

## **Transparency in our dealings with the Healthcare and Patient Communities**

GW is passionate about advancing medicines in the fields of cannabinoid science and developing therapies to address serious medical needs with limited treatment options, particularly in neurological conditions. Healthcare professionals, healthcare organisations and the patient community all have an important role to play in this work. There is much that we can learn from them about the conditions they treat or live with as well as the everyday practical use of our medicines. This collaboration is essential to drive innovation in the development of medicines and to deliver the best treatment options for patients.

We also recognise that it is crucial that patients and the public have faith in our industry and the important work we are carrying out to advance research and treatment in areas of high unmet need. That is why we are committed to transparency in our financial dealings with the healthcare and patient communities.

To the extent we are permitted by law, we disclose our payments to healthcare professionals and healthcare organisations in the United Kingdom on the ABPI Disclosure platform annually on or around 30th of June covering the previous calendar year. Payments in France are published on the DGS (Department of the Ministry of Health) National Public Platform by 1st September of each year for the benefits granted during the first half of the year and by 1st March of each year for the benefits granted during the second half of the preceding year. All other payments to HCPs in other countries outside of the United States are uploaded to our websites.

### **Patient Organisation Support**

GW was founded to respond to the needs of patients. As a result, we understand the importance of partnering with Patient Organisations, patients and carers in order to deliver better medicines and better outcomes for patients. GW has a legitimate interest in supporting initiatives and organisations related to the advancement of research, disease awareness, education, and patient programs. GW also relies upon the expert knowledge provided by patient organisations. As such, the principle purpose of exchange with Patient Organisations is to enhance patient care, improve the lives of patients and improve the health of communities which we serve. Increasing transparent engagements with Patient Organisations has led to better trials, better engagements and greater communication throughout the life cycle of medicines. This has resulted in better patient outcomes.

GW is committed to disclosing transparently all financial support and significant non-financial support given to Patient Organisations. Below are the details of patient organisations we have supported across Europe.

### **2020**

Patient Organisation	Finance (GBP)	Description
AEER - Associazione Epilessia Emilia Romagna	4,542.28	Contribution to support services in the following areas: One-On-One support to families who have a child with epilepsy. Adult Social Support for adults who have epilepsy including their families and caregivers. Community Services social worker who can provide additional information on epilepsy, advocacy,

		<p>support, community resource referrals, and linkage to appropriate Epilepsy Association services.</p> <p>Seminar Series- The Seminar Series are focused events that address issues and concerns voiced by the agency's clients and other individuals affected by epilepsy and their families. These seminars feature presentations by neurologists on relevant epilepsy related topics.</p>
Alliance Syndrome de Dravet	45,195.85	To fund a pilot study for ASD to understand support required by families and medical staff who support or care for ASD patients.
Apoyodravet - Asociacion de Voluntarios en Favor	8,823.37	To support a series of initiatives: International Day of Dravet Syndrome 2020. Informational services to affected people and families
Associazione Famiglie LGS Italia	13,844.04	To support the association with its work linked to clinical, scientific and social aspects of Lennox-Gastaut
Associazione Sclerosi Tuberosa - APS	9,195.00	To provide support in financing research on TSC, creation of the "Register for scientific and clinical research on Tuberous Sclerosis", in-training through scientific events, activities for people affected by the disease, communication projects for the awareness and diffusion of knowledge on TSC, advice and relief to caregivers.
Deutsche Epilepsievereinigung e V	4,549.15	The donation will be used to support a project for the creation of a nationwide network of counselling centres (NNCC) for patients with epilepsy who are newly diagnosed and their relatives as well as public, teachers, and educators.
Dravet Italia Onlus	35,871.00	<p><b>Monies were provided to support three separate projects. Project details are below:</b></p> <p><b>PROJECT DAND - Dravet Associated Neuropsychiatric Disorders Scale.</b> The project foresees the realization and validation of an ad hoc scale for Dravet's Syndrome - DAND for the evaluation of the different clinical aspects of DRAVET SYNDROME aimed at defining the areas of need so as to develop a multidisciplinary care approach adapted to the individual subject.</p> <p><b>CONVENTION 10 YEARS DRAVET</b> Summarize the 10 years of activity, and presenting the future needs for developments on Dravet Syndrome with the round table "Horizons</p>

		<p>for Dravet Syndrome"</p> <p><b>S.A.D.S. Sleep and Cicadian Rhythm in Dravet Syndrome</b></p> <p>To record the sleep of patients suffering from Dravet's syndrome for 7 days, in particular quantity and quality, with a device with actigraphy recording, based on a validated and certified algorithm.</p>
Dravet Syndrome European Federation	76,272.71	<p>To support a patient registry.</p> <p>Project Deliverables:</p> <ol style="list-style-type: none"> <li>1) Increase the number of accredited Centers and their geographical distribution</li> <li>2) Reach within the next two years the number of approximately 1.000 patients (including patients with DS, SCN1A mutation carriers with other phenotypes, and patients with PCDH19 mutations)</li> <li>3) Realize an international network allowing to input data of patients provided by physicians that are not members of the centres</li> <li>4) Create a collection of subjects with highly characterized genetic, clinical, neurophysiological, neuropsychological, treatment, evolutive aspects, useful for subsequent analysis and researches</li> <li>5) Implement the Registry contributing to a more uniform and homogenous optimal clinical management of the patients.</li> </ol>
Dravet Syndrome Foundation Delegacion en Espania	8,487.73	<p>European Dravet experts and other Dravet patient advocacy associations from Italy, France and Croatia launched on April 14, 2020 a worldwide online survey targeted to families and/or caregivers of diagnosed Dravet patients. The survey was open for 1 month, and it aimed at understanding whether and how the COVID-19 pandemic affected their daily lives. This included, health status and the Dravet symptoms of the patients, access to Dravet-specific medicines, doctors and emergency rooms, and the continuity of therapies and clinical trials followed by the patients. The results of this global survey will be released in an international publication with authorship (Dravet Syndrome Foundation) shared by all the collaborators who participated in the design, analysis and draft of the survey and its results.</p>

	4,507.95	Support for 'Dravet Pillows' to help reduce the risk of SUDEP.
Dravet Syndrome UK	26,563	Family event for children with DS at Centre Parc Grant covers carers costs. DSUK provide two medical teams onsite to support the families should the need arise whilst they are at the event with other DS families.
	15,000	Update existing educational materials. DSUK's leaflets were last updated in 2013.
	30,000.00	Additional grant in relation to the Updating of existing educational materials.
Dravet-Syndrom eV.	2,946.77	To support the production of personalised safety belts for Dravet patients "Emergency Dravet belts" and to support annual family conferences.
Epilepsiforeningen	9,060.24	For the development of support materials for patients and families living with Dravet Syndrome. These materials will contain both peer to peer and medical information.
Epilepsy Research UK	250.00	Charity donation for participation in London Marathon and Great north run.
FEDERACIÓN ESPAÑOLA DE ENFERMEDADES RARAS	9,195.00	FEDER fund helps families it represents through projects such as "Aid Fund" which offers direct assistance to families living with rare diseases.
Fondation Française pour la Recherche sur l'Epilep	6,306.90	Cette subvention nous permettra de financer les actions conformes aux missions statutaires de la Fondation Française pour la Recherche sur l'Epilepsie. Cette subvention est destinée à un usage collectif par la Fondation.
Fundacion 29 de Febrero	36,563.07	Improving the lives of patients living with rare diseases the goal is to <u>end</u> diagnostic odyssey for children with rare disease.
	36,513.11	The grant will be used to for the following including: Beta version, platform usage, communication, website, AI benchmarking, and guidance.
Gruppo Famigile Dravet Associazione Onlus	9,015.99	To support Occupational therapy tele-rehabilitation: the project will involve 15 families and will deliver a scientific publication about the efficacy of Occupation Therapy realized by tele-

		<p>rehabilitation. The goal is to pave the way to use this method also in further years, in order to facilitate the rehabilitation in people with Dravet syndrome.</p> <p>2. Support families in the COVID-19 Emergency: families members provided with PPE to be used with their Dravet kids to restart their activities.</p> <p>3. Support and improve social skills of children and people with Dravet syndrome organizing videocall activities for small groups of people (4/5 persons) supported by the mediation of a neuropsychologist.</p>
International Brain Tumour Alliance IBTA	1,975.00	To support projects including Brain Tumour Magazine, Production and dissemination of the monthly IBTA e-News, Complete the revised version of the Brain Tumour Patients Charter Support also goes towards Brain tumour support groups and other IBTA activities.
Norsk Epilepsiforbund	8,494.92	Separate Website area development for information about rare epilepsy and Norsk dedicated networks for rare epilepsy. This website will help provide information to improve the lives of individuals and families affected by these symptoms.
Tuberous Sclerosis Association	10,000.00	The TSA provides support to people affected by TSC and the professionals who support them through the pandemic.
Tuberous Sclerosis Australia	7,726.92	To support the TSA Connect Conference 2020, Sydney TSC Community Stream Saturday 2 and Sunday 3 May 2020
Young Epilepsy	15,000.00	To support 'The Channel' (co-created with Young Reps and is a place where young people with epilepsy (aged 13 – 25) can get information, advice and guidance when they want it, where they want it (online) and written in a language that they can relate to.) Support creation of content. Conduct independent survey Live streaming activities

**2019**

Patient Organisation	Finance	Description
AEER - Associazione Epilessia Emilia Romagna	£4,341.76	Supporting activities related to 'Transitional Care' of epileptic patients from childhood to adulthood
ASD Alliance Syndrome de Dravet	£12,129.71	The beneficiary's purpose to provide support to patients and caregivers. They are involved in representing patients in several official instances including ANSM & HAS commission de Transparence.
Association ESEFNP	£3,483.93	A 3 year cycle of independent annual training courses, developed specifically for child neurologists and physicians treating children with epilepsy. Plenary interactive sessions and training modules will cover different aspects of treatment strategies in children with all types of epilepsies, including topics relating to pharmacology, pharmacodynamics, pharmacokinetics, analysis of AED trials, design of clinical trials, pre-surgical evaluation, optimal choice of AEDs, review of existing guidelines (ILAE, NICE, other), ketogenic diet, neurostimulation, indications for resective and palliative surgery, pharmacological treatments other than AEDs development of educational therapy programs. Training modules, where one will be in Spanish. A pre- and post-assessment will allow evaluation of background and acquired knowledge.
Associazione Famiglie LGS Italia	£13,353.76	The objective of this project "La voce dei pazienti...AL TEMPO DEL COVID-19" is to arise patients with epilepsy and Lennox-Gastaut Syndrome (LGS) needs and expectations, that have been further accentuated by COVID-19 infection. Needs like prompt treatment access, home therapies, support to adults as well as infant patients and caregivers in case of hospitalization, health care professional (HCP) available to provide support for intellectual disabilities.
Dravet Italia Onlus	£4,324.23	Meeting with the 23 accredited Italian centres of the Registry (scheduled for 04 June 2019). This meeting features presentations by neurologists on relevant epilepsy related topics and new medical data presentation. Currently there are 418 patients (275 Dravet syndromes) With the aim of reaching 500 by the end of the year.
Dravet Italia Onlus	£8926.88	Grant to assist with Dravet diary /App Update
Dravet Syndrome European Federation		Project: International Registry for Dravet Syndrome and other syndromes related to mutation of SCN1A and PCDH19 genes (PLATFORM-RESIDRAS). International network consolidation and Subjects Inclusion increase. Project Deliverables: 1) Increase of the number of accredited Centers and their geographical distribution 2) Reach within the next two years the number of approximately 1.000 patients (including patients with DS, SCN1A mutation carriers with other phenotypes, and patients with PCDH19 mutations)

		<p>3) Realize an international network allowing to input data of patients provided by physicians that are not members of the centers</p> <p>4) Create a collection of subjects with highly characterized genetic, clinical, neurophysiological, neuropsychological, treatment, evolutive aspects, useful for subsequent analysis and research</p> <p>5) Implement the Registry contributing to a more uniform and homogenous optimal clinical management on the patients shared.</p>
FUNDACIÓN SÍNDROME DE WEST	£4,218.45	The Grant is to support the organisation of a Multidisciplinary forum meeting in Madrid on 22 <sup>nd</sup> of November 2019. The key participants in this forum are those affected by these diseases: patient organisations, HCPs, governmental institution and all of them will discuss about Epilepsy and Rare diseases. Estimated attendees approximately 200 in this session, in addition to the possibility of transmitting online via streaming for those who cannot attend.
FUNDACIÓN SÍNDROME DE WEST	£4,218.45	VIII Congress West Syndrome, November 20 <sup>th</sup> – 21 <sup>st</sup> 2019: “Update on epilepsy and autism” Place: CaixaForum. Paseo del Prado, 36. Madrid. Spain Duration: Two days, November 20 <sup>th</sup> -21 <sup>st</sup> 2019 Format: International conference of experts and scientists in the area of epilepsy, rare diseases and autism.
Gruppo Famigile Dravet Associazione Onlus	£6,930.46	Various activities related to the scope of the charity, that include, initiatives aimed at improving disease knowledge and management, and the support of people affected by Dravet Syndrome.
International Brain Tumour Alliance IBTA	£1,750	To support the International Brain Tumour Alliance’s work programme for 2019. Including: Encourage the establishment of brain tumour patient groups in countries where they don’t yet exist. Promote collaboration on programmes and projects to benefit the brain tumour community. Highlight the challenges and needs of brain tumour patients and caregivers. Disseminate knowledge, information and best practices. Help shape health and research policies at national and international levels. Honour the courage and achievements of brain tumour survivors and caregivers.
Landesverband fur Epilepsie Selbsthilfe NRW E V	£861.63	To provide financial support to one of the largest epilepsy patient events in Germany, Bonn.
LGS Foundation UK	£6,830.00	Enable two members of the LGS Foundation UK organisation to attend the LGS Foundation International Conference between 12-14 May 2019 by contributing towards their travel and expenses. <ul style="list-style-type: none"> <li>• Ensure latest information and research about LGS is available to UK members.</li> <li>• Ensure UK members can track the conference live via daily Twitter and Facebook updates</li> <li>• Ensure UK website is updated with conference presentations and feature articles drawing on key conference themes</li> </ul>

		<p>Add value to the visit and reach a wider membership by meeting with and recording multi-media interviews / case studies of families impacted by LGS to share with the UK membership.</p> <ul style="list-style-type: none"> <li>☑ Provide accessible insights and experiences into how LGS effects families.</li> <li>☑ Demonstrate how the latest research has a real-life impact on families.</li> </ul>
Ring14 International Onlus	£9,179.04	The grant will help Ring14 International extend the international community of scientists engaged in Ring14 research when they host the 3rd International Scientific Workshop in October 2019.
Ring20 Research and Support UK CIO	£2000	Produce an information booklet for families living with r(20)so that they can understand the impact of their diagnosis and receive practical , useful information for the future of their care and support needs. The aim is to provide these booklets (1 per family) free of charge. Ideally, we would like to produce 2 booklets: 1x aimed at younger children and 1x aimed at young adults/parents/carers.
Young Epilepsy	£5,000	<p>UK MSL Training Preceptorship Program supplied by Young Epilepsy.</p> <p>To help UK team better understand diagnosis, needs, treatment goals and management / care of children and young people with epilepsy.</p> <p>This includes a tour of The Neville Childhood Epilepsy Centre in Lingfield to understand the diagnostic and treatment facilities at Young Epilepsy.</p>

**Payments Include VAT where applicable**