

# Annual Disability-State Transition Probabilities Estimated from a Large Sample (>6,000) of Australians Living with Multiple Sclerosis from MS-Base







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# BACKGROUND AND AIMS

Multiple sclerosis (MS) is a chronic, neurodegenerative disease characterised by progressing disability and it is most frequently diagnosed in females between the ages of 20 and 40 [1]. MS is high cost and high burden for Australian society, costing \$2.5 billion Australian dollars (AUD) in 2021. Cost increases concomitantly with disability severity; the cost per person living with MS in Australia in 2021 increased from \$32,829 for no disability to \$123,333 for severe disability [2]. MS prevalence is increasing. There are now 33,335 people living with MS in Australia, an increase of over 4,000 people from 2017 [2,3]. The *Atlas of MS* reveals there are 2.9 million people living with MS worldwide. This represents a 26% increase from 2013 (Figure 1).

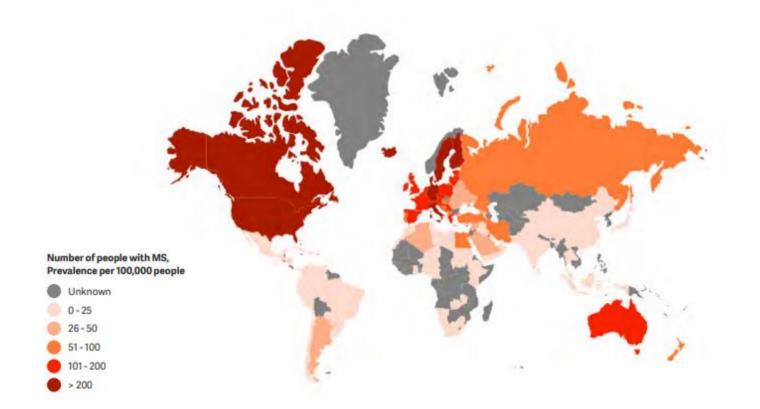


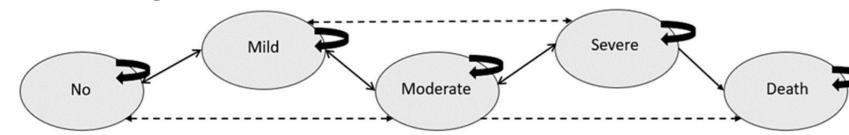
Figure 1: Source
Atas of MS that
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Transition probabilities are a vital input for health economics models, including those used to inform healthcare resourcing decisions relating to MS-specific disease-modifying therapies (DMT) reimbursement decisions.

## **METHODS**

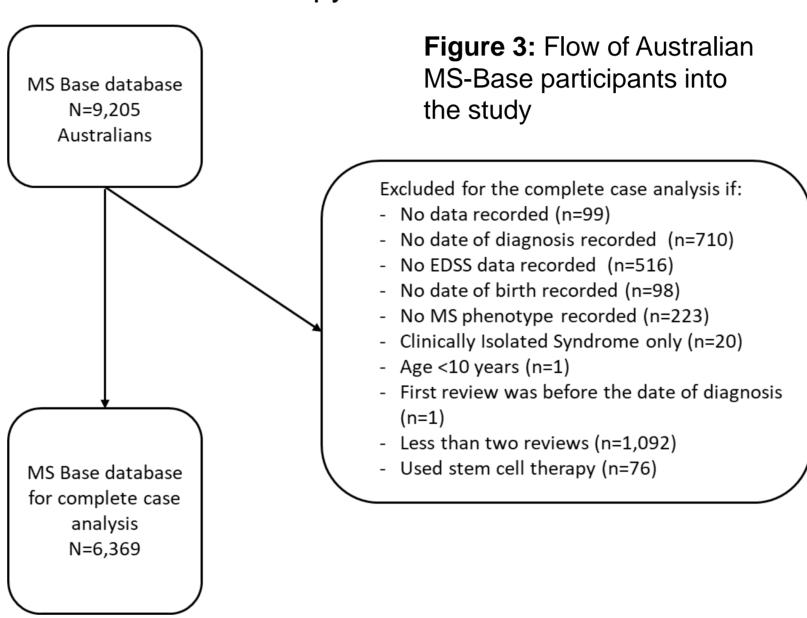
Data was obtained from Australian participants of MS-Base, the largest international registry collecting observational data as part of routine clinical care [4]. We fitted a five-state continuous-time Markov model to describe how people living with MS transition between disability milestones or health states. These states were, as defined by the Expanded Disability Status Scale: no disability (EDSS=0), mild (EDSS=1-3.5), moderate (EDSS=4-6), severe (EDSS=6.5-9.5), and death (EDSS=10). **Figure 2** comprises a diagram of the allowable transitions between health states for the final model. Model covariates included sex, DMT usage, MS-phenotype, and disease duration.

Figure 2: Allowable transitions between health states



## PRELIMINARY RESULTS

**Figure 3** displays the flow of participants into the study. Participants were retained if they represented a complete case, had more than one review, and did not utilise stem cell therapy.



**Table 1** provides a summary of the sociodemographic and clinical characteristics of the study participants. **Table 2** shows summary statistics pertaining to observation times and numbers of clinical reviews.

 Table 1: Participant characteristics

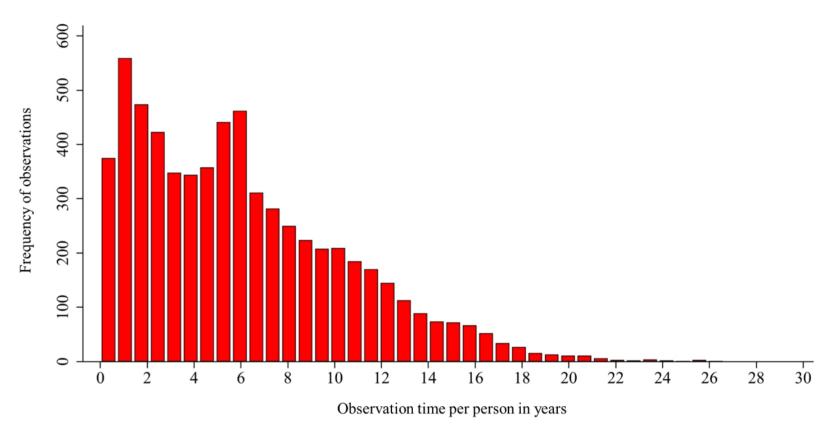
Characteristic	Value					
Number of included participants: n	6,369					
Age: mean (SD)	42.51 (12.37)					
Sex: n (%) female	4,777 (75.00)					
Disability severity* n (%)						
No disability	959 (15.06)					
Mild disability	3554 (55.80)					
Moderate disability	1274 (20.00)					
Severe disability	582 (9.14)					
Number of deaths						
Death	102 (1.60)					
Disease duration: mean (SD)	5.69 (7.40)					
Multiple sclerosis phenotype: n (%)						
Relapsing-remitting	5566 (87.39)					
Secondary-progressive	460 (7.22)					
Progressive-onset	343 (5.39)					
Disease-modifying therapy usage: n (%)						
None	2010 (31.57)					
Category 1	539 (8.46)					
Category 2	514 (8.07)					
Category 3	3306 (51.91)					
Progressive-onset Disease-modifying therapy usage: n (%) None Category 1 Category 2	343 (5.39) 2010 (31.57) 539 (8.46) 514 (8.07)					

**Table 2:** Observation time and clinical reviews

	Observation Time	Number of	
	in Years	<b>Clinical Reviews</b>	
Summary Statistic			
Mean	6.12		
Standard deviation	4.47		
Interquartile Range			
0%	0.02	2	
25%	2.43	4	
50%	5.40	7	
75%	8.87	11	
100%	26.62	48	
Mean		8.58	
Standard Deviation		6.66	
Total	38,992.78	54,669	
Total	Person Years	Clinical Reviews	

# PRELIMINARY RESULTS

**Figure 4** supports Table 2 and shows the frequency of observation times in years; the mean (SD) observation time was 6.12 (4.47) years.



**Table 3** provides the frequency of transitions between the five health states of no disability, mild, moderate and severe disability and death. Rows are the 'from' states and columns are the 'to' states.

	No	Mild	Moderate	Severe	Death
No	4073	1815	41	10	1
Mild	1736	21818	1807	81	11
Moderate	36	1380	8288	1003	25
Severe	5	26	526	5553	65
Death	0	0	0	0	0

Table 3: Total number of transitions between health states

When covariates were set to mean values, one-year transition probabilities included (reported as "value [95% CI]"): remaining disability-free (54.0% [43.6%, 56.2%]) remaining in the mild state (82.0% [60.4%, 82.6%]); transitioning from the no to mild state (42.1% [40.1%, 43.9%]); from the mild to moderate state (11.4% [10.9%, 11.9%]; from the moderate to severe state (9.5% [8.8%, 10.3%]); and from the severe to death state (0.5% [0.3%, 0.7%]) (detailed data not shown; provided in our forthcoming publication)

### **CONCLUSIONS**

Preliminary results indicate that MS progresses quickly from no disability. Estimated transition probabilities will be applied in a health economics simulation model for Australia intended to support reimbursement decisions.

### REFERENCE

[1] Taylor BV. What causes multiple sclerosis? Getting closer to the answers. Med J Aust. 2022 Aug 15;217(4):180-182.; [2] Campbell JA, Simpson S Jr. Ahmad H, Taylor BV, van der Mei I, Palmer AJ. Change in multiple sclerosis prevalence over time in Australia 2010-2017 utilising disease-modifying therapy prescription data. Mult Scler. 2020 Oct;26(11):1315-1328; [3] Campbell JA, van der Mei I, Taylor BV, Palmer AJ. Health economic impact of Multiple Sclerosis in Australia in 2021: an interim update on prevalence, costs and cost of illness. Commissioned Report for MS Australia https://www.msaustralia.org.au/wp-content/uploads/2023/02/health-economic-impact-of-multiple-sclerosis-in-australia-in-2021\_final.pdf

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