



APPLICATION FORM

Expert Patients and Researchers EURORDIS Summer School

ExPRESS

Barcelona, Spain June 1-5, 2015

Instructions for applicants:

- 1. The deadline for applications is December 8, 2014
- 2. This application form is divided into 3 Sections: A, B and C.
- 3. If you are applying as a patient representative, please complete sections A and B.
- 4. If you are a applying as a researcher, please complete sections A and C.
- 5. Please write very clearly or type in your responses ensuring that your answers are complete and legible.
- 6. Please sign and date your application on page 4.
- 7. Only complete, signed and dated applications will be reviewed by the evaluation committee.
- 8. If you are applying as a patient, please send your completed form to nancy.hamilton@eurordis.org or fax to +33 1 56 53 52 15
- 9. If you are applying as a researcher, please send your completed form to a.m.aartsma-rus@lumc.nl

Relation				Postal code			
City					Country		
c. English language skills							
Tick the appropriate boxes							
Please rate your English	Rea	ding ve/Advanced	Writing Native/Advanced		Speaking Native/Advar	nced	Overall Native/Advanced Good
language skills		Good Intermediate Basic None	Good Intermediate Basic None		Intermediate Basic None	2	Intermediate Basic None
d Your expe		ce/knowledge (5					
			nent in clinical trials a	and n	nedicines developmo	ent:	
e. Medicines	De	velopment (10 li	ines max.)				
	s, pati	ient organisations ar			· · · · · · · · · · · · · · · · · · ·		n between other stakeholders dicines development in which
f. Expectatio	ns (1	.0 lines max.)					
Please describe activities.	<i>your</i> (expectations related	to this summer scho	ool ar	nd how you think th	is trair	ning will help you in your

g. Please indicate your experience in:					
Design and objectives of clinical trials and the roles of	Patients' roles & responsibilities in innovative medicines development	Interaction with stakeholders (regulators, industry, etc.) in drug			
all stakeholder		development processes			
Advanced (5+ years of experience) Good (3-5 years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None	Advanced (5+ years of experience) Good (3-5 years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None	Advanced (5+ years of experience) Good (3-5 years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None			
Medicines development	Drug safety and risk/benefit	Pharmaco-economics, health			
process from pre-clinical research to approval	assessment of medicines	economics and health technology assessment			
Advanced (5+ years of experience) Good (3-5 of years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None	Advanced (5+ years of experience) Good (3-5 of years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None	Advanced (5+ years of experience) Good (3-5 years of experience) Intermediate (1-3 years of experience) Basic (under 1 year of experience) None			
h. Commitment					
Please evaluate your level of commitment to potentially be involved in clinical trial and medicines development policies and procedures, to share knowledge and exchange experience and to represent rare disease patients at the national and European level. If I am selected to attend the Expert Patient and Researcher EURORDIS Summer School 2015: I am interested in attending future capacity building sessions and seminars in orphan drug development					
organised by EURORDIS or third parties in conjunction with EURORDIS. As one of the selected participants, I agree to: Attend the full 4.5 day programme					

□ Share my experience □ Be included in EURORDIS list of potential volunteers so as to act as a rare disease patient representative for activities related to clinical trials and drug development. (for patients only) □ Be appointed as "patient expert" in particular for meetings at the EMA such as Protocol Assistance for my disease or review of European Public Assessment Reports (for patients only) □ To review EURORDIS' position papers, contributions and statements for advocacy purposes in the clinical trial and drug development area (for patients only) □ Participate in other conferences and workshops Section B for Expert Patients only (Researchers, please go to section C) 3. Your patient organisation (V.B. if your application is accepted, your organisation must be a member of EURORDIS; if it is not a member at the time of application, you will need to make a request for membership well in advance of the start of the Summer School.) Name: Disease(s) represented: (if applicable) Type of organisation (please tick only one). □ Governmental Non-governmental/not-for-profit Educational/research institution Transnational/Intergovernmental Informal Other (please specify): E-mail Website Elephone Fax Address Postal code City Country Country D. Your role in a patient organisation: Are you? Patient Staff Please tick × all that apply Family of patient Volunteer How long have you been active in the organisation? What are your roles or activities? Do you represent your organisation in any European Committee/Task Force/ Working Group? If yes, please specify	☐ Share my knowledge					
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Are you? Patient Staff Please tick \(\sim \) all that apply How long have you been active in the organisation? What are your roles or activities? Do you represent your organisation in any European Committee/Task Force/ Working Group? If yes, please specify	City				Country	
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European Committee/Task Force/ Working Group? If yes, please specify	What are your roles or activities?					
Group? If yes, please specify	Do you represent your organisation in any					
	European Committee/Task Force/ Working					
What is your professional background?	Group? If yes, please specify					
What is your professional background?						
	What is your professional background?					

Section C (For researchers only) If you are a patient representative, please sign and date your application on page 4.						
c. Your research	organisation:					
Name:						
Your area of rese	Your area of research:					
Type of organisat	ion (please tick only one	e):				
Governmental Non-governmental/not-for-profit Educational/research institution Transnational/Intergovernmental Informal Other (please specify):						
E-mail			Website			
Telephone			Fax			
Address			Postal code			
City			Country			
d. Your role in y	our research organisation	on:				
What is your current position?						
Do you have a M	D and/or a PhD?					
Are you an early	stage researcher (you					
obtained your PhD le	ss than 5 years ago)?					
What is your professional background? (10 lines max.)						
How long have you been involved in your current research field? What are your current roles and responsibilities? (10 lines max)						

Do you represent your organisation in any European Committee/Task Force/Working Group? (If yes, please specify e.g. IRDiRC working groups, European Networks or Consortia)	
Are you a member of COST Action BM1207? "	

Please sign and date your application form:

Signature Date