

New Tests and Test Updates

Modified Date: 12/14/2011

In our continuing effort to provide you with the highest quality toxicology laboratory services available, we have compiled important changes regarding a number of tests we perform. Listed below are the types of changes that may be included in this notification, effective Monday, February 06, 2012

New Tests - Tests recently added to the NMS Labs test menu. New Tests are effective immediately.

Test Changes - Tests that have had changes to the method/ CPT code, units of measurement, scope of analysis, reference comments, or specimen requirements.

Discontinued Tests - Tests being discontinued with alternate testing suggestions.

Please use this information to update your computer systems/records. These changes are important to ensure standardization of our mutual laboratory databases.

If you have any questions about the information contained in this notification, please call our Client Support Department at (866) 522-2206. Thank you for your continued support of NMS Labs and your assistance in implementing these changes.

The CPT Codes provided in this document are based on AMA guidelines and are for informational purposes only. NMS Labs does not assume responsibility for billing errors due to reliance on the CPT Codes listed in this document.

Effective Date:

Monday, February 06, 2012



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Test Code	Test Name	New Test	Test Name	Method / CPT Code	Specimen Req.	Stability	Scope	Units	Reference Comments	Discontinue
0395SP	Armodafinil, Serum/Plasma	•								
2660B	Mercaptopurine, Blood			•	•	•				
3430TI	Perchloroethylene, Tissue									•
4281U	Synthetic Cannabinoid Metabolites (Qualitative) - Expanded, Urine					•				
9562U	Synthetic Cannabinoid Metabolites Screen - Expanded, Urine (Forensic)					•				
2320B	Volatiles Panel, Blood		•							
2320SP	Volatiles Panel, Serum/Plasma		•							
2320TI	Volatiles Panel, Tissue		•							
2320U	Volatiles Panel, Urine		•							
1355U	o-Toluidine, Urine	•								



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New Tests

0395SP A	Armodafin	il, Serum/Plasma			Effective Immediate	ly	
Scope o	of Analysis:	Armodafinil [HPLC]					
	Method(s):	High Performance Lie	quid Chromatogra	aphy (HPLC)			
	Purpose:	Therapeutic Drug Mo	nitoring				
	Category:	Stimulant					
Specimen Req	•••	2 mL Serum or Plasn	na				
	Im Volume:	0.7 mL					
Specia	l Handling:	Serum: Collect samp Plasma: Collect samp Promptly centrifuge a guidelines.	ole in Lavender to	op tube (EDTA	.) or Pink top tube. into a plastic screw capped vial using approved		
Specimen	Container:	Plastic container (pre	servative-free)				
Transport Ter	mperature:	Refrigerated					
Light	Protection:	Not Required					
Rejectio	on Criteria:	Polymer gel separation tube (SST or PST).					
·	Stability:	Room Temperature: ⁻ Refrigerated: 14 day(Frozen (-20 °C): 14 d	s)				
Meth	od: High	Performance Lic	uid Chromat	tography (ł	HPLC)		
Set-Up Days /	TAT: Tuesd	ay Thursday 2nd Shift	3 days (after set-	·up)			
CPT Co	ode: 82491						
Compound Na	ame / Alias	3	Units	RL	Reference Comment		
Armodafinil Nuvigil®			mcg/mL	0.2	The following peak plasma concentrations were reported following daily administration of armodafinil for 7 days: 50 mg: 1.8 +/- 0.2 mcg/mL 100 mg: 4.0 +/- 0.7 mcg/mL 250 mg: 9.2 +/- 0.7 mcg/mL 300 mg: 11 +/- 1.3 mcg/mL 400 mg: 13 +/- 5.3 mcg/mL		
1355U c	o-Toluidine	e. Urine			Effective Immediate	lv	

15550 0-1010101	ne, onne	
Scope of Analysis	s: o-Toluidine [LC-MS/MS]	
Method(s	: High Performance Liquid Chromatography/Tandem Mass Spectrometry (LC-MS/MS)	
Purpose	Exposure Monitoring	
Category	r: Industrial chemical	
Specimen Requirements	s: 7 mL Urine	
Minimum Volume	e: 3.2 mL	
Special Handling	g: Collect urine in a 50 mL plastic bottle containing 5 grams of citric acid. Collect sample a	at end of shift.
Specimen Containe	r: Plastic container	
Transport Temperature	e: Refrigerated	
Light Protectior	n: Not Required	
Rejection Criteria	a: None	
Stability	r: Room Temperature: 30 day(s) Refrigerated: 30 day(s) Frozen (-20 °C): 30 day(s)	



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Method: High Performance Liquid Chromatography/Tandem Mass Spectrometry (LC-MS/MS)

Set-Up Days / TAT: Wednesday 3 days (after set-up)

CPT Code: 83789

Compound Name / Alias	Units	RL	Reference Comment
o-Toluidine 1-Amino-2-methylbenzene; 1-Methyl-2- aminobenzene; 2-Amino-1-methbenzene; 2- Aminotoluene; 2-Methyl-1-aminobenzene; 2- Methylaniline; 2-Methylbenzenamine; Methyl-2-aminobenzene; o-Aminotoluene; o- Methylaniline; o-Methylbenzenamine; o- Tolyamine	ng/mL	0.25	The reported urinary o-toluidine concentration in non-occupationally exposed populations was approximately 2 ng/mL. These population studies included both smokers and non-smokers. Higher values have been reported in smokers than non-smokers. Post-shift urinary o-toluidine levels from workers in a rubber chemical plant averaged 98 +/- 119 ng/mL.

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2660B Mercap	opurine, Blood		
Summary of Ch	Specimen Requirements were changed. Stability was changed. Methods/CPT Codes were changed [LC-MS/MS (83789)]		
Specimen Require			
Transport Tempe			
Specimen Cor			
Light Pro			
Special Ha	•		
Rejection (S Scope of Ar Method (CPT	ability: Room Temperature: 21 day(s) Refrigerated: 30 day(s) Frozen (-20 °C): 30 day(s) alysis: LC-MS/MS (83789): Mercaptopurine		
•	ic Cannabinoid Metabolites (Qualitative) - Expanded, Urine		
Summary of Ch	inges: Stability was changed.		
S	ability: Room Temperature: 30 day(s) Refrigerated: 30 day(s) Frozen (-20 °C): 30 day(s)		
9562U Synthe	c Cannabinoid Metabolites Screen - Expanded, Urine (Forensic)		
Summary of Ch	inges: Stability was changed.		
S	ability: Room Temperature: 30 day(s) Refrigerated: 30 day(s) Frozen (-20 °C): 30 day(s)		
2320B Volatile	s Panel, Blood		
Summary of Ch	inges: Test Name was changed.		
2320SP Volatile	s Panel, Serum/Plasma		
Summary of Ch	inges: Test Name was changed.		
2320TI Volatile	s Panel, Tissue		

Summary of Changes: Test Name was changed.



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2320U Volatiles Panel, Urine

Summary of Changes: Test Name was changed.



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Discontinued Tests

Test Code	Test Name	Alternative Test
3430TI	Perchloroethylene, Tissue	No Alternate Tests Available