

## **FOI 22/1209 - Risk Management Plan for Eylea**

**12 January 2023**

### **MHRA RESPONSE**

Thank you for your email, dated 20<sup>th</sup> December 2022, in which you requested:

**“a copy of the Risk Management Plan for Eylea (afibercept)”**

We can confirm that the MHRA holds a copy of the requested RMP.

Information that has been redacted is exempt under Section 40 (Personal Information) or Section 43 (Commercial Interests) of the Freedom of Information (FOI) Act and is therefore withheld.

Section 40 provides that personal information may be exempt from release where to do so would contravene data protection principles. Section 43 provides that information will be exempt from release where to do so would or would be likely to prejudice commercial interests. Furthermore, we do not believe that there is an overriding public interest in disclosing the redacted information in this instance.

We hope the information provided is helpful, but if you are dissatisfied with the handling of your request, you have the right to ask for an internal review. Internal review requests should be submitted within two months of the date of this response; and can be addressed to this email address.

Please remember to quote the reference number above in any future communications.

Yours sincerely,

#### **FOI Team,**

Safety & Surveillance Group

Medicines and Healthcare Products Regulatory Agency

10 South Colonnade, Canary Wharf, London E14 4PU

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Medicines and Healthcare products Regulatory Agency, (via this email address). After that, if you remain dissatisfied, you may ask the Information Commissioner at:

The Information Commissioner's Office  
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