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-101468-PIP01-24

Scope of the Application

Active Substance(s)

DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE; baxdrostat

Condition(s)

Treatment of chronic kidney disease

Pharmaceutical Form(s)

Tablet

Route(s) of Administration

ORAL USE

Name / Corporate name of the PIP applicant

AstraZeneca UK Ltd

Basis for the Decision

Pursuant to the Human Medicines Regulations 2012, AstraZeneca UK Ltd submitted to the licensing authority on 26/04/2024 14:42 BST an application for a Waiver

The procedure started on 04/06/2024 14:59 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 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-101468-PIP01-24

Of 16/08/2024 15:31 BST

On the adopted decision for DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE; baxdrostat (MHRA-101468-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 DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE; baxdrostat, Tablet , ORAL USE .

This decision is addressed to AstraZeneca UK Ltd, 2 Pancras Square, London, UNITED KINGDOM, N1C 4AG

ANNEX I

1. Waiver

1.1 Condition:

Treatment of chronic kidney disease 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): Tablet Route(s) of administration: ORAL USE Reason for granting waiver: From birth to less than 2 years of age - on the grounds that the specific medicinal product is likely to be unsafe. From 2 years to less than 18 years of age - 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

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

Not Applicable

2.4 Pharmaceutical Form(s):

Not Applicable

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:	