

MS DATA ALLIANCE

Welcome

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There are many gaps in our knowledge about MS that need to be answered using real-world data



How can we diagnose better and earlier?

What affects disease evolution?



How does memory loss relate with disability?

Does an increase of 0.5 in EDSS mean that this person is deteriorating?



How effective are treatments in a real-world setting? (e.g. pregnancy, aging, treatment switch, combined treatments, long-term, ...)

Is one treatment more effective compared to another?



How safe and tolerable are our treatments in a real-world setting? (e.g. pregnancy, aging, treatment switch, combined treatments, long-term, ...)



Objectives of the meeting



To inform different stakeholders about our mission, vision and strategic focus areas of the MS Data Alliance initiative



To engage different stakeholders to be involved in the MS Data Alliance initiative



MSDA Stakeholders – Participants Stakeholder Meeting

e.g. RIMS, International MS Genetic Consortium, EAN, ACTRIMS, RUCTRIMS, Progressive MS Alliance, MS BrainHealth, Consortium of MS Centers, ParadigMS, Big MS Data Network, European Health Data and Evidence Network...

Researchers

Regulatory decision-makers

e.g. European Medicine Agency, BeNeLuxA initiative (=Health Technology Assessment)

Clinicians

People with MS / patient societies

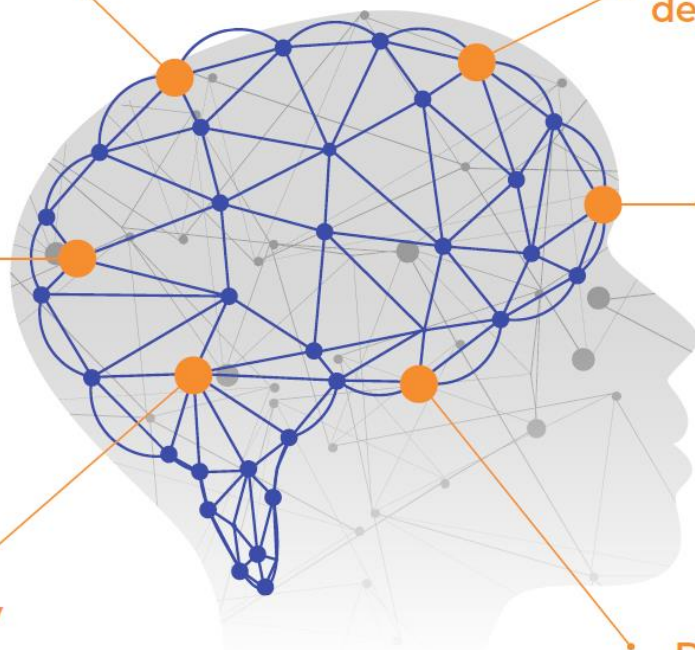
e.g. European MS Platform, MS International Federation, European Patients Forum, United States, Italy, Ireland, Spain, Serbia, Portugal, Denmark, Greece, Belgium, Romania, France, ...

Industry

e.g. European Federation of Pharmaceutical Industries and Associations, Novartis, Merck, Biogen, Icometrix, Quanterix, Qmenta, Canopy Growth Corporation, Mylan, Janssen, Celgene, ...

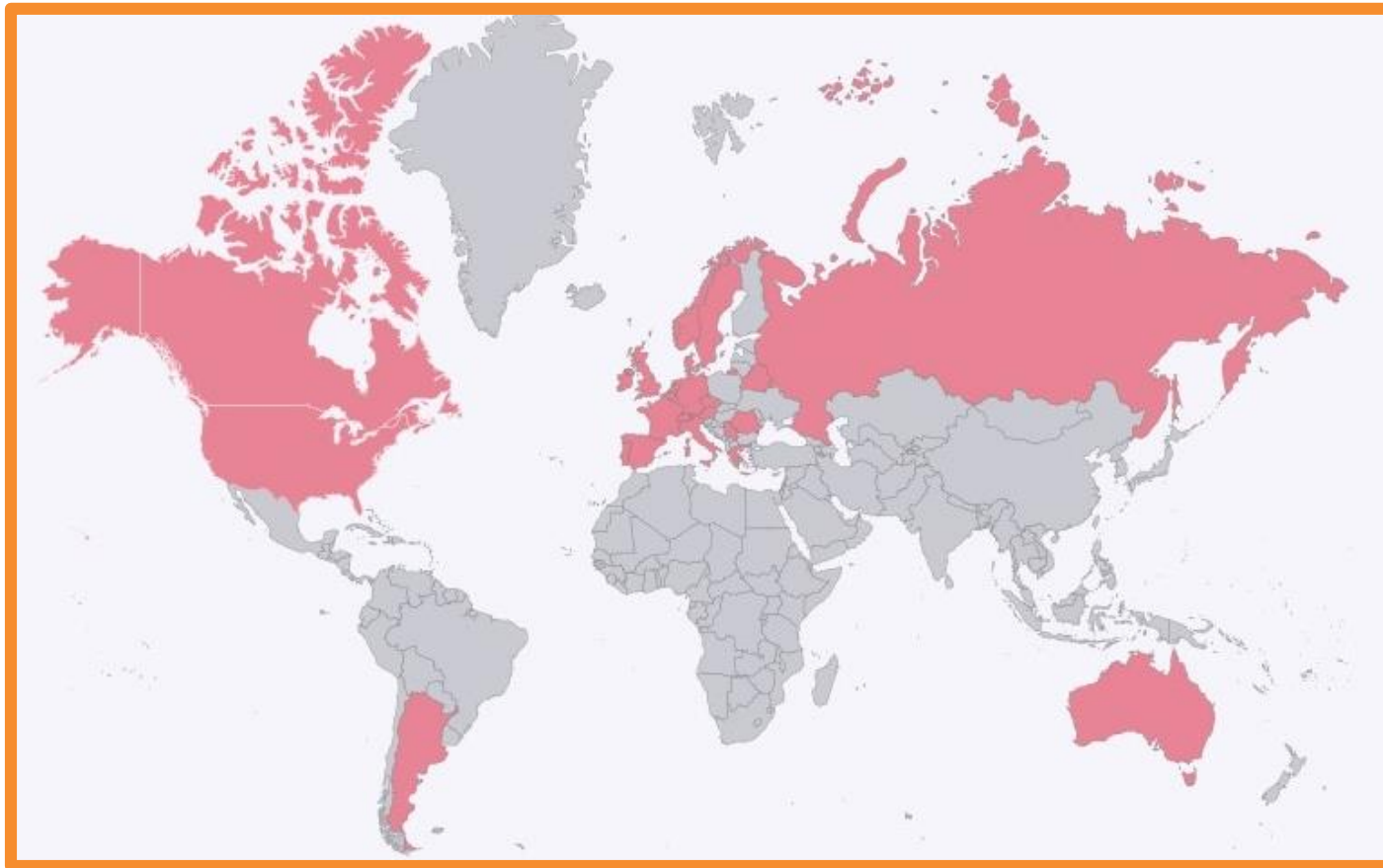
Data custodians

e.g. German MS Registry, UK MS Registry, MSBase Registry, Swedish MS Registry, Danish MS Registry, Italian MS Registry, OptimiseMS, RevelarEM, IconquerMS, Accelerated Cure Project for MS, NeuroTransData, PROMOPROMS, Spanish MS Registry, MultipleMS, German Pregnancy Registry, NARCRMS, CEMCAT CIS Cohort, Canadian Progression Cohort in MS, Greek Registry, NARCOMS, British Columbia MS Database, Australian MS Longitudinal Study, Swiss Cohort Study, Swiss MS Registry, Norwegian MS Registry, BELTRIMS, MSDS3D, ...





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MS DATA ALLIANCE

Because data can
transform the care of people
with Multiple Sclerosis

