DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
11155 Dolfield Boulevard, Suite 117 Owings Mills, MD 21117 (410)779-5455 Fax:(410)779-5707	4/7/2026-4/8/2026
	FEI NUMBER 1276528

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Danny Rivera, Team Supervisor

FIRM NAME	STREET ADDRESS
The American National Red Cross	381 Plantation St Ste 110
CITY, STATE, ZIP CODE, COUNTRY Worcester, MA 01605-4319	TYPE ESTABLISHMENT INSPECTED Donor Center

This document lists observations made by the FDA representative(s) during the inspection of your facility. They are inspectional observations, and do not represent a final Agency determination regarding your compliance. If you have an objection regarding an observation, or have implemented, or plan to implement, corrective action in response to an observation, you may discuss the objection or action with the FDA representative(s) during the inspection or submit this information to FDA at the address above. If you have any questions, please contact FDA at the phone number and address above.

DURING AN INSPECTION OF YOUR FIRM I OBSERVED:

OBSERVATION 1

Failure to maintain facilities in a clean and orderly manner.

Specifically,
A walkthrough of your storage area on 4/7/2026 showed transport (b) (4) positioned (b) (4) with minimal space for pest control activities to be conducted. (b) (4) of Plasma Kits and transport coolers were stored. The bottoms of the (b) (4) appeared to have cobwebs and remnants of dead insects. The Plasma Kits are used for collecting blood and blood products from donors, while the coolers are used to transport blood and blood products to processing centers.

SEE REVERSE OF THIS PAGE

EMPLOYEE(S) SIGNATURE
Charles D Boison, Investigator

X Charles D Boison
Investigator
Signed By: Charles D. Boison -S
Date Signed: 04-08-2026
09:33:39

DATE ISSUED
4/8/2026

The observations of objectionable conditions and practices listed on the front of this form are reported:

1. Pursuant to Section 704(b) of the Federal Food, Drug and Cosmetic Act, or
2. To assist firms inspected in complying with the Acts and regulations enforced by the Food and Drug Administration.

Section 704(b) of the Federal Food, Drug, and Cosmetic Act (21 USC 374(b)) provides:

"Upon completion of any such inspection of a factory, warehouse, consulting laboratory, or other establishment, and prior to leaving the premises, the officer or employee making the inspection shall give to the owner, operator, or agent in charge a report in writing setting forth any conditions or practices observed by him which, in his judgment, indicate that any food, drug, device, or cosmetic in such establishment (1) consists in whole or in part of any filthy, putrid, or decomposed substance, or (2) has been prepared, packed, or held under insanitary conditions whereby it may have become contaminated with filth, or whereby it may have been rendered injurious to health. A copy of such report shall be sent promptly to the Secretary."