



Details of Study ID 202305_MRS

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Date of Approval: May 2023
Name of study: Congenital micropenis in DSD
Summary of the proposed work to be performed with the registry data: There is a lack of systematic studies that examine the outcome of micropenis in DSD in terms of genital appearance, quality of life and sexual satisfaction. This study will assess in detail the management of micropenis among different Centres.
Lay summary for the public (for the I-DSD/CAH websites and other publicity materials) (maximum 50 words) This study will seek to assess the outcomes of micropenis management in boys with DSD.
Inclusion criteria Consent to be included on the I-DSD Registry Raised as boys Identified as having micropenis Year of birth before 2008, ie 16 years of age
Exclusion criteria Raised as girls.
Data to be collected for all participating cases: Core Data And I-DSD longitudinal assessment



The assessments need to be completed at a maximum of 5 time points – birth or initial presentation, at aged one year, between the age of 10-12 years, at 16 years and at most recent presentation.

For missing data, it will be asked to Centres to complete the DSD Assessments Tab within the Registry- a reminder will be send every 2 months for the first 6 months.

Expected outputs

Presentation – I-DSD Symposium; ESPE

Publication

Publication Plan for authorship in outputs (refer to [guidance](#))

The authors involved in study design, data extraction and analysis and writing the paper will be listed first. Clinical contributors would be listed next according to the alphabetical order The senior authors involved in initial design of the study, data analysis and continuous supervision will be listed last.