

TO THE HONOURABLE THE SPEAKER AND MEMBERS OF THE HOUSE OF REPRESENTATIVES

This petition of Australian Citizens, who are friends, family or supporters of Australian sufferers of Paroxysmal Nocturnal Haemoglobinuria (PNH) disease, a rare and potentially fatal disease of the blood **draws to the attention of the House** the need to publicly fund out of the Federal Government, the only currently available treatment for PNH sufferers, namely Soliris® (Eculizumab) as a matter of urgency.

We therefore ask the House to introduce and pass any legislation or to take any administrative action available to the House that will enable sufferers of PNH disease to urgently receive breakthrough life saving treatment for this very rare and debilitating disease.