

The European Multiple Sclerosis Platform

European Register for Multiple Sclerosis EUReMS: Pooling European Data to Better Understand and Fight Multiple Sclerosis

EMA Workshop on the patient-voice in the evaluation of medicines
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European Register for Multiple Sclerosis (MS)

- Patient oriented project , led by the EMSP
- Partnership with MS Clinical and Academic centres and MS Societies
- Addresses major public health challenges:
 - more than 600,000 people in Europe live with MS
 - access to MS specialists and treatments varies across countries
 - lack of (comparable) data on epidemiology, drug effectiveness and costs and resources in MS
- Utilises and harmonises national and regional MS Registers and databases
- Supported by the European Commission Public Health Program from July 2011 till June 2014

European Register for Multiple Sclerosis in 2011-2014

EUREMS First Studies :

- EPI-1-d *Estimating Prevalence and Incidence of MS in Europe from EUREMS data collection*
- EPI-1-s Study: *Comparison of the effect of the month of birth across Europe*
- DMD-1 Study: **Comparison of access and effectiveness of DMD treatment for people with MS across Europe**
- PRO-1 Study: *Assessment of people with MS' quality of life, the burden of disease and influence of employment from the patient's perspective across European countries*