



ENCePP Checklist for Study Protocols (Revision 3)

Adopted by the ENCePP Steering Group on 01/07/2016

The [European Network of Centres for Pharmacoepidemiology and Pharmacovigilance \(ENCePP\)](#) welcomes innovative designs and new methods of research. This Checklist has been developed by ENCePP to stimulate consideration of important principles when designing and writing a pharmacoepidemiological or pharmacovigilance study protocol. The Checklist is intended to promote the quality of such studies, not their uniformity. The user is also referred to the [ENCePP Guide on Methodological Standards in Pharmacoepidemiology](#), which reviews and gives direct electronic access to guidance for research in pharmacoepidemiology and pharmacovigilance.

For each question of the Checklist, the investigator should indicate whether or not it has been addressed in the study protocol. If the answer is "Yes", the section number of the protocol where this issue has been discussed should be specified. It is possible that some questions do not apply to a particular study (for example, in the case of an innovative study design). In this case, the answer 'N/A' (Not Applicable) can be checked and the "Comments" field included for each section should be used to explain why. The "Comments" field can also be used to elaborate on a "No" answer.

This Checklist should be included as an Annex by marketing authorisation holders when submitting the protocol of a non-interventional post-authorisation safety study (PASS) to a regulatory authority (see the [Guidance on the format and content of the protocol of non-interventional post-authorisation safety studies](#)). The Checklist is a supporting document and does not replace the format of the protocol for PASS as recommended in the Guidance and Module VIII of the Good pharmacovigilance practices (GVP).

Study title: The BRodalumab Assessment of Hazards: A Multinational Safety (BRAHMS) study in electronic healthcare databases

Study reference number: EMEA/H/C/003959

Section 1: Milestones	Yes	No	N/A	Section Number
1.1 Does the protocol specify timelines for				
1.1.1 Start of data collection ¹	X	<input type="checkbox"/>	<input type="checkbox"/>	6
1.1.2 End of data collection ²	X	<input type="checkbox"/>	<input type="checkbox"/>	6
1.1.3 Study progress report(s)	X	<input type="checkbox"/>	<input type="checkbox"/>	6
1.1.4 Interim progress report(s)	X	<input type="checkbox"/>	<input type="checkbox"/>	6
1.1.5 Registration in the EU PAS register	X	<input type="checkbox"/>	<input type="checkbox"/>	6

¹ Date from which information on the first study is first recorded in the study dataset or, in the case of secondary use of data, the date from which data extraction starts.

² Date from which the analytical dataset is completely available.

<u>Section 1: Milestones</u>	Yes	No	N/A	Section Number
1.1.6 Final report of study results.	X	<input type="checkbox"/>	<input type="checkbox"/>	6

Comments:

<u>Section 2: Research question</u>	Yes	No	N/A	Section Number
2.1 Does the formulation of the research question and objectives clearly explain:	X	<input type="checkbox"/>	<input type="checkbox"/>	8
2.1.1 Why the study is conducted? (e.g. to address an important public health concern, a risk identified in the risk management plan, an emerging safety issue)	X	<input type="checkbox"/>	<input type="checkbox"/>	8
2.1.2 The objective(s) of the study?	X	<input type="checkbox"/>	<input type="checkbox"/>	8
2.1.3 The target population? (i.e. population or subgroup to whom the study results are intended to be generalised)	X	<input type="checkbox"/>	<input type="checkbox"/>	8
2.1.4 Which hypothesis(-es) is (are) to be tested?	X	<input type="checkbox"/>	<input type="checkbox"/>	8
2.1.5 If applicable, that there is no <i>a priori</i> hypothesis?	<input type="checkbox"/>	<input type="checkbox"/>	X	

Comments:

<u>Section 3: Study design</u>	Yes	No	N/A	Section Number
3.1 Is the study design described? (e.g. cohort, case-control, cross-sectional, new or alternative design)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1
3.2 Does the protocol specify whether the study is based on primary, secondary or combined data collection?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1, 9.2
3.3 Does the protocol specify measures of occurrence? (e.g. incidence rate, absolute risk)	<input type="checkbox"/>	<input type="checkbox"/>	X	
3.4 Does the protocol specify measure(s) of association? (e.g. relative risk, odds ratio, excess risk, incidence rate ratio, hazard ratio, number needed to harm (NNH) per year)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.7
3.5 Does the protocol describe the approach for the collection and reporting of adverse events/adverse reactions? (e.g. adverse events that will not be collected in case of primary data collection)	<input type="checkbox"/>	<input type="checkbox"/>	X	11

Comments:

<u>Section 4: Source and study populations</u>	Yes	No	N/A	Section Number
4.1 Is the source population described?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2
4.2 Is the planned study population defined in terms of:				
4.2.1 Study time period?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2

<u>Section 4: Source and study populations</u>	Yes	No	N/A	Section Number
4.2.2 Age and sex?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2
4.2.3 Country of origin?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2
4.2.4 Disease/indication?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2
4.2.5 Duration of follow-up?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2
4.3 Does the protocol define how the study population will be sampled from the source population? (e.g. event or inclusion/exclusion criteria)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.2

Comments:

<u>Section 5: Exposure definition and measurement</u>	Yes	No	N/A	Section Number
5.1 Does the protocol describe how the study exposure is defined and measured? (e.g. operational details for defining and categorising exposure, measurement of dose and duration of drug exposure)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.3
5.2 Does the protocol address the validity of the exposure measurement? (e.g. precision, accuracy, use of validation sub-study)	<input type="checkbox"/>	X	<input type="checkbox"/>	
5.3 Is exposure classified according to time windows? (e.g. current user, former user, non-use)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1, 9.3, 9.7
5.4 Is exposure classified based on biological mechanism of action and taking into account the pharmacokinetics and pharmacodynamics of the drug?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.3

Comments:

<u>Section 6: Outcome definition and measurement</u>	Yes	No	N/A	Section Number
6.1 Does the protocol specify the primary and secondary (if applicable) outcome(s) to be investigated?	X	<input type="checkbox"/>	<input type="checkbox"/>	8, 9.1, 9.3, 9.7
6.2 Does the protocol describe how the outcomes are defined and measured?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.3
6.3 Does the protocol address the validity of outcome measurement? (e.g. precision, accuracy, sensitivity, specificity, positive predictive value, prospective or retrospective ascertainment, use of validation sub-study)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4, 9.9
6.4 Does the protocol describe specific endpoints relevant for Health Technology Assessment? (e.g. HRQoL, QALYs, DALYS, health care services utilisation, burden of disease, disease management)	<input type="checkbox"/>	X	<input type="checkbox"/>	

Comments:

<u>Section 7: Bias</u>	Yes	No	N/A	Section Number
7.1 Does the protocol describe how confounding will be addressed in the study?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1, 9.7, 9.9
7.1.1. Does the protocol address confounding by indication if applicable?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1, 9.7, 9.9
7.2 Does the protocol address:	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
7.2.1. Selection biases (e.g. healthy user bias)	<input type="checkbox"/>	X	<input type="checkbox"/>	
7.2.2. Information biases (e.g. misclassification of exposure and endpoints, time-related bias)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.9
7.3 Does the protocol address the validity of the study covariates?	<input type="checkbox"/>	X	<input type="checkbox"/>	

Comments:

<u>Section 8: Effect modification</u>	Yes	No	N/A	Section Number
8.1 Does the protocol address effect modifiers? (e.g. collection of data on known effect modifiers, sub-group analyses, anticipated direction of effect)	<input type="checkbox"/>	X	<input type="checkbox"/>	

Comments:

<u>Section 9: Data sources</u>	Yes	No	N/A	Section Number
9.1 Does the protocol describe the data source(s) used in the study for the ascertainment of:				
9.1.1 Exposure? (e.g. pharmacy dispensing, general practice prescribing, claims data, self-report, face-to-face interview)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.1.2 Outcomes? (e.g. clinical records, laboratory markers or values, claims data, self-report, patient interview including scales and questionnaires, vital statistics)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.1.3 Covariates?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.2 Does the protocol describe the information available from the data source(s) on:				
9.2.1 Exposure? (e.g. date of dispensing, drug quantity, dose, number of days of supply prescription, daily dosage, prescriber)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.2.2 Outcomes? (e.g. date of occurrence, multiple event, severity measures related to event)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.2.3 Covariates? (e.g. age, sex, clinical and drug use history, co-morbidity, co-medications, lifestyle)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.3 Is a coding system described for:				
9.3.1 Exposure? (e.g. WHO Drug Dictionary, Anatomical Therapeutic Chemical (ATC) Classification System)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.3.2 Outcomes? (e.g. International Classification of Diseases (ICD)-10, Medical Dictionary for Regulatory Activities (MedDRA))	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4
9.3.3 Covariates?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4

<u>Section 9: Data sources</u>	Yes	No	N/A	Section Number
9.4 Is a linkage method between data sources described? (e.g. based on a unique identifier or other)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.4

Comments:

<u>Section 10: Analysis plan</u>	Yes	No	N/A	Section Number
10.1 Is the choice of statistical techniques described?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.7
10.2 Are descriptive analyses included?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.7
10.3 Are stratified analyses included?	<input type="checkbox"/>	<input type="checkbox"/>	X	
10.4 Does the plan describe methods for adjusting for confounding?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.1, 9.7
10.5 Does the plan describe methods for handling missing data?	<input type="checkbox"/>	X	<input type="checkbox"/>	
10.6 Is sample size and/or statistical power estimated?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.5

Comments:

<u>Section 11: Data management and quality control</u>	Yes	No	N/A	Section Number
11.1 Does the protocol provide information on data storage? (e.g. software and IT environment, database maintenance and anti-fraud protection, archiving)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.8
11.2 Are methods of quality assurance described?	X	<input type="checkbox"/>	<input type="checkbox"/>	9.8
11.3 Is there a system in place for independent review of study results?	<input type="checkbox"/>	X	<input type="checkbox"/>	

Comments:

<u>Section 12: Limitations</u>	Yes	No	N/A	Section Number
12.1 Does the protocol discuss the impact on the study results of:				
12.1.1 Selection bias?	X	<input type="checkbox"/>	<input type="checkbox"/>	
12.1.2 Information bias?	<input type="checkbox"/>	X	<input type="checkbox"/>	
12.1.3 Residual/unmeasured confounding? (e.g. anticipated direction and magnitude of such biases, validation sub-study, use of validation and external data, analytical methods)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.9
12.2 Does the protocol discuss study feasibility? (e.g. study size, anticipated exposure, duration of follow-up in a cohort study, patient recruitment)	X	<input type="checkbox"/>	<input type="checkbox"/>	9.5

Comments:

<u>Section 13: Ethical issues</u>	Yes	No	N/A	Section Number
13.1 Have requirements of Ethics Committee/ Institutional Review Board been described?	X	<input type="checkbox"/>	<input type="checkbox"/>	10
13.2 Has any outcome of an ethical review procedure been addressed?	<input type="checkbox"/>	X	<input type="checkbox"/>	
13.3 Have data protection requirements been described?	X	<input type="checkbox"/>	<input type="checkbox"/>	10

Comments:

<u>Section 14: Amendments and deviations</u>	Yes	No	N/A	Section Number
14.1 Does the protocol include a section to document amendments and deviations?	X	<input type="checkbox"/>	<input type="checkbox"/>	5

Comments:

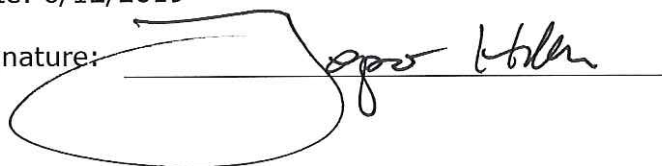
<u>Section 15: Plans for communication of study results</u>	Yes	No	N/A	Section Number
15.1 Are plans described for communicating study results (e.g. to regulatory authorities)?	X	<input type="checkbox"/>	<input type="checkbox"/>	12
15.2 Are plans described for disseminating study results externally, including publication?	X	<input type="checkbox"/>	<input type="checkbox"/>	12

Comments:

Name of the main author of the protocol: Jesper Hallas

Date: 6/12/2019

Signature:



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