
View Now | Addressing the Gaps in Clinical Trial Readiness for FSHD

Facioscapulohumeral muscular dystrophy (FSHD) is one of the most common myopathies affecting anywhere between 1:8,000 to 1:20,000 individuals. There are currently no effective therapies for this indication. The FSHD Society is a research-focused patient advocacy nonprofit organization dedicated to finding a treatment for this devastating disease while also empowering the patient community.



2022 WEBINAR SERIES

Addressing the gaps in clinical trial
readiness for FSHD

presented by FSHD Society

Aug. 17, 2022 | 12 PM ET



REGISTER: https://bit.ly/RDCADAP_Aug_Webinar

With increasing interest by biopharmaceuticals in this indication, the FSHD Society is leveraging existing clinical data to help chart the natural history of the disease, promote the identification of suitable clinical endpoints and enhance patient stratification for upcoming trials. In this webinar, we will discuss the perceived gaps in clinical trial readiness and the various efforts the FSHD Society is taking to help accelerate clinical development.

[View Now](#)

[fda_acknowledgement]