

The Enroll-HD Clinical Trial Committee (CTC) provides recommendations and guidance to clinical trial sponsors for clinical development and/or review of trial protocols for operational support using Enroll-HD resources

- \succ Coordinates the prolnitial contact point for sponsors planning interventional therapeutic HD clinical programs
- > Offers expert advice to partners in pharma, biotech, device companies and academic consortia.
- > Access to CHDI clinical expertise
- > Panel of independent expert advisors in HD and all aspects of clinical trial design and methodology
- > Interface between the Enroll-HD platform and partners
- > Provision of advice and platform support and provides an opportunity for consultation in the early stages of protocol/program development
- > Oversight of HD Clinical Trial Site Certification Program

Chair: Cristina Sampaio; Management: Juliana Bronzova, David Howland, Tim McLean, Swati Sathe, Jenny Townhill



Enroll-HD Clinical Trial Committee (CTC): Support for Clinical Development of HD Therapeutics

The Enroll-HD Platform Team

How can we help? Identify the needs of clinical trial Sponsor







What can we do? Access to expertise and operational support

Support

Advisory activity

CTC acceptance for Enroll-HD operational support

Maintenance of acceptance

Access to HD clinical expertise to advise on clinical questions; study protocols and programs

> CHDI Clinical Experts Independent expert panel

Support provided for HD Phase I-III trial development

Access to Enroll-HD resources enables support during:

Study execution Study planning and

- ✓ Issue resolution
 - Recruitment -
 - potential participants and referrals

Review of protocol amendments to continue provision of operational support

HD Clinical Trial Site Certification Program



Protocol development

■ Site Identification/Feasibility Participant recruitment

Training

Operational support

Number of companies accessing different types of support

Informs Enroll-HD platform support for site feasibility and identification

Open to all HD clinical sites worldwide: assesses fundamental criteria for HD clinical trial participation

HD patient population

• Site staff expertise and training (e.g. GCP, HD rating scales)

• Facilities

start up

✓ In silico feasibility

✓ Site identification

Certified Sites)

(inc. HD Trial

Training and

certification

138 clinical trial ready certified sites

To learn more and initiate discussions contact us Jenny. Townhill@Enroll-HD.org

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