Curriculum vitae

Date of birth **83.06.24**
Professional Address: LINK Medical Research AS

EDUCATION (most current date first)		
Degree	**Institution/School, Country**	**Year**
Master/Pharmaceutical Bioscience (with a focus on clinical trials)	University of Gothenburg Sweden	2002-2007

CURRENT AND PREVIOUS RELEVANT POSITIONS (most current date first)		
Title	**Institution, Company, Country**	**Period**
Regulatory Associate	Link Medical Research AS, Sweden (in-house and out-sourced to Allergan (former Actavis))	Jan 2015-Ongoing
Clinical Research Associate	Link Medical Research AS, Sweden (out-sourced to Bayer)	Jan 2014- Jan 2015
Regulatory Associate	Link Medical Research AS, Norway-Sweden	April 2013-Dec 2013
Clinical Research Associate	Link Medical Research AS, Norway	Aug 2011-April 2013
Clinical Research Associate	TrialFormSupport AS, Norway (out-sourced to Sanofi)	April 2008-June 2011

PROFESSIONAL EXPERIENCE (most current date first)		
Function / Responsibility	**Therapy area and phase** (if relevant)	**Period**
Allergan • Responsible for a product portfolio (24 products) • Content proof reading • Print proof reading • Art work requests • Variation applications to all 5 countries • Update of PI	Nordic responsibility (Sweden, Denmark, Finland, Norway and Iceland)	May 2015-ongoing
Readability test	Responsible for a readability test performed in Sweden.	2015
Variation application	To the Swedish Authorities	2015
Information search	Regulatory authorities (SE/DK/NO)	2015
Update of SmPC and PIL	Swedish/Norwegian	2013,2015
Translations of SmPC and PIL	To Swedish/Norwegian/English	2013, 2015
Review of SmPC and PIL	Swedish/Norwegian/English	2013, 2015
Proof-reading of Mock-ups	Swedish/Norwegian	2015
Country lead Monitor (monitoring, submissions to EC, budget handling, an overall management of the study)	Lymphoma II	June 2014-Jan 2015
Monitoring/Close-out	Diabetes nephropathy II	Jan 2014-Nov 2014
Applications sent to the Danish and Norwegian Data Protection Authorities	Hypertension/ non-interventional study with medical device	2013

Initiation/Monitoring	Pancreatic cancer I/II	Nov2012-Jun2013
Monitoring	Intestinal cancer III	Feb 2011 to June 2011
Selection/initiation/close out	Atrial fibrilliation III	Aug 2008 to May 2011
Monitoring/close out	Pulmonary embolism III	Aug 2008 to April 2011
Monitoring	Diabetes III	Feb 2009 to June 2011
Monitoring/data management	Hypertension III	April 2008 to Nov 2008
Contract negotiation/Amendments	Lung cancer III	Oct 2011-Sep2012
Feasibilities and capabilities		2013
Data management/initiation/monitoring/close out	PB observational study IV	Apr 2008-Oct 2008

COURCES / CERTIFICATIONS (most current date first) **Description**	**Date Institution, Country**	**Time & Duration**
Basic course in Regulatory Affairs (with certification) A 3 days course covering all the basic needs.	Läkemedelsakademin Sweden	3-5 Sep 2013
Excel course	Sweden	2014 (1 workingday)
"Val av och samarbete med CRO/konsultföretag inom Regulatory Affairs"	Sweden/Läkemedelsakademin	2013 (1 workingday)
Different GCP courses over the years, most current one was provided by Barnett	Norway-Sweden	2008-2014
Word course	Norway	2010
Registered and approved as Clinical Research Associate (CRA), by the LMI (pharmaceutical industry), arranged by the Norwegian Association of Pharmaceutical Manufacturers Association of Foreign Pharmaceutical Companies in Norway.	LMI, Norway	Jan 2010, 06 Days

LANGUAGE SKILLS	<<mother tongue, fluent, intermediate, beginner level>>
English	Intermediate
Swedish	Fluent
Norwegian	Intermediate
Farsi	Mother Tongue

COMPUTER SKILLS	<<advanced, intermediate, basic>>
Word, intermediate	
Excel, intermediate	
Power Point, intermediate	
eCRF system, intermediate	

PUBLICATION LIST

EXTERNAL LECTURING

OTHER

Date	Signature