# GODIF – log for site master file for new documents and/or new updated versions

**Date: 10.01.2022**: Added the annual safety report for GODIF trial (site master file #4f).

**Date: 14.12.2021:** Added: a revised document for case-money (site master file #6a), revised document for trial medication dispensing system (site master file #9aiii) and a revised powerpoint show for trial medication and dispensing (site master file #9ciii).

**Date: 29.11.2021:** Added a new document from The Capital Region Knowledge Center for Data Compliance. (site master file #4c).

**Date: 27.10.2021**: Added an updated Trial synopsis (site master file #1c and #9biii).

**Date: 02.09.2021: Added an updated document of “The rights of the patient in clinical research” (site master file #7c), former GODIF newsletters are up-loaded (site master file #13f), SOP for follow-up and MoCA mini test and instructions are updated (site master file #17a and #17c).**

**Date: 27.05.2021:** Added a new slide show for initiation for investigators and for primary personel (site master file #9ci).

**Date: 25.05.2021:** Added an updated eCRF overview for screening, baseline and dayform (site master file #2), an updated document of primary data source template and for Sundhedsplatformen (site master file #12) and finally an updated plan for data-verification (site master file #15d).

**Date: 10.05.2021**: Added an approved GODIF protocol version 2.6 (site master file #1a), approval from The Committees on Health Research Ethics (#4c) and the Danish Medicine Agency (#4a), new EudraCT (#4b), consent forms (#7b) and trial information with up-dated protocol version (#7a), updated SOP for trial medication (#9avi), pocket cards (#9biv), leaflet for clinical staff (#9bv), and eligibility (#9ai), GODIF algorithms (#9avii), GODIF trial medication for notice board (#9bi). Table for ideal body weight and 5% fluid overload according to height (#9bviii) - has changed name to: Table for minimum fluid overload on inclusion and goal for fluid removal - an up-dated document is added. Bedside worksheet is withdrawn/obsolete (#9bvii).

**Date: 21.01.2021:** Added: annual safety report 2020

**Date: 08.01.2021: Added: updated Trial medication dispensing system (site master file #9aiii).**

**Date: 05.01.2021:** Added: data verification plan (site master file #15d) and Front page of protocol signed (site master file #1d).

**Date: 04.01.2021:** Added: protocol 2,5 (site master file #1a), Approval from The Danish Medicine Agency (site master file #4a), EudraCT (site master file 4b), Approval from The Committees on Health Research Ethics (site master file #4c), Trial information (site master file #7a), Consent forms (site master file #7b), Eligibility (site master file #9ai), SAE,SAR,SUSAR and unblinding (site master file #9av), SOP for trial medication (site master file #9avi), Algorithms for trial medication and resuscitation (site master file #9avii), Inclusion and exclusion criteria for notice board (site master file #9bii), Pocket cards (Danish) (site master file #9biv), Leaflet for clinical staff (Danish) (site master file #9bv), Initiation for investigators and primary trial personnel (slides) (site master file #9ci), Primary data source (template and for Sundhedsplatformen) (site master file #12).

**Date: 10.12.2020**: Added: updated co-enrolment list (site master file #8a).

**Date: 13.10.2020**: Added a new plan for data verification from the GCP-unit and a new template for the day form to the eCRF. (site master file #15d and #2).

**Date: 28.09.2020**: Added SOP for trial medication, SOP for escape procedures, and GODIF algorithms in Danish (site master file #9vi and 9vii).

**Date: 15.09.2020**: Added an updated co-enrolment list (site master file #8a)

**Date: 27.08.2020**: Added: new EudraCT file and up-dated approvals and correspondence from The Danish Medicine Agency and The Committees on Health Research Ethics (site master file #4b, #4a, #4c).

**Date: 19.08.2020:** Added: Instructions in the eCRF (site master file #9aiv).

**Date: 04.01.2021:** Added: protocol 2,5 (#1a), Approval from The Danish Medicine Agency (#4a), EudraCT (#4b), approval from The Committees on Health Research Ethics (#4c), Trial information (#7a), Consent forms (#7b).