

Summary of Health Consumer Organisations supported by Medicines Australia Member Companies
For the period: 01 Jan 2017 - 31 Dec 2017

Company: CSL Behring

Name of Health Consumer Organisation	Description and/or purpose of support	Nature of support - monetary value (or equivalent) or description of non-financial support
Myositis Association Australia	The Myositis Association Australia is an organisation dedicated to supporting myositis patients and their families. Support of the Myositis Association Australia 2016 patient conference was provided to assist patient education with medical specialists from Australia and the USA as well as allied health practitioners, carers and lifestyle experts.	\$4,000.00
Haemophilia Foundation of Australia (HFA)	The Haemophilia Foundation of Australia (HFA) is an organisation that represents people with haemophilia, von Willebrand disorder and other inherited bleeding disorders. Support was provided to assist the HFA raise community awareness of haemophilia by developing educational video stories of patients with haemophilia.	\$20,000.00
Haemophilia Foundation of Australia (HFA)	The Haemophilia Foundation of Australia (HFA) is an organisation that represents people with haemophilia, von Willebrand disorder and other inherited bleeding disorders. Support of the 18th Australian and New Zealand bi-annual Conference on Haemophilia and Rare Bleeding Disorders in 2017 was provided to assist with patient and carer education with medical specialists and allied health.	\$30,000.00
Immune Deficiencies Foundation Australia (IDFA)	The Immune Deficiencies Foundation Australia (IDFA) is a not-for-profit organisation raising awareness and caring for people who have Primary (Genetic) and Secondary (due to chemotherapy and other treatments/ or autoimmune illness). Support was provided to assist the IDFA's educational meetings to help patients and their carers manage their disease and move from hospital to home treatment and care.	\$31,590.91
Inflammatory Neuropathy Support Group of Victoria Inc (InGroup)	The Inflammatory Neuropathy Support Group of Victoria Inc (InGroup) is a non profit organisation that supports patients with Guillain-Barré Syndrome (GBS) or Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) or other Inflammatory Neuropathies. Support was provided to assist the InGroup's educational website platform that helps to educate patients and their carers around their disease.	\$200.00
Myasthenia Alliance Australia (MAA)	The Myasthenia Alliance Australia (MAA) is an organisation that supports patients with myasthenic syndrome and represents the needs of myasthenics from all states in Australia. Support was provided for the MAA 2017 patient conference to assist with patient education with medical specialists.	\$3,719.86. Includes provision of food and beverages
Australian Primary Immunodeficiency Patient Support (AusPIPS)	The Australian Primary Immunodeficiency Patient Support (AusPIPS) group is an organisation that supports patients with Primary Immune Deficiency (PID) and associated chronic illnesses in Australia. Funding was provided to support the AusPIP's educational patient meetings regarding their disease, improve patient advocacy and facilitate patient outcomes through educational meetings with health care professionals.	\$14,100.00
Australian Primary Immunodeficiency Patient Support (AusPIPS)	The Australian Primary Immunodeficiency Patient Support (AusPIPS) group is an organisation that supports patients with Primary Immune Deficiency (PID) and associated chronic illnesses in Australia. Support was provided to AusPIPs to allow representation of the group at the Australasian Society for Clinical Immunology and Allergy (ASCI) 28th Annual Conference to raise awareness of PID amongst healthcare professionals.	\$2,430.59
HAE Australasia	HAE Australasia is a not for profit patient advocacy organisation, dedicated to providing support to Australian and New Zealand hereditary angioedema (HAE) patients and their families, as well as raising awareness of HAE resulting from C1-inhibitor deficiency. Support was provided for the HAE Australasia 2017 National Patient and Carers Conference to assist patient education with medical specialists from Australia.	\$10,000.00
HAE Australasia	HAE Australasia is a not for profit patient advocacy organisation, dedicated to providing support to Australian and New Zealand hereditary angioedema (HAE) patients and their families, as well as raising awareness of HAE resulting from C1-inhibitor deficiency. Support was provided to HAE Australasia to help increase Australian and New Zealand community awareness and understanding of hereditary angioedema through their awareness program.	\$20,000.00
HAE Australasia	HAE Australasia is a not for profit patient advocacy organisation, dedicated to providing support to Australian and New Zealand hereditary angioedema (HAE) patients and their families, as well as raising awareness of HAE resulting from C1-inhibitor deficiency. Support was provided for the HAE Australasia's State Patient Meetings and Psychology Workshops to help patients living with HAE and who may be experiencing depression and anxiety due to their disease.	\$13,000.00
HAE Australasia	HAE Australasia is a not for profit patient advocacy organisation, dedicated to providing support to Australian and New Zealand hereditary angioedema (HAE) patients and their families, as well as raising awareness of HAE resulting from C1-inhibitor deficiency. Support was provided to assist with HAE Australasia's patient database project.	\$20,000.00