

To the House of Representatives

NZ Patients with Rare Diseases need treatment too

The Petition of Samantha Lenik

I request that the House of Representatives ensure PHARMAC has sufficient resources to deliver on its rare disease policy, and provide urgent access to the eight treatments it signalled when it set up this policy over two and a half years ago.

We want patients with Pompe disease, Fabry disease, Morquio disease, Hurler disease, Hunter disease, aHUS, PNH and Cystic Fibrosis to get the treatments they need in a fair and equitable way. Most of these diseases have no other treatment options available, and Samantha and these other Kiwis need these proven treatments to live a better quality of life with their families, for longer.

We want no patient left behind - no family forgotten

Name	Signature

***Please return your signed forms to
167 Hollister Lane, Ohauti, Tauranga before 11th November 2016***