

FEDERAL DRUG TESTING CUSTODY AND CONTROL FORM



SPECIMEN ID NO. 0000001

ACCESSION NO.

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

A. Employer Name, Address, I.D. No.

B. MRO Name, Address, Phone No. and Fax No.

C. Donor SSN or Employee I.D. No.

We are adding CDL State and No. as an option for donor identification here

D. Specify Testing Authority: HHS NRC Specify DOT Agency: FMCSA FAA FRA FTA PHMSA USCG

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 _____

Fax _____

Other _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate).

URINE

ORAL FLUID

COLLECTION: Split Single None Provided, Enter Remark.

We are adding a horizontal line to separate COLLECTION and URINE entries here

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

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.

SPECIMEN BOTTLE(S)/TUBE(S) RELEASED TO:

X

Signature of Collector

AM

PM

(PRINT) Collector's Name (First, MI, Last)

Date (Mo/Day/Yr)

Time of Collection

Name of Delivery Service

RECEIVED AT LAB OR IITF:

X

Signature of Accessioner

(PRINT) Accessioner's Name (First, MI, Last)

Date (Mo/Day/Yr)

Primary Specimen Seal Intact

YES NO

If NO, Enter remark in Step 5A.

SPECIMEN BOTTLE(S)/TUBE(S) RELEASED TO:

Primary/Single Specimen Device Expiration Date: _____
(Mo/Day/Yr)

Split Specimen Device Expiration Date: _____
(Mo/Day/Yr)

STEP 5A: PRIMARY SPECIMEN REPORT - COMPLETED BY TEST FACILITY

NEGATIVE REJECTED FOR TESTING ADULTERATED SUBSTITUTED INVALID RESULT

DILUTE

POSITIVE for: _____

Analyte(s) in ng/mL

REMARKS:

Test Facility (if different from above): _____

I certify that the specimen identified on this form was examined upon receipt, handled using chain of custody procedures, analyzed, and reported in accordance with applicable federal requirements.

X

Signature of Certifying Technician/Scientist

(PRINT) Certifying Technician/Scientist's Name (First, MI, Last)

Date (Mo/Day/Yr)

STEP 5b: COMPLETED BY SPLIT TESTING LABORATORY

RECONFIRMED FAILED TO RECONFIRM - REASON _____

I certify that the split specimen identified on this form was examined upon receipt, handled using chain of custody procedures, analyzed, and reported in accordance with applicable federal requirements.

X

Signature of Certifying Scientist

(PRINT) Certifying Scientist's Name (First, MI, Last)

Date (Mo/Day/Yr)

Laboratory Name

Laboratory Address

Date (Mo/Day/Yr)

Donor's Initials

PLACE OVER CAP



0000001
SPECIMEN A



0000001
SPECIMEN B

Date (Mo/Day/Yr)

Donor's Initials

PLACE OVER CAP

OMB No. 0930-0158

PLEASE HARD - YOU ARE MAKING MULTIPLE COPIES

Version C 11 December 2019

80308

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SPECIMEN ID NO. **0000001**

ACCESSION NO.

OMB No. 0930-0158

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C. Donor SSN or Employee I.D. No. _____

D. Specify Testing Authority: HHS NRC Specify DOT Agency: FMCSA FAA FRA FTA PHMSA USCG

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 _____
 Fax _____
 Other _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate). URINE ORAL FLUID

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

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.

X _____
 Signature of Collector

_____ AM
 _____ PM

(PRINT) Collector's Name (First, MI, Last) _____ Date (Mo/Day/Yr) _____ Time of Collection _____

SPECIMEN BOTTLE(S)/TUBE(S) RELEASED TO: _____
 Name of Delivery Service

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.

X _____
 Signature of Donor

(PRINT) Donor's Name (First, MI, Last) _____ Date (Mo/Day/Yr) _____

Email address: _____ Daytime Phone No. () _____ Evening Phone No. () _____ Date of Birth _____
 (Mo/Day/Yr)

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.

STEP 6: COMPLETED BY MEDICAL REVIEW OFFICER - PRIMARY SPECIMEN URINE ORAL FLUID

In accordance with applicable federal requirements, my verification is:

NEGATIVE POSITIVE for: _____
 DILUTE

REFUSAL TO TEST because – check reason(s) below: TEST CANCELLED

ADULTERATED (adulterant/reason): _____

SUBSTITUTED

OTHER: _____

REMARKS: _____

X _____
 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: _____

X _____
 Signature of Medical Review Officer

(PRINT) Medical Review Officer's Name (First, MI, Last) _____ Date (Mo/Day/Yr) _____

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.

We are moving the Public Burden Statement to be on the back of Copies 1-5 and moving the Privacy Act Statement from the front of Copy 5 to the back of the page.

FEDERAL DRUG TESTING CUSTODY AND CONTROL FORM

SPECIMEN ID NO. **0000001**

ACCESSION NO.

OMB No. 0930-0158

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

A. Employer Name, Address, I.D. No. _____	B. MRO Name, Address, Phone No. and Fax No. _____
C. Donor SSN or Employee I.D. No. _____	
D. Specify Testing Authority: <input type="checkbox"/> HHS <input type="checkbox"/> NRC Specify DOT Agency: <input type="checkbox"/> FMCSA <input type="checkbox"/> FAA <input type="checkbox"/> FRA <input type="checkbox"/> FTA <input type="checkbox"/> PHMSA <input type="checkbox"/> USCG	
E. Reason for Test: <input type="checkbox"/> Pre-employment <input type="checkbox"/> Random <input type="checkbox"/> Reasonable Suspicion/Cause <input type="checkbox"/> Post Accident <input type="checkbox"/> Return to Duty <input type="checkbox"/> Follow-up <input type="checkbox"/> Other (specify) _____	
F. Drug Tests to be Performed: <input type="checkbox"/> THC, COC, PCP, OPI, AMP <input type="checkbox"/> THC & COC Only <input type="checkbox"/> Other (specify) _____	
G. Collection Site Address: _____	Collector Contact Info: Phone _____ Fax _____ Other _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate). URINE ORAL FLUID

COLLECTION: <input type="checkbox"/> Split <input type="checkbox"/> Single <input type="checkbox"/> None Provided, Enter Remark.
URINE: Collector reads urine temperature within 4 minutes. Temperature between 90° and 100° F? <input type="checkbox"/> Yes <input type="checkbox"/> No, Enter Remark <input type="checkbox"/> Observed, Enter Remark
ORAL FLUID: Split Type: <input type="checkbox"/> Serial <input type="checkbox"/> Concurrent <input type="checkbox"/> Subdivided Each Device Within Expiration Date? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> 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

<p><i>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.</i></p> <p>X _____ Signature of Collector</p> <p>_____/_____/_____ (PRINT) Collector's Name (First, MI, Last) Date (Mo/Day/Yr) AM PM</p>	<p>SPECIMEN BOTTLE(S)/TUBE(S) RELEASED TO:</p> <p>_____ Name of Delivery Service</p>
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STEP 5: COMPLETED BY DONOR

<p><i>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.</i></p>			
<p>X _____ Signature of Donor</p>	<p>_____ (PRINT) Donor's Name (First, MI, Last)</p>	<p>_____/_____/_____ Date (Mo/Day/Yr)</p>	
Email address: _____	Daytime Phone No. () _____	Evening Phone No. () _____	Date of Birth _____ (Mo/Day/Yr)
<p>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.</p>			

We are adding Steps 6 and 7 from the MRO Copy (Copy 2) here and moving the Public Burden Statement to the back of the page.

Public Burden Statement

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E. Reason for Test: <input type="checkbox"/> Pre-employment <input type="checkbox"/> Random <input type="checkbox"/> Reasonable Suspicion/Cause <input type="checkbox"/> Post Accident <input type="checkbox"/> Return to Duty <input type="checkbox"/> Follow-up <input type="checkbox"/> Other (specify) _____	
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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.

X _____ (PRINT) Donor's Name (First, MI, Last) _____ Date (Mo/Day/Yr) _____

Signature of Donor _____

Email address: _____ Daytime Phone No. () _____ Evening Phone No. () _____ Date of Birth _____ (Mo/Day/Yr)

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.

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 is provided in 5 U.S.C. Sec. 3301 (2), 5 U.S.C. Sec. 7301, 5 U.S.C. Sec. 12564 and 5 U.S.C. 7301, test results may be disclosed to the 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.

We are adding Steps 6 and 7 from the MRO Copy (Copy 2) here and moving the Privacy Act Statement and Public Burden Statement to the back of the page.

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