

SOUTHERN HEALTH & SOCIAL CARE TRUST

NUMBER OF INPATIENT HOSPITAL ADMISSION WITH A DIAGNOSES OF (ATTR-CM) OR (ATTR-PN)

ADMISSION DATE BETWEEN 01/09/2025 TO 31/12/2025

Notes:

The Information Team was only able to provide responses to Questions 3 and 4. As no date range was specified, the data has been based on the last four fully clinically coded months (September–December 2025), in line with the timeframe used for Questions 1 and 2.

Coding completeness for the period is as follows:

- September 2025: 99.9% coded
- October 2025: 100% coded
- November 2025: 100% coded
- December 2025: 100% coded

Wild type Transthyretin Amyloid Cardiomyopathy (ATTR CM) was identified using the following ICD 10 codes:

- E85.4 – Organ limited amyloidosis
- I43.1 – Cardiomyopathy in metabolic diseases

Hereditary Transthyretin Amyloid Polyneuropathy (ATTR PN) was identified using the following ICD 10 codes:

- G63.3* – Polyneuropathy in other endocrine and metabolic diseases
- E85.1 – Neuropathic hereditary amyloidosis

Codes confirmed by Clinical Coding 22/05/2026.

Regular Attenders have been excluded.

Patient Class = Outpatient. Community bedded, home treatment and inpatient mental health have also been excluded.

3. What is the total number of patients with wild type Transthyretin Amyloid Cardiomyopathy (ATTR-CM) being treated in your Trust?

16

4. What is the total number of patients with hereditary Transthyretin Amyloid Polyneuropathy (ATTR-PN) being treated in your Trust?

0