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Regulating Medicines and Medical Devices

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11/03/2015

Dear Mr S Purnell

THE MEDICINES FOR HUMAN USE (CLINICAL TRIALS) REGULATIONS 2004 S.I. 2004/1031

Our Reference:	20363/0273/001-0013
Eudract Number:	2009-012717-22
Product:	rituximab
Protocol number:	UCL/08/0167
Substantial Amendment Code Number:	Code Number: Substantial Amendment 03.02.2015
Version: 6.1	
Date: 2015/01/27	

NOTICE OF ACCEPTANCE OF AMENDMENT

I am writing to inform you that the Licensing Authority accepts the proposed amendment to your clinical trial authorisation (CTA), received on 10/02/2015.

This amendment may therefore be made.

You are reminded that where it is appropriate, the Ethics Committee should also be notified of amendments.

Yours sincerely,

Clinical Trials Unit
MHRA

Medicines and Healthcare Products Regulatory Agency

RECEIVED 13 MAR 2015
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