DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
19701 Fairchild Irvine, CA 92612-2445 (949)608-2900 Fax:(949)608-4417	5/11/2026-5/14/2026
	FEI NUMBER 3015775917

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Julie D. Thompson, Center Director

FIRM NAME	STREET ADDRESS
Octapharma Plasma Inc	9124 E Apache Trl Ste 1
CITY, STATE, ZIP CODE, COUNTRY Mesa, AZ 85207-8735	TYPE ESTABLISHMENT INSPECTED Source plasma donation 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

Equipment used in the storage and distribution of blood and blood components is not observed, standardized and calibrated with at least the frequency required.

Specifically, an installation qualification to verify label accuracy was not performed on the (b)(4) printer (Equipment ID: (b)(4)) following its installation on (b)(4). This printer is used to generate (b)(4) labels for plasma unit distribution.

SEE REVERSE OF THIS PAGE	EMPLOYEE(S) SIGNATURE		DATE ISSUED
	Garrad R Poole, Investigator	X Garrad R Poole Investigator Signed By: GARRAD R. POOLE - S Date Signed: 05-14-2026 11:20:41	5/14/2026

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."