

## Who are IPPOSI?

We are a patient-led organisation that works with patients, science & industry to put patients at the heart of health innovation





## What do we do, and how?

**EDUCATION** 

**ADVOCACY** 

**INFORMATION** 

## Our Patient-led, Public Private Partnership

105

patient organisations\*

122

individual patients

\*represents the largest network of patient organisations in the country

210

individual scientists/researchers

27

healthcare companies

- In-person/virtual conferences
- Training programmes
- Digital discussions & podcasts
- Matchmaking
- Member resources
- Coalitions and networks
- Citizens' juries





## We have three priority areas



ACCESS TO INNOVATION



PATIENT & PUBLIC INVOLVEMENT



DIGITAL HEALTH & INFORMATION





## 'Patient Centricity' in Irish Healthcare

Individual patient experience of healthcare

Patient co-design of care services

Patient co-production of service delivery

Patient representation +
high-level involvement
in strategy, policy,
oversight

Meaningful PPI Partner Opportunities + Training





## 'Learning Framework of Involvement' for Health

#### **POLICY-LEVEL**

- National Patient Involvement Unit in Dept. of Health
  - Chief Patient Officer akin to Chief Medical/Nursing/Clinical Officers

#### **SERVICES**

- Co-designed Roadmap for Healthcare Services
  - Establish a vision for what 'success' looks like and a 'mission' to achieve it
  - Mechanisms, supports, education, skills-training for patients, professionals
  - Resourced to coordinate at national, regional, local levels

#### **RESEARCH + INNOVATION**

Embed PPI in ideation and early-stage development of new innovations

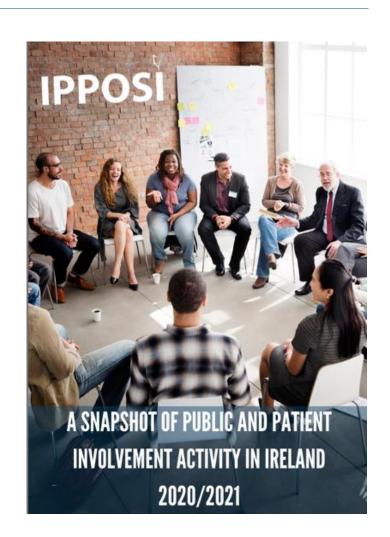




### **Building a new Irish healthcare + research environment**

#### In 2022, Patient Advocates are members of the

- Board of the Health Service Executive (HSE)
- HSE Rare Disease Tech Review Committee
- National Clinical Effectiveness Committee
- Steering Group of Health Innovation Hub
- Health Products Regulatory Authority Patient Forum
- PPI reviewers in Health Research Funder (HRB)
- University-based National PPI Network
- National Research Ethics committees
- Many, many more...

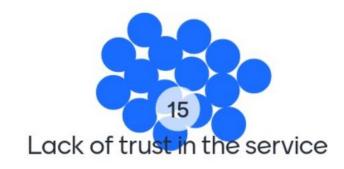


#### Mentimeter

## Having heard these presentations, what do you believe are the main challenges/barriers to PPI in general?









Difficult to find the 'right patient'





Lack of evidence of its impact



## The many Ps of patient & public involvement

**PUBLIC** and **PATIENT** involvement requires **PEOPLE** with a PASSION and a PURPOSE with a PLACE to meet and a PAYMENT for their involvement. It requires a willingness to pursue POLICY CHANGE with PERSEVERANCE and PERSISTENCE while mindful of the need for PRACTICAL APPLICATION and in PARTNERSHIP with the health system.



# IPPPOS

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