

23 August 2023

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) 52
- 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) (ICD10 code E76.1) 0

Paediatric Services

- Fabry Disease (ICD10 code E75.21) 6
- 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) (ICD10 code E76.1) 0

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) <5
- Pompe Disease (ICD10 Code E74.02) <5
- Pompe Disease (ICD10 Code E74.02) Infantile-onset (Patients Diagnosed before age 1) 0
- 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 this could make patients personally identifiable. Disclosure would constitute a breach of the principles of the General Data Protection Regulation 2018.

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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) 14
- Galafold (migalastat) 10
- Elfabrio (pegunigalsidase alfa) 0
- VPRIV (velaglucerase alfa) <5
- Cerezyme (imiglucerase) <5
- Cerdelga (eliglustat) 0
- Zavesca (miglustat) <5
- Myozyme (alglucosidase alfa) 0
- 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) 0
- Myozyme (alglucosidase alfa) 0
- Nexviazyme (avalglucosidase alfa) 0
- AT-GAA (cipaglucosidase alfa/miglustat) <5

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

No