



Medicines & Healthcare products
Regulatory Agency



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Ms Heather House
UNIVERSITY OF OXFORD
JOINT RESEARCH OFFICE, 1ST FLOOR BOUNDARY BROOK HOUSE,
CHURCHILL DRIVE, HEADINGTON
OXFORD
OX3 7GB
UNITED KINGDOM

15/03/2022

Dear Ms Heather House,

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

Our Reference:	CTA 21584/0423/001-0025
Eudract Number:	2020-001113-21
Product:	Hydrocortisone, RoActemra, Kineret , Dexamethasone , Prednisolone, Empagliflozin, Oseltamivir, Baloxavir, Sotrovimab, Molnupiravir, Nirmatrelvir/ritonavir
Protocol number:	NDPHRECOVERY
Substantial Amendment Code Number:	Substantial Amendment 25

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 10/03/2022.

MEDICAL - Remarks: Clinical remark:

The Paxlovid SmPC provided with this amendment is the one that was reviewed in December 2021. The Sponsor is reminded to submit the Paxlovid SmPC reviewed in February 2022 at the time of the next substantial amendment. The Feb 2022 SmPC will be reviewed and the Sponsor will decide whether changes to the trial protocol are warranted.

This amendment may therefore be made.

If applicable, you should ensure your trial details have been updated on the database where you have registered your trial.



You are reminded that from 1 January 2022 you will need to comply with the requirements specified in the following guidance, where applicable:

o Import of IMPs from listed countries to GB:

<https://www.gov.uk/government/publications/importing-investigational-medicinal-products-into-great-britain-from-approved-countries>

o Supply of IMPs to Northern Ireland:

<https://www.gov.uk/guidance/supplying-investigational-medicinal-products-to-northern-ireland>

o Substantial amendments to clinical trials:

<https://www.gov.uk/guidance/guidance-on-substantial-amendments-to-a-clinical-trial>

Any required substantial amendment to your Clinical Trial Authorisation should be submitted and approved as soon as possible and before 1 January 2022.

Yours sincerely,

**Clinical Trials Unit
MHRA**