REPORTING PERIOD: 01 Jan 2020 - 31 December 2020
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
Alpha-1 Association 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) 2020 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 support the HAE Australasia Patient Survey to gather burden of illness and current treatment information to support access to modern treatments in Australia and New Zealand.	$6,000.00
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.	$4,000.00
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 support patient education with healthcare professionals and allied health professionals regarding hereditary angioedema and pandemics to help improve patient outcomes.	$20,000.00
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 HFA to support training for youth leaders and mentors in the bleeding disorders community, and advancing HFA's digital footprint with youth.	$25,000.00
Immune Deficiency Foundation 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 to promote awareness of Primary Immunodeficiency (PID) to improve diagnosis and treatment and support access to services to support and improve patient quality of life.	$27,272.73
Myeloma Australia	Myeloma Australia is a not-for-profit organisation that supports, educates and advocates for people who are living with myeloma. Support was provided for to improve diagnosis and treatment for Secondary Immune Deficiencies (SID) and support access to services to improve patient quality of life.	$20,000.00
Rare Voices Australia	Support of the Rare Voices Australia's Table of Companies for the remainder of calendar year 2020	$7,500.00