

**Pfizer France : Partenariats et prestations avec les associations de patients
pour l'année civile 2020**

ASSOCIATIONS	ACRONYMES	PARTENARIAT ET PRESTATION € Pfizer SAS	DESCRIPTIF
Action Leucémies		1 050 €	Board enquête AVNIR
Associatio Laurette Fugain	ALF	10 000 €	Partenariat RDV Laurette Fugain 2020
Association des sclérodermiques de France	ASF	1 050 €	Board enquête AVNIR
Association française contre l'amylose	AFCA	17 500 €	Partenariat mini-série amylose cardiaque
ASSOCIATION FRANCAISE DE LUTTE ANTI-RHUMATISMALE	AFLAR	15 000 €	Partenariat pour realiser une enquete Franco - Belge sur la douleur arthrosique
Association Francaise Des Polyarthritiques et des Rhumatismes Inflammatoires Chroniques	AFPric	10 832 €	Salon AFPric
Association France Spondyloarthrites	AFS	2 400 €	Consultant - Groupe de travail sur l'antibioresistance
Association Nationale de défense contre la polyarthrite rhumatoïde	ANDAR	1 050 €	Board enquête AVNIR
Association pour la Recherche sur les Tumeurs du Rein	ARTUR	3 000 €	Partenariat enquête cancer du rein
Association Santé Respiratoire France		1 050 €	AVNIR
Etincelle IDF		5 000 €	Partenariat cancer du sein métastatique
Europa Donna forum France		5 000 €	Partenariat cancer du sein métastatique
France AVC		2 500 €	Consultant - enquête parcours de soin fibrillation atriale
France Lymphome Espoir	FLE	1 050 €	Board enquête AVNIR
HTaP France		3 450 €	Consultant - Groupe de travail sur l'antibioresistance Board enquête AVNIR
Juris Santé		5 000 €	Partenariat cancer du sein métastatique
Life is Rose		5 000 €	Partenariat cancer du sein métastatique
Patients en reseau		16 500 €	Partenariat cancer du sein métastatique et enquête collectif 1310 charge mentale + Interventions
Renaloo		1 050 €	Board enquête AVNIR
Rose-Up		75 000 €	Partenariat Maisons Rose Paris - Bordeaux

**Pfizer International Operation - Pfizer Inc
Soutien des associations de patients basées en France en 2020**

Organisation full name	2020 funding	Description	Organisation description
Amyloidosis Alliance	60 000 EUR	Amyloidosis Alliance- Toolkit	The Amyloidosis Alliance is the global alliance of patient groups working in Amyloidosis. The alliance has a mission to promote diagnosis, awareness, research and coordination between organizations in order to improve outcomes for patients and their families suffering from Amyloidosis.
Amyloidosis Alliance	90 000 EUR	Amyloidosis Alliance-International day of amyloidosis	The Amyloidosis Alliance is the global alliance of patient groups working in Amyloidosis. The alliance has a mission to promote diagnosis, awareness, research and coordination between organizations in order to improve outcomes for patients and their families suffering from Amyloidosis.
Europa Donna France	1381 EURO	Participating and providing input into the Pfizer-hosted Global Breast Cancer Advocacy Impact Summit	Europa Donna France is the essential representative of women affected by breast cancer. The association's mission is to raise awareness, inform, support and facilitate access to the best care.
EURORDIS	17 000 EURO	EURORDIS 9th European Conference and Black Pearl Awards	EURORDIS is a non-governmental patient-driven alliance of patient organisations representing 667 rare disease patient organisations in 61 countries.
EURORDIS	65 000 EURO	EURORDIS International Initiatives 2020	EURORDIS is a non-governmental patient-driven alliance of patient organisations representing 667 rare disease patient organisations in 61 countries.
Patients En Reseau	13 933 EURO	Participating and providing input into the Pfizer-hosted Global Breast Cancer Advocacy Impact Summit	The association Patients en réseau was created in February 2014 based on lived experiences. It brings together patients and relatives and relies on multidisciplinary scientific committees. Its mission is to develop social networks for people affected by severe diseases and their loved ones to facilitate their daily lives in the face of the ordeal of the disease.
Rare Diseases International	10 000 EURO	RDI's capacity – building efforts in Asia Pacific, to empower patient leaders to advocate for national rare disease frameworks	Rare Diseases International. Rare Diseases International (RDI) is the global alliance of people living with a rare disease of all nationalities across all rare diseases
Rare Diseases International	20 000 EURO	RDI Membership Renewal 2020	Rare Diseases International. Rare Diseases International (RDI) is the global alliance of people living with a rare disease of all nationalities across all rare diseases
Vitiligo International Patient Organizations Committee	20 000 EURO	VIPOC-2nd VIPOC conference	The purpose of the association: to have the reality of vitiligo and the diseases associated with it recognized, to bring together the organizations of vitiligo patients to make the voice of the sick and the voice of their organization