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-100316-PIP01-21

Scope of the Application

Active Substance(s)

latozinemab

Condition(s)

Treatment of clinically symptomatic or pre-symptomatic frontotemporal dementia (FTD)

Pharmaceutical Form(s)

Solution for infusion

Route(s) of Administration

Intravenous use

Name / Corporate name of the PIP applicant

Alector Inc.

Basis for the Decision

Pursuant to the Human Medicines Regulations 2012, Alector Inc. submitted to the licensing authority on 14/12/2021 16:34 GMT an application for a Paediatric Investigation Plan

The procedure started on 19/05/2022 15:04 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-100316-PIP01-21

Of 20/06/2022 16:51 BST

On the adopted decision for latozinemab (MHRA-100316-PIP01-21) 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 Paediatric Investigation Plan for latozinemab , Solution for infusion , Intravenous use .

This decision is addressed to Alector Inc., 131 Oyster Point Boulevard, Ste 600, South San Francisco, United States, CA94080

ANNEX I

1. Waiver

1.1 Condition:

Treatment of clinically symptomatic or pre-symptomatic frontotemporal dementia (FTD) The waiver applies / applied to: Paediatric Subset(s): The paediatric population from birth to less than 18 years of age Pharmaceutical form(s): Solution for infusion Route(s) of administration: Intravenous 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):

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	0	Not applicable
Non-Clinical Studies	0	Not applicable
Clinical Studies	0	Not applicable
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:	
Date of completion of the paediatric investigation plan:	
Deferral of one or more studies contained in the paediatric investigation plan:	