

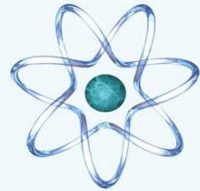
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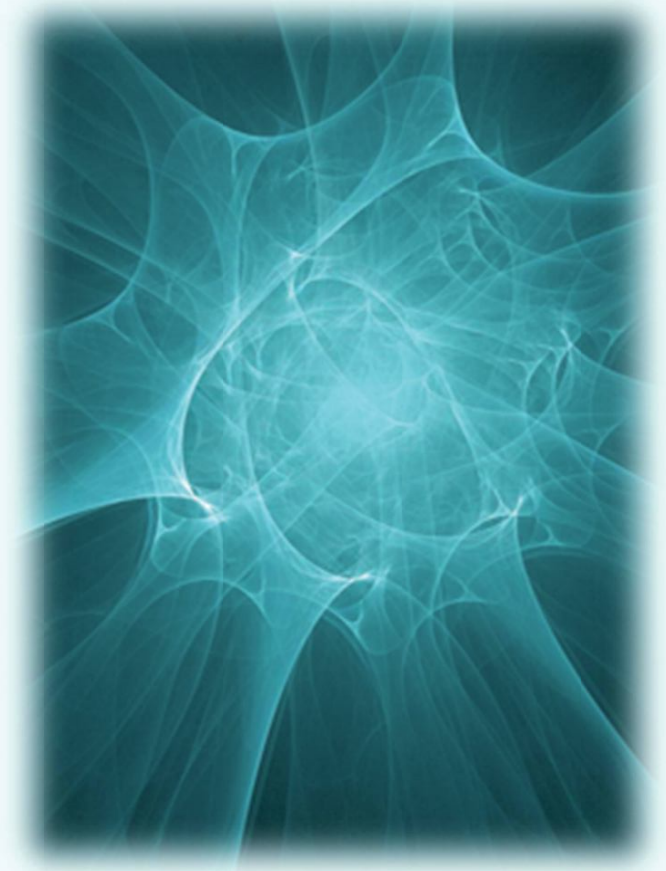
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Pharmaceutical
Service Network



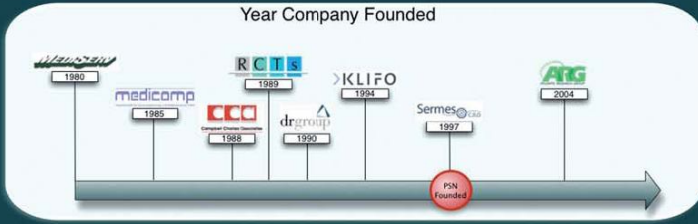
Pharmaceutical
Service Network



Pharmaceutical
Service Network

THINK Global  ACT Local

THINK Global  ACT Local



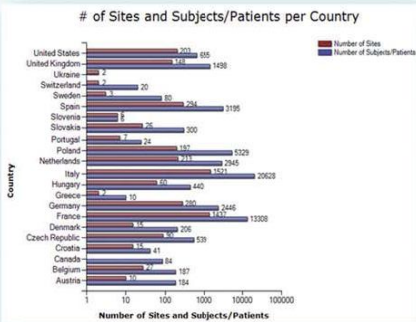
The psn advantage

The **Pharmaceutical Service Network** is a global CRO managed by a group of owner-driven Contract Research Organizations. The concept was born in 1997 to provide a better solution for pharmaceutical and biotech companies looking to conduct multinational trials.

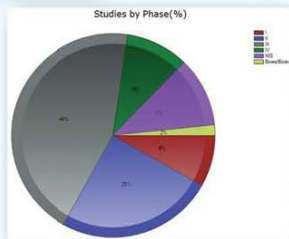
- ☑ Full-service CRO covering the EU and North America
- ☑ Flat management structure and personalized service
- ☑ Lower overhead results in better cost effectiveness
- ☑ Low staff turnover results in a consistent study team
- ☑ Quick contract negotiation and implementation of service
 - ☑ Services supported by common Standard Operating Procedures
 - ☑ Extensive knowledge of local clinical and regulatory requirements

psn has proven experience and expertise in assisting the drug and medical device development process.

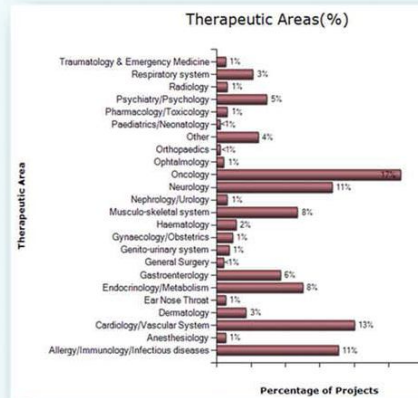
psn figures



CLIENTS: More than 100 active clients and 1000 studies performed

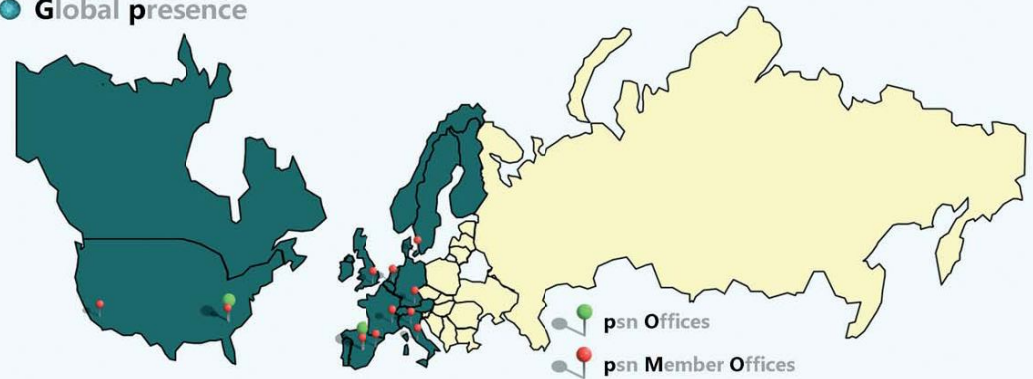


- TYPES OF STUDIES:**
- Clinical trials (all phases)
 - Post-authorization studies
 - Bioeq./Bioav. studies
 - Public health benefit studies



EMPLOYEES: Around 250 employees (20 PMs, 8 QAs ... and over 70 CRAs)

Global presence



COUNTRIES DIRECTLY COVERED BY psn MEMBERS: AUSTRIA, BELGIUM, CANADA, DENMARK, FINLAND, FRANCE, GERMANY, IRELAND, ITALY, NETHERLANDS, NORWAY, PORTUGAL, SPAIN, SWEDEN, SWITZERLAND, UNIDED KINGDOM, UNITED STATES

COUNTRIES COVERED BY psn PARTNERS: BULGARIA, CZECH REPUBLIC, CROATIA, ESTONIA, HUNGARY, LATVIA, LITHUANIA, POLAND, ROMANIA, RUSSIA, SERBIA, SLOVAKIA, SLOVENIA, UKRAINE

Services offered

Project Management

Broad experience managing international projects using a Customizable Clinical Trial Management System called Trial Vista.

Clinical Trial Supply

GMP authorisation for import, stock, randomisation, labelling, repackaging and worldwide distribution of clinical trials supplies.

QA/QC/Auditing

Quality checks during all the stages of a study as well as experience auditing vendors, partners and clients.

SAMM/Post Marketing Surveillance

Safety Assessment of Marketed Medicines and post marketing surveillance studies in various therapeutic areas.

Monitoring Clinical Trials

Local monitors coordinated by a study manager located in your country in accordance with local regulations and ICH requirements.

Medical Writing

Qualified and experienced medical writers available throughout EU and the US.

Training & Education

Global and national courses, workshops and seminars in several areas and languages.

Scientific & Drug Development Counseling

Panel of well-recognized experts that work cross-functionally on several disciplines.

Data Management & Statistics

Quality databases on time and within budget. Compliant with ICH and GCP Guidelines, including FDA 21 CFR Part 11. Experience with EDC systems.

Regulatory Affairs

From a full registration package (RAS and ECs) to special assignments.

Clinical Staff Outsourcing

Qualified staff covering a full range of positions (PMs, CRAs, CTAs, DMs, etc).

Pharmacoeconomics Studies & Evaluation of Product Pricing

Evaluation of efficiency and pricing of pharmaceutical products.

