

Medicines & Healthcare products Regulatory Agency

> 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-101718-PIP01-24

Scope of the Application

Active Substance(s)

Human alpha-1-proteinase inhibitor immunoglobulin G fusion protein, recombinant

Condition(s)

Treatment of emphysema secondary to congenital deficiency of alpha-1 antitrypsin

Pharmaceutical Form(s)

All pharmaceutical forms

Route(s) of Administration

All routes of administration

Name / Corporate name of the PIP applicant

Sanofi AATD Inc.

Basis for the Decision

Pursuant to the Human Medicines Regulations 2012, Sanofi AATD Inc. submitted to the licensing authority on 29/11/2024 16:11 GMT an application for a Waiver

The procedure started on 13/01/2025 14:14 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-101718-PIP01-24

Of 01/04/2025 14:30 BST

On the adopted decision for Human alpha-1-proteinase inhibitor immunoglobulin G fusion protein, recombinant (MHRA-101718-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 Human alpha-1-proteinase inhibitor immunoglobulin G fusion protein, recombinant, All pharmaceutical forms, All routes of administration.

This decision is addressed to Sanofi AATD Inc., 450 Water Street, Cambridge, UNITED STATES OF AMERICA, 02141

ANNEX I

1. Waiver

1.1 Condition:

Treatment of emphysema secondary to congenital deficiency of alpha-1 antitrypsin 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): All pharmaceutical forms Route(s) of administration: All routes of administration 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 | | |
| 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: | |