

## 2015 Alexion Financial Support to Patient Advocacy Organization

Alexion is committed to transparency and developing and maintaining trust-based relationships with stakeholders, the public and patients. Accordingly, Alexion is publicly disclosing, on an annual basis, a report listing the names of all patient advocacy organizations supported, the nature of funding received by each and the financial amount per activity. This chart represents a good faith attempt and best efforts by Alexion to identify and report this information from all available sources and to provide a full and transparent reporting of all financial support provided by Alexion to patient advocacy organizations around the world in calendar year 2015. This chart may be amended if and when new information is identified for inclusion. The funding listed in this chart is reported in the respective local currencies of the recipients. Alexion's Global Policy for Interacting with Patient Advocacy Organizations defines PAOs as not-for-profit organizations (including umbrella organizations to which they belong), mainly composed of patients and/or caregivers that represent and/or support the needs of patients and/or caregivers.

Organization	Description	Country	Funding Amount (in local currency)
APEC (Asociación de pacientes con Enfermedades del Complemento)	To support patients with complement disorders, diagnosis, transportation and Association maintenance.	Argentina	500,000 ARS
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APELRA (Asociacion de Pacientes de Enfermedades Lisosomales de la Republica Argentina)	To support patients with complement disorders, diagnose, transportation and Association maintenance.	Argentina	600,000 ARS
Fundación ENHUE	To support and improve education, access to diagnosis, and treatment of patients suffering from rare diseases.	Argentina	10,000 USD
Fundacion ENHUE*	Support for ENHUE's rare disease patient advocacy activities.	Argentina	10,000 USD
Rare Voices Australia	Membership of Industry Roundtable Group.	Australia	13,636.36 AUD
aHUS Patient Support Group Australia	Unconditional grant to aHUS Patient Support Group Australia.	Australia	40,000 AUD
Associacao Dos Familiares, Amigos E Portadores De Doencas Graves (AFAG)	To support the organization's program and education activities in 2015.	Brazil	1,672,000 BRL
Associadao Paulista dos Familiares e Amigos dos Portadores de Mucopolissacaridose (APMPS)	To support the organization's program and education activities in 2015.	Brazil	320,000 BRL
Associacao Braslieira dos Portadores de Doenca de Hunter e outras Doencas Raras (Casa Hunter)	To support the organization's program and education activities in 2015.	Brazil	50,000 BRL

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Associação Brasileira dos Portadores de Doença de Hunter e outras Doenças Raras (Casa Hunter)	To support organization's efforts for case management and support services for patients.	Brazil	100,000 BRL
Associação Brasileira dos Portadores de Doença de Hunter e outras Doenças Raras (Casa Hunter)	Project awareness raising program for the diseases and continuing education.	Brazil	200,000 BRL
Associação Sergipana de Pessoas com Doenças Raras*	Support for Annual Symposia.	Brazil	1,200 USD
Associação Paulista dos Familiares e Amigos dos Portadores de Mucopolissacaridoses e DOENÇAS RARAS*	Support for APMSP's rare disease day for 2015.	Brazil	2,800 USD
Canadian Association of PNH Patients	To support the organization's disease awareness activities.	Canada	148,500 CAD
The Kidney Foundation of Canada	General education of renal diseases in SW Ontario.	Canada	2,500 CAD
The Kidney Foundation of Canada	Sponsorship of gala fundraiser.	Canada	3,000 CAD
The Kidney Foundation of Canada	Silver level sponsorship of kidney transplant and organ donation summit.	Canada	3,500 CAD

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The Kidney Foundation of Canada	Sponsorship for SW Ontario celebrity event.	Canada	10,000 CAD
The Kidney Foundation of Canada	Sponsorship of Shine a Light Kidney Gala.	Canada	3,500 CAD
aHUS Canada	To support the organization's disease awareness activities.	Canada	100,000 CAD
aHUS Canada	To support the organization's disease awareness activities.	Canada	50,000 CAD
CORD (Canadian Organization for Rare Disorders)	Support for educational and awareness activities regarding the implementation of the Rare Disease Strategy.	Canada	50,000 CAD
CORD (Canadian Organization for Rare Disorders)	Support for 2015 Rare Disease Day, Conference Sponsorship and Gala.	Canada	30,000 CAD
Canadian Association of PNH Patients	To support the organization's disease awareness activities.	Canada	148,500 CAD

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Aplastic Anemia Myelodysplasia Association of Canada	Patient education and support projects.	Canada	40,000 CAD
Network of Rare Blood Disorder Organizations	Sponsor rare blood disorder education day.	Canada	10,000 CAD
Canadian Liver Foundation*	Support for program activities.	Canada	998.28 USD
Fundación de Apoyo Solidario a Pacientes con Enfermedades Raras (FUNDAPER)	To support the organization's program and education activities in 2015.	Colombia	1,401,251,777 COP
Colombian Federation for Rare Diseases (FECOER)	To support the organization's rare disease advocacy efforts.	Colombia	60,000,000 COP
Asociacion Colombiana de Pacientes Con Enfermedades de Deposito Lisomal (ACOPEL)	2016 Support of the organization's patient programs.	Colombia	27,592,600 COP
Asociacion Colombiana de Pacientes Con Enfermedades de Deposito Lisomal (ACOPEL)	Support the organization's patient programs.	Colombia	3,000,000 COP
Asociacion Colombiana de Pacientes Con Enfermedades de Deposito Lisomal (ACOPEL)*	To support the organization's program.	Colombia	4,000 USD

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Croatian Alliance for Rare Diseases	Patient networking and awareness raising in 2015.	Croatia	10,000 EUR
Association pour l'Information et la recherche sur les Maladies Rénales Génétiques (AIRG)	Support the association's communications activities.	France	15,000 EUR
Hypothosphatasise Europe	Support for the organizations activities in particular the translation and dissemination of material on HPP from French into English, set up of a HPP Day in France, translation of website from French into other languages (e.g. English, Spanish).	France	22,000 EUR
Association HPN France	Patient communication on HPN & medular aplasia.	France	15,000 EUR
Association HPN France	Patient communication on HPN & medular aplasia.	France	20,000 EUR
EURORDIS	To provide unrestricted funding for EURORDIS - supporting EURORDIS' work to improve conditions for people throughout Europe who are living with a rare disease; To support the preparation and delivery of a multi-stakeholder conference on rare diseases.	France	20,000 EUR
EURORDIS*	Sponsorship of Gala Dinner.	France	10,000 EUR

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Association de Recherche sur les Microangiopathies Thrombotiques et autres maladies rares hématologiques	To support organization's research activities.	France	726 EUR
Renaloo	To support a patient meeting.	France	1,136 EUR
Sifting Lichterzellen	T-shirts donation to the patient group for the rare disease day events.	Germany	953 EUR
Hypophosphatasie Deutschland e.V.	T-shirts donation to the patient group for the rare disease day events.	Germany	953 EUR
ProRare Austria MPGN & aHUS Selbsthilfe	T-shirts donation to the patient group for the rare disease day events.	Germany	953 EUR
HPP Deutschland	Support Patient Association for an Annual Patient Meeting.	Germany	10,000 EUR

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Osteogenesis Imperfecta Federation Europe (OIFE)	Support for the organization's annual meeting and seminar bringing together patients and clinicians.	Germany	10,000 EUR
Associazione A.M.E.N.A. Onlus	To support the association.	Italy	5,000 EUR
Associazione Culturale Onlus "Giuseppe Dossetti": Valori-Sviluppo e Tutela del Diritti	To support the initiatives of the National Leading Civil Rights and Rare Diseases Advocacy Association.	Italy	8,000 EUR
Associazione Onlus DOMOS Sez Campania	To support the organization's 2015 activities.	Italy	7,999 EUR
Associazione Siciliana Bambini Nefropatici	To support the organization.	Italy	3,000 EUR

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A.S.A.E. A.I.L. Cagliari	To support the organization.	Italy	4,000 EUR
Associazione Sud Italia Trapiantati	Purchase laboratory equipment for supporting the research.	Italy	4,000 EUR
Associazione Donatori di Midollo Osseo e di Cellule Staminali Emopoietiche	Support the scientific research on diseases that currently benefit from Bone Marrow Transplantation.	Italy	7,000 EUR

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<p>Associazione Italiana contro le Leucemie-Linfomi-Mieloma Salerno sez. "Marco Tulimieri" Onlus</p>	<p>Support the Hematology Department of the hospital located in Pagani by supporting the volunteers, purchasing equipment, grant the staff for qualified training and providing support to patients and their families.</p>	<p>Italy</p>	<p>5,000 EUR</p>
<p>AIL - Associazione Italiana contro le Leucemie-Linfomi e Mieloma Onlus - Sez. Avellino</p>	<p>Support the patients affected by hematological malignancies and improve the quality of their lives.</p>	<p>Italy</p>	<p>5,000 EUR</p>
<p>Associazione Italiana contro le Leucemie, Linfoma e Mieloma AIL Salento - Sez. AIL della provincia di Lecce Onlus</p>	<p>Support the volunteers, purchase equipment, grant the staff for qualified training and provide support to patients and their families.</p>	<p>Italy</p>	<p>15,000 EUR</p>

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<p>AIL COSENZA Fondazione "Amelia Scorza" ONLUS</p>	<p>Support the scientific research in the field of hematological malignancies.</p>	<p>Italy</p>	<p>2,000 EUR</p>
<p>Associazione Italiana contro le Leucemie, Linfoma e Mieloma Onlus - Sez. di Parma</p>	<p>Support scientific activity and epidemiological research.</p>	<p>Italy</p>	<p>4,000 EUR</p>
<p>Associazione Sud Italia Trapiantati</p>	<p>Support scientific research at the kidney and transplantation center.</p>	<p>Italy</p>	<p>20,000 EUR</p>

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<p>Associazione Italiana Contro le Leucemie-Linfomi e Mieloma - Milano e Provincia Onlus</p>	<p>Support the scientific research and organize the assistance for people affected by leukemia and other blood cancers.</p>	<p>Italy</p>	<p>15,000 EUR</p>
<p>Associazione Italiana contro le Leucemie-Linfomi e Mieloma Sez. di Bologna Onlus</p>	<p>Support scientific and epidemiological research.</p>	<p>Italy</p>	<p>5,000 EUR</p>
<p>Progetto ALICE Onlus - Associazione per la lotta alla sindrome emolitico uremica</p>	<p>Support all statutory activities.</p>	<p>Italy</p>	<p>15,000 EUR</p>

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Associazione Umbra Studio e Terapia Leucemie e Linfomi Onlus	Support the scientific research in patients with hematological neoplasia.	Italy	8,000 EUR
Saisel Tsubasa no Kai	Support the organization's disease education activities in 2015.	Japan	500,000 JPN
NPO PNH Club	Grant for the 2015 program activities.	Japan	5,000,000 YEN
NPO PNH Club (PNH Patient Advocacy Organization)	Support for program activities in Q3 and Q4.	Japan	10,000,000 JPY

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HIPPSA-JP	HIPPSA-JP - Hypophosphatasia Support Association of Japan - Support organization for 2016.	Japan	1,000,000 JPY
HIPPSA-JP	Travel and speaker fee for HPP meeting.	Japan	62,200 JPY
Organization Mexicana de Enfermedades Raras (OMER)	To support event on June 29 - July 3 2015.	Mexico	70,000 USD
Proyecto PideunDeseo*	Support for the organization's activities.	Mexico	476 USD
Dutch Kidney Association	Support for the organizations activities in particular supporting a working group for aHUS patients, activities to mark Kidney Donor week.	Netherlands	15,000 EUR
Association of Patients with MPS and Rare Diseases	Support for Association of Patients for general activity and organization of Annual Conference: "Rare Disease Crossing Borders Together."	Poland	40,000 PLN
Interregional patient organization "Another Life"	Support for patient meeting.	Russia	310,000 RUB

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Interregional patient organization "Another Life"	Support for Round table meeting.	Russia	320,000 RUB
Interregional patient organization "Another Life"	Third all-Russia colloquium "Live differently".	Russia	800,000 RUB
Interregional patient organization "Another Life"	Support for aHUS Day.	Russia	200,000 RUB
Interregional patient organization "Another Life"	Support for patient meeting.	Russia	300,000 RUB
Center of patients care "Genom"	Round table on Rare Diseases-"Rare diseases yesterday, today, tomorrow" (Saint-Petersburg)	Russia	30,000 RUB
Center of patients care "Genom"	Round table on Rare Disease ( "Rare but equal") (Arkhangelsk)	Russia	50,000 RUB
Center of patients care "Genom"	Round table on Rare Diseases (Crimea)	Russia	150,000 RUB
Center of patients care "Genom"	Round table on Rare Diseases (Blagoveschensk)	Russia	250,000 RUB

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Center of patients care "Genom"	Round table ( in Nizhny Novgorod)	Russia	270,000 RUB
Center of patients care "Genom"	Support for school for patients with metabolic disorders.	Russia	465,000 RUB
Union of patients and patient organizations on rare diseases	International rare diseases day and patient meeting.	Russia	244,681 RUB
All Russia society of orphan diseases	International rare diseases day and patient meeting.	Russia	200,000 RUB
National Association "Genetics"	Round table in Saint-Petersburg to mark international Rare diseases day.	Russia	100,000 RUB
National Association "Genetics"	Support for research project.	Russia	300,000 RUB
National Association "Genetics"	Participation at Mass Media in Moscow.	Russia	200,000 RUB
National Association "Genetics"	Placement of information materials in 'Bulletin/newsletter'.	Russia	222,000 RUB

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National Association "Genetics"	Provision of information on rare diseases.	Russia	260,000 RUB
National Association "Genetics"	Participation at congress "Right for a drug".	Russia	500,000 RUB
Nephro-league	Action "Healthy kidney to everybody" devoted to the Kidney day.	Russia	150,000 RUB
League of patient defenders	Participation at congress "Right for a drug" Moscow, May.	Russia	400,000 RUB
All Russia union of patients	Participation at congress of patients November 12-14, 2015.	Russia	200,000 RUB
Interregional patient organization "Road to life"	2 <sup>nd</sup> Conference of Privlozhsky federal district "Road to Life" October 2-3, 2015.	Russia	60,000 RUB
Asociacion Sindrome Hemolitico Uremico Atipico (ASHUA)	Grant supporting awareness activities including national and regional patient meetings and aHUS wristbands.	Spain	20,000 EUR
AE LALD Asociacion Espanola Deficit De Lipasa Acida Lisosomal	Travel expenses for an event.	Spain	171 EUR

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Asociación Hemoglobinuria Paroxística Nocturna de España (HPNE)	Grant supporting activities including the journal, national and regional patient meetings.	Spain	35,000 EUR
D'GENES ASOCIACIÓN	Sponsorship of VIII National Congress of Rare Diseases.	Spain	3,000 EUR
Federación Española de Enfermedades Raras (FEDER)	Annual sponsorship for disease education activities including a Rare Disease Awareness Campaign.	Spain	25,650 EUR
Fundacion Hipercolesterolemia Familiar*	Support for the Foundation's conference.	Spain	6,050 EUR
Asociacion MPS Espana*	Support for program activities.	Spain	5,514.50 USD
International Osteoporosis Foundation	Support Skelton Rare Disease Capitation of Endorsing Partners Meeting.	Switzerland	35,000 CHF
International Osteoporosis Foundation	Support workshop rare congenital metabolic bone disorders.	Switzerland	38,000 CHF
PNH aHUS Hasta Dernegi Patient Association	Support for vaccination project.	Turkey	190,000 TRY

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PNH-AHUS ve Diger Kompleman Hastaliklari Hasta Dernegi -	To support 2015 Activities of aHUS PA, Turkey.	Turkey	50,000 TRY
aHUS UK/aHUS Alliance	To support Global aHUS Meeting.	United Kingdom	15,000 GBP
PNH Alliance	To support their planned 2015 activities.	United Kingdom	50,000 GBP
aHUS-United Kingdom	Support aHUS patient family conference.	United Kingdom	25,000 GBP
PNH Support England and Wales	To support the organization.	United Kingdom	15,000 GBP
CLIMB (National Information Centre for Inherited Metabolic Diseases)	HPP Patient Support.	United Kingdom	10,000 GBP

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NMO - Research UK	Support for annual meeting.	United Kingdom	5,000 GBP
UK Society for Mucopolysaccharide Diseases	Grant to support the UK MPS Annual Conference June 26-28, 2015 in Coventry, UK.	United Kingdom	5,000 GBP
Transplant 2013	Activities to increase the consent for Organ donation by increasing public awareness.	United Kingdom	65,000 GBP
National Organization for Rare Disorders, Inc. (NORD)	2015 Corporate Council Dues.	United States	25,000 USD
National Organization for Rare Disorders, Inc. (NORD)	Support 2015 PNH Patient Education Program.	United States	502,630 USD

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National Organization for Rare Disorders, Inc. (NORD)	2015 NORD aHUS Patient Education Program.	United States	135,852 USD
National Organization for Rare Disorders, Inc. (NORD)	Sponsorship for 2015 Portraits of Courage Gala.	United States	15,000 USD
National Organization for Rare Disorders, Inc. (NORD)*	Support for program activities.	United States	23,250 USD
The Atypical HUS Foundation	The Atypical HUS Foundation - 2015 Program Support.	United States	125,000 USD
The MAGIC Foundation	Patient scholarship for 50 HPP patients/caregivers during the 2015 Patient Convention.	United States	50,000 USD
The Transverse Myelitis Association	To support the monthly Podcast Series and the Rare Neuro Immune Disorders Symposium on October 23-24 2015 in Dallas TX.	United States	25,000 USD

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Soft Bones, Inc.	2014 HPP Awareness Activities (paid in 2015 - although 2014 project).	United States	20,000 USD
Soft Bones, Inc.	Request for funds to support capacity building for establishment of Soft Bones Canada.	United States	100,000 USD
Soft Bones, Inc.	To support regional patient meetings in conjunction with ASBMR 2015.	United States	5,000 USD
Soft Bones, Inc.	Support organization's Patient Meetings to provide HPP disease education and peer support.	United States	80,000 USD
Global Genes	2015 Global Genes aHUS Patient Education Program.	United States	705,000 USD
Global Genes	Corporate Alliance Membership Dues.	United States	25,000 USD
Global Genes	Sponsorship of 2015 Patient Summit and Tribute of Champions - Gold Sponsorship.	United States	100,000 USD

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Global Genes	Support Teen Summit on October 9-11, 2015 in Orlando, FL.	United States	245,000 USD
Global Genes*	Sponsorship for Tribute of Champions and Corporate Alliance	United States	15,000 USD
National MPS Society	Grant to support the National MPS Society 29th annual Family Conference September 17-19, 2015 in Salt Lake City.	United States	3,000 USD
American Liver Foundation	To educate patients and providers about Lysosomal Acid Lipase Deficiency (LAL D).	United States	36,000 USD
American Liver Foundation*	Support for program activities.	United States	5,000 USD
Aplastic Anemia & MDS International Foundation	Support for 2015 PNH Program.	United States	295,000 USD
Aplastic Anemia & MDS International Foundation	Support for PNH Communities of Hope.	United States	90,000 USD

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Aplastic Anemia & MDS International Foundation	Sponsorship for PNH Walk.	United States	18,000 USD
Aplastic Anemia & MDS International Foundation	Sponsorship for Patient Education Meeting.	United States	35,000 USD
Aplastic Anemia & MDS International Foundation	PNH Webinar Series.	United States	90,000 USD
Aplastic Anemia & MDS International Foundation	PNH Patient Survey (paid in 2015 – although 2014 project).	United States	40,000 USD
Every Life Foundation	Sponsorship of 2015 Education and Awareness Activities.	United States	30,000 USD

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Rare Disease United Foundation	Support for 2015 Rare Disease Day Activities.	United States	6,000 USD
National Kidney Foundation	Disease awareness and education activities.	United States	25,000 USD
Utah Rare	Support for 2015 Rare Disease Day Activities.	United States	1,000 USD
National Tay Sachs and Allied Diseases Association	Support for 2015 Gala Event.	United States	2,500 USD
National Tay Sachs and Allied Diseases Association*	Support for program activities.	United States	1,000 USD

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Guthy-Jackson Charitable Foundation	2015 Industry Council Dues.	United States	25,000 USD
Myasthenia Gravis Foundation of America	Support for 2015 Disease Awareness and Education Initiatives.	United States	25,000 USD
American Kidney Fund	Support for 2015 Gala Event.	United States	10,000 USD
LAL Solace*	Support for program activities.	United States	35,932 USD
LAL Solace*	Support for program activities.	United States	2,000 USD

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National MPS Society*	Support for program activities.	United States	2,322.75 USD
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\*Financial support to Patient Advocacy Organizations by Synageva BioPharma Corp. prior to Alexion acquisition.