

SPECIMEN ID NO. _____

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

ACCESSION NO. _____

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 <input type="checkbox"/> DOT – 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 Phone No. _____	
Collector Fax No. _____	

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate) Collector reads specimen temperature within 4 minutes.

Temperature between 90° and 100° F? <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No, Enter Remark _____	Collection: <input type="checkbox"/> Split <input type="checkbox"/> Single <input type="checkbox"/> None Provided, Enter Remark _____	<input type="checkbox"/> Observed, Enter Remark _____
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REMARKS _____

STEP 3: Collector affixes bottle seal(s) to bottle(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.

<p>X _____ Signature of Collector</p> <p style="text-align: right;">AM PM</p> <p>_____ (PRINT) Collector's Name (First, MI, Last)</p> <p>_____/_____/_____ Date (Mo./Day/Yr)</p> <p>_____/_____/_____ Time of Collection</p>	<p style="text-align: center;">SPECIMEN BOTTLE(S) RELEASED TO:</p> <p>_____ Name of Delivery Service</p>
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<p>RECEIVED AT LAB OR IITF:</p> <p>X _____ Signature of Accessioner</p> <p>_____ (PRINT) Accessioner's Name (First, MI, Last)</p> <p>_____/_____/_____ Date (Mo./Day/Yr)</p>	<p style="text-align: center;">SPECIMEN BOTTLE(S) RELEASED TO:</p> <p>Primary Specimen Bottle Seal Intact <input type="checkbox"/> YES <input type="checkbox"/> NO</p> <p>If NO, Enter remark in Step 5A. _____</p>
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STEP 5A: PRIMARY SPECIMEN REPORT - COMPLETED BY TEST FACILITY

<input type="checkbox"/> NEGATIVE	<input type="checkbox"/> POSITIVE for:	<input type="checkbox"/> Marijuana Metabolite (Δ9-THCA)	<input type="checkbox"/> 6-Acetylmorphine
<input type="checkbox"/> DILUTE	<input type="checkbox"/> Cocaine Metabolite (BZE)	<input type="checkbox"/> Morphine	<input type="checkbox"/> Methamphetamine
	<input type="checkbox"/> PCP	<input type="checkbox"/> Codeine	<input type="checkbox"/> Amphetamine
<input type="checkbox"/> REJECTED FOR TESTING	<input type="checkbox"/> ADULTERATED	<input type="checkbox"/> SUBSTITUTED	<input type="checkbox"/> INVALID RESULT
			<input type="checkbox"/> MDMA
			<input type="checkbox"/> MDA
			<input type="checkbox"/> MDEA

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

<p>_____ Laboratory Name</p> <p>_____ Laboratory Address</p>	<p><input type="checkbox"/> RECONFIRMED <input type="checkbox"/> FAILED TO RECONFIRM - REASON _____</p> <p>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.</p> <p>X _____ Signature of Certifying Scientist</p> <p>_____ (PRINT) Certifying Scientist's Name (First, MI, Last)</p> <p>_____/_____/_____ Date (Mo./Day/Yr.)</p>
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<p style="text-align: center;">A</p> <p style="text-align: center;">SPECIMEN ID NO. _____</p>	<p>↑ HINGE ↑</p> <p>PLACE OVER CAP</p> <p>↓ LATCH ↓</p> <p>↑ HINGE ↑</p> <p>PLACE OVER CAP</p> <p>↓ LATCH ↓</p>	<p style="text-align: center;">_____/_____/_____ Date (Mo./Day/Yr.)</p> <p style="text-align: center;">_____ Donor's Initials</p> <hr/> <p style="text-align: center;">_____/_____/_____ Date (Mo./Day/Yr.)</p> <p style="text-align: center;">_____ Donor's Initials</p>
<p style="text-align: center;">B</p> <p style="text-align: center;">SPECIMEN ID NO. _____</p>	<p>↑ HINGE ↑</p> <p>PLACE OVER CAP</p> <p>↓ LATCH ↓</p>	<p style="text-align: center;">_____/_____/_____ Date (Mo./Day/Yr.)</p> <p style="text-align: center;">_____ Donor's Initials</p>

SPECIMEN ID NO. _____

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

ACCESSION NO. _____

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: HHS NRC DOT – 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 Phone No. _____

Collector Fax No. _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate) Collector reads specimen temperature within 4 minutes.

Temperature between 90° and 100° F? Yes No, Enter Remark _____ Collection: Split Single None Provided, Enter Remark _____ Observed, Enter Remark _____

REMARKS _____

STEP 3: Collector affixes bottle seal(s) to bottle(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) 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

STEP 5: COMPLETED BY DONOR

I certify that I provided my urine specimen to the collector; that I have not adulterated it in any manner; each specimen bottle 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 is correct.

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

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

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)

SPECIMEN ID NO. _____

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

ACCESSION NO. _____

A. Employer Name, Address, I.D. No. _____ C. Donor SSN or Employee I.D. No. _____ D. Specify Testing Authority: <input type="checkbox"/> HHS <input type="checkbox"/> NRC <input type="checkbox"/> DOT – 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 Phone No. _____ Collector Fax No. _____	B. MRO Name, Address, Phone No. and Fax No. _____
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STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate) Collector reads specimen temperature within 4 minutes.

Temperature between 90° and 100° F? <input type="checkbox"/> Yes <input type="checkbox"/> No, Enter Remark _____	Collection: <input type="checkbox"/> Split <input type="checkbox"/> Single <input type="checkbox"/> None Provided, Enter Remark _____	<input type="checkbox"/> Observed, Enter Remark _____
REMARKS _____		

STEP 3: Collector affixes bottle seal(s) to bottle(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) RELEASED TO: _____ Name of Delivery Service
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STEP 5: COMPLETED BY DONOR

I certify that I provided my urine specimen to the collector; that I have not adulterated it in any manner; each specimen bottle 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 is correct.

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

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)

SPECIMEN ID NO. _____

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

ACCESSION NO. _____

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: HHS NRC DOT – 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 Phone No. _____

Collector Fax No. _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate) Collector reads specimen temperature within 4 minutes.

Temperature between 90° and 100° F? Yes No, Enter Remark _____ Collection: Split Single None Provided, Enter Remark _____ Observed, Enter Remark _____

REMARKS _____

STEP 3: Collector affixes bottle seal(s) to bottle(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) 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

STEP 5: COMPLETED BY DONOR

I certify that I provided my urine specimen to the collector; that I have not adulterated it in any manner; each specimen bottle 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 is correct.

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

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

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)

SPECIMEN ID NO. _____

STEP 1: COMPLETED BY COLLECTOR OR EMPLOYER REPRESENTATIVE

ACCESSION NO. _____

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: HHS NRC DOT – 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 Phone No. _____

Collector Fax No. _____

STEP 2: COMPLETED BY COLLECTOR (make remarks when appropriate) Collector reads specimen temperature within 4 minutes.

Temperature between 90° and 100° F? Yes No, Enter Remark _____ Collection: Split Single None Provided, Enter Remark _____ Observed, Enter Remark _____

REMARKS _____

STEP 3: Collector affixes bottle seal(s) to bottle(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) 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 _____

STEP 5: COMPLETED BY DONOR

I certify that I provided my urine specimen to the collector; that I have not adulterated it in any manner; each specimen bottle 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 is correct.

X

Signature of Donor _____

(PRINT) Donor's Name (First, MI, Last) _____

Date (Mo/Day/Yr) _____

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

In accordance with applicable Federal requirements, my verification is:

NEGATIVE **POSITIVE** for: _____
 DILUTE

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

ADULTERATED (adulterant/reason): _____

SUBSTITUTED

OTHER: _____

TEST CANCELLED

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) _____

COPY 5 - DONOR COPY

REORDER #0782 REV. 01/14

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, 1 Choke Cherry Road, Room 7-1044, Rockville, Maryland, 20857.



B-1A

Instructions for Completing the Federal Drug Testing Custody and Control Form

When making entries use black or blue ink pen and press firmly

Collector ensures that the name and address of the HHS-certified Instrumented Initial Test Facility (IITF) or HHS-certified laboratory are on the top of the CCF and that the Specimen I.D. number on the top of the CCF matches the Specimen I.D. number on the labels/seals.

STEP 1:

- Collector ensures that the required information is in STEP 1. Collector enters a remark in STEP 2 if Donor refuses to provide his/her SSN or Employee I.D. number.
- Collector gives collection container to Donor and instructs Donor to provide a specimen. Collector notes any unusual behavior or appearance of Donor in the remarks line in STEP 2. If Donor conduct at any time during the collection process clearly indicates an attempt to tamper with the specimen, Collector notes the conduct in the remarks line in STEP 2 and takes action as required.

STEP 2:

- Collector checks specimen temperature within 4 minutes after receiving the specimen from Donor, and marks the appropriate temperature box in STEP 2. If temperature is outside the acceptable range, Collector enters a remark in STEP 2 and takes action as required.
- Collector inspects the specimen and notes any unusual findings in the remarks line in STEP 2 and takes action as required. Any specimen with unusual physical characteristics (e.g. unusual color, presence of foreign objects or material, unusual odor) cannot be sent to an IITF and must be sent to an HHS-certified laboratory for testing as required.
- Collector determines the volume of specimen in the collection container. If the volume is acceptable, Collector proceeds with the collection. If the volume is less than required by the Federal Agency, Collector takes action as required, and enters remarks in STEP 2. If no specimen is collected by the end of the collection process, Collector checks the *None Provided* box, enters a remark in STEP 2, discards Copy 1 and distributes remaining copies as required.
- Collector checks the Split or Single specimen collection box. If the collection is observed, Collector checks the Observed box and enters a remark in STEP 2.

STEP 3:

- Donor watches Collector pour the specimen from the collection container into the specimen bottle(s), place the cap(s) on the specimen bottle(s), and affix the label(s)/seal(s) on the specimen bottle(s).
- Collector dates the specimen bottle label(s)/seal(s) after placement on the specimen bottle(s).
- Donor initials the specimen bottle label(s)/seal(s) after placement on the specimen bottle(s).
- Collector turns to Copy 2 (Medical Review Officer Copy) and instructs Donor to read and complete the certification statement in STEP 5 (signature, printed name, date, phone numbers, and date of birth). If Donor refuses to sign the certification statement, Collector enters a remark in STEP 2 on Copy 1.

STEP 4:

- Collector completes STEP 4 on Copy 1 (signature, printed name, date, time of collection, and name of delivery service), places the sealed specimen bottle(s) and Copy 1 of the CCF in a leak-proof plastic bag, seals the bag, prepares the specimen package for shipment, and distributes the remaining CCF copies as required.

Privacy Act Statement: (For Federal Employees Only)

Submission of the information on the attached form is voluntary. However, incomplete submission of the information, refusal to provide a urine 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 urine 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, 1 Choke Cherry Road, Room 7-1044, Rockville, Maryland, 20857.