

STRATEGY ON A PAGE 2023–2028

MISSION
To provide a global governance framework, database tools and research support to enable worldwide collaborative outcomes and epidemiologic research for the benefit of people with multiple sclerosis and other neuro-immunological diseases.

VISION
To be the global leader in supporting collaborative, real-world, quality research in multiple sclerosis and other neuro-immunological diseases.

OUR VALUES
Respect, Collaboration, Accountability, Quality

The MSBase Foundation Ltd (“MSBase”) is a not-for-profit charitable organisation incorporated as a company in Australia. In collaboration with participating neurologists around the world, the MSBase Foundation has established the MSBase Registry - a unique, web-based platform dedicated to sharing, tracking and evaluating outcomes data in Multiple Sclerosis and other Neuro-immunological diseases.

The Foundation is governed by an international Board of Directors who are responsible for the oversight of the charity. The Board integrates and prioritises the scientific initiatives as determined by the MSBase Scientific Leadership Group (SLG).

STRATEGIC OBJECTIVES:

GOALS

OUTCOMES

IT INNOVATION & SCALABILITY



Modernise systems and applications through a holistic design approach, ensuring MSBase is at the forefront of cloud-based technology

- Successfully deliver a complex redevelopment project of the Foundation’s platforms and systems, by ensuring a robust RFP process and a staff cohort that has the necessary skills and expertise.
- Secure financial support from industry partners and seek expert specialist advice to support the project.

SAFETY & COLLABORATION



Continue systematic collection of Safety outcomes in the MSBase Registry and active participation in the Big MS Data Network

- Successfully recruit centres for the Safe MS Care sub-study and secure additional safety registry funding from pharmaceutical partners.
- Achieve European Medicines Agency certification of the Big MS Data Network and create and support common data models in Big MS.
- Contribute to the policy frameworks to improve safety collection and reporting from registries.

COMMITMENT TO WOMEN'S HEALTH



Popularise pregnancy data and women’s health outcomes data in the MSBase Registry resulting in improved care for women with MS

- Increased awareness of the Women’s Health, Pregnancy, and Neonatal Outcomes Registry amongst MSBase members and the importance of entering pregnancy outcomes in the pregnancy module in iMed and MDS.
- Increased international awareness of the registry via the submission of abstracts to international conferences and by producing a descriptive paper annually on data collection and methods to date.

MRI INTEGRATION



Propel clinical-imaging research by making available to investigators a global MRI repository with real-time imaging analytics and encourage collaborations utilizing this data.

- Greater awareness of the MSBase Imaging Repository (MSBIR) amongst the global membership base, observed by an increase in participating sites and submission of project proposals requesting access to quantitative and raw imaging data.
- Encourage collaborative opportunities with external registries such as MAGNIMS (Magnetic Resonance Imaging in MS).

QUALITY & ASSURANCE



Enhance the quality of MSBase Registry data and systematically report on key quality indicators.

- Defined centre specific and overall data quality indicators.
- Integration of data quality and visualization tools in the redeveloped cloud-based MSBase Foundation IT platform, including centre reports and potentially data quality scores and ranking to enable centres to track and improve.

DISEASE EXPANSION



Champion the growth of specific disease registries such as Neuromyelitis Optica (NMO) and Myasthenia Gravis (MG)

- A coveted resource of high quality, international data on patients with NMO and MG available for investigator-initiated research and to industry partners.
- MSBase will secure financial support from pharmaceutical industry partners to support these registries through extended service agreements.