

CURRENT PROCEDURAL TERMINOLOGY CODES (CPT[®] CODES)

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Healthy Pregnancy (Screening, Testing, Counseling, Preventive Medication, and Treatment)	
Prenatal Diagnosis of Chromosomal Abnormalities and Neural Tube Defects (NTDs) (Screening)	
82105	Alpha-fetoprotein; serum
82106	Amniotic fluid
84702	Gonadotropin, chorionic (hCG); quantitative
84703	Gonadotropin, chorionic (hCG); qualitative
86336	Inhibin A
83632	Lactogen, human placental (HPL) human chorionic somatomammotropin
59000	Amniocentesis
76946	Ultrasound guidance for amniocentesis
59015	Chorionic villus sampling
76945	Ultrasound guidance for chorionic villus sampling
59012	Cordocentesis
76941	Ultrasonic guidance for cordocentesis
89290	Biopsy, oocyte polar body or embryo blastomere, microtechnique (for pre-implantation genetic diagnosis); less than or equal to 5 embryos
89291	Greater than 5 embryos
11100-1	Skin Biopsy
83013	Acetylcholinesterase (ACHE)
82677	Quantitative Estriol
83020	Hemoglobin Electrophoresis, fractionation and quantitation
83896	Nucleic Acid Probe (each)
83898	Nucleic Acid Probe w/Amplification (PCR)
83912	DNA Interpretation and Report

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86316	Cancer Antigen Immunoassay
88230	Cell Culture, Lymphocytes
88233	Cell Culture, Tissue
88235	Cell Culture, Amnio/CVS
88237	Cell Culture, Bone Marrow
88239	Cell Culture, Solid Tumor
88245	Chromosome analysis for breakage syndromes; baseline Sister Chromatid Exchange, 20-25 cells
88248	Chromosome analysis for breakage syndromes; baseline breakage, score 50-100 cells, count 20 cells, 2 karyotypes (eg, for ataxia telangiectasia, Fanconi anemia, fragile X)
88261	Chromosome Analysis, 5 Cell, Karyotype, 15-20 cells
88262	Chromosome analysis; count 15-20 cells, 2 karyotypes, with banding
88267	Chromosome analysis, amniotic fluid or chorionic villus, count 15 cells, 1 karyotype, with banding
88269	Chromosomal Analysis, in situ for amniotic fluid cells, 1Karyotype with banding
88280	Chromosome Analysis, Additional Karyotype
88283	Additional Banding
88285	Additional Cells Counted
88289	Additional High Resolution
88271	Molecular cytogenetics, DNA probe, each (code applied 5 times, once for each probe in the assay)
88275	Interphase in situ hybridization, count 100 to 300 cells
88291	Molecular cytogenetics, interpretation and report

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99401	Preventive medicine counseling/risk factor reduction, 15 minutes
99402	Preventive medicine counseling/risk factor reduction, 30 minutes
99403	Preventive medicine counseling/risk factor reduction, 45 minutes
99404	Preventive medicine counseling/risk factor reduction, 60 minutes
S3828*	Complete gene sequence analysis; MLH1 gene
S3830*	Complete mlh1 and mlh2 gene sequence analysis for hereditary nonpolyposis colorectal cancer (HNPCC) genetic testing
S3831*	Single-mutation analysis (in individual with a known MLH1 and MLH2 mutation in the family) for hereditary nonpolyposis colorectal cancer (HNPCC) genetic testing
S3833*	Complete APC gene sequence analysis for susceptibility to familial adenomatous polyposis (FAP) and attenuated FAP
S3834*	Single-mutation analysis (in individual with a known APC mutation in the family) for susceptibility to familial adenomatous polyposis (FAP) and attenuated FAP
S3835*	Complete gene sequence analysis for cystic fibrosis genetic testing
S3837*	Complete gene sequence analysis for hemochromatosis genetic testing
S3840*	DNA analysis for germline mutations of the ret proto-oncogene for susceptibility to multiple endocrine neoplasia type 2
S3841*	Genetic testing for retinoblastoma
S3842*	Genetic testing for von Hippel-Lindau disease
S3843*	DNA analysis of the F5 gene for susceptibility to Factor V Leiden thrombophilia
S3844*	DNA analysis of the connexin 26 gene (GJB2) for susceptibility to congenital, profound deafness
S3845*	Genetic testing for alpha-thalassemia

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Prenatal Diagnosis of Chromosomal Abnormalities and Neural Tube Defects (NTDs) (Screening)	
S3846*	Genetic testing for hemoglobin E beta-thalassemia
S3847*	Genetic testing for Tay-Sachs disease
S3848*	Genetic testing for Gaucher disease
S3849*	Genetic testing for Niemann-Pick disease
S3850*	Genetic testing for sickle cell anemia
S3851*	Genetic testing for Canavan disease
S3853*	Genetic testing for myotonic muscular dystrophy
Prenatal Diagnosis of Chromosomal Abnormalities and Neural Tube Defects (NTDs) (Testing)	
59000	Amniocentesis
76946	Ultrasound guidance for amniocentesis
59015	Chorionic villus sampling
76945	Ultrasound guidance for chorionic villus sampling
76801	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation, first trimester (14 weeks 0 days), transabdominal approach; single or first gestation
76802	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation, first trimester (14 weeks 0 days), transabdominal approach; each additional gestation (List separately in addition to code for primary procedure)
76805	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation, after first trimester (> or = 14 weeks 0 days), transabdominal approach; single or first gestation
76810	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation, after first trimester (> or = 14 weeks 0 days), transabdominal approach; each additional gestation

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76811	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation plus detailed fetal anatomic examination, transabdominal approach; single or first gestation
76812	Ultrasound, pregnant uterus, real time with image documentation, fetal and maternal evaluation plus detailed fetal anatomic examination, transabdominal approach; each additional gestation
76815	Ultrasound, pregnant uterus, real time with image documentation, limited (eg, fetal heart beat, placental location, fetal position, and/or qualitative amniotic fluid volume), one or more fetuses
76816	Ultrasound, pregnant uterus, real time with image documentation, follow-up (e.g., re-evaluation of fetal size by measuring standard growth parameters and amniotic fluid volume, re-evaluation of organ system(s) suspected or confirmed to be abnormal on a previous scan), transabdominal approach, per fetus
76817	Ultrasound, pregnant uterus, real time image documentation, transvaginal
76818	Fetal biophysical profile; with non-stress testing
76819	Fetal biophysical profile; without non-stress testing

*“S” codes are national Permanent Level II HCPCS Codes that are maintained by the HCPCS National Panel, a group comprised of representatives from the Blue Cross/Blue Shield Association (BCBSA), the Health Insurance Association of America (HIAA), and the Centers for Medicare and Medicaid Services (CMS). Permanent Level II HCPCS Codes provide a standardized coding system that is managed jointly by public and private insurers, thus providing a stable system for claims processing. These codes can be used by all private and public insurers.