

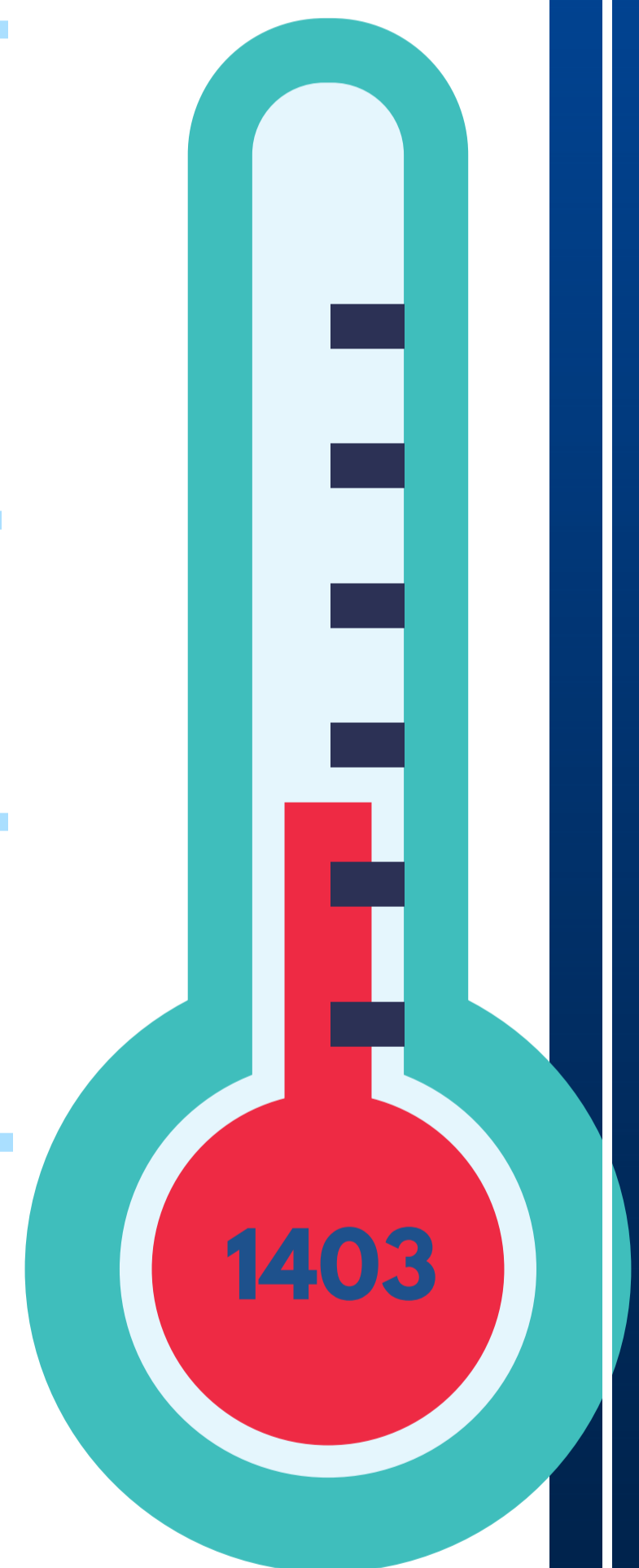
# Australian Epilepsy Research Registry

The primary aim of the registry is to learn valuable information about epilepsy, and its social and psychological impacts that can be used to improve the lives of people affected by this condition.



Year	2010	2013	2016
Enrollments by year	621	883	1415
Participants/ Response rate per Wave survey	343 55%	318 36%	N/A N/A
Average age (years)	43	40	38
Gender (Male/Female)	43%/57%	30%/63%	21%/79%

**Target 2000**



**Current Total**

In 2006 the number of Australians living with epilepsy was estimated at 133,700 or 1 in every 147.

When underreporting was taken into account this number increased to 268,473 people or 1 in every 73 Australians\*

**Currently only estimates are available, hence the real number of people with epilepsy is still unknown.**

\*Source: Brown, K. Indicators of the social consequences of epilepsy in Society. Behaviour, ed. J Pinikahana and C Walker, Nova Biomedical, 2011: 17-28.