

Migraine Foundation Aotearoa New Zealand

Strategic Plan - 2023/24

Review date: February 2025

Migraine Foundation Aotearoa New Zealand

Strategic Plan - 2023/24



To raise awareness of the impact of migraine disease and support people living with migraine in Aotearoa New Zealand.

Vision

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To minimise the health, economic and social burden of migraine disease in Aotearoa New Zealand.



Values/culture:

Migraine Foundation Aotearoa New Zealand Incorporated (MFANZ) is an inclusive, open, active and action-orientated organisation. We value respectful partnerships and relationships and evidence-based information. Supporting people with migraine is at the core of everything we do.

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Strategic objectives

- Connect and support people living with migraine disease.
- Increase awareness of migraine disease and advocate for positive change.
- Create and disseminate robust information about migraine.
- Support New Zealand-focused research on migraine and collaborate with researchers in planning, recruitment and dissemination of findings.
- Build a sustainable, trusted and respected national migraine organisation.

Why do we exist?

Globally, migraine disease affects approximately 1 in 7 people and is one of the most disabling medical conditions. Migraine disease is estimated to affect 642,000 people in Aotearoa New Zealand, although more research is needed for a more accurate picture of the prevalence and impact of migraine in Aotearoa New Zealand.

Migraine disease is under-researched, under-funded and under-diagnosed. Treatments and knowledge about migraine have improved in recent years; sadly Aotearoa New Zealand has been left behind, with poor access to migraine specialists, newer migrainespecific treatments and basic migraine care.

Migraine Foundation Aotearoa New Zealand exists for all people in Aotearoa New Zealand living with migraine. We're dedicated to connecting people living with migraine, supporting positive change and ensuring our voice is heard.



Pillars of service