

Internal Review of FOI 23/882

Dear

As you will be aware, we've logged your email of 13 December 2023 regarding the response to FOI 23/882 as an internal review.

You have stated in your email below that the highlighted text on page 3 of the UKHSA document shows "*UKHSA recommending Oseltamir treatment immediately if the human patient meets case definition for avian influenza.*"

However, having checked the document that you attached, your statement is not correct, as this is not what the highlighted text on page 3 says. It says, "*Start oseltamivir treatment immediately if the patient meets case definition for avian influenza.*"

To confirm, the UKHSA document you've attached says '**oseltamivir**' and not, as you've said in your email, '**Oseltamir**'. I have included a screenshot of the wording you highlighted to clarify this:

Start oseltamivir treatment immediately if the patient meets case definition for avian influenza. For guidance on dosage refer to [UKHSA guidance \(https://www.gov.uk/government/publications/influenza-treatment-and-prophylaxis-using-anti-viral-agents\)](https://www.gov.uk/government/publications/influenza-treatment-and-prophylaxis-using-anti-viral-agents) on the use of antiviral agents for the treatment and prophylaxis of seasonal influenza.

The online version of the UKHSA document also refers to oseltamivir, not oseltamir:

<https://www.gov.uk/government/publications/avian-influenza-guidance-and-algorithms-for-managing-human-cases/investigation-and-initial-clinical-management-of-possible-human-cases-of-avian-influenza-with-potential-to-cause-severe-human-disease>

As stated in our response to FOI 23/882, the MHRA has granted Marketing Authorisations for Oseltamivir. Please find a link to the Public Assessment Report for Oseltamivir 30 mg, 45 mg and 75 mg hard capsules (PL 00142/1278-128) below: [5d2c770dfac07e0c8f8ffa0d3c1452690356945e \(windows.net\)](https://www.mhra.gov.uk/assessments/5d2c770dfac07e0c8f8ffa0d3c1452690356945e)

We hope that this provides the clarification you seek.

If you remain dissatisfied, you may ask the Information Commissioner (ICO) to make a decision on whether or not we have interpreted the FOIA correctly in dealing with the request and subsequent internal review. The ICO's address is:

The Information Commissioner's Office
Wycliffe House
Water Lane
Wilmslow
Cheshire
SK9 5AF

Yours sincerely,

MHRA Customer Experience Centre

Communications and engagement team

Medicines and Healthcare products Regulatory Agency

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