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Research Paper

The natural history of secondary progressive MS

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Abstract

Background: The onset of secondary progression is a pivotal event in the course of relapsing-remitting (RR) multiple sclerosis (MS). Patients with secondary progressive MS (SPMS) experience continuous worsening of symptoms independent of the occurrence of relapses. Possible risk factors associated with the onset of SPMS remain under-investigated in natural history studies of MS disease course.

Methods: We used Kaplan-Meier survival analyses and Cox regression models to investigate the influence of gender, onset age and onset symptoms on time to and age at SPMS in British Columbia (BC) MS patients with a RR disease onset who were unexposed to immunomodulatory drugs.

Results: Of 5778 patients in the BCMS database with definite MS, 5207 (90%) had a RR onset. Median time to SPMS was 21.4 years (95%CI:20.6-22.2), reached at a median age of 53.7 years (95%CI:53.1-54.3). Male gender and motor onset symptoms were associated with a shorter time to and a younger age at SPMS. A younger age at disease onset was associated with a longer time to SPMS, but also with a younger age at secondary progression. Other onset symptoms were not associated with time to, or age at, SPMS.

Conclusions: We identified three factors influencing the onset of SPMS in untreated patients with RRMS: motor onset symptoms and male gender were associated with both a shorter time to and a younger age at SPMS. A younger age at disease onset should not be viewed as indicating a better prognosis.

Introduction

Natural history studies of multiple sclerosis (MS) often focus on the time from disease onset until patients reach certain landmark disability scores (such as EDSS 6.0, when a cane is required for walking, or EDSS 8.0, when a patient is mostly restricted to a wheelchair). Besides the accumulation of disability, patients with relapsing-remitting MS are also at risk of experiencing a critical change in their disease course, the development of secondary progression.¹

A secondary progressive disease course has developed when a patient's clinical condition starts to deteriorate steadily and independently of relapses. Patients with secondary progressive MS (SPMS) have an unfavorable prognosis and can expect a slow and unremitting worsening of their ability to function.² The onset of secondary progression is therefore another important outcome to examine in natural history studies.

The onset of secondary progression has been addressed by several small studies that may have had limited power, and have often focused on patients who had already reached SPMS.³⁻⁵ Tremlett and co-workers have previously investigated a subset of the British Columbia MS (BCMS) cohort,⁶ but when examining predictors of progression, were unable to fully adjust findings for the different patient characteristics considered. Here, we expand findings by examining the entire available cohort and by fitting a multivariable model when considering possible predictors of progression. For the present study, we investigate the natural history of secondary progression in IMD-naïve patients by examining factors associated with the time from onset to the development of

SPMS in the whole BCMS cohort of relapsing at onset patients. Because MS is a chronic, life-long disease, the age at which patients reach the progressive phase is important and relevant to patients and their physicians; we have therefore also examined the factors that influence the age at which SPMS develops.

Methods

Clinical information

The clinical information was taken from the British Columbia MS (BCMS) database, a longitudinal database linking the four MS clinics in British Columbia, Canada, and capturing prospective data for over 80% of the MS population in the province.^{7,8} We selected all patients with a diagnosis of clinically definite MS according to the Poser diagnostic criteria,⁹ with a relapsing-at onset disease course who were enrolled in the database from September 1st 1980 to July 31st, 2003. SPMS disease course was defined clinically by the treating neurologist as a progressive course following a relapsing-remitting onset.¹⁰ Follow-up visits typically occurred annually. In order to investigate the natural history of the disease without the possible influence of immunomodulatory drugs (IMDs), patients who received such treatments were censored at the first day of treatment. Gender, age at disease onset and onset symptoms (motor, sensory, optic neuritis, and cerebellar/ataxia/brainstem), recorded at the patient's first BCMS clinic visit, were also extracted from this database. Age at onset was divided into four categories: <30 years, 30 to <40, 40 to <50 and \geq 50 years. The main outcome measures were the time from disease onset and from birth to the onset of secondary progression.

Statistics

The time from disease onset to onset of secondary progression and the age at secondary progression (i.e. time from birth to SPMS) were estimated from Kaplan-Meier

survival analyses. Group differences in survival time were compared with log-rank tests. In addition we built multivariable Cox regression models with the time to secondary progression and age at secondary progression as the dependent variables. Covariates included in these models were gender, onset symptoms and age at disease onset for the time from onset to secondary progression, and gender and onset symptoms for the age at secondary progression. The age at disease onset was not included as a covariate in the model for age at secondary progression because its value is necessarily smaller than the outcome in every case. The latter model was stratified by age at disease onset instead. We tested for possible violations of the proportional hazards assumption in these multivariable models with log-minus-log plots and Schoenfeld residuals.

All statistical analyses were performed with the SPSS statistical software package version 16 (SPSS Inc, Chicago, Ill., USA).

Standard Protocol Approvals, Registrations, and Patient Consents

The study was approved by the Clinical Research Ethics Board of the University of British Columbia.

Results

Patient cohort

Of 5778 patients with definite MS, 5207 (90%) had a relapsing-remitting onset. IMDs had been used by 1249 (24%) patients; these patients were censored at the first day of treatment. The date of onset of secondary progression was unavailable for 38 patients (0.7%), these patients were excluded from all analyses. The time of disease onset was unknown for a further 7 patients (0.1%), these patients were excluded from the analyses of the time to secondary progression from onset of MS and from the stratified Cox regression analysis of predictors of age at SPMS. Of the 5169 available patients, 1821 (35%) had reached SPMS by the study end.

Kaplan-Meier survival analyses

The results of the Kaplan-Meier survival analyses are shown in the table. The median time to secondary progression was 21.4 years from onset (95% CI: 20.6 - 22.2 years) and the median age at secondary progression was 53.7 years (95% CI: 53.1 - 54.3 years). There was wide variability in survival estimates in both the time to and the age at secondary progression: while the 25% of patients with the quickest disease progression reached secondary progression in less than 11.4 years and at a younger age than 45.1 years (first quartile estimated from the Kaplan-Meier analysis), the 25% with the slowest progression reached secondary progression after more than 32 years and at an age of more than 63.1 years (last quartile). Men and patients with motor onset symptoms developed SPMS after a shorter time and at a younger age (See Table, Figures 1 and

2). Patients with a younger age at onset took a longer time to SPMS, but reached SPMS at a younger age ($p < 0.0005$, Figure 3).

Multivariable regression models

The results of the multivariable Cox regression models are shown in the table. Male gender and the presence of motor onset symptoms were independently associated with a shorter time to, and a younger age at, SPMS (table). Younger age at disease onset was associated with a longer time to SPMS; the hazard rate of reaching SPMS increased by 27.1% (95%CI:23.4-30.9%) for every 5-year increase in onset age (hazard ratio per one year increase: 1.05 (95%CI: 1.04-1.06)).

Other onset symptoms were not associated with these outcomes. Our investigation of all models with log-minus-log plots and Schoenfeld residuals showed no evidence of a violation of the proportional hazards assumption

Discussion

We report on the long-term follow-up of a cohort of relapsing at onset MS patients from a population-based MS database in British Columbia, Canada. We found the onset of secondary progression to be a gradual process in the cohort as a whole, with a steady increase of conversions to SPMS observed over follow-up. However, the window was wide, with SPMS being reached within one year of MS onset, through to more than 50 years post-onset, and in individuals from 11 to 82 years of age.

Overall, the median time to SPMS was 21.4 years, which is similar to that reported for the few other cohorts in which survival analyses have been applied, including 19.1 years in Lyon¹¹ and 20.0 years in Lorraine, France.¹²

Three factors were associated with the onset of secondary progression in our cohort: gender, age at onset and motor onset symptoms.

Male gender predicted a shorter time to, and a younger age at, SPMS in our cohort. The influence of male gender on the time to onset of SPMS has been controversial, while some previous studies suggested that men have a worse prognosis than women,^{6,11,13} other studies have found no such association.¹⁴⁻¹⁸ Sex hormones may play a role in this gender difference - an increase in estrogen levels in particular is believed to reduce CNS

inflammation.¹⁹ Relapse frequency is also reduced during pregnancy and shows a rebound after delivery. This phenomenon has been associated with the rise and fall of estrogen and progesterone levels during pregnancy and the postpartum period.²⁰ Furthermore, the administration of oral estriol in women with MS resulted in a decrease in the number and volume of gadolinium enhancing lesions on cranial MRI in one preliminary study.²¹ A further trial investigating the potential therapeutic value of sex hormones for MS is currently ongoing.²²

Motor or 'long tract' onset symptoms have reportedly been associated with the time to SPMS in some studies,^{11,14,16} while others have found no association between onset symptoms and the timing of secondary progression.^{13,15,17} It is not clear why a motor onset should impact the disease course more than other onset symptoms. The steady worsening of neurological signs may be more apparent in patients with a motor disease onset, and this could lead to the earlier detection of secondary progression in this group. It is also possible that a significant portion of those patients with motor onset symptoms have an early involvement of the spinal cord, which is believed to carry a worse prognosis.²³ This merits further investigation.

The age at disease onset is considered to be the strongest predictor of the time to SPMS.^{11,15-17} A young age at the onset of MS has long been viewed as a relatively good prognostic sign, because a longer delay to specific disability outcomes has typically been observed in these patients.²⁴ In terms of the pathophysiology of MS, a younger

age has been viewed as favorable, because older age is associated with worse recovery from experimental demyelination in animal models due to impaired recruitment and differentiation of oligodendrocyte precursor cells.²⁵ Our investigation of the influence of MS onset age on conversion to SPMS shows that a younger age at MS onset should not be viewed as favorable: patients with a young age at disease onset may take longer until they convert to SPMS, but they are also younger at the onset of SPMS. This concurs with recent findings from other natural history studies examining time to disability (EDSS) milestones in both adult and pediatric MS.²⁶⁻²⁸

The main strength of this study is the fact that we have investigated a large population-based cohort followed for a long time period with prospectively collected data. A limitation of our study, however, is the fact that an estimated 20% of the MS patients in BC have not been captured by the database; we are unable to determine if these other MS patients in BC would have progressed to SPMS faster or slower than the patients included in this study. On the other hand, there is no reason to believe that factors such as gender, onset symptoms and age at onset would have any different effect on disease course amongst those not included in the database.

Recent research suggests that the use of IMDs may influence the timing of secondary progression.^{15,29} In this study we concentrated on untreated patients to evaluate which factors are relevant in the natural history of the disease and to provide a treatment-naïve comparison for future cohort studies. This information may also be helpful for the design of randomized controlled trials of the effect of IMDs on conversion to SPMS.

UBC MS Neurologists (in alphabetical order)

D. Adams, D. Craig, L. Daly, V. Devonshire, S. Hashimoto, O. Hrebicek, J. Hooge, B. Jones, L. Kastrukoff, S. Meckling, J. Oger, D. Parton, D. Paty, P. Smyth, W. Shtybel, T. Traboulsee.

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Peter Rieckmann serves as Research Chair of the MS Society of Canada; serves on scientific advisory boards for the Germany MS Society, and receives research support from the MS Society of Canada, the US National MS Society, and the CIHR.

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Competing interests

Peter Rieckmann serves on scientific advisory boards for Merck Serono, Novartis, Teva Pharmaceutical Industries Ltd., and Bayer-Schering Pharma; serves on the editorial advisory board of Therapeutics in Neurology; has received speaker and/or consulting honoraria from Bayer-Schering Pharma, Biogen Idec, Merck Serono, Novartis, and Teva Pharmaceutical Industries Ltd.

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Table

Kaplan-Meier estimates and multivariable hazard ratios of time to and age at secondary progression by potential risk factors

Factor	n	Median time to SPMS (95% CI)	p	Time to SPMS Hazard ratio** (95% CI)	p	n	Median age at SPMS (95% CI)	p	Age at SPMS Hazard ratio** (95% CI)	p
Overall	5162	21.4, 20.6 - 22.2	---			5169	53.7, 53.1 - 54.3			
Gender										
Women	3792	22.7, 21.8 23.7	< 0.0005	1.0 (reference)	<0.0005	3796	54.4, 53.7 - 55.2	<0.0005	1.0 (reference)	<0.0005
Men	1370	17.1, 15.4 - 18.7		1.43, 1.30 – 1.58		1373	51.2, 49.9 - 52.5		1.43, 1.30 – 1.58	
Age at onset/ years*						*				
<30	2533	24.9, 23.8 - 26.0	< 0.0005	1.05, 1.04 – 1.06	<0.0005	2533	48.6, 47.6 - 49.6	<0.0005	---	---
30 to <40	1680	18.5, 17.3 - 19.7				1680	52.8, 51.7 - 53.8			
40 to <50	761	12.2, 11.0 - 13.4				761	56.5, 54.9 - 58.0			
50 and above	188	8.1, 7.0 - 9.3				188	64.8, 63.0 - 66.5			
Onset symptoms										
Motor										
absent	4131	22.0, 21.1 - 22.9	< 0.0005	1.0 (reference)	0.02	4137	54.1, 53.3 - 54.9	0.31	1.0 (reference)	0.004
present	1031	18.6, 17.1 – 20.0		1.03 – 1.31		1032	52.9, 51.7 - 54.1		1.12, 1.06 – 1.35	
Sensory										
absent	2597	22.7, 21.8 - 23.5	0.22	1.0 (reference)	0.95	2603	53.4, 52.6 - 54.1	0.29	1.0 (reference)	0.68
present	2565	23.1, 21.9 - 24.2		1.0, 0.89 – 1.13		2566	54.7, 53.3 - 55.2		1.03, 0.91 – 1.16	
CAB										
absent	4511	23.1, 22.3 - 23.9	0.60	1.0 (reference)	0.3	4518	53.9, 53.2 - 54.5	0.18	1.0 (reference)	0.15
present	651	22.0, 20.5 - 23.6		1.08, 0.93 – 1.26		651	52.8, 50.6 - 54.9		1.12, 0.96 – 1.30	
ON										
absent	4251	21.4, 20.5 - 22.3	0.71	1.0 (reference)	0.34	4258	53.9, 53.2 - 54.6	0.02	1.0 (reference)	0.15
present	911	21.4, 19.9 – 23.0		1.08, 0.93 – 1.25		911	52.6, 51.1 - 54.2		1.03, 0.96 – 1.30	

CAB: cerebellar, ataxia or brainstem; ON: optic neuritis; * Seven patients for whom date at disease onset was unknown were excluded from all analyses of time to SPMS and from the analysis of the association between age at onset and age at SPMS. ** derived from Cox regression analyses

\$ Hazard ratios represent a per year increase in risk

Figure Legends

Figure 1

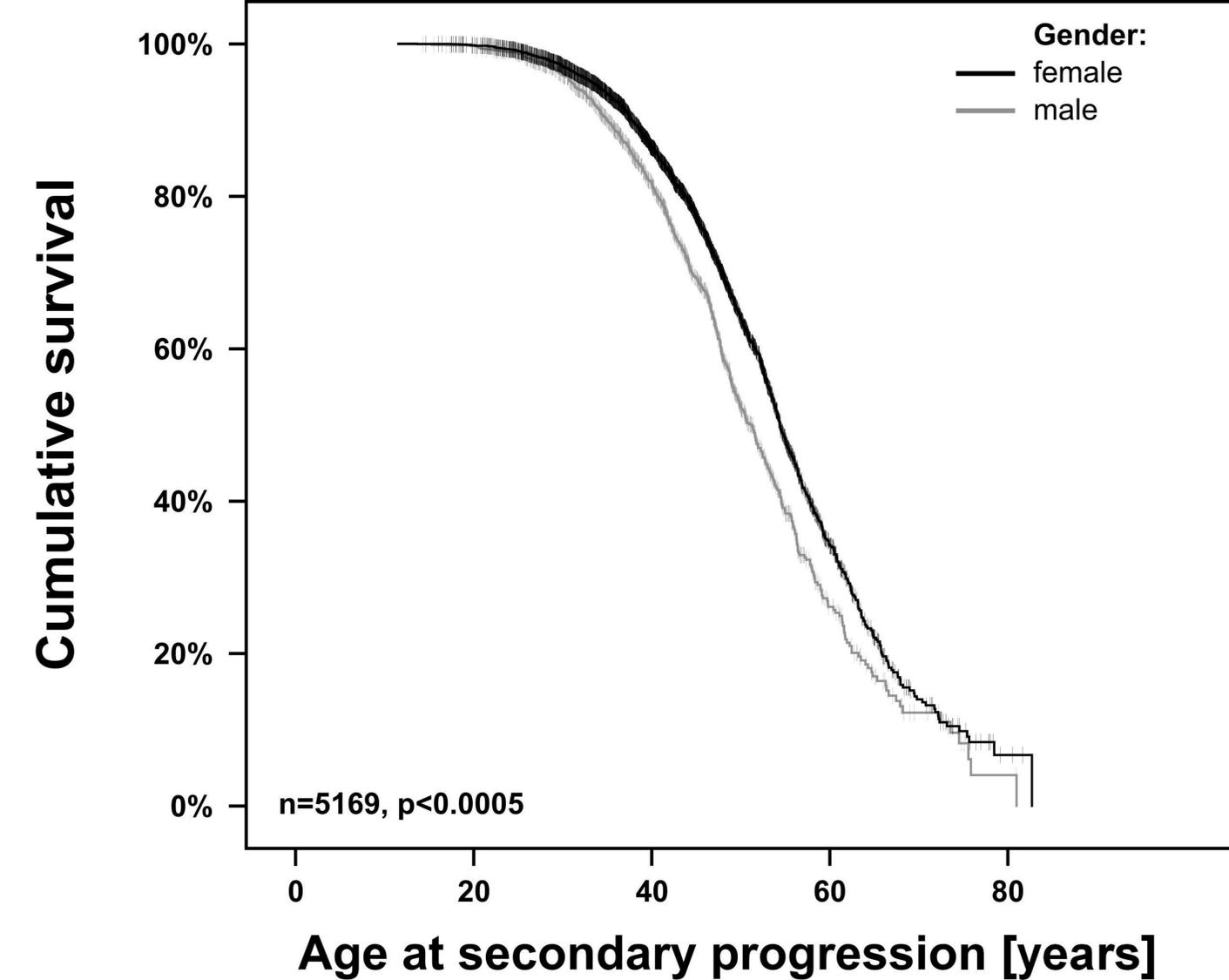
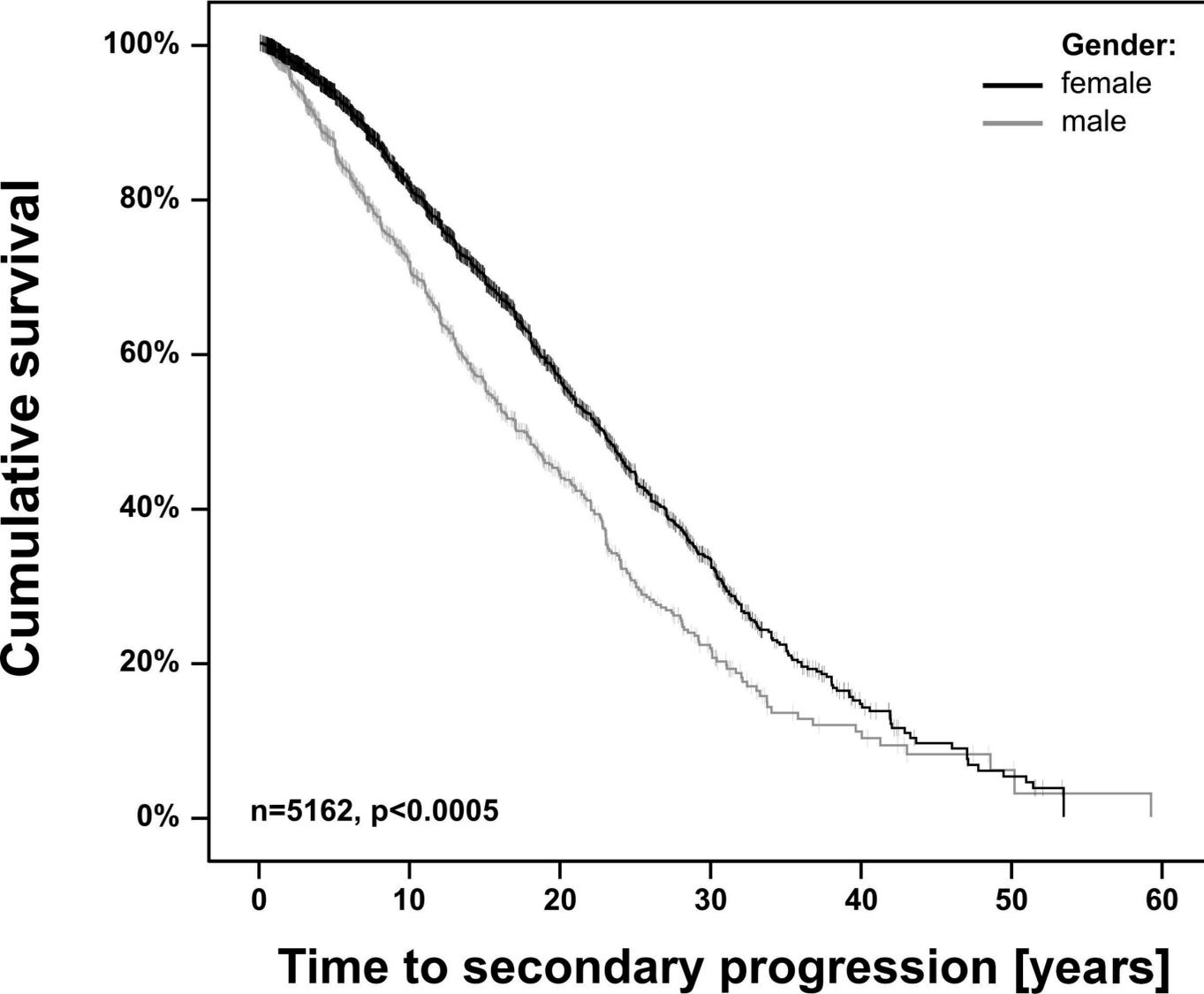
Kaplan-Meier curves showing time to (left) and age at (right) secondary progression by gender.

Figure 2

Kaplan-Meier curves showing time to (left) and age at (right) secondary progression by motor onset symptoms.

Figure 3

Kaplan-Meier curves showing time to (left) and age at (right) secondary progression by age at disease onset.

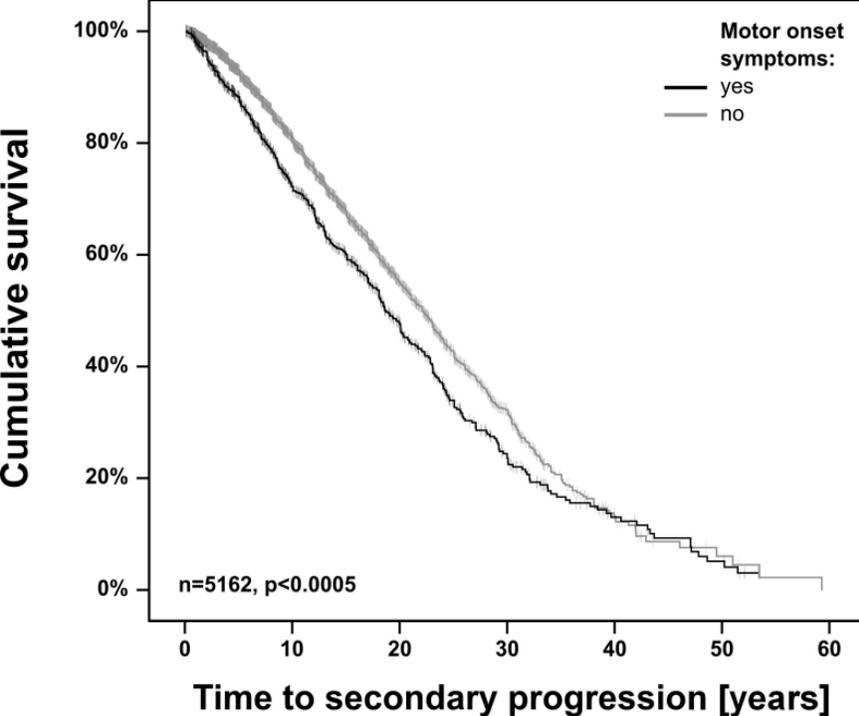


Patients at risk of secondary progression at selected time points:

Gender:	0	10	20	30	40	50	60	Censored patients:
female	3792	1841	672	182	32	7	0	2552 (67%)
male	1370	580	199	52	13	3	0	790 (58%)

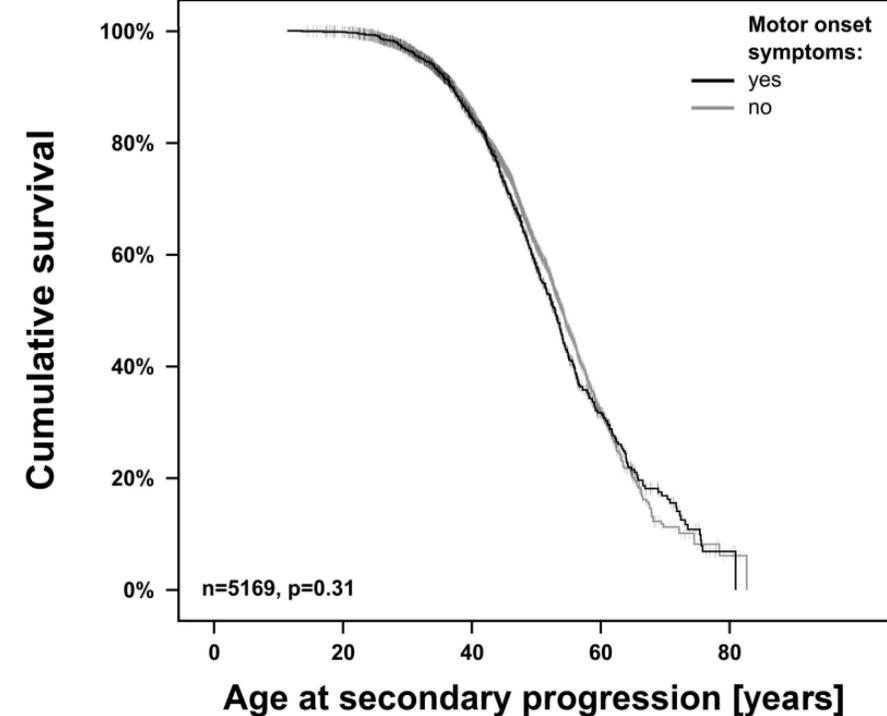
Patients at risk of secondary progression at selected time points:

Gender:	0	20	30	40	50	60	70	80	Censored Patients:
female	3796	3769	3380	2246	910	196	36	3	2555 (67%)
male	1373	1363	1218	800	304	69	12	2	793 (58%)



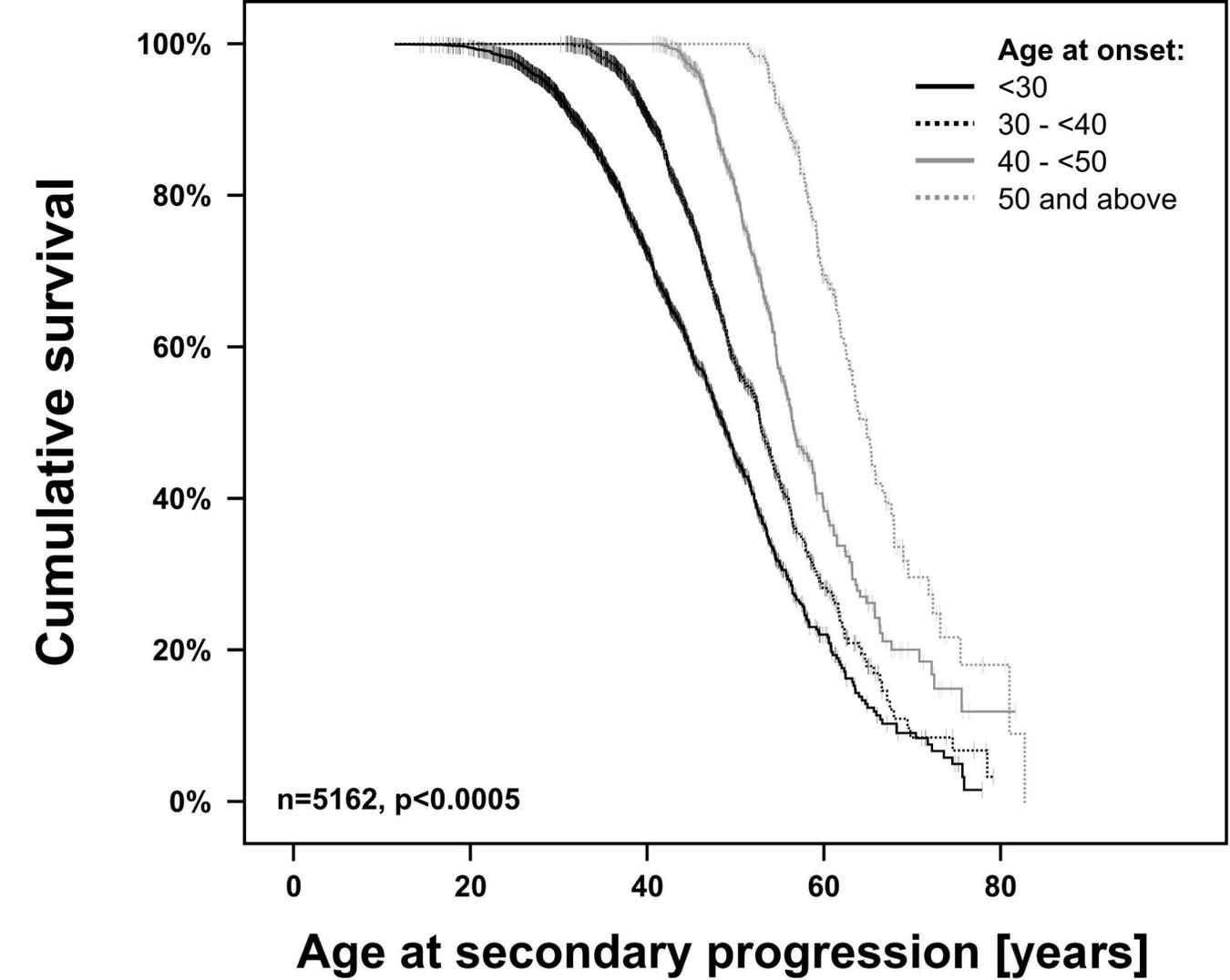
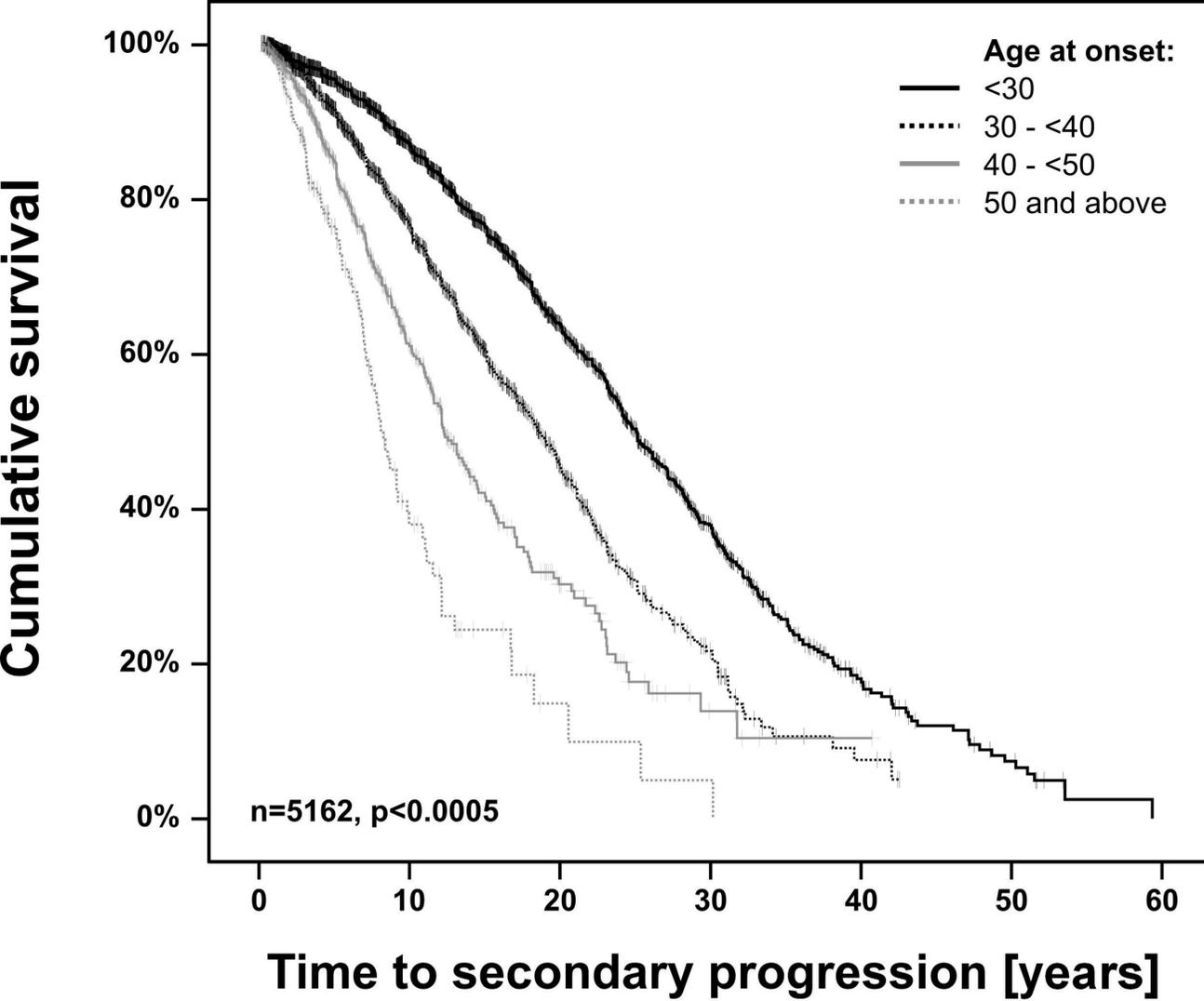
Patients at risk of secondary progression at selected time points:

Motor onset symptoms:	0	10	20	30	40	50	60	Censored patients:
yes	1031	483	198	58	19	6	0	555 (54%)
no	4131	1938	673	176	26	4	0	2787 (67%)



Patients at risk of secondary progression at selected time points:

Motor onset symptoms:	0	20	30	40	50	60	70	80	Censored Patients:
yes	1032	1022	936	675	316	92	26	3	555 (54%)
no	4137	4110	3662	2371	898	173	22	2	2793 (68%)



Patients at risk of secondary progression at selected time points:

	0	10	20	30	40	50	60
Age at onset:							
<30	2533	1514	659	197	39	10	0
30 to <40	1680	698	174	31	5	0	0
40 to <50	761	184	35	5	1	0	0
50 and above	188	25	3	1	0	0	0

Censored patients:

<30	1635 (65%)
30 to <40	1113 (66%)
40 to <50	490 (64%)
50 and above	104 (55%)

Patients at risk of secondary progression at selected time points:

	0	20	30	40	50	60	70	80	Censored Patients:
Age at onset:									
<30	2533	2496	1962	936	299	61	14	0	1635 (65%)
30 to <40	---	---	1680	1158	341	62	7	0	1113 (66%)
40 to <50	---	---	---	761	385	64	13	2	490 (64%)
50 and above	---	---	---	---	188	78	14	3	104 (55%)