

9 January 2024

## Treatment of Rare Diseases

I have a Freedom of Information request - I would be grateful if you could please answer the following questions:

### 1. Within your Trust, how many patients currently have a diagnosis for:

#### Adult Services

- Fabry Disease (ICD10 code E75.21) 53
- Gaucher Disease (ICD10 code E75.22) <5
- Pompe Disease (ICD10 Code E74.02) <5
- Pompe Disease (ICD10 Code E74.02) 0
- Infantile-onset (Patients Diagnosed before age 1)
- MPS II (Hunter Syndrome) (ICD10 code E76.1) <5

**Use of <5 (less than five):** We are unable to provide an exact figure - exempt from release under Section 40(2) of the Freedom of Information Act, as the small numbers involved could make patients personally identifiable. Disclosure would constitute a breach of the principles of the General Data Protection Regulation 2018.

#### Paediatric Services

- Fabry Disease (ICD10 code E75.21) 7
- Gaucher Disease (ICD10 code E75.22) 0
- Pompe Disease (ICD10 Code E74.02) 0
- Pompe Disease (ICD10 Code E74.02) 0
- Infantile-onset (Patients Diagnosed before age 1)
- MPS II (Hunter Syndrome) 0
- (ICD10 code E76.1)

### 2. Of the patients above, how many patients have been newly diagnosed within the past 3 months for:

#### Adult Services

- Fabry Disease (ICD10 code E75.21) <5
- Gaucher Disease (ICD10 code E75.22) 0
- Pompe Disease (ICD10 Code E74.02) 0
- Pompe Disease (ICD10 Code E74.02) Infantile-onset 0
- (Patients Diagnosed before age 1)
- MPS II (Hunter Syndrome) (ICD10 code E76.1) 0

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Paediatric Services

- Fabry Disease (ICD10 code E75.21) 0
- Gaucher Disease (ICD10 code E75.22) 0
- Pompe Disease (ICD10 Code E74.02) 0
- Pompe Disease (ICD10 Code E74.02) Infantile-onset (Patients Diagnosed before age 1) 0
- MPS II (Hunter Syndrome) (ICD10 code E76.1) 0

3. How many patients have been treated in the last 3 months with the following products:

Adult Services

- Replagal (agalsidase alfa) <5
- Fabrazyme (algalsidase beta) 13
- Galafold (migalastat) 12
- Elfabrio (pegunigalsidase alfa) 0
- VPRIV (velaglucerase alfa) <5
- Cerezyme (imiglucerase) <5
- Cerdelga (eliglustat) 0
- Zavesca (miglustat) 0
- Myozyme (alglucosidase alfa) <5
- Nexviazyme (avalglucosidase alfa) 0
- AT-GAA (cipaglucosidase alfa/miglustat) 0

Paediatric Services

- Replagal (agalsidase alfa) 0
- Fabrazyme (algalsidase beta) 0
- Galafold (migalastat) 0
- Elfabrio (pegunigalsidase alfa) 0
- VPRIV (velaglucerase alfa) 0
- Cerezyme (imiglucerase) 0
- Cerdelga (eliglustat) 0
- Zavesca (miglustat) <5 (NP Type C)
- Myozyme (alglucosidase alfa) 0
- Nexviazyme (avalglucosidase alfa) 0
- AT-GAA (cipaglucosidase alfa/miglustat) 0

4. Do you participate in any clinical trials for Fabry Disease? If so, can you please provide the name of each trial along with the number of patients taking part?

Adult - No

Paediatrics - No

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5. Do you participate in any clinical trials for Gaucher Disease? If so, can you please provide the name of each trial along with the number of patients taking part?

Adult - No

Paediatrics - No