

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

gov.uk/mhra

Decision Cover Letter

Decision of the licensing authority to:

grant a product specific waiver

MHRA-102403-PIP01-26

Scope of the Application

Active Substance(s)

mebufotenin benzoate

Condition(s)

Treatment of major depressive disorder

Pharmaceutical Form(s)

Dry powder formulation

Route(s) of Administration

Intranasal use

Name / Corporate name of the PIP applicant

Beckley Psytech Limited

Basis for the Decision

Pursuant to the Human Medicines Regulations 2012, Beckley Psytech Limited submitted to the licensing authority on 18/03/2026 16:54 GMT an application for a Waiver

The procedure started on 30/03/2026 09:51 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-102403-PIP01-26

Of 09/07/2026 09:29 BST

On the adopted decision for mebufotenin benzoate (MHRA-102403-PIP01-26) 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 mebufotenin benzoate, Nasal powder , Intranasal use .

This decision is addressed to Beckley Psytech Limited, Beckley Park, Beckley, Oxford, United Kingdom, OX3 9SY, Oxford, UNITED KINGDOM, OX3 9SY

ANNEX I

1. Waiver

1.1 Condition:

Treatment of major depressive disorder 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): Nasal powder Route(s) of administration: Intranasal use Reason for granting waiver: For the paediatric subset from birth to less than 7 years of age: • The disease or condition for which the specific medicinal product is intended does not occur in the specified paediatric subset For the paediatric subset from 7 years to less than 18 years of age: • The specific medicinal product does not represent a significant therapeutic benefit as clinical studies are not feasible

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:	