PIP01-24

### **Scope of the Application**

#### **Active Substance(s)**

Volrustomig

#### **Condition(s)**

Treatment of head and neck epithelial malignant neoplasms

#### **Pharmaceutical Form(s)**

Concentrate for solution for injection/infusion

#### **Route(s) of Administration**

INTRAVENOUS USE

#### **Name / Corporate name of the PIP applicant**

AstraZeneca UK Limited

#### **Basis for the Decision**

Pursuant to the Human Medicines Regulations 2012, AstraZeneca UK Limited submitted to the licensing authority on 18/10/2024 20:34 BST an application for a Waiver

The procedure started on 02/12/2024 08:28 GMT

1. The licensing authority, having assessed the application in accordance with the Human Medicines Regulations 2012, decides, as set out in the appended summary report:

to grant a product specific waiver

2. The measures and timelines of the paediatric investigation plan are set out in the Annex I.

This decision is forwarded to the applicant, together with its annex and appendix.

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## Final Decision Letter

MHRA-101654-PIP01-24

Of 19/02/2025 17:59 GMT

On the adopted decision for Volrustomig (MHRA-101654-PIP01-24) in accordance with the Human Medicines Regulations 2012.

The licensing authority, in accordance with the Human Medicines Regulations 2012, has adopted this decision:

Granting a waiver in all age groups for the listed condition(s)

This decision applies to a Waiver for Volrustomig , Concentrate for solution for injection/infusion ,  
INTRAVENOUS USE .

This decision is addressed to AstraZeneca UK Limited, 2 Pancras Square, 8th Floor, London, UNITED KINGDOM, N1C 4AG

## ANNEX I

### 1. Waiver

#### 1.1 Condition:

Treatment of head and neck epithelial malignant neoplasms The waiver applies / applied to:  
Paediatric Subset(s): All subsets of the paediatric population from birth to less than 18 years of age  
Pharmaceutical form(s): Concentrate for solution for injection/infusion Route(s) of administration:  
INTRAVENOUS USE Reason for granting waiver: on the grounds that the specific medicinal  
product does not represent a significant therapeutic benefit over existing treatments

### 2. Paediatric Investigation Plan:

#### 2.1 Condition(s):

Not Applicable

**2.2 Indication(s) targeted by the PIP:**

Not Applicable
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**2.3 Subset(s) of the paediatric population concerned by the paediatric development:**

Not Applicable
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**2.4 Pharmaceutical Form(s):**

Not Applicable
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**2.5 Studies:**

Study Type	Number of Studies	Study Description
Quality Measures		
Non-Clinical Studies		
Clinical Studies		
Extrapolation, Modeling & Simulation Studies		
Other Studies		
Other Measures		

**3. Follow-up, completion and deferral of a PIP:**

Concerns on potential long term safety and efficacy issues in relation to paediatric use:	
Date of completion of the paediatric investigation plan:	
Deferral of one or more studies contained in the paediatric investigation plan:	