SPECIALTY LABORATORIES

2006 CPT Coding Change Recommendations

These recommendations for 2006 coding changes, additions and revisions to our recommended codes are based on information in the 2006 CPT and AMA instructions. All coding recommendations are subject to change at any time. It is the client's responsibility to verify the accuracy of the codes and to assign values based on the reimbursement for your area. If you have any questions, please refer to the Current Procedural Terminology (CPT) Manual published by the American Medical Association. To verify reimbursement or if you have questions regarding usage of a CPT code, please contact your local Medicare carrier. Please contact Client Services at 800-421-4449 if you have questions regarding CPT coding changes to custom panels.

***Changes in **bold type.**

TEST CODE	NAME OF TEST	2005 CPT CODES	2006 CPT CODE CHANGES
1410	Acetylcholine Receptor Binding AutoAbs	84238	83519
1412	Acetylcholine Receptor Blocking AutoAbs	84238	83519
1413	Acetylcholine Receptor Modulating AutoAbs	84238	83519
1518	Alpha-1-Antitrypsin Deficiency Fetal Study	83891, 83892x4, 83896x10, 83903x2, 83912	83891, 83892x4, **83908x10**, 83903x2, **83912x2**
1515	Alpha-1-Antitrypsin GenotypR™	83891, 83892x4, 83896x10, 83903x2, 83912	83891, 83892x4, **83908x10**, 83903x2, **83912x2**
A48457	ALT-LDL GGE (Gradient Gel Electrophoresis)	83715 but should have been 83716	83701
4565	AmpliChip™ CYP450	83891, 83892, 83896x29, 83901x2, 83912	83891, 83892, **83900**, **88384**
1900	Apolipoprotein Evaluation	82172x3	**82172x2**, **83695**
5206	Ashkenazi Jewish GenotypR™-Carrier Panel	83891, 83892x2, 83896x31, 83901, 83912	83891, **83892**, **83914x31**, **83900**, **83901x16**, 83912
5216	Bloom Syndrome GenotypR™-Carrier Testing	83891, 83892x2, 83896, 83901, 83912	83891, **83892**, **83914**, **83898**, 83912
5217	Bloom Syndrome GenotypR™-Diagnostic Testing	83891, 83892x2, 83896, 83901, 83912	83891, **83892**, **83914**, **83898**, 83912
5218	Bloom Syndrome GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896, 83901, 83912	83891, **83892**, **83914**, **83898**, 83912
5991	Cadiovascular Thrombotic Risk AssessR™	82172, 85246, 85384, 85420	**83695**, 85246, 85384, 85420
5226	Canavan Disease GenotypR™-Carrier Testing	83891, 83892x2, 83896x5, 83901, 83912	83891, **83892**, **83914x5**, **83900**, **83901**, 83912
5227	Canavan Disease GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x5, 83901, 83912	83891, **83892**, **83914x5**, **83900**, **83901**, 83912
5228	Canavan Disease GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x5, 83901, 83912	83891, **83892**, **83914x5**, **83900**, **83901**, 83912
1078	Celiac Disease GenotypR™	83891, 83894x2, 83896x5, 83901x2, 83912	83891, 83894x2, 83896x5, **83900**, 83912
1650	Cellular Immune Dysfunction Evaluation	85048, 86064, 86359, 86360, 88185x3	85048, **86355**, 86359, 86360, **88184**, **88185x2**
7230	Connexin 26 GenotypR™	83891, 83892x4, 83896x10, 83903x2, 83912	83891, 83892x4, **83908x10**, 83903x2, **83912x2**

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TEST CODE	NAME OF TEST	2005 CPT CODES	2006 CPT CODE CHANGES
3133	Cyclic Citrullinated Peptide	83520	86200
5355	Cystic Fibrosis 40 GenotypR™: Carrier Study	83890, 83892x2, 83896x40, 83901x2, 83912	83891, 83892, 83900, 83901x3, 83914x40, 83912
5356	Cystic Fibrosis 70 GenotypR™: Carrier Study	83890, 83892x2, 83896x70, 83901x2, 83912	83891, 83892x2, 83900, 83901x19, 83914x70, 83912
5357	Cystic Fibrosis 70 GenotypR™: Diagnostic Study	83890, 83892x2, 83896x70, 83901x2, 83912	83891, 83892x2, 83900, 83901x19, 83914x70, 83912
5358	Cystic Fibrosis 70 GenotypR™: Fetal Study	83890, 83892x2, 83896x70, 83901x2, 83912	83891, 83892x2, 83900, 83901x19, 83914x70, 83912
1966	Factor V (Leiden) GenotypR™	83891, 83892x2, 83896x5, 83903, 83912	83891, 83892x2, 83908x5, 83903, 83912
5371	Factor II (Prothrombin) GenotypR™	83891, 83892x2, 83896x5, 83903, 83912	83891, 83892x2, 83908x5, 83903, 83912
5236	Familial Dysautonomia GenotypR™-Carrier Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5237	Familial Dysautonomia GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5238	Familial Dysautonomia GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5246	Fanconi Anemia GrrpC GenotypR™-Carrier Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5247	Fanconi Anemia GrrpC GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5248	Fanconi Anemia GrrpC GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5363	Fragile X Fetal Study	83890, 83892x2, 83894x2, 83896, 83897, 83898, 83912	83891, 83892x2, 83894, 83896, 83897, 83898, 83909, 83912
5362	Fragile X GenotypR™	83890, 83894, 83898, 83912	83891, 83892x2, 83894, 83896, 83897, 83898, 83909, 83912
5256	Gaucher Disease GenotypR™-Carrier Testing	83891, 83892x2, 83896x8, 83901, 83912	83891, 83892, 83914x8, 83900, 83901x2, 83912
5257	Gaucher Disease GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x8, 83901, 83912	83891, 83892, 83914x8, 83900, 83901x2, 83912
5258	Gaucher Disease GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x8, 83901, 83912	83891, 83892, 83914x8, 83900, 83901x2, 83912
S48642	HDL GGE	83715 but should have been 83716	83701
5369	Hemochromatosis GenotypR™	83891, 83892x4, 83896x10, 83903x2, 83912	83891, 83892x4, 83908x10, 83903x2, 83912x2
8144	Hepatitis B Virus Core/Precore Mutant DetectR	83890, 83892x2, 83894x2, 83898x2, 83912	83890, 83894, 83896, 83898x2, 83912
8134	Hepatitis B Virus GenotypR™	83890, 83898, 83904x6, 83912	83890, 83894, 83896, 83898x2, 83904x7, 83912
7420	HIV Phenoscript	87903, 87904x2	87903, 87904x6
7420NY	HIV Phenoscript-New York	87903, 87904x2	87903, 87904x5
1376	HLA-A DetectR™	83891, 83894, 83896x66, 83901, 83912	83891, 83894, 83896x66, 83900, 83912
1368	HLA-A, B, C DetectR™	83891, 83894x3, 83896x156, 83901x3, 83912	83891, 83894x3, 83896x156, 83900, 83901, 83912

TEST CODE	NAME OF TEST	2005 CPT CODES	2006 CPT CODE CHANGES
1369	HLA-A, B, C, DR DetectR™	83891, 83894x4, 83896x199, 83901x4, 83912	83891, 83894x4, 83896x199, 83900, 83901x2, 83912
1377	HLA-B DetectR™	83891, 83894, 83896x28, 83901, 83912	83891, 83894, 83896x28, 83900, 83912
1364	HLA-B27 GenotypR™	83891, 83894, 83896x25, 83901, 83912	83891, 83894, 83896x25, 83900, 83912
1378	HLA-C DetectR™	83891, 83894, 83896x62, 83901, 83912	83891, 83894, 83896x62, 83900, 83912
1379	HLA-DR DetectR™	83891, 83894, 83896x43, 83901, 83912	83891, 83894, 83896x43, 83900, 83912
5921	LDL (Low Density Lipoprotein) Subfractions	82465, 83716	82465, 83701
3446	Lipoprotein (a)	82172	83695
3445	Lipoprotein Electrophoresis	82465, 83715, 84478	82465, 83700, 84478
S47985	Lipoprotein Electrophoresis Fluid	83715x4 but should have been 83715 only once	83700
1671	Lymphocyte Enumeration, Basic	85048, 86064, 86359, 86360	85048, 86355, 86359, 86360
1668	Lymphocyte Enumeration, Basic & NK Cells	85048, 86064, 86359, 86360, 86379	85048, 86355, 86359, 86360, 86357
1658	Lymphocyte Enumeration, T & B cell	85048, 86064, 86359	85048, 86355, 86359
5365	Maternal Cell Contamination (MCC) Detection Fetal/Cord Blood	83894x2, 83901x4, 83912	83891, 83900, 83901x3, 83909, 83912
4562	MTHFR C677T/A1298C GenotypR™	83891, 83892x4, 83896x10, 83903x2, 83912	83891, 83892x4, 83908x10, 83903x2, 83912x2
5266	Mucolipidosis Type IV GenotypR™-Carrier Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5267	Mucolipidosis Type IV GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
5268	Mucolipidosis Type IV GenotypR™-Fetal Study w/ reflex to MCC	83891, 83892x2, 83896x2, 83901, 83912	83891, 83892, 83914x2, 83898, 83912
1025	Myasthenia Gravis Evaluation	84238, 86256	83519, 86256
1026	Myasthenia Gravis Evaluation, Plus	84238x3, 86256	83519x3, 86256
1705	Narcolepsy EvaluatR™	83891, 83894x4, 83901x4, 83912	83891, 83894x4, 83900, 83901x2, 83912
5422	Natural Killer Cell EvaluatR™	83519, 85048, 86379,	83519, 85048, 86357
1872	Natural Killer Cell Quantitation	85048, 86379	85048, 86357
5276	Niemann-Pick Disease GenotypR™-Carrier Testing	83891, 83892x2, 83896x4, 83901, 83912	83891, 83892, 83914x4, 83900, 83912
5277	Niemann-Pick Disease GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x4, 83901, 83912	83891, 83892, 83914x4, 83900, 83912
5278	Niemann-Pick Disease GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x4, 83901, 83912	83891, 83892, 83914x4, 83900, 83912
S50671	NMR Lipo-Profile	83716	83701
2363	Ova & Parasite: *Coccidia* Evaluation	87015, 87206x3, 88313	87015, 87206x3, 87209
2362	Ova & Parasite: Comprehensive Exam with *Coccidia* Evaluation	87177, 87206x3, 88313x2	87177, 87206x3, 87209x2
2361	Ova & Parasite: Routine Exam	87177, 88313	87177, 87209
5375	Plasminogen Activator Inhibitor (PAI-1) GenotypR™	83891, 83892x2, 83896x5, 83903, 83912	83891, 83892x2, 83908x10, 83903, 83912

TEST CODE	NAME OF TEST	2005 CPT CODES	2006 CPT CODE CHANGES
RHV	Reflex HIV Phenoscript	87903, 87904x2	87903, 87904x6
1013	Rheumatoid Arthritis EvaluatR™	83520, 86431x3	86200, 86431x3
4940	Sirolimus	80299	80195
5286	Tay-Sachs Disease GenotypR™-Carrier Testing	83891, 83892x2, 83896x7, 83901, 83912	83891, 83892, 83914x7, 83900, 83901x2, 83912
5287	Tay-Sachs Disease GenotypR™-Diagnostic Testing	83891, 83892x2, 83896x7, 83901, 83912	83891, 83892, 83914x7, 83900, 83901x2, 83912
5288	Tay-Sachs Disease GenotypR™-Fetal Study reflex to MCC	83891, 83892x2, 83896x7, 83901, 83912	83891, 83892, 83914x7, 83900, 83901x2, 83912
5990	Thrombotic Risk AssessR™	82172, 83090, 85384	83090, 83695, 85384
5973	Thrombotic Risk Evaluation 3	82172, 83090, 85300, 85301, 85302, 85303, 85305, 85306x2, 85307, 85384, 85410, 85420	83090, 83695, 85300, 85301, 85302, 85303, 85305, 85306x2, 85307, 85384, 85410, 85420
5353	TPMT GenotypR™	83890, 83892x6, 83896x15, 83903x3, 83912	83891, 83892x6, 83908x15, 83903x3, 83912x3
S50365	VAP Cholesterol	83716, 84478	83701, 84478