

# **Declaration of Interests for ENCePP SEAL Studies**

# INTRODUCTION

This document includes your personal details and your declaration of interests to be made public in line with the provisions of the ENCePP Code of Conduct. All parts must be duly completed. Your declaration will not be accepted if any fields are left empty. You are responsible for the accuracy and completeness of the submitted information.

The form is designed to be filled in electronically and to be transmitted to the ENCePP Secretariat by email; in parallel, a copy of the form should be uploaded to the <u>EU PAS Register</u>.

# SECTION 1: PERSONAL DETAILS

First Name:	Lisbeth Munksgaard	
Last Name:	Baggesen	
Organisation / Research Centre :	Department of clinical epidemiology, Aa	rhus University Hospital
Country:	Denmark	
Contact e-mail Address:	lisbeth@clin.au.dk	
	Intravenous Iron Postauthorisation Safe Severe Hypersensitivity Reactions	ety Study (PASS): Evaluation of the Risk of
Study Reference Number:	EUPAS 2 0 7 2 0	
Are you the (Primary) Lead	Investigator of the above study?	No 🖌 Yes
Are you an investigator/res	earcher contributing to the above study	No 🗌 Yes 🖌

### SECTION 2: DECLARATION OF INTERESTS RELATED TO PHARMACEUTICAL INDUSTRY

In this section you must declare any interests in the pharmaceutical industry that you currently have or had within the past 3 years. If you have interests to declare please tick 'Yes' to the relevant questions. All questions in this part must be answered.

## 2.1 Employment

No 🕢 Yes 🔾

Yes ()

### Employment in a pharmaceutical company during past 3 years of study application?

Pharmaceutical company includes supply or service companies which contribute to research, development, production and maintenance of a medicinal product. Employment relates to salaries currently being directly paid to you by a pharmaceutical company.

## 2.2 Financial Interest

### Financial interests in the capital of a pharmaceutical company?

Financial interests relate to current holding of shares of a pharmaceutical company with the exclusion of independently managed investment funds/pensions schemes that are not exclusively based on the pharmaceutical sector.

No

# 2.3 Patent

### Patent for a medicinal product?

Relates to a patent for a medicinal product currently owned by either you as individual or your organisation/ research centre, and you as individual are the beneficiary.

# 2.4 Consultancy



Yes (

No 🕢 Yes 🔿

### Consultancy for a pharmaceutical company during the past 3 years of study application?

Consultancy refers to provision of advice or services to a pharmaceutical company excluding the concerned study and including but not limited to reviewing activities, data monitoring, statistical analysis, end point committees, regardless of contractual arrangements or any form of remuneration such as consulting fees or honoraria. Note that conference/seminar attendance is not considered a consultancy but should be indicated if subject to fee or honorarium.

No

 $(\mathbf{\Lambda})$ 

### 2.5 Strategic Advisory Role

Strategic Advisory role on activities of a pharmaceutical company during the past 3 years of study application?

Participation with the right to vote on/influence the output in a (scientific) advisory board/steering committee with the role of providing advice/expressing opinions on the future strategy, direction or development activities of a pharmaceutical company either in terms of general or product-related strategy, regardless of contractual arrangements or any form of remuneration.

# 2.6 Grant / Funding

Grant/funding from a pharmaceutical company other than funds contemplated in the concerned study contract?

Refers to a grant or funding from a pharmaceutical company which is currently being received by your research group, irrespective of whether you are employed or a volunteer, and you receive no personal gain.

### SECTION 3: ANY OTHER INTERESTS

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Any other relationships/conditions/circumstances that present a potential conflict of interest, including matters relating to members of your family?

Further to the interests declared above, I do hereby declare on my honour that I do not have any further interests or facts that should be made public in relation to the conduct of the study.

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FULL NAME:	Lisbeth Munksgaard Baggesen	Date:	13/03/2019	
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No 🕢 Yes 🔾

No 🕢 Yes 🔿

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# **SECTION 1: PERSONAL DETAILS**

First Name:	Cécile
Last Name:	Droz-Perroteau
Organisation / Research Centre :	Bordeaux PharmacoEpi - Bordeaux CIC 1401
Country:	France
Contact e-mail Address:	cecile.droz@u-bordeaux.fr
	Intravenous Iron Postauthorisation Safety Study (PASS): Evaluation of the Risk of Severe Hypersensitivity Reactions
Study Reference Number: E	UPAS 2 0 7 2 0
Are you the (Primary) Lead	Investigator of the above study? No $\checkmark$ Yes
Are you an investigator/rese	earcher contributing to the above study No $\square$ Yes $\checkmark$

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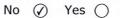
Please specify the pharmaceutical company:



No 🕢 Yes 🔿

No 🕢 Yes 🔿

No () Yes ()



No 🕢 Yes 🔿

No () Yes ()



Name of Pharmaceutical Company	Comments		
ALK-Abello	Impact of a desensitization with OSIRIS® APSI on healthcare resource use for the treatment of allergies and asthma		
Aptalis	Drug utilisation study, safety and pharmacokinetic study of Pylera® in real-life		
AstraZeneca	Benefit-risk of antiplatelet agents during secondary prevention of acute coronary syndrome in France		
AstraZeneca	Health outcomes, resource use, costs in patients with stable coronary artery disease in France		
AstraZeneca	Assessment of the high risk and unmet need in patients with coronary artery disease and type 2 diabetes in France		
Bayer	Benefit-risk of arterial thrombotic prevention with rivaroxaban for atrial fibrillation in France		
Biogen France	Effectiveness of Tecfidera® in multiple sclerosis in France		
Biogen USA	Multiple sclerosis natural history study in France		
BMS	Benefit-risk of antithrombotic treatments after orthopedic surgery in France		
Boehringer	Benefit-risk of anticoagulants in nonvalvular atrial fibrillation in France		
Boehringer	Educational grant for research activities in COPD disease management		
Janssen Cilag	Therapeutic strategy in metastatic castration-resistant prostate cancer in France		
Janssen France	Assessment of the treatment-resistant depression in France		
Lundbeck	Assessment of Selincro® in real-life		
Merck KGaA	Malignancies in multiple sclerosis in France		
Merck Serono	Assessment of targeted therapies in patients with colorectal cancer treated with Erbitux® in first line treatment		
Novartis	Post inscription studies of ranibizumab for visual impairment due to diabetic macular edema and retinal vein occlusion		
Novartis	Assessment of the inhalation systems handling in patients with COPD in real-life		
Novartis Pharma AG	A Phase IIIb, multicentre, open-label study of nilotinib in adult patients with newly diagnosed Philadelphia chromosome positive and/or BCR-ABL positive CML in chronic phase		
Pierre Fabre	Observational cohort study of myocardial infarction with long-term follow-up in France		
Sanofi-Aventis	Assessment of cabazitaxel in real-life		
Stallergenes	Assessment of Oralair® in real-life		
Shire	Incidence, prevalence and burden of dry eye disease in France		

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No 🕢 Yes 🔿

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FULL NAME: Cécile Droz-Perroteau	Date:	07/02/2019	
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First Name:	Uffe					
Last Name:	Heide-Jørgensen					
Organisation / Research Centre :	Department of Clinical Epidemiology, Aa	irhus Univ	ersity ł	lospital		
Country:	Denmark			1		
Contact e-mail Address:	uhj@clin.au.dk					
	Intravenous Iron Postauthorisation Safe Severe Hypersensitivity Reactions	ty Study	(PASS):	Evaluat	ion of the	Risk of
Study Reference Number:	EUPAS 2 0 7 2 0					
Are you the (Primary) Lead	Investigator of the above study?	No 🗸	Yes			
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No

 $\bigcirc$ 

No 🕢

No 🔿

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Department of Clinical Epidemiology carries out research studies funded by several pharmaceutical companies through institutional research grants to and administered by Aarhus University.

No 🕢 Yes ()

No 🕢 Yes ()

No 🕢

Yes 🔘



Yes 🔘

Yes 🔘

Yes 🕢

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FULL NAME: Uffe Heide-Jørgensen Date: 11/02/2020

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