



Evaluating the Long-Term Outcomes and Durability of Effect Following Treatment with Cladribine Tablets for Multiple Sclerosis

14/08/2025 09:13:29

Main Information

Primary registry identifying number

LBCTR2020030215

Protocol number

MS700568_0026

MOH registration number

Study registered at the country of origin

Yes

Study registered at the country of origin: Specify

Type of registration

Prospective

Type of registration: Justify

N/A

Date of registration in national regulatory agency

25/07/2019

Primary sponsor

Merck KGaA

Primary sponsor: Country of origin

Germany

Date of registration in primary registry

18/03/2020

Date of registration in national regulatory agency

25/07/2019

Public title

Evaluating the Long-Term Outcomes and Durability of Effect Following Treatment with Cladribine Tablets for Multiple Sclerosis

Acronym

Scientific title

An Exploratory Phase IV Ambispective Study of Patients Who Previously Participated in the CLARITY/CLARITY-EXT and ORACLE MS Clinical Trials

Acronym

Brief summary of the study: English

The purpose of this study is to explore the long-term outcomes, durability of effect, and real-world treatment patterns in patients previously participating in the Phase III ORACLE MS and CLARITY/CLARITY-EXT clinical trials (i.e. parent studies). The results from this study may be of benefit to patients with multiple sclerosis (MS) and clinicians by helping to inform future treatment approaches and treatment decision-making.

Brief summary of the study: Arabic

تقييم النتائج طويلة الأمد و مدة تأثير العلاج بأقراص Cladribine عند مرضى مصابين بتصلب المتعدد

Health conditions/problem studied: Specify

Multiple Sclerosis

Interventions: Specify

No Intervention, except:

*Optional blood sample: Patients willing to consent to provide an optional blood sample and who are seen at a site with available capabilities to store and ship samples, will have a blood draw taken at Study Visit 1 for pharmacogenetics testing.

Key inclusion and exclusion criteria: Inclusion criteria

1. Patients with MS randomised in CLARITY/CLARITY-EXT clinical trial(s) who have received ≥ 1 course of IMP (Cladribine Tablets or





placebo).

or

Patients with their FCDE randomised in ORACLE MS clinical trial who have received ≥ 1 course of IMP (Cladribine Tablets or placebo).

2. Informed Consent

Key inclusion and exclusion criteria: Gender

Both

Key inclusion and exclusion criteria: Specify gender

Key inclusion and exclusion criteria: Age minimum

18

Key inclusion and exclusion criteria: Age maximum

65

Key inclusion and exclusion criteria: Exclusion criteria

1. Medical Conditions:

Any condition, including any uncontrolled disease state other than MS, that in the Investigator's opinion, constitutes an inappropriate risk or a contraindication for participation in the study or that could interfere with the study objectives, conduct, or evaluation.

For the MRI sub-study:

1. Female Participants Who are pregnant

2. Patient taking Cladribine Tablets as part of another study at the time of the start of this study

Type of study

Observational

Type of intervention

N/A

Type of intervention: Specify type

N/A

Trial scope

N/A

Trial scope: Specify scope

N/A

Study design: Allocation

N/A

Study design: Masking

N/A

Study design: Control

N/A

Study phase

N/A

Study design: Purpose

N/A

Study design: Specify purpose

N/A

Study design: Assignment

N/A

Study design: Specify assignment

N/A

IMP has market authorization

IMP has market authorization: Specify

Name of IMP

Year of authorization

Month of authorization

Type of IMP

Pharmaceutical class

Therapeutic indication

Therapeutic benefit

Study model

Cohort

Study model: Specify model

N/A

Time perspective

Other

Time perspective: Specify perspective

Ambispective

Target follow-up duration

6

Number of groups/cohorts

4

Biospecimen retention

Samples with DNA**

Target sample size

8

Date of first enrollment: Type

Actual

Date of study closure: Type

Actual

Recruitment status

Recruiting

Date of completion

IPD sharing statement plan

No

Additional data URL

Admin comments

Study model: Explain model

NA

Time perspective: Explain time perspective

Ambispective: Retrospective and Prospective

Target follow-up duration: Unit

Weeks

Biospecimen description

8ml blood sample will be collected for DNA analysis from consenting participants

Actual enrollment target size

Date of first enrollment: Date

15/08/2019

Date of study closure: Date

28/02/2021

Recruitment status: Specify

IPD sharing statement description

NO Individual Patient Data Sharing

**Trial status**

Approved

Secondary Identifying Numbers

Full name of issuing authority	Secondary identifying number
European Clinical Trials Database	EudraCT number: 2019-000069-19

Sources of Monetary or Material Support

Name
Merck KGaA Germany

Secondary Sponsors

Name
N/A

Contact for Public/Scientific Queries

Contact type	Contact full name	Address	Country	Telephone	Email	Affiliation
Public	Dr. Bassem Yamout	Hamra, Cairo Street	Lebanon	+9613221 222	-	American University of Beirut Medical Center
Scientific	Kristin Gabriel	EMD Serono, Inc. One Technology Place, Rockland MA 02370	United States of America	+1 781 427 1502	Kristin.Gabriel@emdserono.com	EMD Serono, Inc.

Centers/Hospitals Involved in the Study

Center/Hospital name	Name of principles investigator	Principles investigator speciality	Ethical approval
Bellevue Medical Center	Dr. Souheil Gbeily	Neurology	Approved
American University of Beirut Medical Center	Dr. Bassem Yamout	Neurology	Not approved



Ethics Review

Ethics approval obtained	Approval date	Contact name	Contact email	Contact phone
Bellevue Medical Center	10/09/2019	Prof. Souheil Jbeily	souheil.gebeily@gmail.com	9611682666

Countries of Recruitment

Name
Australia
Austria
Belgium
Bulgaria
Canada
Croatia
Czech Republic
Estonia
Finland
France
Georgia
Germany
Greece
Italy
Republic of Korea
Latvia
Lithuania
Morocco
Norway
Poland
Portugal



Romania
Russian Federation
Republic of Serbia
Spain
Sweden
Switzerland
Tunisia
Ukraine
United Kingdom
United States of America
Lebanon

Health Conditions or Problems Studied

Condition	Code	Keyword
Multiple Sclerosis	Multiple sclerosis (G35)	MS, Multiple Sclerosis

Interventions

Intervention	Description	Keyword
Optional Blood Sampling	Purpose: Pharmacogenetics Testing	blood sampling



Primary Outcomes

Name	Time Points	Measure
To evaluate long-term mobility after treatment with an investigational medicinal product (IMP; Cladribine Tablets or placebo) as part of the Phase III ORACLE MS and CLARITY/CLARITY-EXT clinical trials.	Proportion of study participants using a wheelchair (defined as unable to walk beyond approximately 5 meters even with aid, essentially restricted to wheelchair; wheels self in standard wheelchair and transfers alone; up and about in wheelchair some 12 hours a day) the majority of the time in the 3 months prior to Study Visit 1 for the CLARITY/CLARITY-EXT and ORACLE MS populations,	<ul style="list-style-type: none">• Expanded Disability Status Scale (EDSS) score of 7.0 or higher (if available), or• Alternative clinical description data in medical records.



Key Secondary Outcomes

Name	Time Points	Measure
To assess the long-term disability status after treatment with IMP as part of the Phase III ORACLE MS and CLARITY/CLARITY-EXT clinical trials for the CLARITY/CLARITY-EXT and ORACLE MS populations.	Proportion of study participants with 3-month sustained (i.e. ambulatory disability consistent with EDSS on at least 2 clinic visits no less than 3 months apart)	EDSS of 6.0 or higher in the last year prior to enrollment or corresponding clinical description in medical records
To evaluate differences in clinical characteristics between long-term responders and study participants requiring alternate therapies following treatment with IMP for the CLARITY/CLARITY-EXT and ORACLE MS populations.	Clinical characteristics at Study Visit 1 of long-term responders (defined as study participants who did not demonstrate any evidence of disease reactivation based on Investigator assessment of clinical and imaging outcomes until Year 4 or later following their last dose of IMP and who did not receive disease modifying treatment until Year 4 or later following their last dose of IMP) compared to those of other study participants who started on alternate therapy less than 4 years following their last dose of IMP for the CLARITY/CLARITY-EXT and ORACLE MS populations.	-
To evaluate differences in magnetic resonance imaging (MRI) characteristics between long-term responders and study participants requiring alternate therapies following treatment with IMP for the CLARITY/CLARITY-EXT and ORACLE MS populations.	MRI characteristics at Study Visit 2 of long-term responders (defined as study participants who did not demonstrate any evidence of disease reactivation based on Investigator assessment of clinical and imaging outcomes until Year 4 or later following their last dose of IMP and who did not receive disease modifying treatment until Year 4 or later following their last dose of IMP) compared to those of other study participants who started on alternate therapy less than 4 years following their last dose of IMP for the CLARITY/CLARITY-EXT and ORACLE MS populations.	-



Trial Results

Summary results

Study results globally

Date of posting of results summaries

Date of first journal publication of results

Results URL link

Baseline characteristics

Participant flow

Adverse events

Outcome measures

URL to protocol files