Country	Year	Name of Organisation	Nature of Organisation	Reason for Support	Amount Paid	Currency
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	Sponsor a BPA Connect (UK Porphyria PAG Meeting)	2000 GBP	
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	Sponsor a BPA Connect 2021 event	2000	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	UK Porphyria PAG Talk	210	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	AHP Patient Symptom Tracker	280	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	AHP Family Follow Up, Meeting in a a box (MIAB)	210	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	PEGASUS Virtual Training - Social Media	560	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	AHP MIAB VAB	280	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	AHP Family Follow Up, Meeting in a a box (MIAB)	280	GBP
United Kingdom	2021	The British Porphyria Association	Aims to support and educate patients, relatives and medical professionals about the porphyrias, so as to improve the lives of those living with its effects.	UK Porphyria PAG Talk	210	GBP
United Kingdom	2021	Cardiomyopathy UK	The specialist national charity for people affected by cardiomyopathy, a condition that affects the heart muscle. Their vision is that everyone affected by cardiomyopathy should live a long and fulfilling life.	Donation towards an improvement programme for patient care	140	GBP
United Kingdom	2021	Cardiomyopathy UK	The specialist national charity for people affected by cardiomyopathy, a condition that affects the heart muscle. Their vision is that everyone affected by cardiomyopathy should live a long and fulfilling life.	Consultancy on Patient Advocacy and Engagement projects incl PAG Summit	10000	GBP
United Kingdom	2021	Cardiomyopathy UK	The specialist national charity for people affected by cardiomyopathy, a condition that affects the heart muscle. Their vision is that everyone affected by cardiomyopathy should live a long and fulfilling life.	Donation towards an improvement programme for patient care	1820	GBP
United Kingdom	2021	Metabolic Support UK	The leading patient organisation for Inherited Metabolic Disorders supporting thousands of patients worldwide.	PH1 Meeting	770	GBP
United Kingdom	2021	UK ATTR Amyloidosis Patients	A place where patients with amyloidosis, family, friends and carers from all over the UK can connect, communicate and help each other.	One Step Further is a campaign that aimed to highlight the impact conditions such as hATTR amyloidosis have on people and carers affected by the condition, and more widely provide support to those in need. The campaign materials will be used by UKATPA on their website	3500	GBP
United Kingdom	2021	UK ATTR Amyloidosis Patients	A place where patients with amyloidosis, family, friends and carers from all over the UK can connect, communicate and help each other.	The National Amyloidosis Centre developed this website as a public service to provide patients and their family and friends with a comprehensive, accurate and up to date resource on all types of amyloidosis.	5000	GBP