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Highlights

- Head-to-head comparison of the effectiveness of natalizumab vs fingolimod
- Pooled data from three large international registries
- Uniform inclusion criteria and analyses

Journal Pre-proof

Title:**The effectiveness of natalizumab vs fingolimod – a comparison of international registry studies**

Authors: Johanna B Andersen*, Sifat Sharmin *, Mathilde Lefort *, Nils Koch-Henriksen, Finn Sellebjerg, Per Soelberg Sørensen, Claudia C Pflieger, Peter V Rasmussen, Michael B Jensen, Jette L Frederiksen, Stephan Bramow, Henrik K Mathiesen, Karen I Schreiber, Dana Horakova, Eva K Havrdova, Raed Alroughani, Guillermo Izquierdo, Sara Eichau, Serkan Ozakbas, Francesco Patti, Marco Onofri, Alessandra Lugaresi, Murat Terzi, Pierre Grammond, Francois Grand'Maison, Bassem Yamout, Alexandre Prat, Marc Girard, Pierre Duquette, Cavit Boz, Maria Trojano, Pamela McCombe, Mark Slee, Jeannette Lechner-Scott, Recai Turkoglu, Patrizia Sola, Diana Ferraro, Franco Granella, Vahid Shaygannejad, Julie Prevost, Olga Skibina, Claudio Solaro, Rana Karabudak, Bart V Wijmeersch, Tunde Csepany, Daniele Spitaleri, Steve Vucic, Romain Casey, Marc Debouverie, Gilles Edan, Jonathan Ciron, Aurélie Ruet, Jérôme D Sèze, Elisabeth Maillart, Hélène Zephir, Pierre Labauge, Gilles Defer, Christine Lebrun, Thibault Moreau, Eric Berger, Pierre Clavelou, Jean Pelletier, Bruno Stankoff, Olivier Gout, Eric Thouvenot, Olivier Heinzlef, Abdullatif Al-Khedr, Bertrand Bourre, Olivier Casez, Philippe Cabre, Alexis Montcuquet, Abir Wahab, Jean-Philippe Camdessanché, Aude Marousset, Ivania Patry, Karolina Hankiewicz, Corinne Pottier, Nicolas Maubeuge, Céline Labeyrie, Chantal Nifle, Emmanuelle Leray, David A Laplaud, Helmut Butzkueven[^], Tomas Kalincik[^], Sandra Vukusic[^], Melinda Magyari[^]

*, [^] authors contributed equally to the manuscript

Johanna Balslev Andersen, MSc, The Danish Multiple Sclerosis Registry, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen, Denmark

Sifat Sharmin, PhD, CORE, Department of Medicine, University of Melbourne, Melbourne, Australia; Department of Neurology, Royal Melbourne Hospital, Melbourne, Australia

Mathilde Lefort, PhD, Rennes University, EHESP, REPERES – EA 7449, F-35000 Rennes, France; Univ Rennes, CHU Rennes, Inserm, CIC 1414 [(Centre d'Investigation Clinique de Rennes)], F-35000 Rennes, France

Nils Koch-Henriksen, MD, DMSsc, Department of Clinical Epidemiology, Aarhus University Hospital. Aarhus, Denmark

Finn Sellebjerg, MD, PhD, DMSc., The Danish Multiple Sclerosis Center, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen, Denmark

Per Soelberg Sørensen, MD, PhD, DMSc., The Danish Multiple Sclerosis Center, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen, Denmark

Claudia C Pflieger, MD, PhD, Department of Neurology Aalborg University Hospital; Multiple Sclerosis Unit

Peter V Rasmussen, MD, PhD, Aarhus University Hospital, Neurology, PPJ Boulevard, DK-8200 Aarhus N

Michael B Jensen, MD, PhD, Department of Neurology, University Hospital of Northern Sealand

Jette L Frederiksen, MD, PhD, Department of Neurology, Rigshospitalet Glostrup and Institute of Clinical Medicine, University of Copenhagen, Denmark

Stephan Bramow, MD, PhD, Danish Multiple Sclerosis Centre, Dept. of Neurology, Copenhagen University Hospital, Rigshospitalet in Glostrup, 2600 Glostrup, Denmark

Henrik K Mathiesen, MD, PhD, Department of Neurology, Copenhagen University Hospital Herlev, Denmark.

Karen I Schreiber, MD, PhD, The Danish Multiple Sclerosis Center, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen, Denmark

Dana Horakova D, MD, PhD, Department of Neurology and Center of Clinical Neuroscience, First Faculty of Medicine, Charles University in Prague and General University Hospital, Prague, Czech Republic

Eva K Havrdova, MD, PhD, Department of Neurology and Center of Clinical Neuroscience, First Faculty of Medicine, Charles University in Prague and General University Hospital, Prague, Czech Republic

Raed Alroughani, MD, Division of Neurology, Department of Medicine, Amiri Hospital, Sharq, Kuwait

Guillermo Izquierdo, MD, Hospital Universitario Virgen Macarena, Sevilla, Spain

Sara Eichau, MD, Hospital Universitario Virgen Macarena, Sevilla, Spain

Serkan Ozakbas, MD, Dokuz Eylul University, Konak/Izmir, Turkey

Francesco Patti, MD, GF Ingrassia Department, University of Catania, Catania, Italy; Policlinico G Rodolico

Marco Onofri, MD, Department of Neuroscience, Imaging, and Clinical Sciences, University G. d'Annunzio, Chieti, Italy

Alessandra Lugaresi, MD, PhD, Department of Biomedical and Neuromotor Sciences, University of Bologna, Bologna, Italy; IRCCS Istituto delle Scienze Neurologiche di Bologna, Bologna, Italy

Murat Terzi, MD, Medical Faculty, 19 Mayıs University, Samsun, Turkey

Pierre Grammond, MD, CISSS Chaudiere-Appalache, Levis, Canada

Francois Grand'Maison, MD, Neuro Rive-Sud, Quebec, Canada

Bassem Yamout, MD, Nehme and Therese Tohme Multiple Sclerosis Center, American University of Beirut Medical Center, Beirut, Lebanon

Alexandre Prat, MD, PhD, Hopital Notre Dame, Montreal, Canada; CHUM and Universite de Montreal, Montreal, Canada

Marc Girard, MD, Hopital Notre Dame, Montreal, Canada; CHUM and Universite de Montreal, Montreal, Canada

Pierre Duquette, MD, Hopital Notre Dame, Montreal, Canada; CHUM and Universite de Montreal, Montreal, Canada

Cavit Boz, MD, KTU Medical Faculty Farabi Hospital, Trabzon, Turkey

Maria Trojano, MD, Department of Basic Medical Sciences, Neuroscience and Sense Organs, University of Bari, Bari, Italy

Pamela McCombe, MBBS, University of Queensland, Brisbane, Australia; Royal Brisbane and Women's Hospital

Mark Slee, BMBS, PhD, Flinders University, Adelaide, Australia

Jeannette Lechner-Scott, MD, PhD, School of Medicine and Public Health, University Newcastle, Newcastle, Australia; Department of Neurology, John Hunter Hospital, Hunter New England Health, Newcastle, Australia

Recai Turkoglu, MD, Haydarpasa Numune Training and Research Hospital, Istanbul, Turkey

Patrizia Sola, MD, PhD, Department of Neuroscience, Azienda Ospedaliera Universitaria, Modena, Italy

Diana Ferraro, MD, PhD, Department of Neuroscience, Azienda Ospedaliera Universitaria, Modena, Italy

Franco Granella, MD, Department of Medicine and Surgery, University of Parma, Parma, Italy;
Department of Emergency and General Medicine, Parma University Hospital, Parma, Italy

Vahid Shaygannejad, MD, Isfahan University of Medical Sciences, Isfahan, Iran

Julie Prevost, MD, CSSS Saint-Jerome, Saint-Jerome, Canada

Olga Skibina, MD, Monash University, Melbourne, Australia

Claudio Solaro, MD, Department of Neurology, ASL3 Genovese, Genova, Italy; Department of
Rehabilitaiton, ML Novarese Hospital Moncrivello

Rana Karabudak, MD, Hacettepe University, Ankara, Turkey

Bart V Wijmeersch, MD, PhD, Rehabilitation and MS-Centre Overpelt and Hasselt University,
Hasselt, Belgium

Tunde Csepany, MD, Department of Neurology, Faculty of Medicine, University of Debrecen,
Debrecen, Hungary

Daniele Spitaleri, MD, Azienda Ospedaliera di Rilievo Nazionale San Giuseppe Moscati Avellino,
Avellino, Italy

Steve Vucic, MBBS, PhD, Westmead Hospital, Sydney, Australia

Romain Casey, PhD, Service de neurologie, sclérose en plaques, pathologies de la myéline et
neuro-inflammation, Hôpital Neurologique Pierre Wertheimer, Hospices Civils de Lyon, 69677
Lyon/Bron, France; Centre des Neurosciences de Lyon, Observatoire Français de la Sclérose en
Plaques, INSERM 1028 et CNRS UMR5292, 69003 Lyon, France ; Université Claude Bernard
Lyon 1, Faculté de médecine Lyon Est, F-69000 Lyon, France ; Eugene Devic EDMUS
Foundation, 69677 Lyon/Bron, France

Marc Debouverie, MD, PhD, Centre hospitalier régional universitaire de Nancy, Hôpital central, Service de neurologie, Nancy, France

Gilles Edan, MD, PhD, Centre hospitalier universitaire de Rennes, Hôpital Pontchaillou, Service de neurologie, Rennes, France

Jonathan Ciron, MD, Centre hospitalier universitaire de Toulouse, Hôpital Purpan, Service de neurologie inflammatoire et neuro-oncologie, Toulouse, France

Aurélie Ruet, MD, PhD, Centre hospitalier universitaire de Bordeaux, Hôpital Pellegrin, Service de neurologie, Bordeaux, France

Jérôme D Sèze, MD, PhD, Hôpitaux universitaires de Strasbourg, Hôpital de Hautepierre, Service des maladies inflammatoires du système nerveux – neurologie, Strasbourg, France

Elisabeth Maillart, MD, Assistance publique des hôpitaux de Paris, Hôpital de la Pitié-Salpêtrière, Service de neurologie, Paris, France

Hélène Zephir, MD, PhD, Centre hospitalier universitaire de Lille, Hôpital Salengro, Service de neurologie D, Lille, France

Pierre Labauge, MD, PhD, Centre hospitalier universitaire de Montpellier, Hôpital Gui de Chauliac, Service de neurologie, Montpellier, France

Gilles Defer, MD, PhD, Centre hospitalier universitaire de Caen Normandie, Service de neurologie, Hôpital Côte de Nacre, Caen, France

Christine Lebrun, MD, PhD, Centre hospitalier universitaire de Nice, Université Nice Côte d'Azur, Hôpital Pasteur, Service de neurologie, Nice, France

Thibault Moreau, MD, PhD, Centre hospitalier universitaire Dijon Bourgogne, Hôpital François Mitterrand, Service de neurologie, maladies inflammatoires du système nerveux et neurologie générale, Dijon, France

Eric Berger, MD, Centre hospitalier régional universitaire de Besançon, Hôpital Jean Minjoz, Service de neurologie, Besançon, France

Pierre Clavelou, MD, PhD, Centre hospitalier universitaire de Clermont-Ferrand, Hôpital Gabriel-Montpied, Service de neurologie, Clermont-Ferrand, France

Jean Pelletier, MD, PhD, Assistance publique des hôpitaux de Marseille, Centre hospitalier de la Timone, Service de neurologie et unité neuro-vasculaire, Marseille, France

Bruno Stankoff, MD, PhD, Assistance publique des hôpitaux de Paris, Hôpital Saint-Antoine, Service de neurologie, Paris, France

Olivier Gout, MD, Fondation Adolphe de Rothschild de l'œil et du cerveau, Service de neurologie, Paris, France

Eric Thouvenot, MD, PhD, Centre hospitalier universitaire de Nîmes, Hôpital Carémeau, Service de neurologie, Nîmes, France

Olivier Heinzlef, MD, Centre hospitalier intercommunal de Poissy Saint-Germain-en-Laye, Service de neurologie, Poissy, France

Abdullatif Al-Khedr, MD, Centre hospitalier universitaire d'Amiens Picardie, Site sud, Service de neurologie, Amiens, France

Bertrand Bourre, MD, Centre hospitalier universitaire Rouen Normandie, Hôpital Charles-Nicolle, Service de neurologie, Rouen, France

Olivier Casez, MD, Centre hospitalier universitaire Grenoble-Alpes, Site nord, Service de neurologie, Grenoble/La Tronche, France

Philippe Cabre, MD, PhD, Centre hospitalier universitaire de Martinique, Hôpital Pierre Zobda-Quitman, Service de Neurologie, Fort-de-France, France

Alexis Montcuquet, MD, Centre hospitalier universitaire Limoges, Hôpital Dupuytren, Service de neurologie, Limoges, France

Abir Wahab, MD, APHP, Hôpital Henri Mondor, Department of Neurology, F-94000 Créteil, France

Jean-Philippe Camdessanché, MD, PhD, Centre hospitalier universitaire de Saint-Étienne, Hôpital Nord, Service de neurologie, Saint-Étienne, France

Aude Marousset, MD, Centre hospitalier régional universitaire de Tours, Hôpital Bretonneau, Service de neurologie, Tours, France

Ivania Patry, MD, Centre hospitalier sud francilien, Service de neurologie, Corbeil-Essonnes, France

Karolina Hankiewicz, MD, Centre hospitalier de Saint-Denis, Hôpital Casanova, Service de neurologie, Saint-Denis, France

Corinne Pottier, MD, Centre hospitalier de Pontoise, Service de neurologie, Pontoise, France

Nicolas Maubeuge, MD, Centre hospitalier universitaire de Poitiers, Site de la Milétrie, Service de neurologie, Poitiers, France

Céline Labeyrie, MD, Assistance publique des hôpitaux de Paris, Hôpital Bicêtre, Service de neurologie, Le Kremlin-Bicêtre, France

Chantal Nifle, MD, Centre hospitalier de Versailles, Hôpital André-Mignot, Service de neurologie, Le Chesnay, France

Emmanuelle Leray, PhD, Rennes University, EHESP, REPERES – EA 7449, F-35000 Rennes, France; Univ Rennes, CHU Rennes, Inserm, CIC 1414 [(Centre d'Investigation Clinique de Rennes)], F-35000 Rennes, France

David A Laplaud, MD, PhD, CHU de Nantes, Service de Neurologie & CIC015 INSERM, F-44093 Nantes, France; INSERM CR1064, F-44000 Nantes, France

Helmut Butzkueven, MBBS, PhD, Central Clinical School, Monash University, Melbourne, Australia; Department of Neurology, The Alfred Hospital, Melbourne, Australia; Department of Neurology, Box Hill Hospital, Monash University, Melbourne, Australia

Tomas Kalincik, MD, PhD, CORE, Department of Medicine, University of Melbourne, Melbourne, Australia; Melbourne MS Centre, Department of Neurology, Royal Melbourne Hospital, Melbourne, Australia

Sandra Vukusic, MD, PhD, Service de neurologie, sclérose en plaques, pathologies de la myéline et neuro-inflammation, Hôpital Neurologique Pierre Wertheimer, Hospices Civils de Lyon, 69677 Lyon/Bron, France; Centre des Neurosciences de Lyon, Observatoire Français de la Sclérose en Plaques, INSERM 1028 et CNRS UMR5292, 69003 Lyon, France; Université Claude Bernard Lyon 1, Faculté de médecine Lyon Est, F-69000 Lyon, France

Melinda Magyari, MD, PhD, The Danish Multiple Sclerosis Registry, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen, Denmark; The Danish Multiple Sclerosis

Center, Department of Neurology, University of Copenhagen, Rigshospitalet, Copenhagen,
Denmark

Corresponding author:

Johanna Balslev Andersen

Blegdamsvej 9

2100 København

Phone: +45 35 45 78 80

Email: johanna.balslev.andersen@regionh.dk

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ABSTRACT

Background

Natalizumab and fingolimod were the first preparations recommended for disease breakthrough in priorly treated relapsing-remitting multiple sclerosis. Of three published head-to-head studies two showed that natalizumab is the more effective to prevent relapses and EDSS worsening.

Methods

By re-analyzing original published results from MSBase, France, and Denmark using uniform methodologies, we aimed at identifying the effects of differences in methodology, in the MS-populations, and at re-evaluating the differences in effectiveness between the two drugs.

We gained access to copies of the individual amended databases and pooled all data. We used uniform inclusion/exclusion criteria and statistical methods with Inverse Probability Treatment Weighting.

Results

The pooled analyses comprised 968 natalizumab- and 1479 fingolimod treated patients. The on-treatment natalizumab/fingolimod relapse rate ratio was 0.77 ($p=0.004$). The hazard ratio (HR) for a first relapse was 0.82 ($p=0.030$), and the HR for sustained EDSS improvement was 1.4 ($p=0.009$). There were modest differences between each of the original published studies and the replication study, but the conclusions of the three original studies remained unchanged: in two of them natalizumab was more effective, but in the third there was no difference between natalizumab and fingolimod.

Conclusion

The results were largely invariant to the epidemiological and statistical methods but differed between the MS populations. Generally, the advantage of natalizumab was confirmed.

Keywords: Multiple sclerosis; Natalizumab; Fingolimod; Treatment effectiveness; Head-to-head comparison;

1. Introduction

Evidence-based 2018 guidelines (1) for the use of disease modifying drugs (DMDs) in multiple sclerosis (MS) suggest that in patients with relapsing-remitting MS (RRMS) the choice of DMD should be based upon patient characteristics and comorbidities, disease severity, drug safety profile and accessibility of the drug. In RRMS patients with inadequate treatment response it is recommended to switch to a drug with higher efficacy including natalizumab or fingolimod (1). No randomized clinical trial has assessed the comparative efficacy of natalizumab and fingolimod in RRMS patients. Observational studies have shown inconsistent results as to difference in clinical effectiveness in real life settings (2–10) These studies varied in sources of data, sample size, inclusion and exclusion criteria, study design, outcomes, and statistical analyses as well as in the MS populations. This study is based on three published studies from The Danish Multiple Sclerosis Treatment Register (2), the French MS Registry (Observatoire Français de la Sclérose en Plaques) OFSEP (7) and MSBase (8) which led to seemingly discordant results. We hypothesize that these differences are primarily driven by differences in the studied populations rather than the used analytical methodology. This study is the first of a series of three studies which will replicate and combine the observations from the original three analyses, then quantify the effect of clinical and demographic differences between the MS populations on the observed effects of the two DMDs with high efficacy, and, lastly in detail explore the effect of statistical methodology. The present study may by its study design adjust the outcomes of the original studies as well as the robustness and internal validity. Differences in the studied samples may influence external validity and reflect variability in reported response to treatment in different patient subgroups (11).

Kalincik et al. (7) used data from the MSBase (12) and reported a higher effectiveness of natalizumab compared to fingolimod in reducing the annualized relapse rate (ARR) and sustained disability improvement in RRMS. Barbin et al. (8), using the French Multiple Sclerosis Registry (OFSEP) (13), supported the finding of higher effectiveness of natalizumab compared to fingolimod on reducing the proportion of relapse-free patients. Conversely, Koch-Henriksen et al. (2) analysed data from the nationwide Danish Multiple Sclerosis Treatment Register and found no significant differences when comparing the effectiveness of natalizumab and fingolimod in any of the clinical endpoints.

The three original head-to-head studies represented different MS populations, and they differed to some extent in inclusion/exclusion criteria, and there may have been local differences in how clinicians prescribed the two preparations.

The purpose of this study was to compare disease activity after switch from first-line therapy to natalizumab or fingolimod using pooled and extended data from the three databases and to replicate their differences when using uniform methodology.

2. Material and Methods

2.1 Study design

This study is a historical cohort study of prospectively collected data, recorded in three large MS registries, OFSEP, DMSR and MSBase (2,7,8).

The study consists of two parts: 1) the replication *study* in which the same data were subjected to new and uniform selection criteria, definition of endpoints, and statistical analyses, and 2) the *pooled study* in which data from the three cohorts were pooled and subjected to the same methods and analyses as used for the replication study (see below).

Data in MSBase and DMSR had been updated with more patients and longer follow-up presented in this study, whereas the data provided from OFSEP for this study was the same as used in the original study. Table 1 shows the differences in the inclusion/exclusion criteria, statistical methods and clinical endpoints used in the three original studies and in the pooled study.

Table 1. Differences in methods for the original studies and the present replication study and pooled study

	MSBase 2015 (7)	OFSEP 2016 (8)	DMSTR 2016 (2)	Present replication study and pooled study
Number of centers	66	27	14	174
Design	Cohort, longitudinal data.	Cohort, longitudinal data.	Cohort, longitudinal data.	Cohort, longitudinal data.
Inclusion/exclusion	Relapse or disability worsening within 6 months before start; No previous participation in randomized trials	RRMS. Age 18 to 65. EDSS \leq 5.5	RRMS \geq 1 relapse within 12 months before start or, if treatment naïve, else \geq 2 relapses with	RRMS; \geq 90 days of DMD first-line treatment prior to study medication; \geq 3 months of study treatment; no

			residual symptoms	previous participation in randomized trials;
Propensity score: Matching or weighting	Matched by propensity score based on age; sex; number of relapses in 6 or 12 months EDSS; Disease activity under previous treatment (relapses, EDSS-worsening or both). MRI data available from a proportion of patients, multiple imputation used.	Weighted by inverse probability of treatment (IPTW) based on sex; number of relapses in previous year; EDSS; hospital; Gd-enhancing lesions on MRI.	Matched by propensity score based on sex; age; being treatment naïve; ARR during previous treatment; MSSS (derived from EDSS) with ignoring unmatchable cases. No MRI data available for matching.	Weighted by stabilized inverse probability of treatment (IPTW) based on sex; age; MS duration; EDSS; #relapses in 12 months; disease activity in 12 months (relapses, EDSS-worsening or both).
Statistical analyses	Adjusted paired proportional hazards models and weighted negative binomial model	<i>t</i> test; Wilcoxon test; chi-square.	Generalized linear models assuming negative binomial distribution; Kaplan-Meyer analysis; Mann Whitney U test; Pearson chi-square.	Negative binomial model; Cox proportional Hazards; Anderson-Gill model
Follow-up	December 2013	July 2014	October 2015	-
Clinical study endpoints	Freedom from clinical relapses. ARR. Disability worsening. Disability improvement.	Proportion of patients with at least one on-study relapse in the first year and at two years.	ARR; proportion of patients remaining free of relapses; time to 1 st relapse; proportion with worsening or improving EDSS.	ARR; relapse rate ratio; time to 1 st relapse; increase in EDSS sustained for 6 months; improvement of EDSS sustained for 6 months

2.2 Data sources

The MSBase Registry is a large international collaboration database with patient records from 129 participating MS centres located in 34 different countries (12). The MSBase longitudinally collects data most from tertiary MS centres. The inclusion criteria for the MSBase is a diagnosis of MS or clinically isolated syndromes based on the 2005 or 2010 revised McDonald Criteria. The MSBase protocol stipulates update on the minimum data set at least annually, although this was not a required inclusion criterion. The median inter-visit interval is 5 months. The data entry portal was either iMed MS patient record system or the MSBase online data system. An operationalised data quality procedure was applied(14).

The Danish Multiple Sclerosis Registry (DMSR) (15) was founded in 1956. It comprises data on all patients diagnosed with - or suspected of having - MS by a neurologist. The diagnostic criteria applied before 2005 were the Poser criteria (16) and thereafter the current version of the McDonald criteria (17). Since 1996, acquisition of relapses and Expanded Disability Status Scale (EDSS) scores and of the clinical characteristics has been performed in all DMD-treated patients at baseline, after 3 months and thereafter every 6 months during the clinical follow-up with mandatory notification of the DMSR due to reimbursement. Only departments of neurology in public hospitals are authorized to prescribe and dispense the DMDs to the patients, and the treating neurologists are joined in a network enabling use of uniform guidelines. All 14 Danish MS centres contribute, and data collection is done through an online data collection platform, which enables continuous completion of data improving its completeness and validity.

The Observatoire Français de la Sclérose en Plaques (OFSEP) (18) collects information from 40 MS expert centres throughout France, representing more than one half of the French MS population. Clinical data are collected during routine follow-up visits, usually at least once a year, retrospectively at the first visit and prospectively thereafter. Minimum standardized datasets are recorded through the EDMUS database and synchronised with the OFSEP database at 6-month intervals. OFSEP has implemented a strategy to improve the quality of its data and samples. The EDMUS software has an integrated data verification tool to identify missing or incoherent data. Twice a year, a quality report is sent to all centres, with queries on incoherent data entries. Information documents, data quality indicators, training sessions and audits are displayed.

2.3 Inclusion criteria for the replication study and the pooled study

The inclusion criteria and statistical methods used in the replication study and the pooled study were agreed upon by the three registries. They were: 1) RRMS at commencing study treatment; 2) patients have commenced treatment with either natalizumab or fingolimod for the first time on or after 1st of January 2011 (to ensure accessibility of both drugs in Europe and Australia); 3) continuous treatment with either natalizumab or fingolimod for ≥ 3 months; 4) no prior exposure to immunotherapies with extended effect (mitoxantrone, alemtuzumab, ocrelizumab, daclizumab, rituximab, cyclophosphamide, or cladribine); 5) no prior participation in any interventional randomised controlled trials; 6) exposure to DMD treatment for more than 90 consecutive days within the 12 months immediately prior to commencing natalizumab or fingolimod; 7) sufficient EDSS follow-up (consisting of EDSS recorded 6 months to +1 months of baseline; more than one EDSS assessment recorded on study therapy and more than one EDSS assessment recorded ≥ 6 months later (irrespective of the treatment status at that time)). EDSS scores recorded ≤ 30 days after a prior relapse were ignored. Baseline was defined as the day of initiation of natalizumab or fingolimod. Patients' follow-up was censored at discontinuation of the study therapy or the last recorded follow-up. The numbers of eligible patients are presented in Figure 1 and Table 2. All three registries used equivalent definitions of the EDSS score as derived from functional score systems described by Kurtzke (19). Relapses were defined as occurrence of new or worsening neurological symptoms persisting for at least 24 hours in the absence of fever and infection (17) and onset year as the year of first experienced symptom of MS. MSBase and OFSEP had the date and year of onset registered, whereas only year of onset was recorded for some patients in the DMSR (if missing, date was set to 15/6 in the recorded year of onset).

Figure 1. Flowchart presentation of the included patients in the pooled cohort.

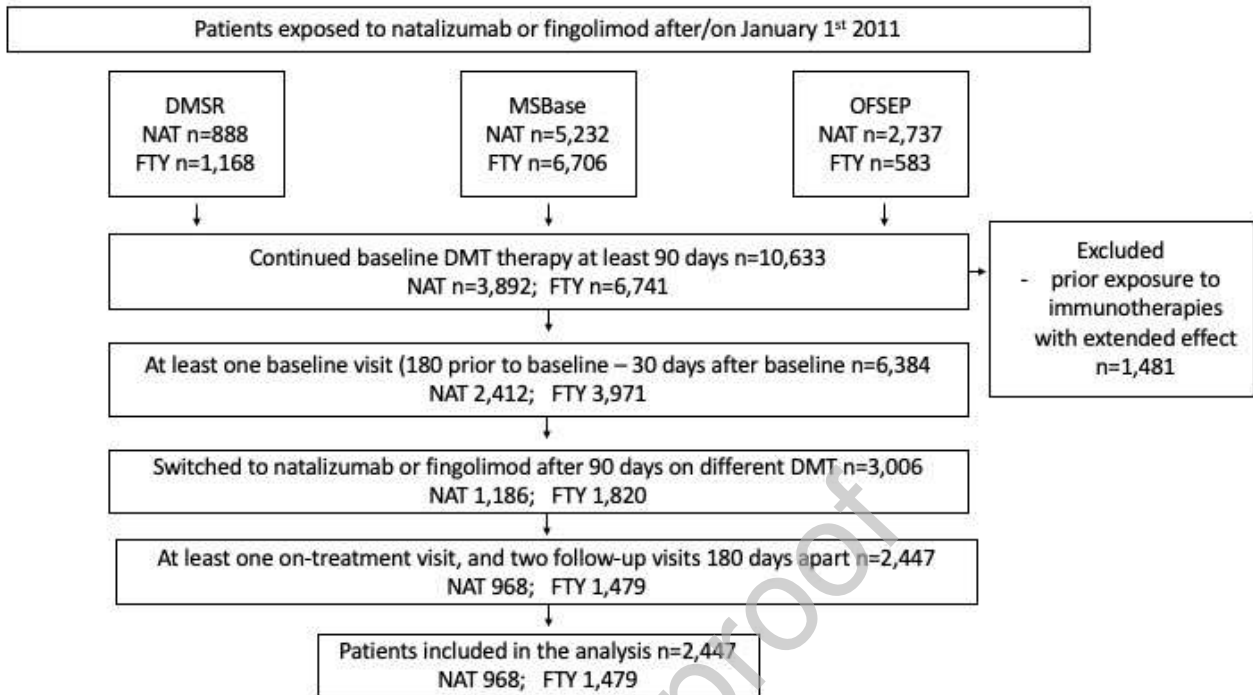


Table 2. Baseline characteristics of the pooled cohort and the three individual cohorts contributing to it before and after stabilized inverse probability of treatment weighting (sIPTW)

	Before sIPTW			After sIPTW		
	Natalizumab	Fingolimod		Natalizumab	Fingolimod	
<i>MSBase (international)</i>	N = 410	N = 792	SMD*			SMD*
Female %	73.2	72.9	0.071	72.1	72.8	0.018
Mean age at baseline, years (sd)	36.2 (10.3)	38.1 (9.6)	0.183	37.6 (11.1)	37.6 (9.5)	0.004
Mean MS duration, years (sd)	8.1 (6.6)	9.2 (7.2)	0.151	9.1 (7.9)	8.9 (7.0)	0.025
Mean EDSS at baseline, (sd)	2.9 (1.52)	2.28 (1.51)	0.411	2.66 (1.42)	2.49 (1.58)	0.114
Mean nr of relapses 12 months prior to baseline	1.35 (0.96)	0.94 (0.84)	0.447	1.14 (0.88)	1.07 (0.90)	0.079
Mean nr of previous DMDs, (sd)	1.59 (0.80)	1.67 (0.88)	0.106	1.67 (0.88)	1.67 (0.87)	0.008
Disease activity 12 months prior to baseline%						
None	13.66	27.02	0.337	18.92	22.86	0.097
Worsening	3.90	5.56	0.078	4.36	5.25	0.042
Relapse	51.95	45.2	0.135	49.06	46.16	0.058
Relapse and worsening	30.49	22.22	0.188	27.66	25.73	0.044
<i>DMSR (Denmark)</i>	N = 399	N = 581				
Female %	70.9	65.1	0.126	67.3	67.4	0.017
Mean age at baseline (sd)	39.2 (9.5)	40.4 (9.2)	0.131	39.9 (9.5)	39.9 (9.3)	0.001
Mean MS duration, years (sd)	8.8 (7.4)	8.9 (6.7)	0.025	8.8 (7.6)	8.8 (6.6)	0.003
Mean EDSS at baseline	2.90 (1.59)	2.63 (1.46)	0.171	2.73 (1.56)	2.74 (1.50)	0.005
Mean nr of relapses 12 months prior to baseline	0.76 (0.84)	0.71 (0.75)	0.072	0.73 (0.80)	0.73 (0.78)	0.001
Mean nr of previous DMDs	1.61 (0.95)	1.51 (0.76)	0.117	1.56 (0.87)	1.55 (0.79)	0.005
Disease activity 12 months prior to baseline%						
None	25.81	30.98	0.115	28.99	28.97	0.001
Worsening	18.30	13.77	0.124	15.36	15.41	0.002
Relapse	28.32	30.98	0.058	30.10	30.02	0.002
Relapse and worsening	27.57	24.27	0.075	25.55	25.60	0.001
<i>OFSEP (France)</i>	N = 159	N = 106				
Female %	76.7	73.6	0.073	74.9	76.7	0.042
Mean age at baseline (sd)	37.1(10.2)	39.1 (9.2)	0.198	37.9(10.4)	37.8 (9.5)	0.023
Mean MS duration (years)	8.0 (5.4)	9.8 (6.9)	0.297	8.7 (5.8)	8.6 (6.3)	0.015
Mean EDSS at baseline	2.82 (1.58)	2.61 (1.67)	0.131	2.77 (1.54)	2.85 (1.66)	0.049
Mean nr of relapses 12 months prior to baseline	1.62 (1.07)	0.99 (0.93)	0.623	1.38 (1.06)	1.41 (1.1)	0.029
Mean nr of previous DMDs	1.69 (0.89)	1.58 (0.87)	0.114	1.66 (0.89)	1.68 (0.9)	0.024
Disease activity 12 months prior to baseline%						
None	6.92	25.47	0.520	14.09	14.03	0.002
Worsening	3.77	7.55	0.164	5.12	5.10	0.001
Relapse	45.91	44.34	0.032	45.29	44.31	0.019
Relapse and worsening	43.4	22.64	0.453	35.49	36.56	0.022
<i>Pooled cohort (MSBase+DMSR+OFSEP)</i>	N = 968	N = 1479				
Female %	72.8	69.8	0.066	70.3	71.0	0.015
Mean age at baseline (sd)	37.6 (10.0)	39.1 (9.5)	0.150	38.8 (10.5)	38.6 (9.5)	0.022
Mean MS duration, years (sd)	8.4 (6.8)	9.1 (6.9)	0.110	9.0 (7.8)	8.9 (6.8)	0.026
Mean EDSS at baseline	2.89(1.56)	2.44 (1.51)	0.289	2.71 (1.50)	2.65 (1.57)	0.036
Mean nr of relapses 12 months prior to baseline	1.15 (0.99)	0.85 (0.82)	0.327	0.99 (0.91)	0.98 (0.91)	0.017
Mean nr of previous DMDs	1.61 (0.88)	1.6 (0.84)	0.011	1.61 (0.84)	1.62 (0.84)	0.004

Disease activity 12 months prior to baseline%						
None	17.56	28.47	0.261	23.12	24.29	0.028
Worsening	9.81	8.92	0.031	8.97	9.41	0.015
Relapse	41.22	39.55	0.034	40.64	39.51	0.023
Relapse and worsening	31.40	23.06	0.188	27.27	26.79	0.011

*Standardized mean difference (difference as fraction of the pooled standard deviation)

2.4 Study endpoints of the replication study and the pooled study

ARR was calculated at the individual level as the number of relapses divided by annualized observed person-time from baseline to treatment discontinuation or censor date in years.

Time to first relapse was calculated as the time from baseline to the date of start of a first relapse.

Worsening of EDSS was defined as an increase by ≥ 1.5 step sustained for 6 months if EDSS at baseline was 0; or ≥ 1 step if EDSS at baseline was ≥ 1 and ≤ 5.5 ; and ≥ 0.5 step if EDSS at baseline was ≥ 6 . Improvement of EDSS was defined as a decrease by ≥ 1 EDSS step if EDSS at baseline was ≤ 6 and ≥ 1.5 ; ≥ 0.5 step if EDSS at baseline was > 6 ; and 1.5 step if EDSS at baseline was 1.5, of which all should be confirmed by EDSS scores recorded over ≥ 6 months.

2.5 Statistical analyses

The replication study

Estimation of propensity scores. To control for treatment indication bias, we used stabilized inverse probability of treatment weighting (sIPTW) calculated from propensity scores. The propensity score is a balanced score representing the probability of being treated with natalizumab (relative to fingolimod) given the patients' baseline clinical and demographic characteristics. In the replication analyses, it was computed separately for each database using a multivariable logistic regression based on sex, age, MS duration, EDSS at baseline, number of relapses in the 12 months prior to baseline, disease activity 12 months prior to baseline (classified as relapse or EDSS progression, or both), and the number of previously DMDs commenced prior to baseline. In the pooled analysis we

computed sIPTW based on the pooled data. For the MSBase cohort and the pooled cohort, the models of sIPTW included country as a random effect.

Using the propensity scores, we calculated sIPTW (20). Each patient who fulfilled the inclusion criteria was assigned a weight. The weight was proportional to the inverse of the probability of receiving the treatment that the subject actually received (21) given the individual patient's baseline characteristics, e.g. a patient treated with natalizumab with a low probability of being treated with natalizumab was assigned a high weight.

Comparison of treatment effectiveness

Demographic and clinical characteristics of patients from either treatment group within each registry as well as the pooled cohort at baseline are reported, including their standardized differences. A difference of $\leq 10\%$ was considered acceptable (20). The propensity score distributions in the two groups were assessed for the degree of overlap, also named the common support.

ARR for natalizumab and fingolimod were reported. The counts of relapses between natalizumab and fingolimod in the treated periods were compared using generalized linear models with weighted negative binomial distribution model and with logarithmic transformed length of treatment period as offset. The regression coefficients were exponentiated to obtain the ratio of relapse rates.

Weighted Cox proportional hazards models were used to evaluate the cumulative hazard of 1st relapse as well as 1st EDSS improvement and 1st EDSS worsening. The weighted Andersen-Gill proportional hazards model was used to evaluate the cumulative hazards of multiple events of EDSS worsening and improvement. Robust estimation of variance was used.

Analyses were performed per protocol using the R-software (R 3.4.0).

3. Results

3.1 The replication analyses

Table 2 shows the baseline characteristics for the cases from the three databases, before and after stabilized inverse probability weighting (sIPTW). The weighting improved the balance between the natalizumab and the fingolimod treated groups which is demonstrated by the reduced standardized mean differences (SMD).

3.2 The MSBase. This part of the study included 1202 patients, 410 treated with natalizumab and 792 treated with fingolimod. The detailed demographic and clinical baseline characteristics are shown in Table 2 before and after sIPTW. The results of the replication analysis from the MSBase using unified methodology showed an ARR of 0.091 for natalizumab and 0.144 for fingolimod. With fingolimod as reference, the weighted ratio of the ARRs was 0.619; $p = 0.0013$. The hazard ratio (HR) for a first relapse was 0.61 ($p = 0.003$). HR for the first sustained EDSS-worsening was close to unity: 1.08 ($p = 0.767$), but the Cox regression analysis of a first sustained EDSS-improvement indicated that natalizumab was associated with a greater chance of decrease in EDSS than fingolimod: HR = 1.89 ($p = 0.003$). The estimates and confidence intervals are shown in Table 3.

3.3 The DMSR cohort. This cohort included 980 patients, 399 treated with natalizumab and 581 with fingolimod. Demographic and clinical baseline characteristics before and after sIPTW are presented in Table 2. In the replication analyses the results were comparable between the treatment groups: ARR was 0.178 for natalizumab and 0.151 for fingolimod. With fingolimod as reference the weighted ratio of the ARRs was 1.115 ($p = 0.397$). HR for a first relapse was 1.12 ($p = 0.359$), for a first sustained EDSS-worsening: 0.97 ($p = 0.91$), and for a first sustained EDSS-improvement: 1.11 ($p = 0.539$). For the full results and confidence intervals see Table 3.

3.4 The OFSEP cohort. This part of the study included 265 patients, 159 treated with natalizumab and 106 with fingolimod. Table 2 presents detailed demographic and clinical baseline characteristics before and after sIPTW. The replication analysis of the OFSEP data showed that the ARR was 0.183 on natalizumab and 0.387 on fingolimod. Treatment with fingolimod as reference the weighted ratio of the ARRs was 0.466 ($p = 0.002$). For the other outcomes there were no statistically significant differences between the treatment groups. HR for a first relapse was 0.66 ($p=0.111$), for a first sustained EDSS worsening: 0.77 ($p=0.645$), and for a first sustained EDSS-improvement: 1.57 ($p=0.342$). For full results of the analysis and confidence intervals see Table 3.

In summary, with some differences, the results of the present replication analyses for each database were roughly the same as those published in the three individual original studies (2,7,8), when using the uniform design and statistical analyses, with a larger cohort and longer follow-up times for two of the study populations.

3.5 The pooled analysis

The pooled cohort from the three databases consisted of 2447 patients, 968 treated with natalizumab and 1479 treated with fingolimod. In patients treated with natalizumab the ARR was 0.138, compared with the ARR of 0.165 in patients treated with fingolimod. With fingolimod as reference the weighted ratio of the ARRs was 0.771 ($p = 0.004$), and HR for a first relapse was 0.82; ($p=0.030$). We found no difference in hazards for a first sustained EDSS-worsening: HR 1.13 ($p = 0.438$), but sustained EDSS improvement was in the favour of natalizumab with a HR of 1.40 ($p=0.009$), and for multiple EDSS-improvement events of 1.37 ($p = 0.011$). A visual presentation of the results is presented in Figure 2. Table 3 shows full results with confidence intervals. Analyses with interaction terms for registry x treatment confirmed the differences in comparative effectiveness presented in the replication analyses above (data not shown).

Table 3. Results of replication analyses based on weighted (IPTW) data in the pooled cohort and the three individual cohorts contributing to it

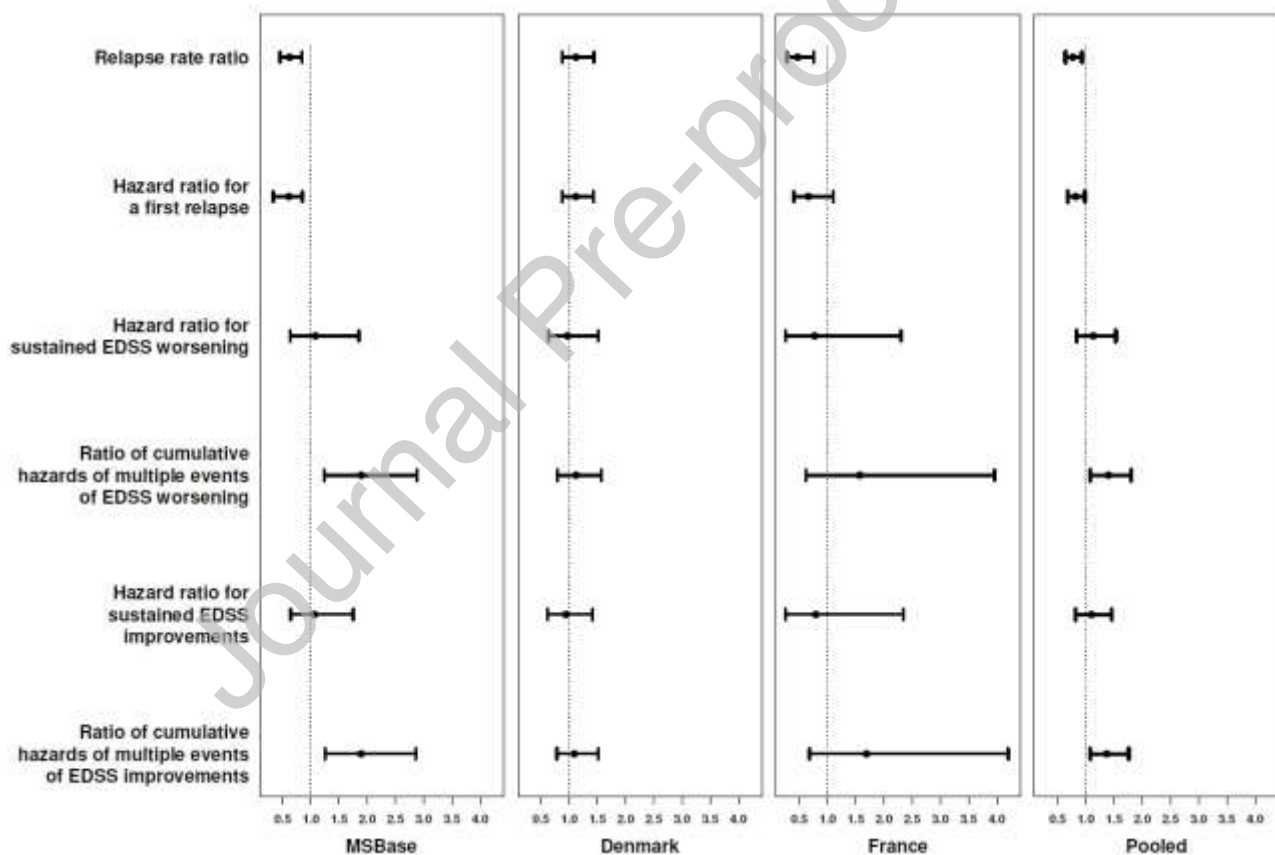
		Pooled cohort N=2447	MSBase N=1202	DMSR N=980	OFSEP N=265
Annualized relapse rate	Natalizumab [95% CI]	0.14 [0.12; 0.16]	0.09 [0.06; 0.12]	0.18 [0.14; 0.22]	0.18 [0.14; 0.23]
	Fingolimod [95% CI]	0.17 [0.14; 0.19]	0.14 [0.12; 0.17]	0.15 [0.12; 0.18]	0.39 [0.21; 0.57]
Difference of means (FTY minus NAT)	[95% CI]	0.026 [-0.004; 0.06]	0.053 [0.02; 0.09]	-0.027 [0.07; -0.02]	0.204 [0.02; 0.39]
Relapse rate ratio*§	[95% CI]	0.77 [0.64; 0.93]	0.62 [0.45-0.84]	1.12 [0.87; 1.44]	0.47 [0.28; 0.76]
	<i>p-value</i>	0.004	0.0013	0.397	0.002
Hazard Ratio* for a first relapse	[95% CI]	0.82 [0.68; 0.98]	0.61 [0.44; 0.85]	1.12 [0.88; 1.43]	0.66 [0.40; 1.10]
	<i>p value</i>	0.030	0.0032	0.359	0.111
Hazard Ratio* for a first sustained EDSS-worsening	[95% CI]	1.13 [0.83; 1.53]	1.08 [0.63; 1.85]	0.97 [0.63; 1.51]	0.77 [0.26; 2.30]
	<i>p value</i>	0.438	0.767	0.910	0.645

Hazard Ratio* for a first sustained EDSS-improvement	[95% CI] <i>p value</i>	1.40 [1.08; 1.80] 0.009	1.89 [1.24; 2.88] 0.003	1.11 [0.79; 1.57] 0.539	1.57 [0.62; 3.96] 0.342
Ratio* of cumulative hazards of multiple events of EDSS-worsening	[95% CI] <i>p value</i>	1.10 [0.82; 1.46] 0.528	1.06 [0.64; 1.75] 0.814	0.94 [0.62; 1.41] 0.745	0.79 [0.26; 2.34] 0.669
Ratio* of cumulative hazards of multiple events of EDSS-improvement	[95% CI] <i>p value</i>	1.37 [1.08; 1.76] 0.011	1.89 [1.25; 2.86] 0.002	1.09 [0.78; 1.51] 0.624	1.69 [0.68; 4.20] 0.259

*with Fingolimod as reference

§calculated as the adjusted exponentiated regression coefficient of count of relapses with logarithmic transformed observation time as offset.

Figure 2. Comparative presentation of study outcomes. Fingolimod is the reference drug in all comparisons.



4. Discussion

Using unified design and methodology, this study reanalysed original and extended clinical data from three different published studies that compared effectiveness of natalizumab and fingolimod in

RRMS. The analyses of the pooled cohort confirmed an advantage of natalizumab over fingolimod in reducing the risk of relapses by 23% and facilitating early recovery from neurological disability by 40%. These results were largely driven by MSBase and OFSEP. However, similar to the original studies, the pooled study found no difference in the risk of EDSS worsening between the two disease modifying therapies.

Also, the original studies from OFSEP and MSBase (7,8) showed that natalizumab was associated with lower risk of relapses than fingolimod. The study in the MSBase cohort also suggested that natalizumab was associated with a higher probability of recovery from disability. On the other hand, there was a certain degree of heterogeneity as the study from DMSR (2) showed no significant differences between the effects of the two drugs.

When we replicated the results from the three contributing databases with the uniform present inclusion criteria and methodology, the results were roughly the same as in the original studies. The heterogeneity between the results of MSBase, OFSEP, and, on the other hand, DMSR can best be explained by differences in the clinical and demographic characteristics of the study populations (11): For example, the OFSEP and the MSBase cohorts were enriched for younger patients with higher prior relapse activity (mean ARR 1.38-1.41 and 1.07-1.14, respectively) and greater exposure to DMDs prior to their treatment with natalizumab or fingolimod than the DMSR in the original studies (mean ARR 0.73). In 12 months prior to treatment switch more of the DMSR patients had experienced worsening compared with patients from MSBase and OFSEP, but fewer of them had recorded relapses in this period. This could also explain some of the differences between the main results from the three databases. In fact, the difference in the effect on relapses between natalizumab and fingolimod was greatest in the cohort with the highest disease activity (OFSEP). This suggests that a 'floor effect' exists when one compares effectiveness among highly potent DMDs, and the differences between fingolimod and natalizumab become apparent in patients with highly active disease. The overall frequency of relapses was higher in the OFSEP dataset than in the DMSR dataset, and the magnitudes of treatment effectiveness were similar or greater in the MSBase and OFSEP datasets than in the combined dataset. We cannot rule out that these differences may be partly driven by differences in reporting methods among the three registries. Confounding by variables that influence the choice of treatment as well as short-term disease outcomes is a major concern when comparing treatment arms in non-randomized open-label studies. The three original studies had dealt with this issue using different statistical methods. The present study used a uniform analytical methodology, based on a consensus among the

investigators, and we used the sIPTW to successfully reduce treatment indication bias. This is reflected by the very close balance of baseline variables between the two treatment arms after weighting. To account for possible heterogeneity, we have included the country of data origin in the estimation of sIPTW in the pooled analyses.

The reported findings were mainly driven by the MSBase and the DMSR cohorts which constitute 49% and 40% of the data in the pooled cohort, respectively. The size of the treatment groups in the individual cohorts (with the exception of the fingolimod group in MSBase) decreased as a result of more rigorous inclusion criteria in the unified analyses. However, our inclusion of data from 183 MS-centers across 36 countries strengthens the generalizability of our pooled data in a real-world setting.

The results of our pooled study are in keeping with a growing body of studies showing the advantage of natalizumab over fingolimod in terms of treatment effectiveness (9,10,22).

Limitations

The inclusion only of patients with sufficient follow-up EDSS is a limitation of this study as this inclusion criterion, which aimed at including a population of patients who became established on their new therapy and with sufficient on-treatment disability information available for the analysis, would limit generalization of the observations for the subset of patients who discontinued their therapy early after only a brief time on treatment.

Furthermore, the lack of magnetic resonance imaging (MRI) data, either as a baseline or as an endpoint parameter is a limitation of this study. A recently published guideline (1) emphasises the advantage of using MRI activity as short- and long-term predictors of disability worsening in RRMS patients. However, two of the original analyses had used (OFSEP) or imputed (MSBase) MRI information in their analyses, without any noticeable effect on the magnitude of the reported difference in the latter. Small numbers in some of the cohorts could have a negative impact on the power of the specific replication analyses, and their results should be interpreted with some caution. Reassuringly, these results confirmed the results of the original studies. Finally, this study did not compare incidence of adverse events, as this information was not available from all combined registries.

In conclusion: This study, conducted in a large combined cohort from three MS registries, reconciles the results of several previous analyses, and shows that natalizumab, after controlling for

indication bias, is associated with a better control of relapse activity and improved chance of early recovery from disability among patients with active RRMS. The different results between the registries are primarily attributable to clinical and demographic differences between the studied cohorts.(23). These characteristics warrant further research as they hold the promise of guiding personalised approach to choosing between different treatment options.

Data availability

DMSR: Anonymized data will be shared on request from any qualified researcher under approval from the Danish Data Protection Agency.

OFSEP: The individual data from the present study can be obtained upon request and after validation from the OFSEP scientific committee (see website: <http://www.ofsep.org/fr/http://www.ofsep.org/en/data-access>)

MSBase: MSBase is a data processor, and warehouses data from individual principal investigators who agree to share their datasets on a project-by-project basis. Each principal investigator will need to be approached individually for permission to access the datasets.

Standard protocol approvals, registrations, and patient consents

The MSBase registry was approved by the Melbourne Health Human Research Ethics Committee and the local ethics committees at participating centres. Enrolled patients provided written informed consent as required. OFSEP was conducted in accordance with the French law relative to clinical noninterventional research according to the French law on Bioethics. Data confidentiality and safety are ensured according to the recommendations of the French Commission Nationale Informatique et Libertés (CNIL). OFSEP has received approval for storing clinical, biological and imaging data for research purpose. Patients gave informed consent for their data to be stored in the database and used for research, in France and abroad (www.ofsep.org/en/cohort/ofsep-consent). The cohort has been registered to clinicaltrials.gov under the number NCT02889965. The Danish study was conducted according to the Danish laws. Non-interventional register-based studies do not require ethical approval in Denmark. Required approvals were obtained with the Center for Data Review applications (j. nr. 2012-58-0004/VD-2018-121 I-suite 6361).

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J.B. Andersen received travel grant and congress participation support from Merck.

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F. Sellebjerg served on scientific advisory boards, been on the steering committees of clinical trials, served as a consultant, received support for congress participation, received speaker honoraria, or received research support for his laboratory from Biogen, Merck, Novartis, Roche, Sanofi Genzyme and Teva.

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J. Frederiksen received no funding to support the presented work. She has served on scientific advisory boards for and received funding for travel related to these activities as well as honoraria from Biogen Idec, Merck Serono, Sanofi-Aventis, Teva, Novartis, Roche and Almirall. She has received speaker honoraria from Biogen Idec, Teva, Roche and Novartis

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M. Magyari served on scientific advisory board for Biogen, Sanofi, Teva, Roche, Novartis, Merck, has received honoraria for lecturing from Biogen, Merck, Novartis, Sanofi, Genzyme, has received research support and support for congress participation from Biogen, Genzyme, Teva, Roche, Merck, Novartis.

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S. Ozakbas did not declare any competing interests.

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A. Lugaresi served as a Bayer, Biogen, Merck, Novartis, Roche, Sanofi/ Genzyme and Teva Advisory Board Member. She received congress and travel/accommodation expense compensations and speaker honoraria from Bayer, Biogen, Merck, Novartis, Sanofi/Genzyme, Teva and Fondazione Italiana Sclerosi Multipla (FISM). Her institutions received research grants from Novartis.

M. Terzi received travel grants from Novartis, Bayer-Schering, Merck and Teva; has participated in clinical trials by Sanofi Aventis, Roche and Novartis.

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B. Yamout did not declare any competing interests.

A. Prat did not declare any competing interests.

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C. Boz received conference travel support from Biogen, Novartis, Bayer-Schering, Merck and Teva; has participated in clinical trials by Sanofi Aventis, Roche and Novartis.

M. Trojano received speaker honoraria from Biogen-Idec, Bayer-Schering, Sanofi Aventis, Merck, Teva, Novartis and Almirall; has received research grants for her Institution from Biogen-Idec, Merck, and Novartis.

P. McCombe did not declare any competing interests.

M. Slee has participated in, but not received honoraria for, advisory board activity for Biogen, Merck, Bayer Schering, Sanofi Aventis and Novartis.

J. Lechner-Scott accepted travel compensation from Novartis, Biogen and Merck. Her institution receives the honoraria for talks and advisory board commitment from Bayer Health Care, Biogen, Genzyme Sanofi, Merck, Novartis and Teva, has been involved in clinical trials with Biogen, Novartis and Teva.

R. Turkoglu did not declare any competing interests.

P. Sola served on scientific advisory boards for Biogen Idec and TEVA, she has received funding for travel and speaker honoraria from Biogen Idec, Merck, Teva, Sanofi Genzyme, Novartis and Bayer and research grants for her Institution from Bayer, Biogen, Merck, Novartis, Sanofi, Teva.

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F. Granella received an institutional research grant from Biogen and Sanofi Genzyme, served on scientific advisory boards for Biogen, Novartis, Merck, Sanofi Genzyme and Roche, received funding for travel and speaker honoraria from Biogen, Merck, and Sanofi-Aventis.

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J. Prevost accepted travel compensation from Novartis, Biogen, Genzyme, Teva, and speaking honoraria from Biogen, Novartis, Genzyme and Teva.

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C. Solaro served on scientific advisory boards for Merck, Genzyme, Almirall, and Biogen; received honoraria and travel grants from Sanofi Aventis, Novartis, Biogen, Merck, Genzyme and Teva.

R. Karabudak did not declare any competing interests.

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S. Vucic did not declare any competing interests.

H. Butzkueven served on scientific advisory boards for Biogen, Novartis and Sanofi-Aventis and has received conference travel support from Novartis, Biogen and Sanofi Aventis. He serves on steering committees for trials conducted by Biogen and Novartis, and has received research support from Merck, Novartis and Biogen.

T. Kalincik served on scientific advisory boards for Celgene, Roche, Sanofi-Genzyme, Novartis, Merck and Biogen, steering committee for Brain Atrophy Initiative by Sanofi-Genzyme, received conference travel support and/or speaker honoraria from WebMD Global, Novartis, Biogen, Roche, Genzyme-Sanofi, Teva, BioCSL and Merck and received research support from Biogen.

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universitaire de Caen Normandie, Service de neurologie, Hôpital Côte de Nacre, Caen, France; Mikaël Cohen, MD, Centre hospitalier universitaire de Nice, Université Nice Côte d'Azur, Hôpital Pasteur, Service de neurologie, Nice, France; Agnès Fromont, MD, Centre hospitalier universitaire Dijon Bourgogne, Hôpital François Mitterrand, Service de neurologie, maladies inflammatoires du système nerveux et neurologie générale, Dijon, France; Sandrine Wiertelowsky, MD, Centre hospitalier universitaire de Nantes, Hôpital nord Laennec, Service de neurologie, Nantes/Saint-Herblain, France; Eric Berger, MD, Centre hospitalier régional universitaire de Besançon, Hôpital Jean Minjoz, Service de neurologie, Besançon, France; Pierre Clavelou, MD, Centre hospitalier universitaire de Clermont-Ferrand, Hôpital Gabriel-Montpied, Service de neurologie, Clermont-Ferrand, France; Bertrand Audoin, MD, Assistance publique des hôpitaux de Marseille, Centre hospitalier de la Timone, Service de neurologie et unité neuro-vasculaire, Marseille, France; Claire Giannesini, MD, Assistance publique des hôpitaux de Paris, Hôpital Saint-Antoine, Service de neurologie, Paris, France; Olivier Gout, MD, Fondation Adolphe de Rothschild de l'œil et du cerveau, Service de neurologie, Paris, France; Eric Thouvenot, MD, Centre hospitalier universitaire de Nîmes, Hôpital Carémeau, Service de neurologie, Nîmes, France; Olivier Heinzlef, MD, Centre hospitalier intercommunal de Poissy Saint-Germain-en-Laye, Service de neurologie, Poissy, France; Abdullatif Al-Khedr, MD, Centre hospitalier universitaire d'Amiens Picardie, Site sud, Service de neurologie, Amiens, France; Bertrand Bourre, MD, Centre hospitalier universitaire Rouen Normandie, Hôpital Charles-Nicolle, Service de neurologie, Rouen, France; Olivier Casez, MD, Centre hospitalier universitaire Grenoble-Alpes, Site nord, Service de neurologie, Grenoble/La Tronche, France; Philippe Cabre, MD, Centre hospitalier universitaire de Martinique, Hôpital Pierre Zobda-Quitman, Service de Neurologie, Fort-de-France, France; Alexis Montcuquet, MD, Centre hospitalier universitaire Limoges, Hôpital Dupuytren, Service de neurologie, Limoges, France; Alain Créange, MD, Assistance publique des hôpitaux de Paris, Hôpital Henri Mondor, Service de neurologie, Créteil, France; Jean-Philippe Camdessanché, MD, Centre hospitalier universitaire de Saint-Étienne, Hôpital Nord, Service de neurologie, Saint-Étienne, France; Justine Faure, MD, Centre hospitalier universitaire de Reims, Hôpital Maison-Blanche, Service de neurologie, Reims, France; Aude Maurousset, MD, Centre hospitalier régional universitaire de Tours, Hôpital Bretonneau, Service de neurologie, Tours, France; Ivania Patry, MD, Centre hospitalier sud francilien, Service de neurologie, Corbeil-Essonnes, France; Karolina Hankiewicz, MD, Centre hospitalier de Saint-Denis, Hôpital Casanova, Service de neurologie, Saint-Denis, France; Corinne Pottier, MD, Centre hospitalier de Pontoise, Service de neurologie, Pontoise, France; Nicolas

Maubeuge, MD, Centre hospitalier universitaire de Poitiers, Site de la Milétrie, Service de neurologie, Poitiers, France; Céline Labeyrie, MD, Assistance publique des hôpitaux de Paris, Hôpital Bicêtre, Service de neurologie, Le Kremlin-Bicêtre, France; Chantal Nifle, MD, Centre hospitalier de Versailles, Hôpital André-Mignot, Service de neurologie, Le Chesnay, France;

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Appendix 1.

Author		Role	Contribution
Melinda Magyari, MD, PhD	Danish Multiple sclerosis Center, Rigshospitalet Glostrup, Copenhagen, Denmark	Principal investigator	Conceptualised and designed the study, recruited patients, contributed data, carried out statistical analysis, interpreted the results, drafted and edited the manuscript.
Johanna B Andersen, MSc	Danish Multiple sclerosis Registry, Rigshospitalet Glostrup, Copenhagen, Denmark	Analyst	Contributed to the study design, interpreted the results, carried out statistical analysis, edited the manuscript
Nils Koch-Henriksen, MD, DMSc	Aarhus University Hospital, Aarhus, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Finn Sellebjerg, MD, PhD, DMSc	Danish Multiple sclerosis Center, Rigshospitalet Glostrup, Copenhagen, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Per Solberg Soerensen, MD, PhD, DMSc.	Danish Multiple sclerosis Center, Rigshospitalet Glostrup, Copenhagen, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Claudia Pfleger, MD	Aalborg University Hospital, Aalborg, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Peter V Rasmussen, MD, PhD	Aarhus University Hospital, Aarhus, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Stephan Bramow, MD, PhD	Danish Multiple sclerosis Center, Rigshospitalet Glostrup, Copenhagen, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Jette L Battisini, MD, PhD	Danish Multiple sclerosis Center, Rigshospitalet Glostrup, Copenhagen, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Henrik K Mathiesen, MD, PhD	Copenhagen University Hospital, Herlev, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Michael B Jensen, MD, PhD	University Hospital of Northern Sealand, Denmark	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Karen I Schreiber, MD	Danish Multiple sclerosis Center, Rigshospitalet Glostrup,	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript

	Copenhagen, Denmark		
Tomas Kalincik, MD, PhD	Royal Melbourne Hospital, Melbourn, Australia	Principal investigator	Conceptualized and designed the study, recruited patients, contributed data, carried out statistical analysis, interpreted the results, drafted and edited the manuscript.
Sifat Sharmin, PhD	Royal Melbourne Hospital, Melbourn, Australia	Analyst	Contributed to the study design, interpreted the results, carried out statistical analysis, edited the manuscript
Dana Horakova, MD, PhD	General University Hospital, Prague, Czech Republic	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Eva Kubala Havrdova, MD, PhD	General University Hospital, Prague, Czech Republic	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Maria Trojano, MD	University of Bari, Bari, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Guillermo Izquierdo, MD	Hospital Universitario Virgen Macarena, Sevilla, Spain	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Alessandra Lugaresi, MD, PhD	University of Bologna, Bologna, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Alexandre Prat, MD, PhD	Hospital Notre Dame, Universite de Montreal, Montreal, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Pierre Duquette, MD	Hospital Notre Dame, Universite de Montreal, Montreal, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Pierre Grammond, MD	CISSS Chaudiere-Appalache, Levis, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Francois Grand'Maison, MD	Neuro Rive-Sud, Quebec, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Patrizia Sola, MD, PhD	Azienda Ospedaliera Universitaria, Modena, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Diana Ferraro, MD, PhD	Azienda Ospedaliera Universitaria, Modena, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Vahid Shaygannejad, MD	Isfahan University of Medical Sciences, Isfahan, Iran	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Raed Alroughani,	Amiri Hospital, Sharq, Kuwait	Investigator	Recruited patients, contributed data, interpreted the results, edited the

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Murat Terzi, MD	19 Mayis University, Samsun, Turkey	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Cavit Boz, MD	KTU Medical Faculty Farabi Hospital, Trabzon, Turkey	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Jeannette Lechner-Scott, MD, PhD	School of Medicine and Public Health, University Newcastle, John Hunter Hospital, Newcastle, Australia	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Franco Granella, MD	University of Parma, Parma University Hospital, Parma, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Daniele Spitaleri, MD	Azienda Ospedaliera di Rilieco Nazionale Dan Giuseppe Moscation Avellino, Avellino, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Mark Slee, BMBS, PhD	Flinders University, Adelaide, Australia	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Steve Vucic, MBBS, PhD	Westmeas Hospital, Sydney, Australia	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Sara Eichau, MD	Hospital Universitario Virgen Macarena, Sevilla, Spain	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Serkan Ozakbas, MD	Dokuz Eylul University, Konak/Izmir, Turkey	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Francesco Patti, MD	University of Catania, Catania, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Marco Onofri, MD	University G. d'Annunzio, Chieti, Italy	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Bassem Yamout, MD	American University of Beirut Medical Center, Beirut, Libanon	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Marc Girad, MD	Hospital Notre Dame, Universite de Montreal, Montreal, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Recai Turkoglu, MD	Haydarpasa Numune Training and Research Hospital, Istanbul, Turkey	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Claudio	ASL3 Genovese,	Investigator	Recruited patients, contributed data,

Solaro, MD	Genova, Italy, ML Novarese Hospital Moncrivello		interpreted the results, edited the manuscript
Rana Karabudak, MD	Hacettepe University, Ankara, Turkey	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Bart V Wijmeersch, MD, PhD	Hasselt University, Hasselt, Belgium	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Pamela McCombe, MBBS	University of Queensland, Brisbane, Royal Brisbane and Women's Hospital, Australia	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Julie Prevost, MD	CSSS Saint-Jerome, Saint-Jerome, Canada	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Olga Skibina, MD	Monash University, Melbourne, Australia	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Tunde Csepany, MD	University of Debrecen, Debrecen, Hungary	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Helmut Butzkueven, MBBS, PhD	Monash University, The Alfred Hospital, Melbourne, Australia	Investigator	Contributed to the study design, recruited patients, contributed data, interpreted the results, edited the manuscript
Sandra Vukusic, MD, PhD	Hospices Civils de Lyon, Lyon/Bron, France	Principal investigator	Conceptualised and designed the study, recruited patients, contributed data, interpreted the results, drafted and edited the manuscript.
Mathilde Lefort, PhD	Rennes University, Rennes, France	Analyst	Contributed to the study design, interpreted the results, carried out statistical analysis, edited the manuscript
Emmanuelle Leray, PhD	Rennes University, Rennes, France	Analyst	Contributed to the study design, interpreted the results, edited the manuscript
David Laplaud, MD, PhD	CHU de Nantes, Nantes, France	Investigator	Contributed to the study design, interpreted the results, edited the manuscript
Romain Casey, PhD	Hospices Civils de Lyon, Lyon/Bron, France	Analyst	Contributed to the study design, interpreted the results, edited the manuscript
Marc Debouverie, MD, PhD	Hôpital Central, Nancy, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Gilles Edan,	Hôpital Pontchaillou,	Investigator	Recruited patients, contributed data,

MD, PhD	Rennes, France		interpreted the results, edited the manuscript
Jonathan Ciron, MD	Hôpital Purpan, Toulouse, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Aurélie Ruet, MD, PhD	Hôpital Pellegrin, Bordeaux, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Jérôme De Seze, MD, PhD	Hôpital de Hautepierre, Strasbourg, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Elisabeth Maillart, MD	Hôpital de la Pitié-Salpêtrière, Paris, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Hélène Zephir, MD, PhD	Hôpital Salengro, Lille, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Pierre Labauge, MD, PhD	Hôpital Gui de Chauliac, Montpellier, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Gilles Defer, MD, PhD	Hôpital Côte de Nacre, Caen, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Christine Lebrun-Frenay, MD, PhD	Hôpital Pasteur, Nice, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Thibault Moreau, MD, PhD	Hôpital Francois Mitterrand, Dijon, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Eric Berger, MD	Hôpital Jean Minjoz, Besancon, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Pierre Clavelou, MD, PhD	Hôpital Gabriel-Montpied, Clermont-Ferrand, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Jean Pelletier, MD, PhD	Centre hospitalier de la Timone, Marseille, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Bruno Stankoff, MD, PhD	Hôpital Saint-Antoine, Paris, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Olivier Gout, MD	Foundation Adolphe de Torhschild de l'œil et du cerveau, Paris, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Eric Thouvenot, MD, PhD	Centre Hospitalier universitaire de Nîmes, Hôpital Carémeau,	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript

	Nîmes, France		
Olivier Heinzlef, MD	Centre Hospitalier intercommunal de Poissy Saint-Germain-en-Laye, Poissy, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Abullatif Al-Khedr, MD	Centre Hospitalier universitaire d'Amiens Picardie, Amiens, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Bertrand Bourre, MD	Hôpital Charles-Nicolle, Rouen, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Olivier Casez, MD	Centre Hospitalier universitaire Grenoble-Alpes, Grenoble/La Tronche, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Philippe Cabre, MD, PhD	Hôpital Pierre Zobda-Quitman, Fort-de-France, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Alexis Montcuquet, MD	Hôpital Dupuytren, Limoges, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Abir Wahab, MD	Hôpital Henri Mondor, Créteil, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Jean-Philippe Camdessanche, MD, PhD	Hôpital Nord, Saint-Étienne, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Aude Maurousset, MD	Hôpital Bretonneau, Tours, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Ivania Patry, MD	Centre Hospitalier sud francilien, Corbeil-Essonnes, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Karolina Hankiewicz, MD	Hôpital Casanoca, Saint-Denis, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Corinne Pottier, MD	Centre Hospitalier de Pontoise, Pontoise, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Nicolas Maubeuge, MD	Centre Hospitalier universitaire de Poitiers, Poitiers, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Céline Labeyrie, MD	Hôpital Bicêtre, Le Kremlin-Bicêtre, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the manuscript
Chantal Nifle, MD	Hôpital André-Mignot, Le Chesnay, France	Investigator	Recruited patients, contributed data, interpreted the results, edited the

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