**Petition of Sue Haldane for Rare Disorders NZ: Develop a National Rare Disorder Framework**

**Petition request**

That the House of Representatives urge the Government to acknowledge the universal challenges faced by people living with a rare disease, and the unfairness within the current system, by committing to the development of a New Zealand National Rare Disorder Framework.

**Petition reason**

There are around 300,000 Kiwis living with a rare disease. NZ lags behind most OECD countries in supporting people living with rare disorders and their families to access the best healthcare. We believe a shift in mind-set is needed for rare disorders to stop being considered in isolation, and instead to be regarded as a significant factor within health policy frameworks. This is in alignment with the global rare disease movement headed by Rare Disease International.

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