

Thinking about joining?

Neurologists treating MS patients:

- interested in participating in a worldwide Registry?
- interested in conducting research studies with colleagues within your own country and worldwide?
- able to benefit from using a free, structured tool for data collection and investigator collaboration?

Join now by going to:

www.msbase.org

Click on '**Register now**'

and complete your details

Nurse, Scientist, or Administrator who works with a Neurologist who is a member of MSBase? Request to join his/her centre as a co-investigator.

Patient, healthcare professional, student, MS industry, interested in keeping informed of the registry activities? Become a Friend of the registry.

Register now by going to:

www.msbase.org

Take part. Make a difference.

MSBase
Multiple Sclerosis dataBase

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MSBase
Free International Online Registry for Multiple Sclerosis Researchers

www.msbase.org

Registry Overview

The MSBase Registry is an international online registry for MS clinical researchers. It is a *free* tool for electronic data collection, sub study management, and online Investigator collaboration with Multiple Sclerosis Researchers worldwide.

The Registry collects and aggregates coded patient data from routine clinic visits for studies of MS regionally, nationally or globally, with a fully customisable sub-study functionality and flexible electronic case report form.

This research tool can be used by MS clinics contributing data to MSBase to participate in global analyses or to run their own studies.

Anonymity of Data

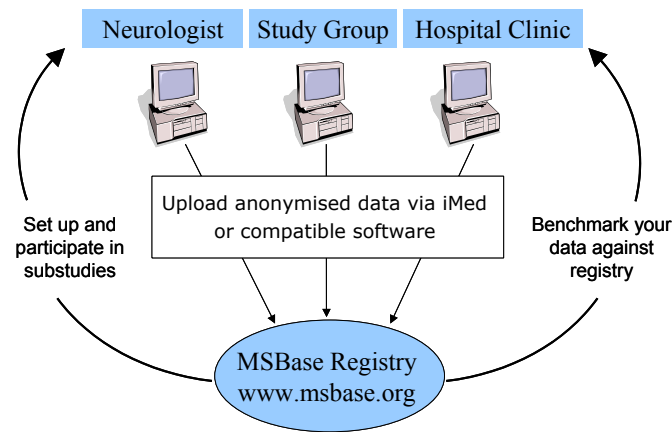
Patients in the MSBase Registry are identified by an assigned identification code (ID). Any identifying details are removed from the record before reaching the registry.

Registry Objectives

The objectives of the Registry are to:

- Provide a platform for the conduct of observational studies on MS 'real world' data
- Minimise the cost, time and setup of studies of MS with an electronic data collection solution
- Encourage global research collaboration

Structure



Ethical Requirements

- Ethics approval or exemption is required
- Patient consent is required

Benefits of membership

- Participation in a worldwide database that allows the benchmarking of your centre's data against the registry
- Communication with neurologists worldwide via email correspondence
- Participation in sub-studies with local and international neurologists
- Opportunity to develop sub-studies using a free system for identification of potential collaborators, communication, data quality assurance and data management
- Participation in MS research without additional case report forms (for those using a compatible local electronic documentation system)

Sponsor

The Registry is sponsored by the MSBase Foundation, a health related charity which is incorporated in Melbourne, Australia.

The Foundation seeks sponsorship from Industry, Government and other granting bodies.