Multiple sclerosis and motherhood choice: an observational study in Portuguese women patients

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Introduction. Multiple sclerosis (MS) is a disabling disease occurring mainly in women of childbearing age. MS may interfere with family planning and motherhood decision.

Aim. To study the influence of MS diagnosis and course of the disease on motherhood decision.

Patients and methods. The cohort of 35 to 45-year-old female patients diagnosed with MS for at least ten years was selected from six Portuguese MS centers. A structured questionnaire was applied to all patients in consecutive consultation days. Clinical records were reviewed to characterize and collect information about the disease and pregnancies.

Results. One hundred women were included; mean age at MS diagnosis was 26.3 ± 5.0 years; 90% of the participants presented with a relapsing-remitting MS; 57% had no pregnancies after the diagnosis. MS type and number of relapses were not significantly different between women with or without pregnancies after the diagnosis (p = 0.39 and p = 0.50, respectively). Seventy-seven percent of the patients did not have the intended number of pregnancies. Main reasons given were fear of future disability and the possibility of having relapses. Forty-three women considered that pregnancy might worsen MS.

Conclusion. In our population, motherhood choice was unrelated to the MS type and the number of relapses. However, a relevant number of women had fewer pregnancies than those intended before MS diagnosis and believed that pregnancy could worsen the disease. An effort to better inform the patients should be made to minimize the impact of MS diagnosis on motherhood decision.


Introduction

Multiple sclerosis (MS) is a chronic disease affecting young adults, particularly women in childbearing age. In Portugal, the estimated prevalence of MS is 46.3/100,000 [1, 2]. In the past, women with MS were discouraged from becoming pregnant due to the possible negative effect of pregnancy on the course of the disease. In addition, the influence of the disease on the course and outcome of pregnancy (and vice versa) was also unknown for many years.

The Pregnancy in Multiple Sclerosis (PRIMS) study was the first prospective study to demonstrate a reduction in the relapse rate during pregnancy, especially in the third trimester, and a subsequent increase in the first three months after delivery [3]. Moreover, it showed that an increased relapse rate in the pre-pregnancy year and during pregnancy and a high Kurtzke's Disability Status Scale score at the beginning of pregnancy were positively related to the occurrence of a relapse in the post-partum period [3]. Other retrospective and prospective studies helped to reach the consensus that the relapse rate actually declines during pregnancy and increases in the first three months post-partum without greater future disability [4-6]. Even though the most important risk factor for a relapse in the post-partum period is the number of relapses before and during pregnancy, additional risk factors, modifiable or not, need yet to be determined. Besides, as the relapse rate prior to pregnancy is nowadays decreased due to the use of immunomodulatory drugs, the relapse rate prior to pregnancy may not represent the activity of the disease [7].

MS treatment is based on immunomodulatory and immunosuppressive drugs, which have a high patient's adequate adherence [8]. However, evidence of the safety of these drugs during pregnancy is still lacking [9]. The reduction in relapses during pregnancy is about the same obtained with the existing drugs. Although women stop MS therapy during pregnancy, the disease is not expected to worsen [10].
Concerning obstetric and neonatal characteristics, no major complications have been described in women with MS [9,11]. In delivery there were reports of an increase of cesarean labor and instrumental labor (caused by a prolonged second stage) but without further complications. In neonates although a higher percentage of low birth weight was observed no major differences were found in Apgar scores, birth defects or birth mortality [9].

As a chronic and disabling disease, MS might have an impact on family planning, especially with respect to the motherhood decision of female patients. However, the literature about this topic is scarce. Few descriptive studies tried to identify the main concerns of women with MS when considering motherhood decision. Uncertainty about the course of MS, the effect of pregnancy on the disease course and conflicting advice were some of the concerns identified [12-14]. A recently published Canadian study concluded that the most common MS-related reasons for not becoming pregnant were: symptoms interfering with parenting, fear of burdening the partner or fear of children inheriting MS [15]. There was also evidence that women needed more information about the effect of MS on pregnancy and the effect of pregnancy on the progression of MS [13].

The aim of this study was to document the influence of MS diagnosis and course of the disease on motherhood decision in a population of Portuguese female patients with MS, which is an issue scarcely addressed in the literature.

**Patients and methods**

We conducted an observational multicentre study in women with MS, involving six hospital MS centers that cover a large geographic area of Portugal, in 2012. The recruitment took place in consecutive MS consultation days, during 3 months (May to July 2012).

The population of the study was selected according to the following inclusion criteria: age between 35 and 45 years; diagnosis of definite MS for at least 10 years according to the revised McDonald criteria from 2005 [16]. Women with co-morbidities imposing additional disability, those taking chronic treatments with teratogenic potential and those who were hysterectomized were excluded from the study. All patients gave written informed consent after the approval by the Ethics Committees, Boards of the participating centres and National Commission for Data Protection was obtained.

A standardized questionnaire including social and demographic data (date of birth, marital status,
educational level and employment status), medical past conditions, MS data (year of clinical onset and definite diagnosis, clinical form, prior and current treatments, number of relapses, relapses occurring during the pregnancies and progression of the disease) and reproductive history (intended and actually occurred pregnancies before and after MS diagnosis, specific pregnancies data and type of delivery) was applied to all patients.

From patients’ answers and clinical records, the authors retrospectively filled out electronic data-collection forms. Data were anonymously transferred to the database to safeguard the privacy of the patients.

Descriptive statistics were calculated for patient data (mean ± standard deviation, minimum, median, maximum for continuous variables, and percentages for categorical variables). Variables were compared using the Mann-Whitney and the Chi-square test as appropriate, and statistical significance was set at $p < 0.05$. Analysis was performed using SPSS software, version 15.0.

Results

Demographics and MS history

The sample was composed by 100 women with MS, evenly recruited in the participating hospitals. The mean age was 40.0 ± 3.3 years and the mean number of school-attended-years was 12.0 ± 4.4. The majority of women was married (61%) and employed (52%). Mean age at the time of MS diagnosis was 26.3 ± 5.0 years. Ninety percent of patients presented a relapsing-remitting MS, 8% had secondary progressive course and 2% suffered from primary progressive MS. The total number of relapses the year before pregnancy was quite variable, with a median of two.

Reproductive history

At the time of MS diagnosis, 57% of women had never been pregnant and 28% had already had the intended number of children. Before MS diagnosis, 62% of women had no children, 28% had already had one child, 8% had had two children and 2% had had three children. Fifty-seven percent of the participants had no pregnancies after MS diagnosis, 31% had one pregnancy, 11% had two and one percent had three pregnancies. Seven percent tried to get pregnant and stopped the MS medication without the physician’s knowledge (in most situations the women and the physicians discussed if the medication should be stopped or changed).

The characteristics of the first pregnancy after MS diagnosis are described in table I. Other pregnancies are not presented because no relevant differences between them and the first pregnancy were observed. We perceived that 93% of the pregnancies were well succeeded (7% had spontaneous miscarriages, which is a smaller prevalence than that of the general population). We also found that during pregnancy no relapses occurred and in the three months post-partum most of the women were relapse-free.

For the women who breastfed (72%) no additional co-morbidity, namely relapses, was observed during this period.

History of motherhood decision

Seventy-seven percent of the patients (53 out of 69 patients) reported that the number of intended pregnancies changed after MS diagnosis (28 women who had already had the intended number of pregnancies before diagnosis and 3 who did not answer were excluded from this analysis). Among these 77% of patients, the number of intended pregnancies decreased in 98% and increased in 2% (one patient). Concerning this matter, we decided to place these women in two different groups, according to their academic qualifications: in one group women had attended school less than nine years and in the other group women had studied more than nine years. There were no significant statistical differences between them ($p = 0.26$), both groups showing a decrease in the number of intended pregnancies.

The main reasons presented for having fewer pregnancies than previously planned were fear of future disability and the possibility of having relapses (Table II).

The number of relapses, MS type and therapies used were not significantly different between women with or without pregnancies after MS diagnosis (Table III).

When asked about the impact of MS on pregnancy, 81% of the patients considered that MS does not affect pregnancy; 70% thought that it does not affect childbirth; 80% believed that the disease causes no fetal malformations. Despite the fact that most considered that MS has no effect on pregnancy and pregnancy does not influence the disease course, still 43% of participants believed that pregnancy might worsen the disease.

Among women who got pregnant after MS diagnosis, 35% decided to delay the pregnancy due to
the disease and its possible consequences, namely due to fear of future disability.

Discussion

Motherhood decision in patients with MS is scarcely addressed in the literature, despite the higher frequency of this disease in childbearing age females. On the other hand, although in the last few years a lot of information has been gathered about MS pathophysiology and clinical course, patients still have many doubts concerning pregnancy, breastfeeding, and their impact in the disease course.

To the best of our knowledge this is the first observational study that addresses the impact of MS on motherhood choice and family planning in female patients, in Europe.

Our sample may be considered to be representative of the female Portuguese MS population, because we included MS women from different regions of the country, with different levels of education and distinct economic and marital status. We decided to study only female patients aged 35 to 45 years and with at least ten years of MS diagnosis, taking into account that women with that age-span would have already had the possibility to become pregnant and could also have enough years of disease evolution, thus being able to have a better concept of the present and future burden that the disease might have upon their lives.

Our study showed that a large percentage of women (77%) changed the number of intended pregnancies (decided to have fewer children) after MS diagnosis. Although not statistically significant, these results suggest important implications of the disease on motherhood decision as well as an impact of diagnosis in these women life planning.

The main reasons pointed out for having fewer children are the possibility of having relapses and fear of future disability. These results don't fit those recently reported by Alwan et al who concluded that the most common MS-related reasons for not becoming pregnant are: symptoms interfering with parenting, fear of burdening the partner or fear of children inheriting MS [15]. This may be explained by epidemiological and cultural differences between Canadian and Portuguese populations. Furthermore, Alwan et al study lacked data on the subtypes of MS, disease duration and disease severity. Also, there was a broad spectrum of age of the patients included, therefore becoming a very heterogeneous group of study. However, both studies denoted a fear about the future.

Borisow et al stated that psychological factors should not be underestimated in family planning, especially in sexual contacts because factors like bladder or bowel dysfunction, inserted catheters, or fear of partner disapproval are relevant features that contribute to diminished sexual interest [14]. Although we did not study these particular issues we consider them to be of special importance.

Our results are in accordance with those previously reported by Twork et al who found no differences regarding demographic data between female patients having children and childless MS women, namely in what concerns academic qualifications, addressed in our study [17]. However, in opposition to our results, Twork et al also found these women to be more commonly on a relationship and unemployed. It has been shown that there are some differences in living conditions, quality of life and coping behavior between mothers and childless women with MS, but not enough to confirm the hypothesis that a better quality of life and coping behavior would be visible in MS mothers.

Regarding MS history in the group with children, we found that the number of pregnancies af-
The diagnosis does not correlate with type of disease, number of relapses or immunomodulating therapies. Moreover, 35% of women who decided to get pregnant after MS diagnosis delayed this decision because of the disease itself. In this population, pregnancies and deliveries after MS diagnosis were devoid of complications and the majority of patients decided to breastfeed, in accordance to what is described in the literature.

Our study revealed that patients in general have the knowledge that MS does not influence pregnancy, delivery and newborn development. These results are in consonance with those of Albrecht et al [18]. However, an important number of patients still believe that pregnancy is harmful to MS and worsens the prognosis of the disease. In fact, nowadays information is easily available worldwide but this does not mean that patients are fully aware of the implications of the disease or understand the information that is being given to them.

Concerning breastfeeding, the main question in deciding whether or not to breastfeed is the fear of postpartum relapses that has been described to be higher in this period. Therefore many MS women decide not to breastfeed in order to initiate immunomodulating therapies to avoid postpartum relapses. This fact may explain why so many patients still believe that pregnancy worsens the disease. In a recent study it was concluded that exclusive breastfeeding, due to the consequent hormonal pattern that ensues, has a protective effect in decreasing the number of relapses [7]. However, there are no guidelines regarding this subject and the options should be discussed case by case.

The main limitation of our study is the possible memory bias, as not only a transversal but also a retrospective collection of data was done. Our sample size limits statistical analysis, but a possible explanation for that was our age-span inclusion criterion.

The present study discloses the need of a general effort to inform MS women in order to reduce the negative impact of MS diagnosis on reproductive issues. We believe that it is essential not to discourage pregnancy in these patients and to accurately inform them. Risks and benefits of breastfeeding should also be discussed with the patients, as well as the concern of concomitantly resuming or not MS therapies. We also believe that prevention of postpartum relapses poses a challenge for the future: a decrease in relapse rate in this period could increase the confidence of MS women and encourage them to fulfill their desire to have a family.

References

Esclerosis múltiple y decisión de la maternidad: estudio observacional en pacientes portuguesas

Introducción. La esclerosis múltiple (EM) es una enfermedad incapacitante que afecta mayoritariamente a mujeres en edad fértil. La EM puede alterar el deseo de crear una familia y concebir hijos.

Objetivo. Estudiar la influencia del diagnóstico de la EM y de su evolución sobre la decisión de ser madre.

Pacientes y métodos. Se seleccionó una cohorte integrada por pacientes de 35-45 años diagnosticadas de EM desde hacía por lo menos 10 años que eran atendidas en seis centros portugueses. Las participantes respondieron a un cuestionario estructurado en días de consulta consecutivos. Se revisaron las historias clínicas para caracterizar y recabar información sobre la enfermedad y los embarazos.

Resultados. Participaron 100 mujeres; la media de edad en el momento del diagnóstico de la EM era de 26,3 ± 5,0 años; el 90% de las participantes presentaba la forma remitente recurrente; el 57% de las pacientes no se habían quedado embarazadas después del diagnóstico. El tipo de EM y el número de recidivas no difirieron de manera significativa entre las mujeres que habían concebido después del diagnóstico y las que no \((p = 0,39 \text{ y } p = 0,50, \text{ respectivamente})\). El 77% no había tenido el número de hijos deseado. Los principales motivos aducidos fueron el temor a la incapacidad futura y la posibilidad de sufrir recidivas. Cuarenta y tres mujeres creían que el embarazo podía agravar la EM.

Conclusión. En la población del estudio, la decisión de ser o no ser madre no guardó relación con el tipo de EM ni con el número de recidivas. No obstante, un número relevante de mujeres tuvieron menos embarazos de los que habían deseado antes de ser diagnosticadas y pensaban que la gestación podía empeorar la enfermedad. Sería conveniente mejorar la información que reciben estas pacientes a fin de minimizar el impacto del diagnóstico de la EM en la decisión de ser madre.