

REPORTING PERIOD: 01 Jan 2018 - 31 Dec 2018

COMPANY NAME: 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
Haemophilia Foundation Australia	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's 2018 Youth Lead Connect (YLC) program to build education and life skills for young people in the bleeding disorders community.	\$25,000.00
Haemophilia Foundation Australia	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 in the development of personal video stories of people living with bleeding disorders to raise awareness of bleeding disorders.	\$20,000.00
Haemophilia Foundation of Australia	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 a continuing education plan to enable members of the Australia and New Zealand Physiotherapy Haemophilia Group (ANZPHG) to develop and enhance the national standard of physiotherapy care.	\$1,335.00
Alpha-1 Foundation of Australia	The Alpha-1 Association of Australia (AAA) is a not for profit organisation representing patients with Alpha-1 Antitrypsin Deficiency (Alpha-1) and their carers throughout Australia. The AAA focuses on increasing Alpha-1 awareness and education, diagnosis, improving patient access to therapy and encouraging research. Support was provided to sponsor attendance to the Annual Scientific Meeting of The Australia and New Zealand Society of Respiratory Science and The Thoracic Society of Australia and New Zealand (TSANZSRS) 2018 conference.	\$5,000.00
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 disease 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.	\$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 assist with HAE Australasia's Global Multicentre Registry.	\$10,000.00

Immune Deficiency Foundation of Australia	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 for sponsorship of the 2018 Young Adult Support conference for patients living with Primary Immune Deficiencies and patient education through the printing of the Secondary Immune Deficiency Patient booklet.	\$40,000.00
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 help increase Australian community awareness and understanding of PID through their disease awareness program including development of a website and patient support materials.	\$36,363.64