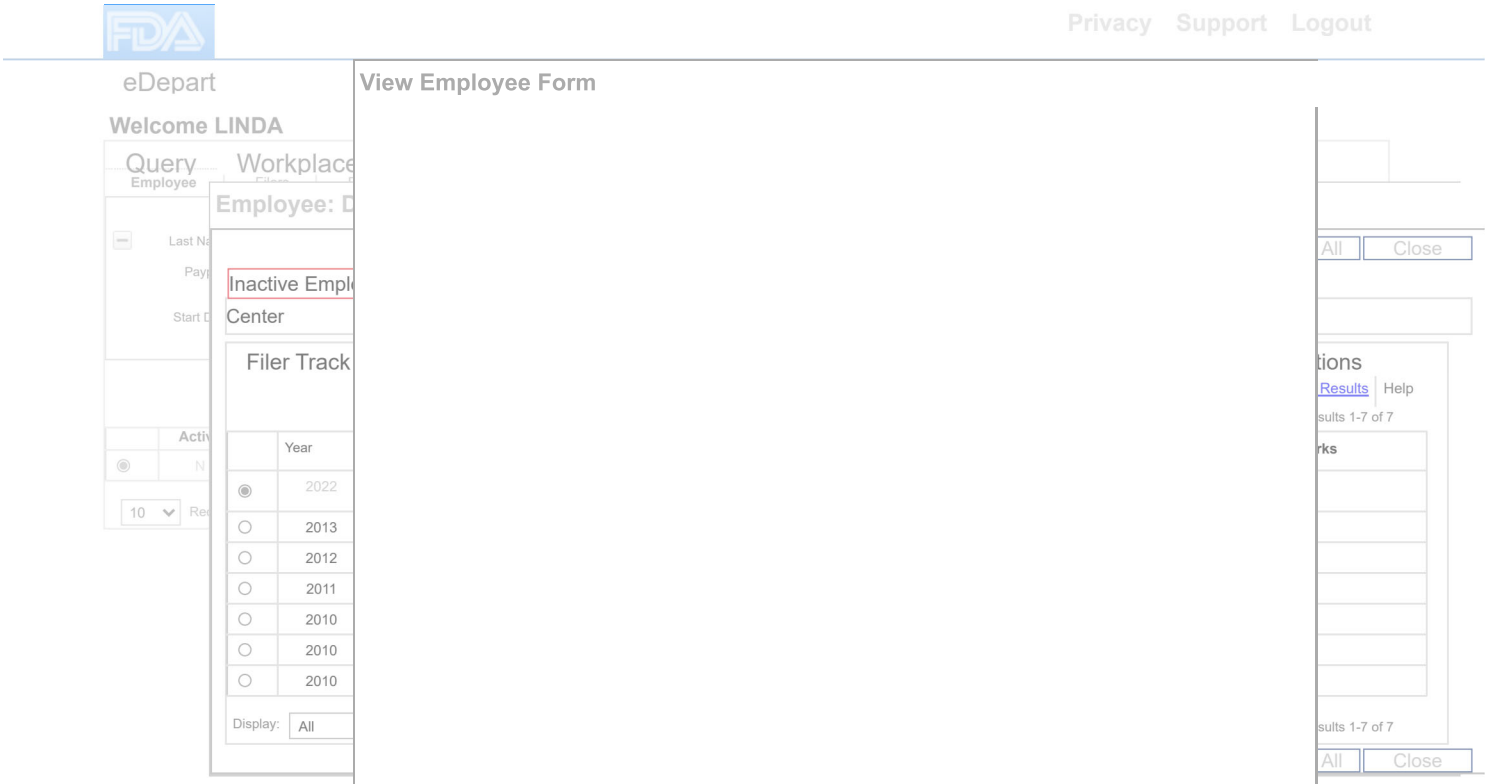
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Employee: DORAN FINK
Form Type: Post Emp Guidance - Regulation Certification for Separating Employee

Non FDA Email	(b) (6)
Employment plans following departure from FDA	**Work for a Non-Federal Entity**
Nature of Self-Employment	
Position at a non-federal entity	**Y**
Entity	**Moderna Therapeutics**
Position	**Executive Director, Infectious Diseases Translational and Early Development**
Kind of Work	**clinical development, vaccines**
Employer Considering is a Law, Accounting or Government related firm	
Supervisory Matters	**N**
Supervisory Matters Description	**In coordination with my supervisor, I have recused myself from all matters with a known or potential conflict of interest with Moderna or any of its competitors.**
Intended Separation Date	**12/(b) (6)/2022**
OGE 278 Public Financial Disclosure Report Filer	**N**
Supervised other FDA employees in past year	**Y**
Procurement Work Participation	**N**
Procurement work at FDA	
Served as the procuring contracting officer, source selection authority or evaluation board member, or chief of a financial or technical evaluation team	
Served as a program manager, deputy program manager, or administrative contracting officer	
Personally made certain decisions such as awarding a contract, subcontract, modification, task or delivery order, establishing overhead, issuing payment, or settling a claim	
Participate in any Trade or Treaty Negotiations	**N**

Non-Federal Employer List:

Employer Name	Communications	Position
No Records Found		

Previous Position Details:

FDA Duties and Responsibilities	Type of Matters	Area of Industry Affected
As a Medical Officer in DVRPA/OVRR/CBER, I was responsible for primary clinical review of regulatory submissions (pre-INDs, INDs, BLAs, and BLA supplements) for vaccines and related biological products regulated by OVRR.	Product reviews	Sponsors developing vaccines and related biological products
As a Clinical Team Leader in DVRPA/OVRR/CBER, I was responsible for primary clinical review and first-level supervisory review of regulatory submissions (pre-INDs, INDs, BLAs, and BLA supplements) for vaccines and related biological products regulated by OVRR.	Product reviews, occasional input on policy development	Sponsors developing vaccines and related biological products
As Deputy Director - Clinical in DVRPA/OVRR/CBER, I was responsible for managing clinical and toxicology review and policy for vaccines and related biological products regulated by OVRR.	Senior-level management oversight of clinical and toxicology reviews, policy development, interactions with external public health stakeholders in the US government and internationally.	Sponsors developing vaccines and related biological products
As Acting Deputy Director, OVRR/CBER, I was responsible for managing regulatory review and policy of vaccines and related biological products regulated by OVRR.	Senior-level management oversight of product reviews, policy development, interactions with external public health stakeholders in the US government and internationally.	Sponsors developing vaccines and related biological products

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