

BESPOKE PAEDIATRIC AND PARENT INVOLVEMENT PLANS TO ENHANCE RECRUITMENT AND RETENTION IN CLINICAL TRIALS



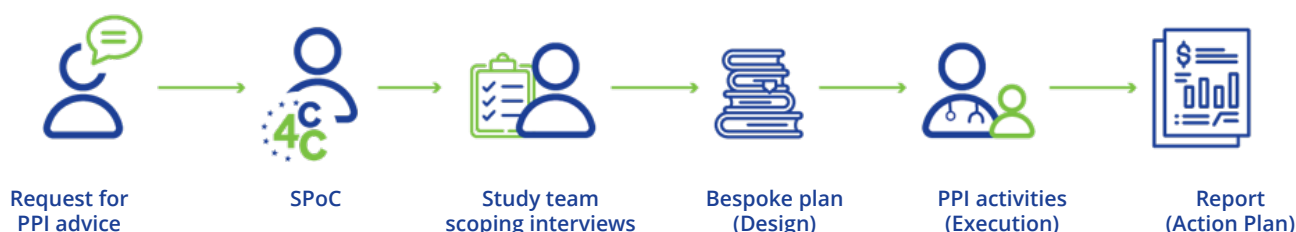
Being able to gather the views of paediatric patients and parents in the design and delivery of clinical trials is essential to ensure they are acceptable to both patients and their families. European Young Persons Advisory Group Network (eYPAGnet) has extensive experience working with children, young people and caregivers to provide advice to sponsors of European and international paediatric clinical trials.

BENEFIT FROM:

- **Expertise in all paediatric specialties** including rare diseases.
- **Strong collaboration** with patient organisations.
- **Partnership** with the c4c National Hubs.
- **Tailored services:** bespoke plan for clinical trials, non-drug interventions and medical device studies.
- Recognised by **Enpr-EMA**.

 PPI SERVICE	EXPERTISE
• Study specific support including co-design of clinical trial protocols, PROMS, PREMS and QoLs • Education and training of key stakeholders • Patient and Parents Involvement (PPI) advice and coordination of involvement activities • Reporting and measuring the impact of involvement	SUBJECT MATTER
	• Multiple languages • Adapted methodologies to fully involve children and parents • Diversity and inclusiveness • Preparing and facilitating focus groups • Qualitative interviews • Surveys • Other methods • Acceptable trial procedures • Endpoints • Recruitment and retention • Patient Journey • Informed Consent • Trial concept design • Other specific study needs

The **Single Point of Contact (SPoC)** acts as the gateway to the **c4c-S network**, providing an efficient and user-friendly way of engaging c4c-S experts with a single request.



CORE GROUP	CORE VALUES
• 4 patient involvement expert leaders in 3 European countries with more than 50 years cumulative experience of including the voices of paediatric patients in the design of clinical trials. • Supported by more than 15 facilitators across Europe.	• Involve patients and caregivers in a meaningful approach. • Respect the needs of patients and caregivers. • Ethical involvement. • Respect children's rights to have an active participation in health research.



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Certain information in this communication contains forward-looking statements, including statements related to the expected scope and timing of launch of services, as well as the number of countries and/or sites at time of launch of related services based upon what we believe are reasonable assumptions.