

Drug testing payment policy

Policy effective date: Sept. 23, 2024

Applies to

Coverage is subject to member's specific benefits. Group specific policy will supersede this policy when applicable.

- **HMO/EPO**: This policy applies to insured HMO/EPO plans.
- **POS**: This policy applies to insured POS plans.
- PPO: This policy applies to insured PPO plans. Consult individual plan documents as state mandated benefits may apply. If there is a conflict between this policy and a plan document, the provisions of the plan document will govern.
- ASO: For self-funded plans, consult individual plan documents. If there is a conflict between this policy and a self-funded plan document, the provisions of the plan document will govern.
- INDIVIDUAL: For individual policies, consult the individual insurance policy. If there is a conflict between this medical policy and the individual insurance policy document, the provisions of the individual insurance policy will govern.
- **MEDICARE**: Coverage is determined by the Centers for Medicare and Medicaid Services (CMS) and/or the Evidence of Coverage (EOC); if a coverage determination has not been adopted by CMS, this policy applies.
- MEDICAID/HEALTHY MICHIGAN PLAN: For Medicaid/Healthy Michigan Plan members, this policy will apply. Coverage is based on medical necessity criteria being met and the appropriate code(s) from the coding section of this policy being included on the Michigan Medicaid Fee Schedule. If there is a discrepancy between this policy and the Michigan Medicaid Provider Manual, the Michigan Medicaid Provider Manual will govern. For Medical Supplies/DME/Prosthetics and Orthotics, refer to the Michigan Medicaid Fee Schedule to verify coverage.

Medical policy

Drug Testing – 91611

Reimbursement information

Reimbursement will be considered for presumptive drug testing codes 80305, 80306 and 80307 and for definitive drug testing codes 80320-80377, 83992, G0480-G0483 and G0659. Third party codes are not reimbursable. Medical records may be requested from the ordering provider. See the Documentation requirements section below.

Note: HMO/EPO plans require prior authorization for all OON labs. POS and PPO plans don't require prior authorization for OON labs; however, all OON labs will be processed at the member's out-of-network benefit level.

Definitions

Presumptive drug testing: A testing methodology to determine the presence or absence of a substance belonging to a general class of drugs. The test result is expressed in non-numerical terms (i.e. positive or negative).

Definitive drug testing: A testing methodology to determine the specific concentration of a drug or drug metabolite. The test result is expressed in numerical terms.

- Review of presumptive results by ordering/treating provider should occur prior to ordering of definitive testing; rationale for ordering definitive testing is detailed in the medical record
- Rationale for ordering definitive drugs should be detailed in the medical record and detail the inconsistent positive finding from qualitative testing (when applicable)
- Rationale for ordering a definitive drug test in lieu of presumptive test should also be documented in the medical record (i.e., Presumptive test to evaluate drug not available)

Qualitative or presumptive drug testing must be medically necessary as defined by medical policy criteria for coverage.

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Documentation requirements

The ordering practitioner's documentation must support the test(s) ordered. Each drug or drug class ordered should be documented in the member's medical record and detailed on the lab order. The medical records should also detail the reasons each test is indicated and ordered to support management of the member's specific medical condition. Such documentation must indicate how the test results will impact clinical care.

- Standing orders or custom panel should not be referenced on the written lab order; only panel tests defined by CMS or CPT are acceptable
- Specific drugs or drug classes should be clearly detailed
- Orders must be signed and dated by the ordering practitioner
- Standard orders and/or routine screenings as part of a practitioner's protocol are not payable without supporting documentation to support member's specific medical assessment and treatment

Medical records may be requested to support accurate coding and support testing ordered. Although we do not expect billing labs to obtain medical records from ordering providers and submit them upon request, it is expected that at a minimum the lab order, requisition and results will be submitted.

This requisition must contain the following:

- Signed, valid requisition from the ordering provider that specifically outlines the tests being ordered
- Specific drugs or drug classes being tested
- Member specific information
- Ordering provider (full name and credentials) and ordering provider NPI
- Legible signature (photocopy, stamp, or signature on file is not accepted)
- Facility/location where specimen was collected
- Sample type (urine, blood, etc.)
- Date sample collected
- Time sample collected
- Individual who collected sample
- Date/time received at the lab facility

Final reports for lab results must contain the following:

- Complete detail for entity performing the lab service (name, address, CLIA)
- Patient full name
- Patient date of birth
- Ordering full name and NPI
- Facility name if different from above
- Date sample was collected
- Date sample was received at facility
- Date results were reported
- Detail of complete test results for each test performed

Claims submitted with insufficient documentation to support lab services will be denied. The provider submitting the claim will receive a denial if there is insufficient documentation to support all services reported.

Submitting orders or requested information alone does not guarantee services will be reimbursed. Supporting documentation from both lab and order provider must support requirements detailed in both payment and medical policy.

Limits

1 qualitative AND/OR 1 quantitative drug test may be billed on a single date of service

Coding information

CPT / HCPCS codes

Presumptive drug testing See CPT guidelines for definitions of drug class A & B		
Code	Description	
80305	Drug test(s), presumptive, any number of drug classes, any number of devices or procedures (e.g., immunoassay); capable of being read by direct optical observation only (e.g., dipsticks, cups, cards, cartridges) includes sample validation when performed, per date of service	
80306	Drug test(s), presumptive, any number of drug classes, any number of devices or procedures (e.g., immunoassay); read by instrument assisted direct optical observation (e.g., dipsticks, cups, cards, cartridges), includes sample validation when performed, per date of service	
80307	Drug test(s), presumptive, any number of drug classes, any number of devices or procedures, by instrument chemistry analyzers (e.g., utilizing immunoassay [e.g., EIA, ELISA, EMIT, FPIA, IA, KIMS, RIA]), chromatography (e.g., GC,HPLC), and mass spectrometry either with or without chromatography, (e.g., DART, DESI, GC-MS, GC-MS/MS, LC-MS, LC-MS/MS, LDTD, MALDI,TOF) includes sample validation when performed, per date of service	

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Definitive d Definitive tes	rug testing sting codes should not be performed routinely as this is not appropriate for every
	ledical records from the ordering provider must support the medical necessity of
The second secon	r drug class ordered.
Code	Description
G0480	Drug test(s), definitive, utilizing (1) drug identification methods able to identify individual drugs and distinguish between structural isomers (but not necessarily stereoisomers), including, but not limited to, GC/MS (any type, single or tandem and LC/MS (any type, single or tandem) and excluding immunoassays (e.g., IA, EIA, ELISA, EMIT, FPIA) and enzymatic methods (e.g., alcohol dehydrogenase)), (2) stable isotope or other universally recognized internal standards in all samples (e.g., to control for matrix effects, interferences and variations in signal strength), and (3) method or drugspecific calibration and matrix-matched quality control material (e.g., to control for instrument variations and mass spectral drift); qualitative or quantitative, all sources(s), includes specimen validity testing, per day, 1-7 drug class(es), including metabolite(s) if performed
G0481	Drug test(s), definitive, utilizing (1) drug identification methods able to identify individual drugs and distinguish between structural isomers (but not necessarily stereoisomers), including, but not limited to, GC/MS (any type, single or tandem) and LC/MS (any type, single or tandem) and excluding immunoassays (e.g., IA, EIA, ELISA, EMIT, FPIA) and enzymatic methods (e.g., alcohol dehydrogenase)); (2) stable isotope or other universally recognized internal standards in all samples (e.g., to control for matrix effects, interferences and variations in signal strength), and (3) method or drugspecific calibration and matrix-matched quality control material (e.g., to control for instrument variations and mass spectral drift); qualitative or quantitative, all sources(s), includes specimen validity testing, per day, 8-14 drug class(es), including metabolite(s) if performed
G0482	Drug test(s), definitive, utilizing (1) drug identification methods able to identify individual drugs and distinguish between structural isomers (but not necessarily stereoisomers), including, but not limited to, GC/MS (any type, single or tandem) and LC/MS (any type, single or tandem) and excluding immunoassays (e.g., IA, EIA, ELISA, EMIT, FPIA) and enzymatic methods (e.g., alcohol dehydrogenase)); (2) stable isotope or other universally recognized internal standards in all samples (e.g., to control for matrix effects, interferences and variations in signal strength), and (3) method or drugspecific calibration and matrix-matched quality control material (e.g., to control for instrument variations and mass spectral drift); qualitative or quantitative, all sources(s), includes specimen validity testing, per day, 15-21 drug class(es), including metabolite(s) if performed
G0483	Drug test(s), definitive, utilizing (1) drug identification methods able to identify individual drugs and distinguish between structural isomers (but not necessarily stereoisomers), including, but not limited to, GC/MS (any type, single or tandem) and LC/MS (any type, single or tandem) and excluding immunoassays (e.g., IA, EIA, ELISA, EMIT, FPIA) and enzymatic methods (e.g., alcohol dehydrogenase)); (2) stable isotope or other universally recognized internal standards in all samples (e.g., to control for matrix effects, interferences and variations in signal strength), and (3) method or drugspecific calibration and matrix-matched quality control material (e.g., to control for instrument variations and mass spectral drift); qualitative or quantitative, all sources(s), includes specimen validity testing, per day, 22 or more drug class(es), including metabolite(s) if performed

G0659	Drug test(s), definitive, utilizing drug identification methods able to identify individual drugs and distinguish between structural isomers (but not necessarily stereoisomers), including but not limited to, GC/MS (any type, single or tandem) and LC/MS (any type, single or tandem), excluding immunoassays (e.g., IA, EIA, ELISA, EMIT, FPIA) and enzymatic methods (e.g., alcohol dehydrogenase), performed without method or drug-specific calibration, without matrix-matched quality control material, or without use of stable isotope or other universally recognized internal standard(s) for each drug, drug metabolite or drug class per specimen; qualitative or quantitative, all sources, includes specimen validity testing, per day, any number of drug classes
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Validity testing codesValidity testing is performed to confirm results are accurate and valid. These codes are included in the above base codes and should not be billed separately. Codes 80320 – 80377 not payable by Priority Medicare and Medicaid - use G codes.

Code	Description
80320	Alcohols
80321	Alcohol biomarkers; 1 or 2
80322	Alcohol biomarkers; 3 or more
80323	Alkaloids, not otherwise specified
80324	Amphetamines; 1 or 2
80325	Amphetamines; 3 or 4
80326	Amphetamines; 5 or more
80327	Anabolic steroids; 1 or 2
80328	Anabolic steroids; 3 or more
80329	Analgesics, non-opioid; 1 or 2
80330	Analgesics, non-opioid; 3-5
80331	Analgesics, non-opioid; 6 or more
80332	Antidepressants, serotonergic class; 1 or 2
80333	Antidepressants, serotonergic class; 3-5
80334	Antidepressants, serotonergic class; 6 or more
80335	Antidepressants, tricyclic and other cyclicals; 1 or 2
80336	Antidepressants, tricyclic and other cyclicals; 3-5
80337	Antidepressants, tricyclic and other cyclicals; 6 or more
80338	Antidepressants, not otherwise specified
80339	Antiepileptics, not otherwise specified; 1-3
80340	Antiepileptics, not otherwise specified; 4-6
80341	Antiepileptics, not otherwise specified; 7 or more
80342	Antipsychotics, not otherwise specified; 1-3
80343	Antipsychotics, not otherwise specified; 4-6
80344	Antipsychotics, not otherwise specified; 7 or more
80345	Barbiturates
80346	Benzodiazepines; 1-12
80347	Benzodiazepines; 13 or more
80348	Buprenorphine
80349	Cannabinoids, natural
80350	Cannabinoids, synthetic; 1-3
80351	Cannabinoids, synthetic; 4-6
80352	Cannabinoids, synthetic; 7 or more
80353	Cocaine

80354 Fentanyl 80355 Gabapentin, non-blood 80356 Heroin metabolite 80357 Ketamine and norketamine 80358 Methadone 80359 Methylenedioxyamphetamines (MDA, MDEA, MDMA) 80360 Methylphenidate 80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80356Heroin metabolite80357Ketamine and norketamine80358Methadone80359Methylenedioxyamphetamines (MDA, MDEA, MDMA)80360Methylphenidate80361Opiates, 1 or more80362Opioids and opiate analogs; 1 or 280363Opioids and opiate analogs; 3 or 4	
80357 Ketamine and norketamine 80358 Methadone 80359 Methylenedioxyamphetamines (MDA, MDEA, MDMA) 80360 Methylphenidate 80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80358 Methadone 80359 Methylenedioxyamphetamines (MDA, MDEA, MDMA) 80360 Methylphenidate 80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80359 Methylenedioxyamphetamines (MDA, MDEA, MDMA) 80360 Methylphenidate 80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80360 Methylphenidate 80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80361 Opiates, 1 or more 80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80362 Opioids and opiate analogs; 1 or 2 80363 Opioids and opiate analogs; 3 or 4	
80363 Opioids and opiate analogs; 3 or 4	
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00264 Onicide and enicte analogo: Flor mare	
80364 Opioids and opiate analogs; 5 or more	
80365 Oxycodone	
80366 Pregabalin	
80367 Propoxyphene	
80368 Sedative hypnotics (non-benzodiazepines)	
80369 Skeletal muscle relaxants; 1 or 2	
80370 Skeletal muscle relaxants; 3 or more	
80371 Stimulants, synthetic	
80372 Tapentadol	
80373 Tramadol	
80374 Stereoisomer (enantiomer) analysis, single drug class	
80375 Drug(s) or substance(s), definitive, qualitative or quantitative, not other	rwise
specified	
80376 Drug(s) or substance(s), definitive, qualitative or quantitative, not other	rwise
specified; 4-6	
80377 Drug(s) or substance(s), definitive, qualitative or quantitative, not other	rwise
specified; 7 or more	
83992 Phencyclidine (PCP)	