

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

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## **Decision Cover Letter**

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

accept change(s) to the agreed paediatric investigation plan (MHRA-101583-PIP01-24-M01)  
MHRA-101583-PIP01-24-M02

### **Scope of the Application**

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

Recombinant varicella Zoster Virus Glycoprotein E

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

Prevention of varicella zoster virus (VZV) reactivation

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

Powder and suspension for suspension for injection; Suspension for injection

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

Intramuscular use

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

GlaxoSmithKline UK Limited

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

Pursuant to the Human Medicines Regulations 2012, GlaxoSmithKline UK Limited submitted to the licensing authority on 14/05/2026 10:19 BST an application for a Modification

The procedure started on 19/05/2026 15:31 BST

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 accept change(s) to the agreed paediatric investigation plan

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-101583-PIP01-24-M02

Of 27/08/2026 11:56 BST

On the adopted decision for Recombinant varicella Zoster Virus Glycoprotein E (MHRA-101583-PIP01-24-M02) in accordance with the Human Medicines Regulations 2012.

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

Agreement on modification of a paediatric investigation plan (including modification of a waiver or deferral included in that paediatric investigation plan)

This decision applies to a Modification for Recombinant varicella Zoster Virus Glycoprotein E, Powder and suspension for suspension for injection; Suspension for injection , Intramuscular use .

This decision is addressed to GlaxoSmithKline UK Limited, 79 New Oxford Street, London, UNITED KINGDOM, WC1A 1DG

## ANNEX I

### 1. Waiver

#### 1.1 Condition:

Prevention of varicella zoster virus (VZV) reactivation The waiver applies / applied to: Paediatric Subset(s): The paediatric population from birth to less than 1 year of age Pharmaceutical form(s): Powder and suspension for suspension for injection Suspension for injection Route(s) of administration: Intramuscular use Reason for granting waiver: on the grounds that the disease or condition for which the specific medicinal product is intended does not occur in the specified paediatric subset(s).

### 2. Paediatric Investigation Plan:

#### 2.1 Condition(s):

Prevention of varicella zoster virus (VZV) reactivation

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

Prevention of herpes zoster in immunocompromised subjects

## 2.3 Subset(s) of the paediatric population concerned by the paediatric development:

The paediatric population from 1 year to less than 18 years of age

## 2.4 Pharmaceutical Form(s):

Powder and suspension for suspension for injection Suspension for injection

## 2.5 Studies:

| Study Type                                   | Number of Studies | Study Description                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            |
|----------------------------------------------|-------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Quality Measures                             | 1                 | Study 3 Development of an AS01E-adjuvanted vaccine formulation.                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |
| Non-Clinical Studies                         | 0                 | Not applicable.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |
| Clinical Studies                             | 2                 | Study 1 (Zoster-047) Open-label, randomised study to assess the safety, reactogenicity and immunogenicity of the paediatric formulation of Herpes Zoster candidate vaccine (HZ/su) for the prevention of zoster in immunocompromised children 1 year to less than 18 years of age. Study 2 (Zoster-040) Open-label, non-randomised clinical study to assess safety, immunogenicity and reactogenicity of the paediatric formulation of HZ/su for the prevention of zoster in immunocompromised children 1 year to less than 18 years of age. |
| Extrapolation, Modeling & Simulation Studies | 0                 | Not applicable.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |
| Other Studies                                | 0                 | Not applicable.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |
| Other Measures                               | 0                 | Not applicable.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |

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

|                                                                                           |            |
|-------------------------------------------------------------------------------------------|------------|
| Concerns on potential long term safety and efficacy issues in relation to paediatric use: | No         |
| Date of completion of the paediatric investigation plan:                                  | 30/09/2032 |

|                                                                                        |     |
|----------------------------------------------------------------------------------------|-----|
| <b>Deferral of one or more studies contained in the paediatric investigation plan:</b> | Yes |
|----------------------------------------------------------------------------------------|-----|