Supplementary Table 2 Adherence to the recommended laboratory monitoring tests in the months before and after initiation of the first oral disease-modifying therapy for the treatment of multiple sclerosis.

Number (n) of people	Proportion of people with a laboratory test							
included, by oral DMT	Biocher	iochemical liver Lymphocyte		phocyte	Urinalysis			
and time period	test ^a		count					
	n	%	n	%	n	%		
Dimethyl fumarate								
Pre-treatment								
Within 6 months, n=567	518	91.4	531	93.7	438	77.2		
On-treatment								
Months 1-6, n=339	302	89.1	305	90.0	253	74.6		
Months 7-12, n=197	175	88.8	180	91.4	141	71.6		
Fingolimod								
Pre-treatment								
Within 6 months, n=253	222	87.8	231	91.3		NA		
On-treatment								
Months 1-3, n=225	138	61.3		NA		NA		
Months 4-6, n=206	139	67.5						
Months 7-9, n=192	118	61.5						
Months 10-12, n=167	92	55.1						

Teriflunomide					
Pre-treatment					
Within 6 months, n=196	179	91.3	183	93.4	NA
On-treatment					
Month 1, n=174	70	40.2		NA	NA
Month 2, n=156	64	41.0			
Month 3, n=147	59	40.1			
Month 4, n=131	45	34.4			
Month 5, n=124	39	31.5			
Month 6, n=117	29	24.8			

Key: DMT, disease-modifying therapy

^aMinimum recommendations, for the biochemical liver tests, according to each DMTs product monograph: either ALT or AST for dimethyl fumarate and fingolimod; and for teriflunomide, either ALT or AST pre-treatment, and ALT while on-treatment.

NA – not applicable as the relevant laboratory test was not part of the routine monitoring recommendations.