

**IN THE UNITED STATES DISTRICT COURT
FOR THE DISTRICT OF COLUMBIA**

JUDICIAL WATCH, INC.,)	
425 Third Street SW, Suite 800)	
Washington, DC 20024,)	
)	
Plaintiff,)	Civil Action No.
)	
v.)	
)	
U.S. DEPARTMENT OF HEALTH)	
AND HUMAN SERVICES,)	
200 Independence Avenue SW)	
Washington, DC 20201,)	
)	
Defendant.)	
_____)	

COMPLAINT

Plaintiff Judicial Watch, Inc. (“Plaintiff”) brings this action against Defendant U.S. Department of Health and Human Services (“Defendant”) to compel compliance with the Freedom of Information Act, 5 U.S.C. § 552 (“FOIA”). As grounds therefor, Plaintiff alleges as follows:

JURISDICTION AND VENUE

1. The Court has jurisdiction over this action pursuant to 5 U.S.C. § 552(a)(4)(B) and 28 U.S.C. § 1331.
2. Venue is proper in this district pursuant to 28 U.S.C. § 1391(e).

PARTIES

3. Plaintiff Judicial Watch, Inc. is a not-for-profit, educational organization incorporated under the laws of the District of Columbia and headquartered at 425 Third Street SW, Suite 800, Washington, DC 20024. Plaintiff seeks to promote transparency, integrity, and accountability in government and fidelity to the rule of law. As part of its mission, Plaintiff

regularly requests records from federal agencies, analyzes the responses it receives, and disseminates its findings and the records to the American public to inform them about “what their government is up to.”

4. Defendant U.S. Department of Health and Human Services is an agency of the U.S. Government and is headquartered at 200 Independence Avenue SW, Washington, DC 20201. Defendant has possession, custody, and control of records to which Plaintiff seeks access.

STATEMENT OF FACTS

5. On June 1, 2022, Plaintiff submitted a FOIA request via email to the National Institutes of Allergy and Infectious Diseases (“NIAID”), a component of Defendant, seeking the following records:

1. All safety studies, data, reports, and analyses produced by the Division of Microbiology and Infectious Diseases (DMID) relating to the safety of ‘vaccines’ and/or gene therapies to treat and/or prevent SARS-CoV-2 and/or COVID-19 made by Pfizer, BioNTech, Moderna, Johnson & Johnson, and Janssen.
2. All emails sent to and from the following DMID officials relating to the safety of ‘vaccines’ and/or gene therapies to treat and/or prevent SARS-CoV-2 and/or COVID-19 made by Pfizer, BioNTech, Moderna, Johnson & Johnson, and Janssen:
 - a. The Director of DMID
 - b. The head of the Office of Genomics & Advanced Technologies
 - c. The head of the Office of International Research in Infectious Diseases
 - d. The head of the Office of Regulatory Affairs
 - e. The head of the Office of Clinical Research Affairs
 - f. The head of the Clinical Trials Management Section
 - g. The head of the Virology Branch
 - h. The head of the Respiratory Diseases Branch
 - i. The head of the Influenza, SARS, and Other Viral Respiratory Diseases Section

The time frame for the records sought is June 1, 2020 to the present.”

6. NIAID acknowledged receipt of the FDA request on June 1, 2022 and assigned the request No. 58421.

7. Between June 21, 2022 and June 27, 2022, Plaintiff and a policy analyst from Defendant's FOIA department agreed to narrow the scope of the request. Defendant never responded to Plaintiff's June 27, 2022 email.

8. As of the date of this Complaint, Defendant has failed to: (i) determine whether to comply with the request; (ii) notify Plaintiff of any such determination or the reasons therefor; (iii) advise Plaintiff of the right to appeal any adverse determination; or (iv) produce the requested records or otherwise demonstrate that the requested records are exempt from production.

COUNT I
(Violation of FOIA, 5 U.S.C. § 552)

9. Plaintiff realleges paragraphs 1 through 8 as if fully stated herein.

10. Defendant is in violation of FOIA.

11. Plaintiff is being irreparably harmed by Defendant's violation of FOIA, and Plaintiff will continue to be irreparably harmed unless Defendant is compelled to comply with the law.

12. Plaintiff has no adequate remedy at law.

13. To trigger FOIA's administrative exhaustion requirement, Defendant was required to make final determinations on Plaintiff's request by August 12, 2022, at the latest. Because Defendant failed to make final determinations on Plaintiff's requests/appeal within the time limits set by FOIA, Plaintiff is deemed to have exhausted its administrative appeal remedies.

WHEREFORE, Plaintiff respectfully requests that the Court: (1) order Defendant to conduct a search for any and all records responsive to Plaintiff's FOIA requests and demonstrate that it employed search methods reasonably likely to lead to the discovery of records responsive to Plaintiff's FOIA requests; (2) order Defendant to produce, by a date certain, any and all non-exempt records responsive to Plaintiff's FOIA requests and *Vaughn* indices of any responsive records withheld under claim of exemption; (3) enjoin Defendant from continuing to withhold any and all non-exempt records responsive to Plaintiff's FOIA requests; (4) grant Plaintiff an award of attorneys' fees and other litigation costs reasonably incurred in this action pursuant to 5 U.S.C. § 552(a)(4)(E); and (5) grant Plaintiff such other relief as the Court deems just and proper.

Dated: October 17, 2022

Respectfully submitted,

/s/ Meredith Di Liberto

MEREDITH DI LIBERTO

D.C. Bar No. 487733

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