

EMA/166242/2018 ENCePP Code of Conduct Revision 4



European Network of Centres for Pharmacoepidemiology and Pharmacovigilance

Declaration of compliance with the ENCePP Code of Conduct

The (primary) lead investigator and a person authorised to sign on behalf of the coordinating study entity hereby declare 1 for the purpose of conducting the study <include here study name and identifier/registration no.>: Cohort Study of Long-term Safety of Upadacitinib for the Treatment of Ulcerative Colitis and Crohn's Disease in a Real-world Setting in Europe.

- to follow the rules and principles for the independent and transparent conduct of pharmacoepidemiological and pharmacovigilance studies of the current version of the ENCePP Code of Conduct²;
- to inform the ENCePP Secretariat, of any change or decision to change that constitutes a deviation from the provisions of this Code.

It is of note that the (primary) lead investigator and the person authorised to sign on behalf of the coordinating study entity may be identical.

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Name of (primary) lead investigator: Johan Reutfors	
Date: 24/10/2023 (dd/mm/yyyy)	
Stamp (if applicable) and signature: Johnn Rendfors	
Name of the coordinating study entity: Centre for Pharmacoepidemic	ology (CPE), Karolinska Institutet
Address:	
Karolinska Institutet, CPE Karolinska University Hospital T2 SE-171 76 Stockholm, Sweden	
Name of person authorised to sign on behalf of the coordinatin (primary) lead investigator]:	g study entity [if different from
Date: (dd/mm/yyyy)	
Stamp (if applicable) and signature:	

The (primary) lead investigator should also complete, sign and date the Checklist of the ENCePP Code of Conduct.

 $^{^1}$ Complete the declaration on screen, then print and scan a signed and stamped (if applicable) copy. 2 Adopted Code and any revision thereof at the time of signature of the declaration.