For: JR103629 Inspector General

Phone Number [redacted] (Mobile)
Email [redacted]

Cover Letter_updated 5_14_2026_added Fidelity Records Verification of cover letter and Cycling logged 313 miles week of June 29_July 4th_26.docxCycling logged 313 miles week of June 29_July 4th_26.pngFidelity Record 704014.07.PNGFidelity record showing 337753.13 to 565037.13 .PNGResume with details included.docx

Phone Number [redacted] (Mobile)
Phone Composite Header Grouping Icon Phone
Email [redacted]
Email Composite Header Grouping Icon Email
Location [redacted], MA [redacted] United States of America
Location Header Grouping Icon Location
Jobs Applied to 1
Documents with Check Header Grouping Icon Documents with Check

Attachments

Attachments

Resume / Cover Letter

Attachment	
Resume with details included.docx	
Cover Letter_updated 5_14_2026_added Fidelity Records Verification of cover letter and Cycling logged 313 miles week of June 29_July 4th_26.docx	
Cycling logged 313 miles week of June 29_July 4th_26.png	
Fidelity record showing 337753.13 to 565037.13 .PNG	
Fidelity Record 704014.07.PNG	

Other Documents

Attachment	Category	

Candidate

Job Application Details Card

Job Application Details

Job Requisition JR103629 Inspector General (Open)
Location Providence
Date Applied 08/04/2026 07:06:25 AM

Source Sources -> Apply.RI.Gov Career Website

Thomas A Verdi
Hiring Manager

Colleen Halloran-Villandr
Recruiter

In Progress

	Step	Awaiting Me	Awaiting
Screen: Sean Tessicini - JR103629 Inspector General (C116091)	Referred	Screen Candidate	1

Active Job Applications

Active Job Applications (1)

Job Application for Active Job Applications
Task Sean Tessicini - JR103629 Inspector General (C116091)
Stage Label Referred
Location: Providence | Date Applied: 08/04/2026

Education

Education

none entered

Work History

Work History

Experience

none entered

Credentials

Languages

none entered

Websites

none entered

Skills

none entered

Sean Tessicini
Email: [redacted]
Cell: [redacted]

Cover Letter

I got fired by the FBI (contractors) on behalf of my manager at the FDA. Mr. Chris Elliott stated "you shouldn't be making more money than me." I made 100k (approx) in the stock/crypto markets. I had to submit my stocks to the ethics dept and they informed my manager. All the stocks were ok to own. Mr. Elliott was even allowed to threaten me with a fist as I was sitting in a chair. Everything was documented through the FDA anti-harassment program (https://www.fda.gov/about-fda/jobs-and-training-fda/fdas-anti-harassment-and-inappropriate-conduct-policy).

In addition, I'm someone who understands the smallest nuances on a social level to make the company function effectively and efficiently. I also have a high enough net worth, along with specialized skills to understand everything. Since March 27th, 2025 with no incoming income, I **traded stocks to increase my net worth over $350K** when unemployment could not process claims with no reason why. I also made 100k in realized gains in 2024. I'm extremely personable, extremely easy to get along with yet have the ability to be stern and hold PEOPLE ACCOUNTABLE when necessary. Ty for consideration!

Final Thought: If it is required of me to be a machine, then I'll be a machine. I also want to say I'm grateful for the specialized training the FDA has provided me over the last decade which made me understand when it is me in life, I'm required to be a machine and do more than everyone else. HIGH QUALITY WORK!

Additional Pertinent Information:

I worked for HHS/FDA for a decade and got fired from my mission critical role for making too much money. **I'm 35 years old with a net worth over 695K**. In addition, I went through the anti-harassment program and Eldridge Coles Jr couldn't contact the POCS identified making it a fake anti-harassment program. That is in fact a crime. US Federal Law (Title VII). **Matthew Noonan** (matthew.noonan@fda.hhs.gov, FDA Compliance Officer and **Subject Matter Expert**). stated I was going to get a job in the Human Food Program. Why can Hr not do their job?

Eldridge Coles, Supervisory HR Specialist
Office of Human Capital Management
Voice: 301-796-0141
E-mail: Eldridge.Coles@fda.hhs.gov

Sean Tessicini
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Cell: [redacted]

The FDA taught me how to build dashboards outside of tableau. I've done complex pharmaceutical component for further processing inspections where along the process flow, I was walking along with vats of acid on either side of an elevated walk way. I have aseptically swabbed your favorite foods processing plants. I have self-taught myself how to trade stocks and built a **portfolio over 700K** at the age of 35 all while not making 100K a year to a few years ago. I'm a machine who looks around at everyone in high paying positions and can't fathom how they don't know what they are doing. If I'm not qualified, I want you to get the treated I get in life. You would crumble. You would be in so much debt that you might be homeless. Because when it is me in life, I AM REQUIRED TO BE A MACHINE ALL WHILE BEING TREATED UNFAIRLY. I'm required to be obsessed to not be homeless.

Below is a regulatory-focused process flow for a bovine-derived pharmaceutical component intended for further processing into gelatin, which is an inspection type I completed for the agency. FYI- I was in foods and primarily did food inspections.

Animal Sourcing
↓
Raw Hide/Bone Receipt
↓
Cleaning & Size Reduction
↓
Alkaline or Acid Pretreatment
↓
Hot Water Extraction
↓
Filtration & Purification
↓
Concentration
↓
Sterilization Step
↓
Drying
↓
Milling
↓
Testing (USP Specs)

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Cell: [redacted]

↓
Batch Release
↓
Use as Pharmaceutical Excipient

In addition, some background about QMIS systems and FDA compliance for HR to be abreast on below. I went through the training for an ISO17025 cert which is also more extensive than the current food staff possess in regard to understanding ISO standards within supply chain audits and QMIS systems. It is just the way FOODS are going. Just like devices and pharmaceuticals.

QMIS Systems and FDA Compliance

1. **Regulatory Requirements:**
    - The FDA has strict regulations for industries it oversees (pharmaceuticals, medical devices, biologics, etc.), and these industries must comply with **Good Manufacturing Practices (GMP)** and other FDA requirements. A QMIS is used to ensure that all quality-related processes comply with these standards.
    - Key regulations like **21 CFR Part 820** (for medical devices) and **21 CFR Part 211** (for pharmaceuticals) require manufacturers to implement robust quality management systems. A QMIS helps track data, manage documentation, and ensure all processes are compliant.
2. **Functions of a QMIS:**
    - **Document Control**: Ensures all product documentation (e.g., SOPs, batch records, certificates of analysis) is compliant and accessible.
    - **Non-Conformance Management**: Tracks deviations from specifications and ensures corrective actions are documented and implemented.
    - **CAPA (Corrective and Preventive Actions)**: Ensures issues are identified, investigated, and that actions are taken to prevent recurrence.
    - **Audit Management**: Facilitates internal and external audits, including FDA inspections, and tracks audit findings and resolutions.
    - **Supplier Quality Management**: Ensures suppliers meet regulatory and quality requirements.
    - **Training Management**: Tracks employee training to ensure personnel are qualified to handle regulatory requirements.

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Cell: [redacted]

3. **FDA Inspections:**
    - During FDA inspections, a QMIS can help provide evidence that a company is maintaining compliance with regulatory requirements. The FDA will typically look for documented evidence of quality processes (e.g., audits, training, CAPA processes).
    - Systems like these also provide detailed reporting features, which can help during inspections to demonstrate ongoing compliance.
4. **Benefits of QMIS for FDA Compliance:**
    - **Data Integrity**: Helps ensure that all data is accurate, traceable, and can be easily retrieved for inspections.
    - **Efficiency**: Streamlines reporting and auditing processes, which is crucial for maintaining continuous FDA compliance.
    - **Risk Mitigation**: Helps identify potential issues early, reducing the risk of regulatory violations or product recalls

I also wanted you to be made aware of what my GS 13 national level work was so I'll provide an outline for your reference in regards to "Performs complex and special project management and other Manufactured Food Regulatory Program Standards (MFRPS) and Animal Food Regulatory Program Standards (AFRPS) related assignments" subsection U.S Food and Drug Administration 6/2022 - 7/2022 Series: 0696 Pay Plan: GS Grade: 13 Consumer Safety Officer within the resume. You will now have a better picture of what the FDA is doing (see "MFRPS and AFRPS outline"). Similar to what the CDC and NIH does funding wise through **distinct funding mechanisms**. The main ones are **grants, cooperative agreements, and contracts**, each with different levels of involvement from the agency. In regards to MFRPs and AFRPS (FDA), both programs are funded primarily through **cooperative agreements**.

MFRPS AND AFRPS Outline

The **U.S. Food and Drug Administration (FDA)** created two related frameworks to help state regulatory programs maintain **consistent, high-quality food safety oversight**:

- **Manufactured Food Regulatory Program Standards (MFRPS)**
- **Animal Food Regulatory Program Standards (AFRPS)**

Both programs are part of FDA's **Integrated Food Safety System (IFSS)** and are commonly implemented by state agriculture or public health agencies.

Sean Tessicini
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Cell: [redacted]

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1. Manufactured Food Regulatory Program Standards (MFRPS)

MFRPS apply to **state programs that regulate human food manufacturing, processing, packing, and holding facilities**.

Purpose

- Ensure state manufactured food programs operate with **uniform standards**
- Improve **inspection quality, training, compliance, and enforcement**
- Align state programs with **federal food safety laws**, including the **FDA Food Safety Modernization Act (FSMA)**.

The 10 MFRPS Standards

1. **Regulatory Foundation** – Legal authority to regulate food facilities
2. **Training Program** – Training for inspectors and staff
3. **Inspection Program** – Risk-based inspections of food facilities
4. **Auditing Program** – Internal audits of inspection quality
5. **Food-Related Illness & Outbreak Response**
6. **Compliance & Enforcement Program**
7. **Industry & Community Relations**
8. **Program Resources** – Staffing, equipment, funding
9. **Program Assessment** – Self-assessment and continuous improvement
10. **Laboratory Support** – Access to food testing labs

---

2. Animal Food Regulatory Program Standards (AFRPS)

AFRPS apply to **state programs that regulate animal food and feed manufacturing**.

Purpose

- Ensure **safe animal feed** and protect **animal and human health**
- Align state feed programs with **FSMA preventive controls for animal food**

The 11 AFRPS Standards

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1. **Regulatory Foundation**
2. **Training Program**
3. **Inspection Program**
4. **Auditing Program**
5. **Feed-Related Illness & Outbreak Response**
6. **Compliance & Enforcement Program**
7. **Industry & Community Relations**
8. **Program Resources**
9. **Program Assessment**
10. **Laboratory Support**
11. **Sampling Program**

In addition, I possess a Boating Safety ID card from the state of Florida, exp: Lifetime. It is recognized by the state of Florida, the National Association of State Boating Law Administrators, and by the U.S COAST GUARD**. I'm also an avid cyclist. I ride a Trek Checkpoint SL5, carbon frame, gravel bicycle.**

In addition, I had a Commissioned Corps of the Public Health Service Officer (manager) "train" me during my tenure with the agency.

In addition, I even made 6 figures in year 2025 (103K) on paper without having any incoming income since March 27th 2025 when I got fired by Mr. Chris Elliott for making too much money. Please feel free to verify with the **IRS (owe 7K in taxes for year 2025: Paid 3/28/2025)**.

In addition, my net worth is **now over 695K**. The **last over 350K made with no incoming income** this past year, trading stocks **(debt of 6k only on a 2022 truck**). I'm done trading stocks as the portfolio is now set up on autopilot. More than happy to verify.

Sean Tessicini
Email: [redacted]
Cell: [redacted]

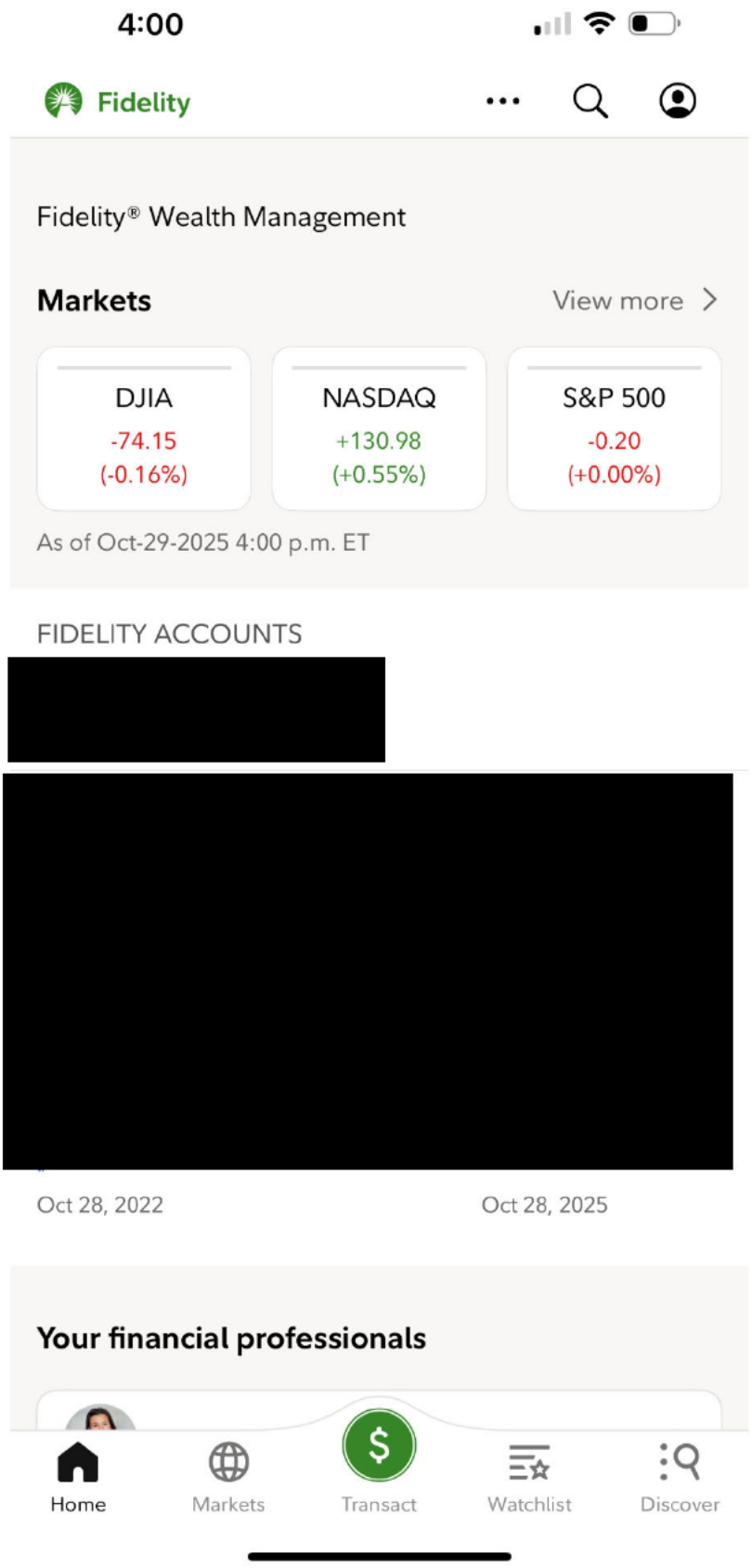
I just need someone to understand how qualified I am. I spend all day every day applying to jobs. I'll never stop and never waver because I understand what is required when it is someone who works hard and does the right things in life! Again, I want to thank the FDA for molding me into this over the last decade.

Final Note: I added the fidelity records for verification purposes (Pages 8-9). I also added my cycling log for the week of Jun 29th-July 4th 2026 (Page 10). The miles logged were over 300 miles in the week. Miles the professional's record. I think it is fitting as it shows the type of person I am. In addition, I also understand some high-level employees prefer hiring the alcoholics instead. That is also ok, m'am. However, I understand what is required when it is me in life. "If i'm required to be a machine, then i'll be a machine." Yes m'am!

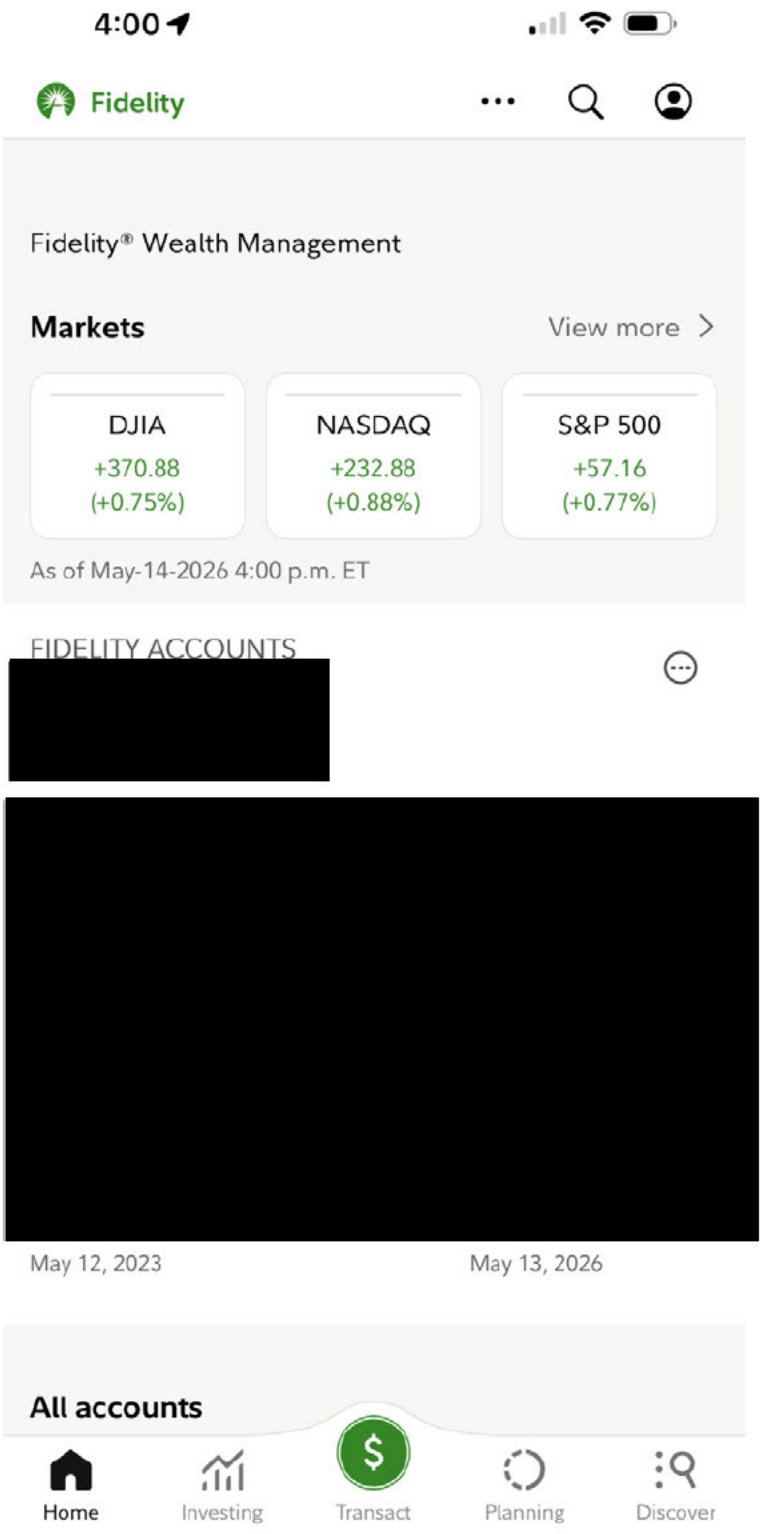
V/R

Sean Tessicini

Sean Tessicini
Email: [redacted]
Cell: [redacted]



Sean Tessicini
Email: [redacted]
Cell: [redacted]



Sean Tessicini
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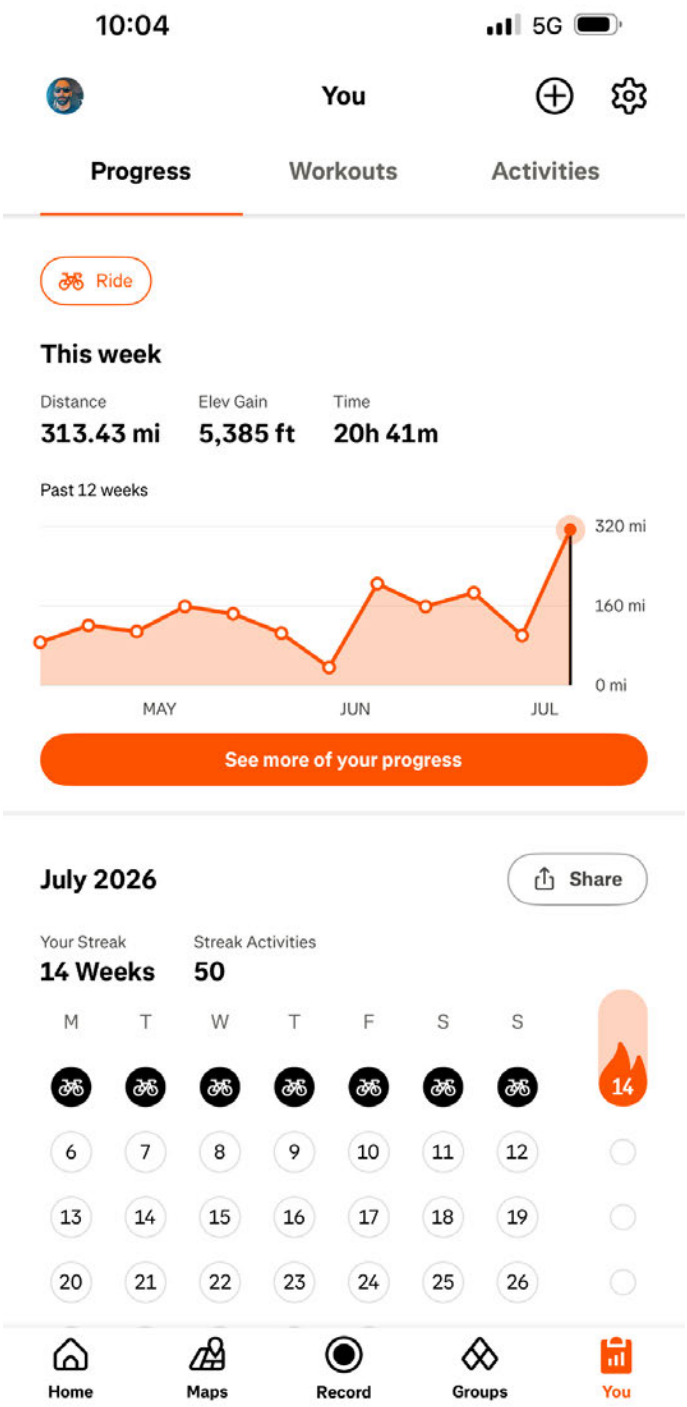
Cell: [redacted]



Sean Tessicini
Phone: [redacted]
Email: [redacted]

Work Experience:

U.S Food and Drug Administration

1 Montvale Ave, Stoneham, MA 02180 United States/109 Holton Street Winchester, MA 01890

07/2018 – 4/2025

Series: 0696 **Pay Plan:** GS **Grade:** 12

Consumer Safety Officer (This is a federal job)

Duties, Accomplishments and Related Skills:

•Primarily plan and conduct regulatory inspections and in-depth investigations of various food facilities. In addition, performed regulatory inspections/investigations of medical device facilities, dietary supplements, and completed an inspection for a pharmaceutical component for further processing.

• Coordinated a recall in the field at the retail setting with the firm owner not speaking English. Something that you would expect the retail food specialist to accomplish. Private labels were done by the supermarket.

• Completed a compliance follow up, following up a class 1 and class 2 recall, importer verification within the HACCP regulation, multiple label reviews, multiple HACCP plan reviews of a 200 million plus business.

• Performed a recall audit check for "Shruumz-brand products" and calmly diffused a high-risk interaction independently in the field.

•Act as a liaison between regulated industry and FDA. Interact with CEOs, CFOs, QA managers, and production floor employees to obtain the necessary information to complete comprehensive inspections of multimillion-dollar companies (potentially billion-dollar companies).

•Writes clear and concise comprehensive establishment inspection reports. Evaluate food safety plans, and HACCP plans

•Educate and provide resources to food industry in the interest of promoting public health.

•Independently carry out investigations, inspections, recall audit checks, and sample collections (surveillance, environmental, and CFSAN directed sample assignments).

•Perform analysis and evaluation on data samples and documented information gathered during inspection and investigations. Interact with various levels of officials representing the establishments subject to regulatory review.

• Help handle/ manage the official establishment inventory of firms. This includes verifying food firms fall under FDA jurisdiction (via phone, email, and in person), performing coding in FMS, entering

investigations within FACTS/eNspect , and addressing any issues with project “B” firms.

U.S Food and Drug Administration
6/2022 - 7/2022
Series: 0696 **Pay Plan:** GS **Grade:** 13
Consumer Safety Officer (This is a time-limited appointment or temporary promotion)
Duties, Accomplishments and Related Skills:
• Plans, delivers, and maintains comprehensive programs for reviewing and analyzing programmatic data of SLTT human and animal food regulatory programs to evaluate progress towards a national integrated food safety system (IFSS).
•Performs complex and special project management and other Manufactured Food Regulatory Program Standards (MFRPS) and Animal Food Regulatory Program Standards (AFRPS) related assignments
• Evaluates return-on-investment and return-on-value of ORA's SLTT/associate investment portfolio
• Evaluates programmatic public health impacts of ORA funding opportunities
• Evaluates SLTT food consumer protection laws, regulations, processes, and procedures to determine equivalency with FDA’s
• Measures and tracks integration of SLTT programs nationally
• Analyzes and evaluates the effectiveness of OP’s programs from an integration viewpoint and advises on metrics, review methods, and funding agreement structures supporting SLTT programs
• Develops and applies new or revised program methods, organizational structures, and technologies to carry out program objectives.
• Prepares summaries and statistical data on integration.
• Performs analysis of the reports generated by financial management systems and subsystems to ensure the timely satisfaction of existing and future reporting requirements.

U.S Department of Commerce: NOAA-Office of International Affairs and Seafood Inspections
55 Great Republic Drive, Gloucester, MA 01930 United States
11/2017 - 07/2018
Series: 0696 **Pay Plan:** ZP
Consumer Safety Officer (This is a federal job)
Duties, Accomplishments and Related Skills:
• Audit seafood processing facilities and vessels to evaluate their conformity with U.S. regulations and industry standards ensuring Current Good Manufacturing Practices (CGMP) are adhered to
•Apply knowledge of the Hazard Analysis and Critical Control Point (HACCP) process in quality system

audits and inspections to identify violations, recommend corrective actions, and work with processing plants and vessels to develop further compliance to their respective HACCP plan.
•Perform inspections of raw and processed seafood to evaluate the safety, quality and health of the products in accordance with U.S. regulations, U.S. grade standards, program policy, and commercial item descriptions (label specifications).
• Provide industry with export health certificates once it is determined seafood product meet the unique requirements of each importing country, and any other specific industry buyer criteria
•Collect samples /perform lab analysis of raw and processed products to determine the quality and condition of seafood. Identify abnormalities (defects) in product characteristics using sensory abilities
• Act as a liaison between the Department of Commerce and industry when performing inspections at approved establishments.

U.S Food and Drug Administration: Winchester Engineering and Analytical Center
109 Holton Street, Winchester, MA 01890-1152 United States
05/2015 - 11/2017
Series: 0303 **Pay Plan:** GS **Grade:** 7
Laboratory Support Technician
Duties, Accomplishments and Related Skills:
•Maintain chain-of-custody of all samples; ensure proper handling, storage and documentation are completed in compliance with applicable FDA procedures
•Identifying Office of Criminal Investigation (OCI) case numbers and making an excel sheet to correspond with the FACTS numbers, among other various OCI support activities.
•CDRH microwave program and setting up all the shipments for commercial microwaves back to industry.
•TLD monitors- scanning them into the system, sending emails to CSO's around the world about sending the Thermoluminescent dosimeter back to WEAC.
•Putting insurance on 100K piece of equipment and setting up the freight service to ship the piece of equipment to Alaska. OP did a project with Alaska dept of public health. WEAC owned the piece of equipment. I dealt with the shipment and insurance. Coordinated everything to make sure that piece of equipment landed safely in Alaska.
• Quality control monitor for sample storage areas and ELIX water purification system. Maintain FV/PM and calibration records; schedule services as indicated by standard operating procedure associated with equipment.
• Complete documentation and proper disposal method associated with each Office of Regulatory Affairs

(ORA) sample at the laboratory

•Inspect samples on arrival at the laboratory. Review records ensuring proper documentation is adequate and complete. Note any discrepancies found. Enter samples into Field Accomplishments and Compliance Tracking System (FACTS).

•Review and analyze scientific QA books associated with regulatory equipment ensuring function verification, calibration, and preventative maintenance are being accomplished within specified time frames.

•Manage GSA Fleet vehicle. Maintain logs and databases associated with vehicle.

•Evaluate problems and make recommendations for possible corrective action.

•Revise SOP's as need. Write sample feedback reports as needed.

•Collaborate on special projects and participate in project teams

Education:

University of Massachusetts Amherst: School of Public Health and Health Sciences Amherst, MA United States

Bachelor's Degree 9 /2013

Major: Bachelor of Science Kinesiology

Job Related Training:

• FD254- Hazard Analysis and Risk Based Preventative Controls for Human Food; 5/12/2023

• FDA Excel Dashboard Training; 7/21/22

•FD249-Conducting Seafood Inspections; 12/07/2020

•FD152-Food Processing and Technology; 7/31/2020

•FD190-Food Current Good Manufacturing Practice, Application, and Evidence Development; 9/20/2019

• USDC/NOAA Seafood HACCP Training Workshop; 4/5/2018

•PJLA-ISO/IEC 17025:2005 Internal Auditor Training; 6/2/2016

10:04 5G

You

Progress Workouts Activities

Ride

This week

Distance	Elev Gain	Time
313.43 mi	**5,385 ft**	**20h 41m**

Past 12 weeks

See more of your progress

July 2026

Share

Your Streak	Streak Activities
14 Weeks	**50**

M	T	W	T	F	S	S	
●	●	●	●	●	●	●	14
6	7	8	9	10	11	12	
13	14	15	16	17	18	19	
20	21	22	23	24	25	26	

Home Maps Record Groups You

4:00

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DJIA	NASDAQ	S&P 500
+370.88	+232.88	+57.16
(+0.75%)	(+0.88%)	(+0.77%)

As of May-14-2026 4:00 p.m. ET

FIDELITY ACCOUNTS

[redacted]

May 12, 2023 May 13, 2026

All accounts

Home Investing Transact Planning Discover

4:00

Fidelity

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Markets

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DJIA	NASDAQ	S&P 500
-74.15 (-0.16%)	+130.98 (+0.55%)	-0.20 (+0.00%)

As of Oct-29-2025 4:00 p.m. ET

FIDELITY ACCOUNTS

[redacted]

Oct 28, 2022 Oct 28, 2025

Your financial professionals

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