ENCePP

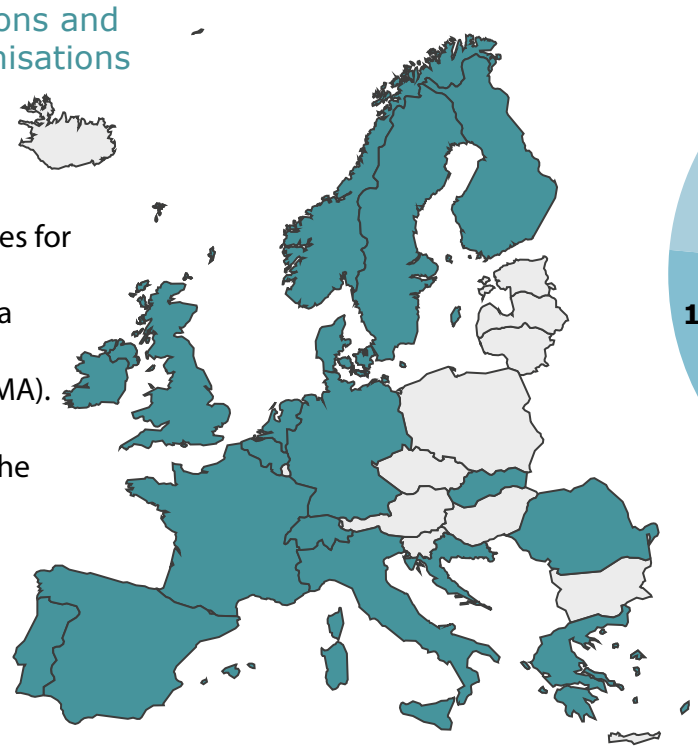
10 years of collaboration

to strengthen the monitoring of benefits and risks of medicines in Europe.

Expertise and collaboration in pharmacovigilance and pharmacoepidemiology in Europe

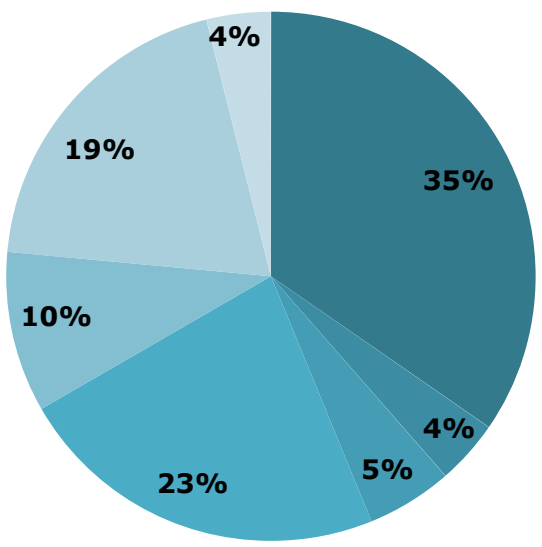
168 Public institutions and research organisations

18 European countries



The European Network of Centres for Pharmacoepidemiology and Pharmacovigilance (ENCePP) is a network coordinated by the European Medicines Agency (EMA).

The members of this network (the ENCePP partners) are public institutions and contract and research organisations (CRO) involved in research in pharmacoepidemiology and pharmacovigilance.



83 Data sources (by type, July 2017)

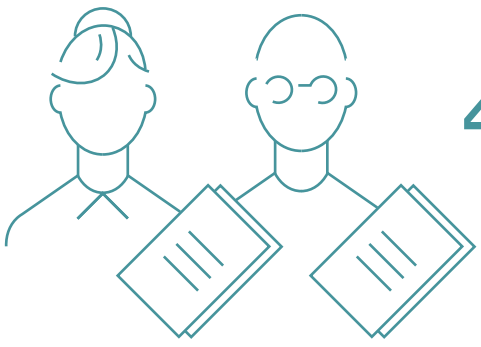
- Disease registry
- Spontaneous reporting database
- Prescription event monitoring
- Claims/administrative databases
- Electronic healthcare records
- Medicine registry
- Other

Methodological guidance

ENCePP Guide on Methodological Standards in Pharmacoepidemiology—a single web resource for methodological English language guidance (6th revision published in July 2017).

5000 views on average per month in 2017 (www.encepp.eu)

850 downloads on average per month in 2017 (www.encepp.eu)



424 references selected, commented and updated

31 authors

Governance principles

ENCePP Checklist for Study Protocols

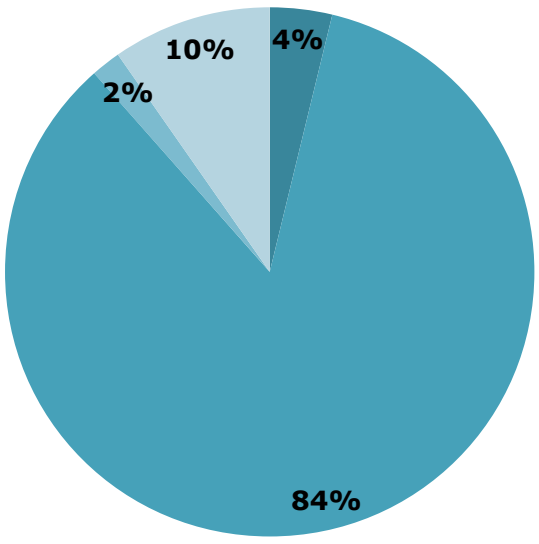
- Supporting best practice in study design
- Promoting transparency on study methods

ENCePP Code of Conduct

- Promoting transparency and scientific independence throughout the research process

Increased transparency in high quality, multi-centre, independent non-interventional post-authorisation studies (PAS)

ENCePP hosts the public **European Union electronic Register of Post-Authorisation Studies (EU PAS Register®)**.



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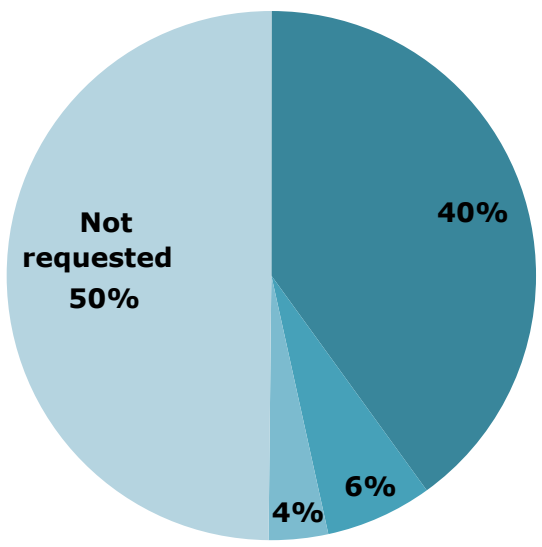
PAS registered (By type, July 2017)

- Active surveillance
- Observational study
- Clinical trial
- Other



50%

of registered studies were requested by a regulator:

- Europe
- United States
- Rest of the World