

SI	PECIMEN ID NO. $00000$	OT AC	CESSION NO.		
STEP 1: COMPLETED BY COLLECTOR OF	R EMPLOYER REPRESENTATIVE	P MPO Nama Addra	as Dhone No and E	av Na	
A. Employer Name, Address, I.D. No.  B. MRO Name, Address, Phone No. and Fax No.					
C. Donor SSN, Employee I.D., or CDL State	and No.				
D. Specify Testing Authority: HHS		] FMCSA   FAA	☐ FRA ☐ FTA	☐ PHMSA ☐ USCG	
E. Reason for Test: Pre-employment Ra	ndom 🔲 Reasonable Suspicion/Cause [	Post Accident Re	turn to Duty  Follow	-up Other (specify)	
F. Drug Tests to be Performed:   THC, C	COC, PCP, OPI, AMP 🔲 THC & COC	Only Othe	r (specify)		
G. Collection Site Address:		Collector Contact	Info: Phone		
STEP 2: COMPLETED BY COLLECTOR (m	nake remarks when appropriate).	URINE	ORAL FLU	IID	
	None Provided, Enter Remark.				
URINE: Collector reads urine temperature					
ORAL FLUID: Split Type: Serial Co	ncurrent Subdivided Each Dev	vice Within Expiration	Date?   Yes   No	Volume Indicator(s) Obser	
REMARKS:					
STEP 3: Collector affixes seal(s) to bottle( STEP 4: CHAIN OF CUSTODY - INITIATED			Donor completes S	TEP 5 on Copy 2 (MRO Copy)	
I certify that the specimen given to me by the			m   SPECIMEN BO	OTTLE(S)/TUBE(S) RELEASED T	
was collected, labeled, sealed and released to the				(0)02_(0)	
X					
	Signature of Collector	Δ.Ν.	-		
	/ /	AN PM			
(PRINT) Collector's Name (First, MI,	Last) Date (Mo/Day/Yr)	Time of Collection		Name of Delivery Service	
RECEIVED AT LAB OR IITF:			Primary Specimen Seal Intact	SPECIMEN BOTTLE(S)/TUBE( RELEASED TO:	
X			☐ YES ☐ NO	KELLAGED TO.	
Si	gnature of Accessioner				
Si	gnature of Accessioner	1 1	If NO, Enter remark		
(PRINT) Accessioner's I	Name (First, MI, Last)	/ Date (Mo/Day/Yr)	If NO, Enter remark in Step 5A.		
	Name (First, MI, Last)  n Date: / /		If NO, Enter remark		
(PRINT) Accessioner's Primary/Single Specimen Device Expiration	Name (First, MI, Last)  n Date: / / (Mo/Day/Yr)		If NO, Enter remark in Step 5A.	Pate: / (Mo/Day/Yr)	
(PRINT) Accessioner's Primary/Single Specimen Device Expiration	Name (First, MI, Last)  n Date: / / (Mo/Day/Yr)  - COMPLETED BY TEST FACILITY	Split Specimer	If NO, Enter remark in Step 5A.	(Mo/Day/Yr)	
(PRINT) Accessioner's I Primary/Single Specimen Device Expiratio  STEP 5A: PRIMARY SPECIMEN REPORT  NEGATIVE DILUTE	Name (First, MI, Last)  n Date: / / (Mo/Day/Yr)  - COMPLETED BY TEST FACILITY	Split Specimer	If NO, Enter remark in Step 5A.  Device Expiration I		
Primary/Single Specimen Device Expiration  STEP 5A: PRIMARY SPECIMEN REPORT  NEGATIVE DILUTE POSITIVE for:	Name (First, MI, Last)  n Date: / / (Mo/Day/Yr)  - COMPLETED BY TEST FACILITY	Split Specimer	If NO, Enter remark in Step 5A.  Device Expiration I	(Mo/Day/Yr)	
(PRINT) Accessioner's I  Primary/Single Specimen Device Expiratio  STEP 5A: PRIMARY SPECIMEN REPORT  NEGATIVE DILUTE POSITIVE for: Analyte(s) in ng/mL	Name (First, MI, Last)  n Date: / / (Mo/Day/Yr)  - COMPLETED BY TEST FACILITY	Split Specimer	If NO, Enter remark in Step 5A.  Device Expiration I	(Mo/Day/Yr)	
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Version C 6May2020

0000001 SPECIMEN ID NO. ACCESSION NO. STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE OMB No. 0930-0158 A. Employer Name, Address, I.D. No. B. MRO Name, Address, Phone No. and Fax No. C. Donor SSN, Employee I.D., or CDL State and No. ☐ NRC Specify DOT Agency: FMCSA FAA FRA FTA PHMSA USCG D. Specify Testing Authority: HHS E. Reason for Test: Pre-employment Random Reasonable Suspicion/Cause Post Accident Return to Duty Follow-up Other (specify) F. Drug Tests to be Performed: THC, COC, PCP, OPI, AMP THC & COC Only Other (specify) G. Collection Site Address: Collector Contact Info: Phone Other ☐ URINE ☐ ORAL FLUID STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate). COLLECTION: Split Single None Provided, Enter Remark. **URINE: Collector reads urine temperature within 4 minutes.** Temperature between 90° and 100° F? ☐ Yes ☐ No, Enter Remark ☐ Observed, Enter Remark ORAL FLUID: Split Type: ☐ Serial ☐ Concurrent ☐ Subdivided Each Device Within Expiration Date? ☐ Yes ☐ No ☐ Volume Indicator(s) Observed REMARKS: STEP 3: Collector affixes seal(s) to bottle(s)/tube(s). Collector dates seal(s). Donor initials seal(s). Donor completes STEP 5 on Copy 2 (MRO Copy) STEP 4: CHAIN OF CUSTODY - INITIATED BY COLLECTOR AND COMPLETED BY TEST FACILITY SPECIMEN BOTTLE(S)/TUBE(S) RELEASED TO: I certify that the specimen given to me by the donor identified in the certification section on Copy 2 of this form was collected, labeled, sealed and released to the Delivery Service noted in accordance with applicable federal requirements. Signature of Collector ΔM PM Name of Delivery Service (PRINT) Collector's Name (First, MI, Last) Date (Mo/Day/Yr) STEP 5: COMPLETED BY DONOR I certify that I provided my specimen to the collector; that I have not adulterated it in any manner; each specimen bottle/tube used was sealed with a tamper-evident seal in my presence; and that the information provided on this form and on the label affixed to each specimen bottle/tube is correct. (PRINT) Donor's Name (First, MI, Last) Signature of Donor Date (Mo/Day/Yr) \_\_\_ Evening Phone No. (\_\_\_)\_ Email address: \_ Daytime Phone No. (\_\_\_)\_ Date of Birth After the Medical Review Officer receives the test results for the specimen identified by this form, he/she may contact you to ask about prescriptions and over-the-counter medications you may have taken. Therefore, you may want to make a list of those medications for your own records. THIS LIST IS NOT NECESSARY. If you choose to make a list, do so either on a separate piece of paper or on the back of your copy (Copy 5). - DO NOT PROVIDE THIS INFORMATION ON THE BACK OF ANY OTHER COPY OF THE FORM, TAKE COPY 5 WITH YOU. ☐ URINE ☐ ORAL FLUID STEP 6: COMPLETED BY MEDICAL REVIEW OFFICER - PRIMARY SPECIMEN In accordance with applicable federal requirements, my verification is: **NEGATIVE** POSITIVE for: REFUSAL TO TEST because – check reason(s) below: ☐ TEST CANCELLED ADULTERATED (adulterant/reason): SUBSTITUTED OTHER: \_\_\_\_\_ REMARKS: Signature of Medical Review Officer (PRINT) Medical Review Officer's Name (First, MI, Last) Date (Mo/Day/Yr) STEP 7: COMPLETED BY MEDICAL REVIEW OFFICER - SPLIT SPECIMEN In accordance with applicable federal requirements, my verification for the split specimen (if tested) is: RECONFIRMED for: \_ \_\_ TEST CANCELLED FAILED TO RECONFIRM for: REMARKS: \_ (PRINT) Medical Review Officer's Name (First, MI, Last) Signature of Medical Review Officer Date (Mo/Day/Yr)

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$\bigcirc$	Paper CCF: Back of Copy 1-4 Electronic CCF: Separate Page	$\bigcirc$
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$\bigcirc$	Public Burden Statement  An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this project is 0930-0158. Public reporting burden for this collection of information is estimated to average: 5 minutes/donor; 4 minutes/collector; 3 minutes/test facility; and 3 minutes/Medical Review Officer. Send comments regarding this	$\bigcirc$
burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to SAMHSA Report Officer, 5600 Fishers Lane, Room 15E57B, Rockville, Maryland, 20852.		$\circ$
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**COPY 5 - DONOR COPY** 

Signature of Medical Review Officer

(PRINT) Medical Review Officer's Name (First, MI, Last)

Date (Mo/Day/Yr)

Paper CCF: Back of Copy 5	
Electronic CCF: Separaté Page	

## Privacy Act Statement: (For Federal Employees Only)

Submission of the information on the Federal Drug Testing Custody and Control Form is voluntary. However, incomplete submission of the information, refusal to provide a specimen, or substitution or adulteration of a specimen may result in delay or denial of your application for employment/appointment or may result in removal from the federal service or other disciplinary action.

The authority for obtaining the specimen and identifying information contained herein is Executive Order 12564 ("Drug-Free Federal Workplace"), 5 U.S.C. Sec. 3301 (2), 5 U.S.C. Sec. 7301, and Section 503 of Public Law 100-71, 5 U.S.C. Sec. 7301 note. Under provisions of Executive Order 12564 and 5 U.S.C. 7301, test results may only be disclosed to agency officials on a need-to-know basis. This may include the agency Medical Review Officer (MRO), the administrator of the Employee Assistance Program, and a supervisor with authority to take adverse personnel action. This information may also be disclosed to a court where necessary to defend against a challenge to an adverse personnel action.

Submission of your SSN is not required by law and is voluntary. Your refusal to furnish your number will not result in the denial of any right, benefit, or privilege provided by law. Your SSN is solicited, pursuant to Executive Order 9397, for purposes of associating information in agency files relating to you and for purposes of identifying the specimen provided for testing. If you refuse to indicate your SSN, a substitute number or other identifier will be assigned, as required, to process the specimen.

## **Public Burden Statement**

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this project is 0930-0158. Public reporting burden for this collection of information is estimated to average: 5 minutes/donor; 4 minutes/collector; 3 minutes/test facility; and 3 minutes/Medical Review Officer. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to SAMHSA Reports Clearance Officer, 5600 Fishers Lane, Room 15E57B, Rockville, Maryland, 20852.