EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Vicky Cann
AsktheEU

team@asktheEU.org

22 July 2014
EMA/446934/2014
Procedure Management & Business Support Division

Dear Ms Cann

Subject: All completed declarations of interest for Jan Regnstrom ASK-4245 – **Release letter to the requester**

Thank you for your request for access to documents received on 14 July 2014 in which you apply for copies of the following documents concerning Jan Regnstrom, an employee of the European Medicines Agency, in particular:

- All completed declarations of interest.

Your request has been handled in accordance with Article 7 of Regulation (EC) No 1049/2001 regarding public access to European Parliament, Council and Commission documents (the Regulation)[1] and Article 6 of the Rules for the Implementation of Regulation (EC) No 1049/2001 on access to European Medicines Agency documents (the Agency rules)[2]. Moreover, it has been assessed pursuant to Article 4 of the Regulation, Article 3 of the Agency rules and the European Medicines Agency policy on access to documents related to medicinal products for human and veterinary use (the Agency policy)[3].

Having assessed your application, the Agency considers that access to the documents requested should be granted. However, the documents have been redacted in accordance with Article 4(1)(b) of the Regulation and the European Union legislation regarding the protection of personal data. All protected personal data was redacted in order to avoid that the disclosure of the document would undermine the privacy and integrity of any individual.

You may submit a confirmatory application in writing against this decision to the European Medicines Agency, within 15 working days. Should you wish to do so, you are kindly invited to provide your reasons against our decision to redact parts of the documents at this stage, which you believe should be taken into account by the Agency in adopting a final decision.

The confirmatory application should be submitted using the on-line request form, available on the European Medicines Agency website, under the following location:

http://www.ema.europa.eu/ema/index.jsp?curl=pages/about_us/landing/ask_ema_landing_page.jsp&mid=WC0b01ac05806499f0

Please use the ASK Procedure Number mentioned in the subject line in any correspondence related to this request.

1 OJ L 145, 31.5.2001, P. 43-48
2 EMEA/MB/203359/2006 Rev 1 Adopted
3 EMA/110196/2006 of 30 November 2010

All the documents concerned will shortly be sent to you via Eudralink. Please note that these documents are made available to you in order to provide you with access in accordance with the Regulation and the Agency rules. In that regard, please visit the Agency's public website to know more about the applicable copyright and limited reproduction notices.

If you have any queries on the enclosed, please do not hesitate to contact the Access to Documents Coordinator for this request, [redacted] e-mail: [redacted] using the ASK Procedure Number mentioned in the subject line.

Yours sincerely,

[redacted]

Dr Anne-Sophie Henry-Eude
Head of Access to Document Service
Business Data & Support Department

[redacted]

Legal Administrator
Legal Department