NEUROLOGICAL CONDITIONS **AT A GLANCE**



CAN AFFECT PEOPLE

ARE ONE OF THE **BIGGEST THREATS** TO OUR HEAI

Are the leading cause of DISABILITY AND SECOND CAUSE OF **DEATH** WORLDWIDE



every year in Aotearoa

IMPACT 1 IN 3 NEW ZEALANDERS in their lifetime

Are the cause of death for 1 IN 5 NEW ZEALANDERS

NUMBERS AFFECTED:



1.3 million New Zealanders suffer from tension-type headaches 642,000 from Migraine

300,000 from Rare Disorders

83,000 from Dementia

70,000 from Myalgic Encephalomyelitis (ME) & Long Covid





ARE RAPIDLY **INCREASING**

AT ANY AGE

as our population grows and ages



ARE **UNDER** RESOURCED

There are significant workforce shortages and underfunded community services

A lack of resources means

LONG WAIT TIMES FOR SPECIALISTS AND DELAYS IN DIAGNOSIS AND TREATMENT



64,000 from Stroke

50,000 from Epilepsy

37,100 from Traumatic Brain

12,000 from Parkinsons; and

4,100 from Multiple Sclerosis

4,000 from Neuro-Muscular Conditions.

NEUROLOGICAL ALLIANCE PRIORITIES

Address health workforce shortages to speed up diagnosis and treatment

There are chronic shortages in neurologists, specialist nurses, physiotherapists, occupational therapists, psychotherapists, speech language therapists and home-carers. NGOs also face challenges with recruitment and retention of specialist staff, as underfunding of the not-forprofit sector by Government undermines our ability to compete for staff within the wider health sector.



Increase funding for pharmaceutical treatment to improve quality of life

Pharmac's complex decision-making processes, lengthy timeframes and strict rationing of pharmaceutical treatments is impacting the quality of life and in some cases, the life expectancy of people with neurological conditions. The Neurological Alliance is calling for full implementation of the recommendations from the Pharmac Review 2021.

Increase funding for community-based care and support services

The majority of people with neurological conditions live at home independently or supported by family whānau carers and rely on services and programmes delivered in their communities. Shortfalls in health funding for community-based workers and historical underfunding of the sector means that our members can only reach a fraction of people who need support. Family whānau carers are also burnt out and struggling to access respite.

Improve data collection on neurological conditions

There is no systematic collection of data on neurological conditions by the Government. Without data on prevalence and incidence of different neurological conditions the Government, Pharmac and Te Whatu Ora are not able to accurately plan service provision, workforce resource requirements, monitor the accessibility of services, or ensure equitable health outcomes.

Data gaps also make it difficult to model the social and economic costs of neurological conditions and creates challenges for neurological research.