

## 00.1.0 Authorisation application form



D591

## 00.2.0 Approved trial forms



D592

## 00.3.0 Swiss SAE-forms



D590

## 01.00 Cover letter



M005

## 02.00 Ethics committees decisions applications



M006

## 03.00 Foreign Competent Authority decisions



M007

## 04.00 CIP



M008

## 05.00 CRF



M009



## 06.00 PI ICF



M010

## 07.00 Compliance EN ISO14155



M011

## 08.1.1 IB



M013

## 08.1.2 instructions for use (IFU)



M014

## 08.1.3 additional device information



M015

## 08.1.4 CE marking information



M016

## 08.1.5 IB information on off-label use



M017

## 08.2.0 Applicable standards list



M018



## 08.3.0 Manufacturer statement



M019

## 08.4.0 Documentation access confirmation



M020

## 09.00 Animal origin information



M021

## 10.00 Devit human tissue information



M022

## 11.00 MRI checklist



D593

## 12.1.0 Key aspects radiation protection



M024

## 12.2.0 Licences required



M025

## 12.3.1 Properties radiopharmaceutical



M026



## 12.3.2 Authorised radiopharmaceutical prescribing information



M027

## 12.3.3 Non authorised radiopharmaceutical information



M028

## 12.3.4 CV responsible radiopharmaceutical use



M029

## 12.3.5 Form SFOPH (BAG OFSP UFSP)



M030

## 13.00 Contract with manufacturer



M031

## 14.00 Trial centres list



M032

## 15.00 Other documents



M033

## 16.00 Continued use predecessor model



M034



## 17.1.0 MEDDEV Tables



M036

## 17.2.0 Interim reports



M037

## 17.3.0 Annual safety reports



M038

## 17.4.0 Final report



M039