

Hoppe Capt Joshua P

From: Hoppe Capt Joshua P
Sent: Friday, June 24, 2022 2:14 PM
To:
Cc:

Subject: [REDACTED]
RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Good Afternoon Gentlemen,

Thank you for the response. The only reason I had a timeline for this request was due to the timelines that many of us have due to what is being offered to us as the "Comirnaty-labeled" product. If the DOD agrees to cease all further actions of the enforcement of this mandate, then I am more than happy to wait as long as necessary for the answers to the legitimate concerns raised from all the appropriate agencies. I have already engaged with the medical clinic and Pfizer directly and they both directed me to contact the DHA (in charge of the vaccine rollout) to answer the questions I have after refusing to answer directly. Please let me know which questions you can answer at this time and I will resubmit the others via FOIA. If there are questions that would be more appropriate for other agencies, please identify which ones and which agencies to prevent this from taking longer than necessary. Thank you again for your time and assistance with these concerns. Have a great weekend!

Semper Fidelis,

Capt Joshua "Hippity" Hoppe
ASO/NATOPS/Education Officer
Office

[REDACTED]

-----Original Message-----

From: [REDACTED]@usmc.mil>
Sent: Friday, June 24, 2022 10:12 AM
To: Hoppe Capt Joshua P <[REDACTED]@usmc.mil>;
[REDACTED]@usmc.mil>;
[REDACTED]@usmc.mil>

Cc: [REDACTED]@usmc.mil>;
[REDACTED]@usmc.mil>; [REDACTED]

[REDACTED]
[REDACTED]
[REDACTED] mail.mil>

Subject: RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Captain Hoppe,

Good deal that you submitted a request to DHA. Copy.

Judge Advocate Division can weigh-in, but here's my recommendation for answering the many questions you have, which, of note, span different entities, echelons, agencies, etc.

1. You can submit additional FOIA requests to additional entities, to include Headquarters Marine Corps.
2. If you have specific questions about informed consent processes/documents, temperatures of particular vaccine doses or labeling at a particular MTF at a particular time, or any other MTF-specific logistics/policy questions, you can engage your local MTF, professionally and politely, and without inappropriate timeline demands. If that MTF cannot answer, they may be able to push appropriate question(s) up through BUMED and/or DHA channels.

At this point, I'll defer all further correspondence from you to HQMC over to Judge Advocate Division.

Thanks very much for your patience, Captain Hoppe, and I wish you the best.

Break...

[REDACTED], over to you all.

V/R - CD [REDACTED]

-----Original Message-----

From: Hoppe Capt Joshua P <[REDACTED]@usmc.mil>

Sent: Friday, June 24, 2022 11:47 AM

To: [REDACTED]@usmc.mil>

Cc: [REDACTED]@usmc.mil>; [REDACTED]@usmc.mil>;

[REDACTED]@usmc.mil>; [REDACTED]

[REDACTED]@usmc.mil>

Subject: RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Good Morning Sir,

I did submit a FOIA request to the DHA on 21 Jun 22 to try and get this ball rolling, but do not agree that this is necessary to answer the RFIs that I sent your office. The additional RFIs that I sent your office yesterday are not on the FOIA request either. Please let me know which ones you can provide answers to as I believe they are all very pertinent medical treatment concerns as it relates to my informed consent as protected by the DoD's Patient Bill of Rights and are of interest for our Command as they

will need these answers in order to lawfully comply with any enforcement of this inoculation order. I have requested that all enforcement ceases until these questions can be appropriately addressed and answered. Thank you for your time and attention to these concerns.

Semper Fidelis,

Capt Joshua "Hippity" Hoppe
ASO/NATOPS/Education Officer
Office
[REDACTED]
[REDACTED]@usmc.mil

-----Original Message-----

From: [REDACTED]@usmc.mil>
Sent: Friday, June 24, 2022 5:46 AM
To: [REDACTED]@usmc.mil>
Cc: [REDACTED]@usmc.mil>; [REDACTED]@usmc.mil>; [REDACTED]@usmc.mil>
Subject: RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Captain Hoppe, did you, or did you not, submit a FOIA request to DHA regarding your questions?

V/R - CDR [REDACTED]

-----Original Message-----

From: Hoppe Capt Joshua P <[REDACTED]@usmc.mil>
Sent: Thursday, June 23, 2022 6:50 PM
To: [REDACTED]@usmc.mil>
Cc: [REDACTED]@usmc.mil>; [REDACTED]@usmc.mil>; [REDACTED]@usmc.mil>
Subject: RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Good Afternoon Sir,

I am following up about the request that I sent you on Tuesday. Attached is the DoD Patient Bill of Rights for reference. In enclosure 2, para 1.f, one of my rights as a patient to Informed Consent states:

"Patients have the right to any and all necessary information in nonclinical terms to make knowledgeable decisions on consent or refusal for treatments, or participation in clinical trials or other research investigations as applicable. Such information is to include any and all complications, risks,

benefits, ethical issues, and alternative treatments as may be available."

My requests below are directly linked to my right to informed consent per this instruction. Please provide the information requested at your earliest availability. I do not believe this is an appropriate avenue for a FOIA request which according to the site provided takes 36-252 days on average for the request to be completed from the DHA. I have already engaged the military treatment facilities and they had directed me to reach out to the DHA for these concerns. Below are a couple additional concerns that I have about the treatment I am being ordered to receive:

- What temperature does a package of the "Comirnaty-labeled" products have to reach before an alarm goes off (while shipping or after), have any temperature alarms been set off at any point, and what actions have occurred as a result?
- Why aren't these vials to be diluted in contrast to some of the other inoculation products already in circulation?
- What studies have been done on the DoD military personnel for the "Warnings and Precautions" section and specifically of Myocarditis and Pericarditis adverse events that are listed in the Comirnaty (Gray Cap) Package Insert?
- What are the current injury rates of these adverse events occurring for service members who have already taken any of the mRNA products?
- From the CDC Pinkbook, there are no COVID-19 vaccines added to the Vaccine Injury Table yet. When will that occur or has there already been a solicitation for public comments on the VIS?
- When did the DoD first have "Comirnaty" in stock and where?
- I was previously told that to have an FDA-approved "Comirnaty" vaccine provided to me would be a "herculean effort". Please explain why this was the case as the DHA have been the ones responsible for the roll out and delivery of the COVID-19 inoculation products.
- Any other supporting documentation that would help verify that these are the FDA-approved products as described in the BLA-Approval/Revision letters previously sent.

Thank you again for your time and assistance with this information.

Semper Fidelis,

Capt Joshua "Hippity" Hoppe
ASO/NATOPS/Education Officer
Office:

@usmc.mil
[REDACTED]

-----Original Message-----

From: Hoppe Capt Joshua P
Sent: Tuesday, June 21, 2022 3:47 PM
To: @usmc.mil>
Cc: @usmc.mil>; R [REDACTED]
[REDACTED]

██████████@usmc.mil>

Subject: RE: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Good Afternoon Sir,

Thanks for the reply. I am unsure why this would fall under a FOIA request as I am simply trying to obtain basic verifiable facts that are required by law for the manufacturing of a "fully licensed and approved vaccine" as required by the SecDef's Memo for mandatory vaccination. My request IS one of medical-specific concerns. Myself and my fellow Marines are being told that this "Comirnaty-labeled" product is the "FDA-approved" product as described in the Approval letters previously attached. I am requesting for the verification of these items as previously requested as it relates to the photos of the product previously attached. The timeliness and accuracy of this information is of utmost importance since many of us have deadlines to comply with this mandate if these are the "FDA-approved" vials. If they are still under EUA, then we need this information to inform my Command so that we can still maintain our informed consent as required by the law and not consent to a product that is not fully licensed and approved with the BLA-approval of Comirnaty. Please provide the answers as soon as you can, but no later than COB tomorrow as this is another deadline at our Command. Thank you for your time and assistance with this in advance.

Semper Fidelis,

Capt Joshua "Hippity" Hoppe

██████████ ASO/NATOPS/Education Officer

Office:

██████████@usmc.mil
██

-----Original Message-----

From: ██████████@usmc.mil>

Sent: Tuesday, June 21, 2022 11:17 AM

To: ██████████@usmc.mil>

Cc: ██████████@usmc.mil>; ██████████

██████████@usmc.mil>

Subject: FW: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

Captain Hoppe,

I am the Director of ██████████ for HQMC Health Services, and the below string was forwarded to me on Friday, asking me to follow-up. I discussed your email with folks from DHA and BUMED on the string below, and with HQMC Judge Advocate Division (Major ██████████ from their office is cc'd, as is Captain ██████████ from the Marine Corps COVID Cell, for their SA on the discussion).

Reading your email below, I am assuming (please correct me if I'm wrong) that you are seeking the information below for personal reasons...i.e., not on behalf of your Command, and not for a "medical"-specific reason.

If I am interpreting your email correctly, then HQMC JAD and DHA have both advised that you submit your RFIs via a FOIA request to DHA. Brief instructions on how to do so, and the link to the submission portal, can be found at:
<https://www.health.mil/Military-Health-Topics/Privacy-and-Civil-Liberties/FOIA>.

If I have interpreted your email incorrectly, please let me (and JAD) know, and we will further discuss how/where your RFIs should be routed.

Thanks very much, Captain Hoppe.

V/R - CDR [REDACTED]

CDR [REDACTED], MC, USN
Director of [REDACTED]
Headquarters Marine Corps, Health Services

DSN:
Office:
Work Cell:
NIPR: [REDACTED]@usmc.mil

-----Original Message-----

From: [REDACTED] DHA PUB HLTH (USA) <[REDACTED].civ@mail.mil>

Sent: Friday, June 17, 2022 10:14 AM

To: [REDACTED]@usmc.mil>

Cc: [REDACTED] SN BUMED FCH VA (USA)
<[REDACTED]@mail.mil>; [REDACTED] CDR USN BUMED FCH VA
(USA) <[REDACTED] mail.mil>; [REDACTED] SAF DHA
IMMUNIZATION (USA) <[REDACTED] mail.mil>

Subject: Request for Assistance: Comirnaty Lot Release (UNCLASSIFIED)

CLASSIFICATION: UNCLASSIFIED

Sir,

Request your assistance to address the questions in an email received below from Capt Hoppe ([REDACTED] ASO/NATOPS/Education Officer).

Thanks.

V/r,

[REDACTED]
Chief, Public Health Communications
Office of the Director
DHA Public Health

[REDACTED].civ@mail.mil
Wingman: Protector #1

-----Original Message-----

From: Hoppe Capt Joshua P <[REDACTED]@usmc.mil>
Sent: Thursday, June 16, 2022 8:33 PM
To: DHA NCR Healthcare Ops Mailbox IHB DoD Vaccines
<[REDACTED]dod-vaccines@mail.mil>
Subject: Comirnaty Lot Release

Good Afternoon,

I have received word that there is a Comirnaty-labeled vaccine available at [REDACTED] (pictures attached). I am requesting verification that this is the FDA-approved product in accordance with the BLA-license as described in the Comirnaty Approval Letter and Supplemental approval letter attached. Please send a copy of the following:

- Notification of lot release from the Director, Center for Biologics Evaluation and Research (CBER) required by the Approval letter.
- Proof of the location and date of manufacture of this Lot number and NDCs.
- How many and which lots were released?
- A copy of the list of differences from what is currently in large circulation (EUA Pfizer-BioNTech) and the Comirnaty.
- If this the FDA-approved version, when will all the EUA in circulation be required to be pulled from the shelves?
- The Vaccine Information Statement for this Comirnaty product that should Replace the EUA FACT sheets once there is an FDA-approved product available.

Please provide the information as soon as you can and no later than Tuesday, 21 June 2022. Please let me know if you have any questions. Thank you for your time and assistance with this matter.

Semper Fidelis,

Capt Joshua "Hippity" Hoppe
ASO/NATOPS/Education Officer
Office
[REDACTED]@usmc.mil
[REDACTED]

CLASSIFICATION: UNCLASSIFIED