

Join the Italian Multiple Sclerosis Register project.

All is needed
is your consent,
to make a contribution
to the research.

And also to your future



WHAT IS THE REGISTER?

It is a multicentric organized infrastructure to collect the data of all people with multiple sclerosis at Italian national level. Developed as a scientific research tool, the data from the Italian Multiple Sclerosis Register will be useful for carried out specific studies and to promote public healthcare planning actions



WHAT IS IT FOR?

The data collected allows monitoring the progression of the disease, assessing the effectiveness of treatments and helping to improve services. Valuable information that increases the knowledge of the disease



WHAT YOU HAVE TO DO?

Participant do not have to undergo any additional examination but only sign the informed consent so that their clinical data can be transmitted to the Register strictly anonymously, and used in an aggregate manner



Over 55 thousand people
have already joined the project.
If you have already done so,
check with your neurologist
to have signed the informed
consent for data processing

Technical Methodological Structure



UNIVERSITÀ
DEGLI STUDI DI BARI
ALDO MORO

Network of the Italian MS Clinical Centers



un mondo
libero dalla SM

AI SM. INSIEME, UNA CONQUISTA DOPO L'ALTRA

www.registroitalianosm.it