

A Phase 2, Randomized, Double-Blind, Placebo-Controlled, Multicenter Study to Evaluate the Safety and Efficacy of Vosoritide in Infants and Young Children with Hypochondroplasia, Aged 0 to < 36 Months

Status: Recruiting

Eligibility Criteria

Sex: Male or Female

Age Group: Up to 18 years old

This study is NOT accepting healthy volunteers

Inclusion Criteria:

- 0 months to less than 36 months old - confirmed genetic diagnosis of Hypochondroplasia (HCH)

•participant's weight must be ≥ 3 kg (6.6 pounds) - see link to clinicaltrials.gov for complete inclusion criteria

Exclusion Criteria:

- short stature condition other than HCH - have an unstable medical condition - see link to clinicaltrials.gov for complete exclusion criteria

Conditions & Interventions

Interventions:

Drug: Placebo, Drug: Vosoritide

Conditions:

Diabetes & Endocrine, Rare Diseases

Keywords:

HCH, Hypochondroplasia

More Information

Description: The purpose of this study is to learn the effects, good or bad, of the study drug, vosoritide, for treating children who have hypochondroplasia. This condition affects the growth of bone and cartilage and in which kids who have it are shorter than other kids of the same age. In this study, your child will get either the study drug or placebo, which is an inactive medicine.

Study Contact: Brad Miller - mille685@umn.edu

Principal Investigator: Brad Miller, MD, PhD

Phase: PHASE2

IRB Number: STUDY00026654

Thank you for choosing StudyFinder. Please visit <http://studyfinder.umn.edu> to find a Study which is right for you and contact sfinder@umn.edu if you have questions or need assistance.