A Study to Evaluate Del-brax (Also Referred to as AOC 1020) in Participants With FSHD

Status: RECRUITING

Eligibility Criteria

Age: 16 years to 70 years old

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

* Clinical and genetic diagnosis of FSHD1 or FSHD2 * Ability to walk independently at pre-specified walking speed (orthoses and ankle braces allowed) for at least 10 meters at screening * Adequate muscle strength based on QMT composite score

Exclusion Criteria:

* Breastfeeding, pregnancy, or intent to become pregnant during the study * Unwilling or unable to comply with contraceptive requirements * Abnormal lab values, conditions or diseases that would make the participant unsuitable for the study * Blood Pressure \> 140/90 mmHg at Screening * Treatment with another investigational drug or biological agent within 1 month of Screening or 5 half-lives of the drug, whichever is longer * Treatment with an oligonucleotide within 9 months of Screening

Conditions & Interventions

Interventions:

DRUG: AOC-1020, DRUG: Placebo

Conditions:

Facioscapulohumeral Muscular Dystrophy, FSHD, FSHD - Facioscapulohumeral Muscular Dystrophy, FSHD1, FSHD2, Fascioscapulohumeral Muscular Dystrophy, Fascioscapulohumeral Muscular Dystrophy Type 1, Fascioscapulohumeral Muscular Dystrophy Type 2, Facioscapulohumeral Muscular Dystrophy 1, Facioscapulohumeral Dystrophy, Facioscapulohumeral Dystrophy, Facioscapulohumeral Muscular Dystrophy 2, Facioscapulohumeral Muscular Dystrophy Type 1 (FSHD1), FSH Muscular Dystrophy, Landouzy Dejerine Dystrophy, Landouzy-Dejerine Syndrome

Keywords:

Avidity, Avidity Biosciences, del-brax, del brax, delbrax, AOC1020, AOC 1020, delpacibart braxlosiran, FORTITUDE-3, FORTITUDE Phase 3, FORTITUDE, FORTITUDE 3

More Information

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Principal Investigator: Phase: PHASE3

IRB Number:

System ID: NCT07038200

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