A Study of Suboptimally Controlled Participants Previously Taking Injectable DMDs for RMS (CLICK-MS)

Status: Recruiting

Eligibility Criteria

Age: 18 years and over

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

• Male or female participants greater than or equal to (>=)18 years

• Signed informed consent

• Have diagnosis of RMS including RRMS and aSPMS and satisfy the approved indication for cladribine tablets as per United States Prescribing Information (USPI)

• Have time since diagnosis of RMS of at least 12 months

• Had received their last previous injectable disease-modifying drug (DMD) for at least 3 months

• Have decided to initiate treatment with cladribine tablets during routine clinical care

• Meet criteria as per the approved USPI

• Have access to a valid e-mail address

• In the opinion of the Investigator, experienced suboptimal response (lack of effectiveness, intolerability, poor adherence) to injectable DMD treatment

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Exclusion Criteria:

• Have been previously treated with cladribine in any dosing form

• Transitioning from previous injectable DMD solely for administrative reasons such as relocation

• Have comorbid conditions that preclude participation

• Have any clinical condition or medical history noted as contraindication on USPI

• Are currently participating in an interventional clinical trial

• Pregnant or breastfeeding women, women who plan to become pregnant or men whose partner plans to become pregnant during the cladribine treatment period

Conditions & Interventions

Interventions:
Drug: Cladribine Tablets
Conditions:
Multiple Sclerosis
Keywords:
Multiple Sclerosis, Cladribine Tablets, Observational, Mavenclad

More Information

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