

How do patients enter the healthcare system after the first onset of multiple sclerosis symptoms? The influence of setting and physician specialty on speed of diagnosis

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Abstract

Background: Diagnosing multiple sclerosis (MS) early is crucial to avoid future disability. However, potentially preventable delays in the diagnostic cascade from contact with a physician to definite diagnosis still occur and their causes are still unclear.

Objective: To identify the possible causes of delays in the diagnostic process.

Methods: We analyzed the data of the Swiss MS Registry. With logistic regression, we modeled the time from the first contact to the first consultation (contact-to-evaluation time, ≤ 1 month/ > 1 month) and the evaluation-to-diagnosis time (≤ 6 months/ > 6 months). Potential factors were health system characteristics, sociodemographic variables, first symptoms, and MS type.

Results: We included 522 participants. Mostly, general practitioners (67%) were contacted first, without delaying the diagnosis. In contrast, first symptoms and MS type were the major contributors to delays: gait problems were associated with longer contact-to-evaluation times, depression as a concomitant symptom with longer evaluation-to-diagnosis times, and having primary progressive MS prolonged both phases. In addition, living in mountainous areas was associated with longer contact-to-evaluation times, whereas diagnosis after 2000 was associated with faster diagnoses.

Conclusion: For a quicker diagnosis, awareness of MS as a differential diagnosis of gait disorders and the co-occurrence of depression at onset should be raised, and these symptoms should be attentively followed.

Keywords: Registries, regression analysis, time to diagnosis, delayed diagnosis, patient-reported outcomes, epidemiology

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Introduction

Diagnosing a patient with multiple sclerosis (MS) is a complex and multifaceted clinical process. However, the past three decades have seen a significant reduction in the time from onset of symptoms to diagnosis.¹ Probable reasons are the constantly refined diagnostic criteria,² an increased magnetic resonance imaging (MRI) availability, and the increasing treatment options, approved in Switzerland since 1993. Nevertheless, a substantial number of patients still experience significant delays before receiving a definite diagnosis. A Swiss study by Kaufmann et al.³ found that—even in recent time periods—40% of all

newly diagnosed persons with MS reported more than 2 years passing between symptom onset and the MS diagnosis. This is even more relevant in light of the mounting evidence on the benefits of early treatment initiation in relapsing remitting multiple sclerosis (RRMS) to prevent long-term disability.^{4–6}

The reasons for delays in the diagnosis are not fully understood, and few studies have attempted more detailed investigations into the time from MS onset to diagnosis (henceforth referred to as diagnostic time).^{7–10} Of these studies, most were unable to investigate the delaying factors using a systematic approach and

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performed in highly specialized MS centers.^{7–9} An exception was a population-based study from Canada,¹⁰ which found that younger age at onset and having primary progressive multiple sclerosis (PPMS) were associated with longer times to diagnosis.

Health system-specific aspects and care access barriers may also play an important role in diagnostic delays. Studies from the United States found that insurance coverage, living in remote rural areas, and public transportation deficiencies can constitute significant barriers to accessing neurological care in MS patients,¹¹ which may equally apply to the diagnostic process.

Therefore, further investigations of delaying factors should not only consider clinical aspects but also possible access barriers and the diagnostic cascade potential MS patients with first symptoms take through the healthcare system.

Using the Swiss Multiple Sclerosis Registry (SMSR), we aimed to analyze the diagnostic cascade from the moment a patient seeks care because of first symptoms until a clinically isolated syndrome (CIS) or an MS diagnosis is confirmed. In addition, we aimed to analyze which setting- and patient-specific features potentially delay the diagnosis at the different cascade steps. This knowledge could lead to an optimization of the diagnostic process and possibly to an earlier start of drug and non-drug (e.g. neurorehabilitation, lifestyle adjustments) treatment of MS patients.

Methods

The conceptual model of the diagnostic cascade

Along the lines of the diagnostic cascade described by Fernández et al.,⁷ we developed a conceptual model of the diagnostic steps in collaboration with MS neurologists, thereby taking into account specific features of the Swiss healthcare system (e.g. free physician choice; no mandatory first visit to the primary care provider unless the health insurance plan requires it). This concept was subsequently used to guide our analysis and was the framework for summarizing our study data. A sketch of the model is outlined in Figure 1 and includes three steps. In step 1, a person notices his or her symptoms and contacts a healthcare provider, which, in Switzerland, can be a general practitioner (GP) or directly a specialist. The contacted physician's office then schedules a visit, which depending on symptom severity could take from a few days to months (step 2, henceforth named "contact-to-evaluation phase"). If, at the actual visit,

the symptoms suggest an MS diagnosis, a neurological referral would follow (step 3, "evaluation-to-diagnosis phase").

In accordance with this conceptual model, separate data analyses were performed to investigate the factors associated with delays in steps 2 and 3, as well as to summarize the variety of observed cascades over both steps.

The data sources

The SMSR is an ongoing, Swiss-based, patient-centered, longitudinal observational study, initiated and funded by the Swiss MS Society (<http://www.ClinicalTrials.gov>; identifier: NCT02980640). Study recruitment is continuous and started in June 2016. Adults with CIS or confirmed MS who live or receive care in Switzerland can voluntarily participate. A wide range of measures have been implemented to motivate and enable Swiss persons with MS to participate in the SMSR (described in detail by Puhani et al.¹²), such as regular information through the news outlets of the Swiss Multiple Sclerosis Society in the three main national languages, presentations to health care professionals and potential participants in clinics and private practices, as well as the possibility to fill in questionnaires on paper or via a specifically designed online platform. At registration, a baseline questionnaire is completed. Afterwards, follow-up questionnaires can be answered on a semi-annual basis. The treating physician provides confirmation of the diagnosis. Details on the study design are described elsewhere.^{12,13}

The Ethics Committee of Zurich approved the study (Study No. PB-2016-00894) and an informed consent was signed by each participant prior to study entry.¹²

Our analysis was restricted to patients diagnosed after 1995 and considered all data collected until 19 March 2018. The study utilized information from the first follow-up questionnaire administered 6 months post baseline. The baseline assessments provide information on age, sex, lifestyle factors, education level, living situation, year of MS diagnosis, current MS form, first symptoms, and family history.¹⁴ The 6-month follow-up questionnaire includes a comprehensive section on the diagnostic process. It covers the healthcare professional first contacted in connection with MS symptoms, details about the setting and specialty of the healthcare provider involved in the diagnosis, as well as the specialty of the healthcare provider in charge of the diagnostic tests. Moreover, for steps 2 and 3 of our conceptual model, the duration of each

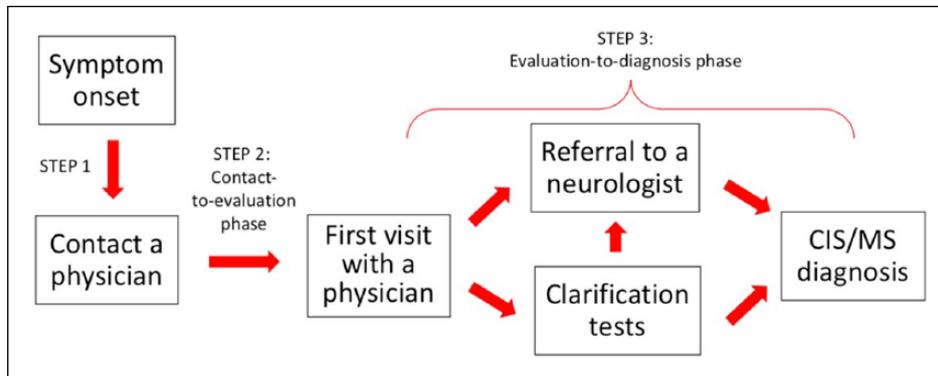


Figure 1. The conceptual model of the diagnostic process.

phase was assessed (contact-to-evaluation phase and evaluation-to-diagnosis phase, respectively).

Description of the cascade

To simplify the different cascades, we created a decision tree on the basis of our conceptual model with the following bifurcations to depict the diagnostic process steps: (1) whether a setting change occurred between the first contact with a physician's office to the first visit (e.g. an immediate referral to a neurology clinic by a GP office), (2) whether a neurologist was present at the first visit, and (3) whether the physician at the first visit led diagnostic tests, referred to another physician, or took no action. Setting changes were defined as any changes in the care setting (private practice, hospital) and/or the specialty of the attending physician between the first contact and the first visit.

Identification of the factors associated with prolonged diagnostic times

In order to assess the influence of different factors on the diagnosis duration, we performed logistic regression models with contact-to-evaluation and evaluation-to-diagnosis durations as outcome variables.

To identify prolonged diagnostic times, the contact-to-evaluation duration was categorized into "within 1 month" and "more than 1 month", whereas the evaluation-to-diagnosis duration was categorized into "within 6 months" and "more than 6 months".

The main factors (variables of interest) in the regression models were the decision tree branches with some modifications due to low patient numbers as follows. For the contact-to-evaluation duration, the

first two bifurcations (setting change, neurologist at first visit) were considered, and for the evaluation-to-diagnosis duration the last two bifurcations (neurologist at first visit, actions on clarification tests) were analyzed. Other fixed regression factors were sex, education level (mandatory education or apprenticeship/school leaving certificate/higher professional education/university degree), living in a mountainous area (No/Yes), MS type (RRMS start/PPMS), age at onset, time period of diagnosis (1996–2000/2001–2010/2011–2018, reflecting the release years of the diagnostic criteria), and first visit setting (private practice/hospital). Potential additional factors were living situation, rural/urban area, language region of residence, Swiss nationality, MS diagnosed in older relatives, and 20 different first symptoms. If a first symptom was rare (in less than 5% of the sample), it was discarded (Supplemental Table S2).

For each logistic regression model, the potential factors were added to the fixed factors in a stepwise approach based on Akaike information criterion (AIC).³ Missing information (ranging from 0.1% to 5% per variable) were imputed by means of the multivariate imputation by chained equations (MICE) algorithm.¹⁴ The entire analysis was performed using R, version 3.4.0.¹⁵

Results

Study population

A total of 522 participants were included (Supplemental Figure S1). Patients were mostly female with RRMS (74%), with a median age of 47 years and an Expanded Disability Status Scale (EDSS) score below 4 (81%). A detailed description of the study population is provided in Table 1.

Table 1. Demographics, clinical information, and diagnostic process features of the included sample.

Variables	Participants included in the regression analyses ^a
<i>N</i>	522
Age at baseline, median (IQR)	47 (38; 54)
Sex—female	384 (73.6%)
Disease duration at baseline (years) ^b , median (IQR)	7 (3; 13)
Disease duration not available	11 (2.1%)
MS form at baseline	
CIS	9 (1.7%)
RRMS	384 (73.6%)
PPMS	49 (9.4%)
SPMS	72 (13.8%)
Not available	8 (1.5%)
EDSS proxy at baseline	
0–3.5	420 (80.5%)
4–6.5	68 (13%)
7–10	32 (6.1%)
Not available	2 (0.4%)
Contacted first doctor	
General practitioner	352 (67.4%)
Ophthalmologist	68 (13%)
Neurologist	44 (8.4%)
Emergency room	34 (6.5%)
Other specialists	23 (4.4%)
Not available	1 (0.2%)
Time from contact with a physician to first symptom evaluation	
Same day/next day	96 (18.4%)
Within 1 week	115 (22%)
1–2 weeks	75 (14.4%)
2–4 weeks	68 (13%)
1–3 months	60 (11.5%)
More than 3 months	94 (18%)
Other	14 (2.7%)
Specialization of doctor at the first visit	
General practitioner/internist	308 (59%)
Neurologist	124 (23.8%)
Other specialist (incl. ophthalmologist)	89 (17%)
Not available	1 (0.2%)
Place of the first visit	
Private practice	349 (66.9%)
Hospital	111 (21.3%)
Emergency room	59 (11.3%)
Not available	3 (0.6%)

Table 1. (Continued)

Variables	Participants included in the regression analyses ^a
Action on evaluation tests by the first physician	
Refer to other MD	335 (64.2%)
Lead the tests	146 (28%)
No action taken	35 (6.7%)
Not available	6 (1.1%)
Next steps and visits sufficiently explained	
Yes	354 (67.8%)
Partially	125 (23.9%)
No	41 (7.9%)
Not available	2 (0.4%)
Setting of further evaluation	
Outpatient	332 (63.6%)
Inpatient	136 (26.1%)
In- and outpatient	47 (9%)
Not available	7 (1.3%)
Setting of diagnosis (from confirmation of diagnosis document)	
Neurologist in a hospital	267 (51.1%)
Neurologist in a private practice	191 (36.6%)
Others (general practitioner, ...)	2 (0.6%)
Not available	62 (11.9%)
Number of doctors visited prior to MS diagnosis	
One	34 (6.5%)
Two	244 (46.7%)
Three	128 (24.5%)
More than three	114 (21.8%)
Not available	2 (0.4%)
Time from first symptom evaluation to MS diagnosis	
Less than 1 week	87 (16.7%)
1–2 weeks	98 (18.8%)
2–4 weeks	78 (14.9%)
1–3 months	101 (19.3%)
3–6 months	48 (9.2%)
6–12 months	34 (6.5%)
More than 1 year	72 (13.8%)
Other	4 (0.8%)

IQR: interquartile range; CIS: clinically isolated syndrome; RRMS: relapsing remitting multiple sclerosis; PPMS: primary progressive multiple sclerosis; SPMS: secondary progressive multiple sclerosis; EDSS: Kurtzke Expanded Disability Status Scale.

^aParticipants at the 6-month follow-up, diagnosed since 1996 and with both dependent variables available.

^bThe MS duration refers to the time difference between the MS diagnosis date and the date of baseline survey. Where available, the diagnosis date was obtained from the diagnosis confirmation sheet provided by the treating physician (available for 89%). For the remainder, diagnosis dates were self-reported in the baseline questionnaires.

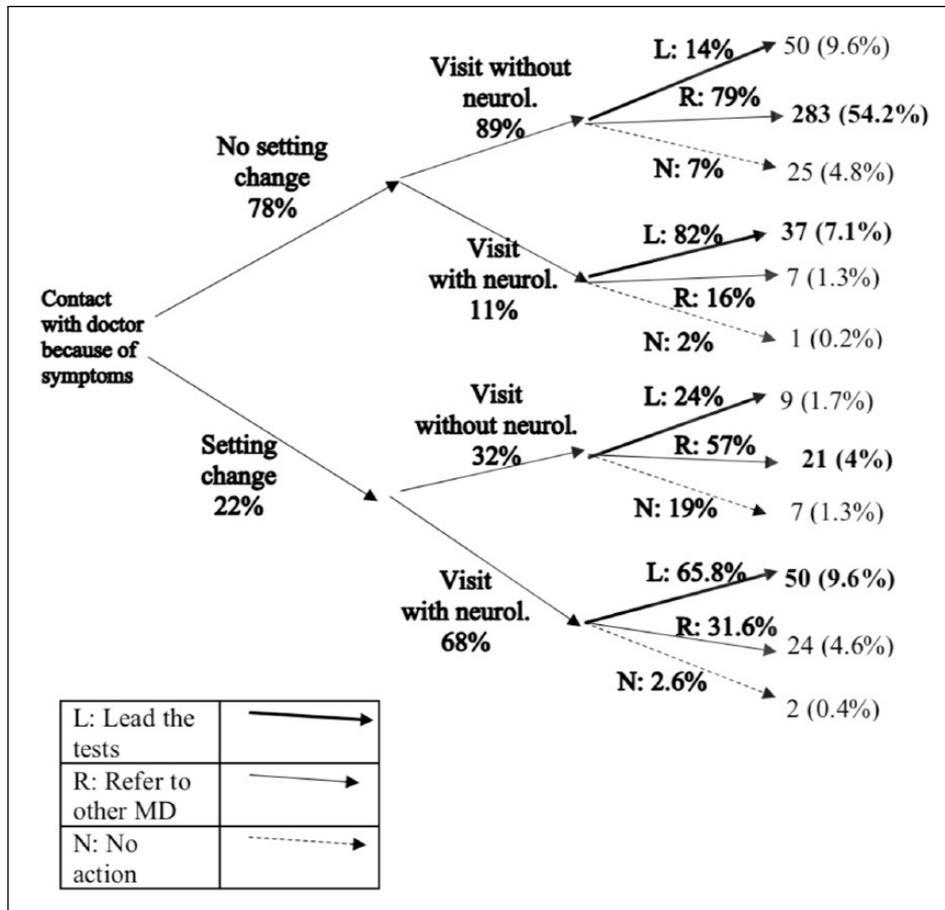


Figure 2. Decision tree, from the first contact until the decisions on the clarification tests are taken.

L: lead the tests; R: refer to other medical doctor; N: no action.

Percentages displayed next to branches illustrate what population fraction within a node entered a specific branch and sum up to 100% for each bifurcation. Numbers indicated at each tree endpoint provide the frequency of participants who have had that combination of events, as well as their share (%) of the entire sample. Numbers in bold, on the right-hand side of the figure, are the most frequent action for each node.

Description of the diagnostic cascades

Table 1 presents the details of the diagnostic process. During the contact-to-evaluation phase, the majority of patients (67%) first contacted a GP because of their symptoms. A minority sought care from ophthalmologists (13%), neurologists (8%), or general emergency services (7%). For 17% of participants, the evaluation-to-diagnosis duration (step 3 in Figure 1) was shorter than a week, for another 53% it occurred within 3 months, for 16% between 3 and 12 months, and for the remaining 14% more than a year.

Figure 2 summarizes the first steps of the diagnostic process with a decision tree. Four out of five participants were seen by the physician they contacted (i.e. no change of setting, 78%). Of these patients, the vast majority (89%) did not see a neurologist at their first visit. By contrast, of those patients who did change

the setting (22%), more than two out of three (68%) were seen by a neurologist. It is further observed from Figure 2 that, regardless of any prior setting change, neurologists were more likely to lead, perform, or organize diagnostic tests, while non-neurologists tended to refer patients to other physicians. Similar patterns can also be observed when the sample is restricted to the participants diagnosed after 2010 (Supplemental Figure S2).

Factors associated with prolonged time from the first contact to symptom evaluation

Table 2 (columns 3 and 4) shows the distribution of all the considered, potential delaying factors for the contact-to-evaluation phase stratified by shorter and more prolonged duration. Persons with PPMS were more likely to wait longer for the first visit (5% in the

Table 2. Stratification of all factors considered for modeling the contact-to-evaluation time (columns 3 and 4) and the evaluation-to-diagnosis time (columns 5 and 6).

Levels		Contact-to-evaluation time: within 1 month	Contact-to-evaluation time: more than 1 month	Evaluation-to-diagnosis time: within 6 months	Evaluation-to-diagnosis time: more than 6 months
<i>N</i>		354	168	412	110
Diagnosis time period	1996–2000	42 (11.9%)	38 (22.6%)	58 (14.1%)	22 (20%)
	2001–2010	150 (42.4%)	69 (41.1%)	171 (41.5%)	48 (43.6%)
	After 2010	162 (45.8%)	61 (36.3%)	183 (44.4%)	40 (36.4%)
Age at onset	6–20	22 (6.2%)	8 (4.8%)	22 (5.3%)	8 (7.3%)
	21–30	120 (33.9%)	41 (24.4%)	131 (31.8%)	30 (27.3%)
	31–40	107 (30.2%)	50 (29.8%)	122 (29.6%)	35 (31.8%)
	41–50	79 (22.3%)	43 (25.6%)	99 (24%)	23 (20.9%)
	51–70	10 (2.8%)	16 (9.5%)	20 (4.9%)	6 (5.5%)
	NA	16 (4.5%)	10 (6%)	18 (4.4%)	8 (7.3%)
	PPMS	No	336 (94.9%)	133 (79.2%)	379 (92%)
	Yes	17 (4.8%)	35 (20.8%)	32 (7.8%)	20 (18.2%)
	NA	1 (0.3%)	–	1 (0.2%)	–
Sex	Female	267 (75.4%)	117 (69.6%)	304 (73.8%)	80 (72.7%)
	Male	87 (24.6%)	51 (30.4%)	108 (26.2%)	30 (27.3%)
Decision tree—first two bifurcations	No setting change, first visit w/o neurologist	246 (69.5%)	115 (68.5%)	–	–
	No setting change, first visit with neurologist	32 (9%)	14 (8.3%)	–	–
	Setting change, first visit w/o neurologist	25 (7.1%)	12 (7.1%)	–	–
	Setting change, first visit with neurologist	51 (14.4%)	27 (16.1%)	–	–
Decision tree—last two bifurcations	First visit w/o neurologist, refer to other MD	–	–	235 (57%)	69 (62.7%)
	First visit w/o neurologist, lead the tests	–	–	49 (11.9%)	10 (9.1%)
	First visit w/o neurologist, no action	–	–	18 (4.4%)	14 (12.7%)
	First visit with neurologist, lead the tests	–	–	76 (18.4%)	11 (10%)
	First visit with neurologist, no action or refer to other MD	–	–	29 (7%)	5 (4.5%)
	NA	–	–	5 (1.2%)	1 (0.9%)
Living in mountainous area	No	310 (87.6%)	132 (78.6%)	354 (85.9%)	88 (80%)
	Yes	41 (11.6%)	34 (20.2%)	54 (13.1%)	21 (19.1%)
	NA	3 (0.8%)	2 (1.2%)	4 (1%)	1 (0.9%)
Place of first visit	Private practice	226 (63.8%)	123 (73.2%)	267 (64.8%)	82 (74.5%)
	Hospital	126 (35.6%)	44 (26.2%)	144 (35%)	26 (23.6%)
	NA	2 (0.6%)	1 (0.6%)	1 (0.2%)	2 (1.8%)
First symptoms (more than one possible)	Gait problems	93 (26.3%)	71 (42.3%)	125 (30.3%)	39 (35.5%)
	Visual problems	135 (38.1%)	49 (29.2%)	145 (35.2%)	39 (35.5%)
	Bladder problems	37 (10.5%)	21 (12.5%)	49 (11.9%)	9 (8.2%)
	Balance problems	90 (25.4%)	52 (31%)	116 (28.2%)	26 (23.6%)
	Bowel problems	24 (6.8%)	14 (8.3%)	29 (7%)	9 (8.2%)
	Dizziness	72 (20.3%)	36 (21.4%)	86 (20.9%)	22 (20%)

Table 2. (Continued)

Levels		Contact-to-evaluation time: within 1 month	Contact-to-evaluation time: more than 1 month	Evaluation-to-diagnosis time: within 6 months	Evaluation-to-diagnosis time: more than 6 months
	Dysarthria	25 (7.1%)	4 (2.4%)	24 (5.8%)	5 (4.5%)
	Paresthesia	216 (61%)	100 (59.5%)	251 (60.9%)	65 (59.1%)
	Weakness	94 (26.6%)	58 (34.5%)	117 (28.4%)	35 (31.8%)
	Paralysis	87 (24.6%)	46 (27.4%)	105 (25.5%)	28 (25.5%)
	Fatigue	113 (31.9%)	61 (36.3%)	137 (33.3%)	37 (33.6%)
	Pain	41 (11.6%)	25 (14.9%)	52 (12.6%)	14 (12.7%)
	Spasticity	22 (6.2%)	25 (14.9%)	36 (8.7%)	11 (10%)
	Sexual dysfunction	17 (4.8%)	18 (10.7%)	27 (6.6%)	8 (7.3%)
	Memory problems	28 (7.9%)	15 (8.9%)	33 (8%)	10 (9.1%)
	Depression	32 (9%)	21 (12.5%)	37 (9%)	16 (14.5%)
Education level	Mandatory education/apprenticeship	141 (39.8%)	83 (49.4%)	180 (43.7%)	44 (40%)
	School leaving certificate	38 (10.7%)	7 (4.2%)	35 (8.5%)	10 (9.1%)
	Higher professional education	54 (15.3%)	30 (17.9%)	59 (14.3%)	25 (22.7%)
	University degree	105 (29.7%)	42 (25%)	121 (29.4%)	26 (23.6%)
	NA	16 (4.5%)	6 (3.6%)	17 (4.1%)	5 (4.5%)
Living situation	With partner only	135 (38.1%)	65 (38.7%)	151 (36.7%)	49 (44.5%)
	With partner and children	134 (37.9%)	55 (32.7%)	157 (38.1%)	32 (29.1%)
	Alone	55 (15.5%)	39 (23.2%)	73 (17.7%)	21 (19.1%)
	Other	28 (7.9%)	9 (5.4%)	29 (7%)	8 (7.3%)
	NA	2 (0.6%)	–	2 (0.5%)	
Language region of Switzerland	German	302 (85.3%)	135 (80.4%)	340 (82.5%)	97 (88.2%)
	French	38 (10.7%)	21 (12.5%)	51 (12.4%)	8 (7.3%)
	Italian	11 (3.1%)	10 (6%)	17 (4.1%)	4 (3.6%)
	NA	3 (0.8%)	2 (1.2%)	4 (1%)	1 (0.9%)
MS diagnosed in older relatives	Yes	39 (11%)	10 (6%)	40 (9.7%)	9 (8.2%)
	No	292 (82.5%)	153 (91.1%)	347 (84.2%)	98 (89.1%)
	NA	23 (6.5%)	5 (3%)	25 (6.1%)	3 (2.7%)
Swiss citizenship	Yes	330 (93.2%)	154 (91.7%)	382 (92.7%)	102 (92.7%)
	No	24 (6.8%)	14 (8.3%)	30 (7.3%)	8 (7.3%)
Typology of residence	Urban	196 (55.4%)	90 (53.6%)	222 (53.9%)	64 (58.2%)
	Urban–rural	99 (28%)	41 (24.4%)	111 (26.9%)	29 (26.4%)
	Rural	56 (15.8%)	35 (20.8%)	75 (18.2%)	16 (14.5%)
	NA	3 (0.8%)	2 (1.2%)	4 (1%)	1 (0.9%)

PPMS: primary progressive multiple sclerosis; NA: not available.

early group, 21% in the late group), as well as those with gait problems at onset (26% early, 42% late), and those residing in a mountainous area (12% early, 20% late). The contact-to-evaluation time was more often shorter after 2000 (88% early, 77% late), or with dysarthria at onset (7% early, 2% late).

Those observations were confirmed by multivariable logistic regression analyzing the contact-to-evaluation

time (Figure 3). The factors associated with a shorter contact-to-evaluation time were diagnosis after 2000 (odds ratio (OR): 2.1, 95% confidence interval (CI): [1.2; 3.8] for 2001–2010, 2.7 and [1.5; 4.8] after 2010), having a school leaving certificate (2.4, [0.97; 5.74]), and dysarthria at onset (3.14, [1.01; 9.77]).

PPMS (0.28, [0.14; 0.56]), gait problems at onset (0.6, [0.39; 0.93]), and living in a mountainous area

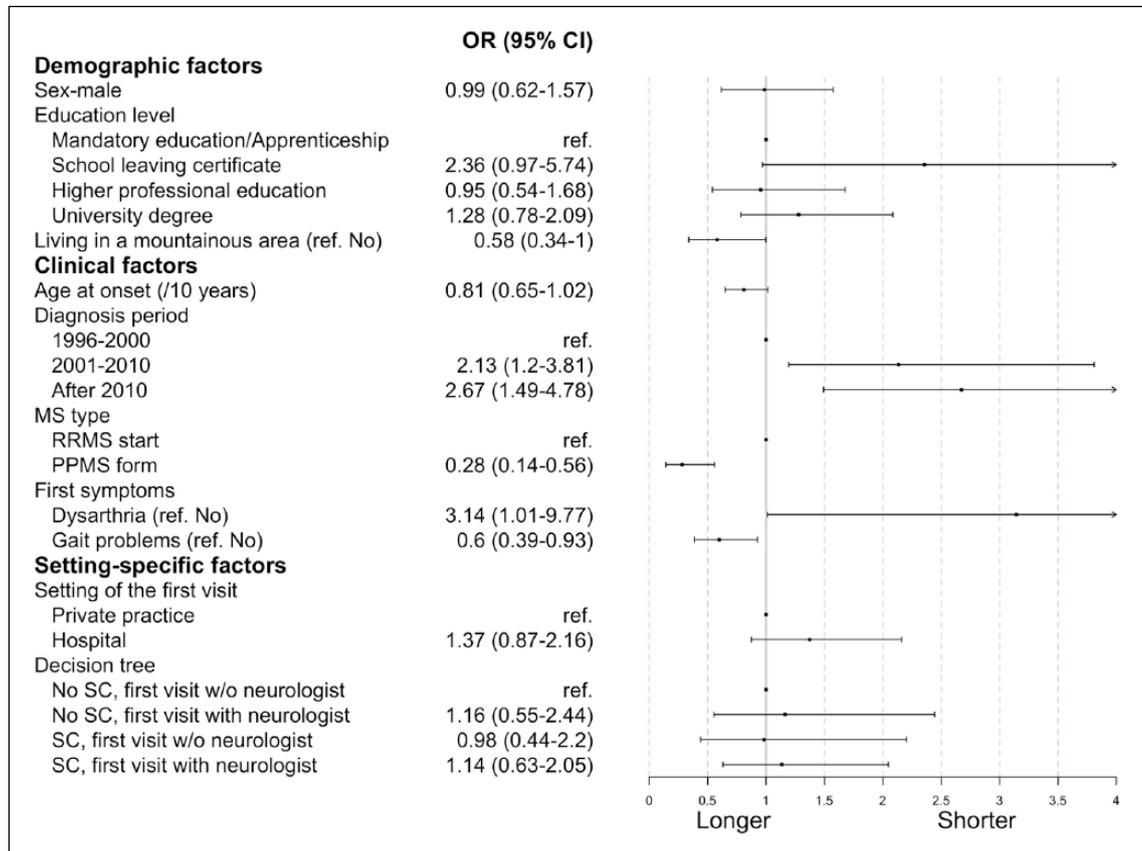


Figure 3. Regression coefficients of the multivariable model on the contact-to-evaluation time.

ref.: reference level; PPMS: primary progressive multiple sclerosis; RRMS: relapsing remitting multiple sclerosis; Priv. Practice: private practice; SC: setting change.

(0.58, [0.34; 1.00]) were instead associated with a longer time. Of note, the tree-derived levels describing the sequence of contacts were not associated with a prolonged time from first contact to first visit. When restricting the analysis to participants without PPMS, gait problems remained significantly associated with a longer contact-to-evaluation time (Supplemental Table S3 and Supplemental Figure S3).

Factors associated with prolonged time from symptom evaluation to MS diagnosis

The last two columns of Table 2 describe all potential delaying factors for the evaluation-to-diagnosis phase. Shorter times were observed more often when the diagnosis was after 2010 (44% early, 36% late), the neurologist led the tests (18% early, 10% late), or the first visit occurred at a hospital (35% early, 24% late). Prolonged durations were observed with depression as a concomitant first symptom (9% early, 15% late), lack of action by a non-neurologist (4% early, 13% late), and PPMS (8% early, 18% late). These pat-

terns also emerged in the multivariable logistic regression (Figure 4).

Based on the different cascades illustrated in Figure 2, we further hypothesized that some cascades were more likely to lead to swifter diagnoses than others, for example, when neurologists are almost immediately involved in diagnosis. Indeed, the multivariable model indicated that the evaluation-to-diagnosis duration was statistically significantly faster when a neurologist led the clarification tests, compared to patients first seeing a GP and then being referred (OR 2.21 [1.07; 4.59]). Moreover, a diagnosis year after 2010, compared with one before 2000, was also associated with faster evaluation-to-diagnosis times (1.96, [1.03; 3.74]). By contrast, having PPMS (0.28, [0.14; 0.56]), depression as a concomitant first symptom (0.46, [0.24; 0.91]), higher professional education (0.49, [0.27; 0.91]), or seeing a (non-neurologist) physician who did not undertake further diagnostic actions (0.28, [0.13; 0.62]) were all associated with a prolonged evaluation-to-diagnosis phase.

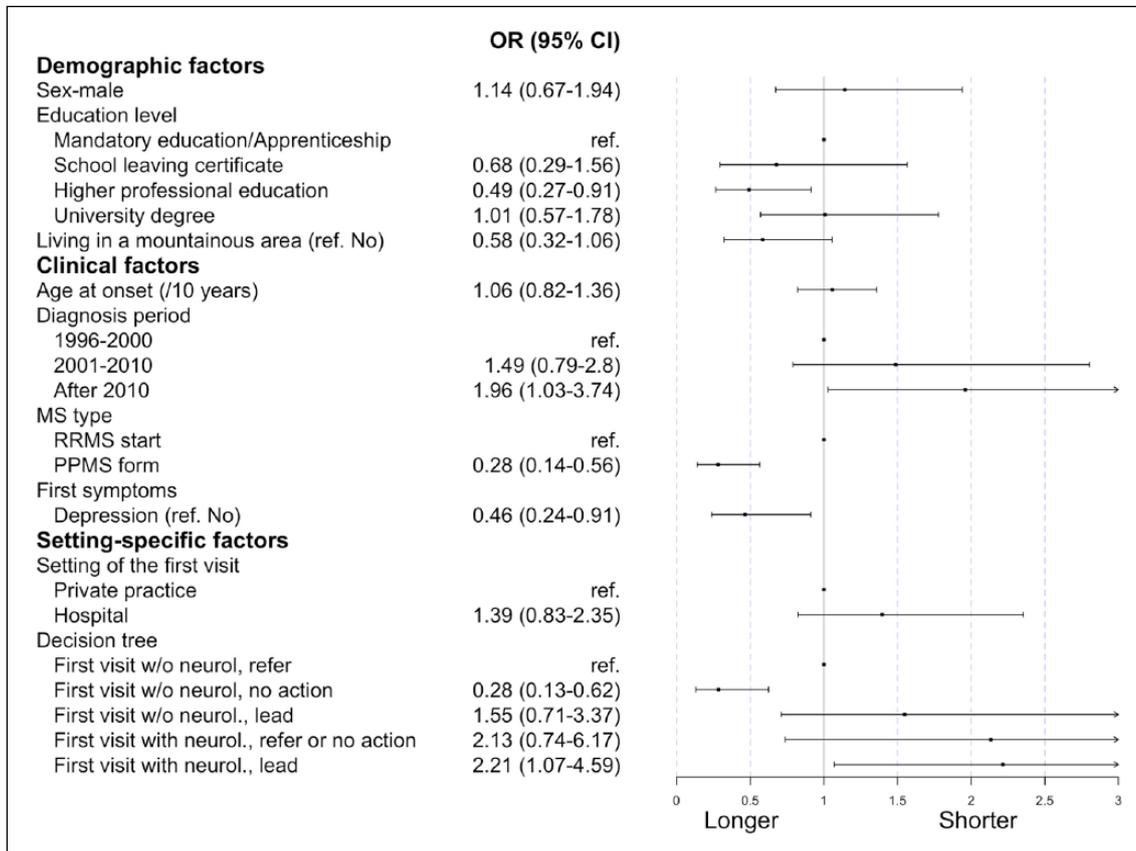


Figure 4. Regression coefficients of the multivariable model on the evaluation-to-diagnosis time.

ref.: reference level; PPMS: primary progressive multiple sclerosis; RRMS: relapsing remitting multiple sclerosis; Priv. Practice: private practice; neurol.: neurologist.

Discussion

Using comprehensive questionnaire data from 522 Swiss MS Registry participants, we described the MS diagnosis process for persons receiving care in Switzerland. Our study analyzed in detail the steps after the first contact and identified the delaying factors.

We observed a broad variety of possibilities on how potential MS patients entered the healthcare system (Supplemental Table S1), but overall the cascade functioned as expected (Figure 2).

GPs were contacted primarily, and they usually referred patients to a neurologist who led further diagnostic tests and finally made the diagnosis. Importantly, this procedure did not prolong the overall diagnostic process compared to patients immediately contacting a neurologist, unless no action was taken. By contrast, the type of MS or first symptoms were more crucial in prolonging the process. Specifically, gait problems were associated with a longer time from the first contact to the first visit and dysarthria with a shorter one.

The concomitance of depression was associated with a longer time from the first visit to the diagnosis and having PPMS was the most influential factor for a prolonged diagnosis duration and negatively influenced both phases.

To our knowledge, few studies have explored the diagnostic pathways and the role of patient-, disease-, and setting-specific aspects in MS diagnosis duration.⁷⁻¹⁰ Fernández *et al.*⁷ identified delays in several steps leading up to an MS diagnosis (onset to first visit, first visit to referral, time to perform the tests) and found the phase from the first symptoms to the first visit to be particularly critical. Kelly *et al.* found that approximately 78% of patients saw a neurologist within 6 weeks from onset, in accordance with UK guidelines. They also provided a list of common reasons for delay.⁸

Detailed examinations of factors delaying the diagnostic process were, however, largely missing. For example, Marrie *et al.*^{1,16} examined the effect of single factors, namely, onset year and comorbidities.

Kingwell et al. pursued a systematic approach to identify the factors associated with the diagnostic time, but did not include sociodemographic factors, access barriers, and features of the diagnostic cascade in their assessment. Recently, Kaufmann et al.³ investigated this question using SMSR data, but without distinguishing between the different diagnostic process phases, instead modeling the whole diagnostic time at once.

This study extends these prior efforts by analyzing the different steps of the cascade in greater detail and testing a broader array of factors potentially delaying the diagnosis. Our findings may add to the literature by emphasizing the role of symptom presentation for a swift MS diagnosis. In particular, the concomitant occurrence of depression at onset (in 10% of the study population) as a delaying factor confirms the results of Marrie et al.¹⁶ There are various explanations: depression appears highly heterogeneous and is not considered a specific MS symptom, the patients themselves might not seek care due to an inability to take action, and the physician might interpret organic symptoms as psychogenic due to depression.

The delaying effect of gait problems might appear to contradict the findings of Kaufmann et al.,³ where they were associated with a quicker diagnostic process. However, in our study, gait problems were associated with a longer contact-to-evaluation phase, but not a longer evaluation-to-diagnosis phase. Gait problems often occur slowly, especially in patients with PPMS, and are therefore not urgent symptoms triggering immediate examinations, despite being distinctive first MS symptoms (Supplemental Table S3).

Furthermore, the fact that the diagnostic cascade was shorter every time the main diagnostic criteria were released (2001, 2010) extends Marrie et al.'s¹ findings on a more recent timeframe.

Our finding of longer diagnostic times in patients with PPMS confirms those reported in other studies.^{3,10} The delaying effect of having PPMS on the time from the first evaluation to diagnosis possibly reflects the requirement of McDonald diagnostic criteria, which require 1 year of confirmed progression for this MS type. However, because delays in PPMS were also statistically significantly longer in the first phase of the diagnostic cascade (i.e. before the first evaluation), this suggests that at least a part of the delay occurs independently of the diagnostic criteria applied. For example, it is possible that the nature of symptoms in PPMS is less disruptive and urgent when compared to those that predominate in RRMS, thus

already leading to delayed care seeking after the first symptom occurrence.

From a healthcare utilization perspective, specific sociodemographic factors could represent care access barriers hindering the diagnostic process.¹¹ However, our findings revealed that most factors potentially constituting a barrier, such as lower education, living in an urban area, or having a migration background, do not lead to a prolonged diagnosis. Only living in mountainous areas was associated with longer diagnosis durations, probably due to a shortage of neurologists in those areas. But overall, our findings suggest that there is a good healthcare access, due to the Swiss mandatory healthcare insurance, a high hospital capacity,¹⁷ and diffuse hospital vicinity,¹⁸ resulting in a reasonably fast diagnostic process.¹⁹

Strengths and limitations

To better understand during which diagnostic cascade phase delays can occur, we differentiated the diagnostic process into two phases. Referring to more objective events (e.g. first contact, first visit) rather than personal interpretations (e.g. onset date) made our outcomes less prone to recall bias and subjectivity.

Moreover, the available information of the SMSR, such as education level, circumstances of life, and family history, is more comprehensive than other similar studies,^{20,21} and adjusting for them contributes to the findings' robustness. In addition, having precise information on a large variety of different first symptoms allowed us to indicate which symptoms should be better monitored to prevent delays.

While some of our findings are setting specific (e.g. the distribution of primary care contacts), others are likely to generalize to other settings. For example, the diffuse nature of some of the MS symptoms associated with delays is likely also of concern in other countries.

This study presents, however, some limitations. First, self-reported data can be affected by biases, such as recall bias and underreporting. Moreover, we did not include information on comorbidities other than depression at the time of the onset that were previously found to contribute to delays.¹⁶

In addition, diagnostic delays can also occur due to financial access barriers, even in countries with social health insurance systems like Switzerland (e.g. due to high out-of-pocket expenses and upfront payments^{22,23}). Unfortunately, we were unable to assess the actual

financial situation of the participants, so we used the education level as a proxy instead.

Prospect for future research

To accelerate the diagnostic process further, future research should examine the reasons why patients may not seek immediate care after the first onset of MS symptoms. Moreover, the potential causes for initial inactivity by first healthcare providers deserve further investigation and may lie in unusual symptom combinations, lack of awareness for MS, or problems in patient–doctor communication.

Conclusion

Our study is among the first to query persons with MS on how they responded after noticing their first symptoms. This effort has provided a comprehensive picture of the patients' first steps in the healthcare system toward an MS diagnosis after symptom onset, which was unknown until now in Switzerland. The observation that the majority of Swiss MS patients were diagnosed in a timely manner regardless of setting and specialty of first physician contact is reassuring, but still too many potentially preventable delays occur in these first steps until diagnosis initiation. For a faster diagnostic process, awareness for MS as a differential diagnosis of gait disorders should be raised, and an attentive follow-up of possible MS cases with depression as a concomitant first symptom is needed.

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