TRICORE REFERENCE LABORATORIES

CPT CODES
GENETIC/MOLECULAR TESTS 2019

81120 - IDH1 Common Variants
81121 - IDH2 Common Variants
81161 - HORIZON 27 PANEL
81162 - BRCA1/2 COMPREHENSIVE
81163 - BREAST AND OVARIAN HEREDITARY
81170 - BCR ABL1 MUTATION ANALYSIS FOR
81200 - ASHKENAZI JEWISH CARRIER TEST
81201 - FAMILIAL ADENOMATOUS POLYPOSIS
81203 - FAMILIAL ADENOMATOUS POLYPOSIS
81205 - ASHKENAZI JEWISH CARRIER TEST
81206 - t(9,22) BCR-ABL
81207 - t(9,22) Minor Breakpt
81208 - BCR/ABL, MRNA DETECTION, RT PR
81209 - ASHKENAZI JEWISH CARRIER TEST
81210 - BRAF gene anal,V600e var
81211 - BRCAVANTAGE COMPREHENSIVE
81212 - BRCA1/2 ASHKENAZI JEWISH
81213 - INTEGRATED BRAC ANALYSIS
81215 - FAMILIAL MUTATION, TARGETED SG
81217 - MLH1 AND BRCA2
81218 - CEBPA Gene Analysis Full
81219 - CALR Gene Analysis
81220 - CF Gene Analysis
81221 - CF Known Fam Variants
81222 - CF Dupl/Deletion Variants
81223 - CYSTIC FIBROSIS SEQ & DEL/DUP
81223 - PANORAMA PRENATAL AND HORIZON
81223 - HORIZON 3 PANEL
81224 - CYSTIC FIBROSIS SEQ & DEL/DUP
81225 - CYTOCHROME P450 PAIN MANAGEMEL

81226 - CYTOCHROME P450 2D6 (CYP2D6)
81227 - WARFARIN SENSITIVITY GENOTYPE
81228 - SIGNATURE CHIPOS
81229 - Cytogenomic Const Analys
81230 - 1A2V,2C19V,2C9GV,2D6CV,3A4V
81231 - 1A2V,2C19V,2C9GV,2D6CV,3A4V
81232 - DIHYDROPYRIMIDINE DEHYDROGENA3
81235 - EGFR gene anal,common var
81238 - HEMOPHILIA B SEQUENCING & DELE
81240 - F2 gene anal,20210G>A var
81241 - F5 gene anal, Lieden var
81242 - ASHKENAZI JEWISH CARRIER TEST
81243 - FMR1 Gene Analysis
81244 - Reflex to Fragile X Meth
81245 - FLT3 Gene Analysis Inter
81247 - GLUCOSE 6 PHOSPHATE DEHYDROGEN
81250 - ASHKENAZI JEWISH CARRIER TEST
81251 - ASHKENAZI JEWISH CARRIER TEST
81252 - PANORAMA PRENATAL/HORIZON 274
81254 - HEARING LOSS,NONSYNDROMIC PANL
81255 - ASHKENAZI JEWISH CARRIER TEST
81256 - Hemochromatosis gene anal
81256 - HEMOCHROMATOSIS 3 MUTATIONS
81257 - Alpha Thal,HBA1,A2
81259 - ERYTHROCYTOSIS EVALUATION
81260 - ASHKENAZI JEWISH CARRIER TEST
81261 - IGH gene rearrangement
81263 - IGH var reg somatic anal
81265 - MATERNAL CELL CONTAMINATION
81266 - MATERNAL CELL CONTAMINATION

TRICORE REFERENCE LABORATORIES

CPT CODES GENETIC/MOLECULAR TESTS 2019, continued

81267 - Chimerism, Post Tx
81270 - JAK2 gene anal,V617F var
81271 - Huntington Disease
81272 - KIT AML Gene Ana Targ Seq
81273 - KIT Gene Ana Mastocytosis
81275 - KRAS Gene Analysis
81276 - KRAS Additonal Variants
81280 - COMPREHENSIVE ARRHYTHMIA PANEL
81281 - SCN5A AND NPHS2 GENE EVAL
81282 - CARDIONEXT, ANALYSIS OF 84 GES
81287 - MGMT Methylation,unlisted
81288 - MLH1 Promoter Methylation
81290 - ASHKENAZI JEWISH CARRIER TEST
81291 - MTHFR Mutation
81292 - HEREDITARY NON POLYPOSIS
81293 - FAMILIAL MUTATION, TARGETED
81294 - HNPCC/LYNCH SYNDROME SEQUENCIG
81295 - HEREDITARY NON POLYPOSIS
81296 - FAMILIAL MUTATION, TARGETED
81297 - BRCA 1/2 ANALYSIS WITH OVANEXT
81298 - HEREDITARY NON POLYPOSIS
81300 - BRCA 1/2 ANALYSIS WITH OVANEXT
81301 - Microsat Instability anal
81302 - Rett Syndrome MECP2
81303 - MECP2 RELATED DISORDERS
81304 - RETT SYNDROME (MECP2), SEQUENC
81305 - MYD88 L265P MUTATION DETECTION
81310 - NPM1 Gene Analysis, Exon
81311 - NRAS Gene Ana Exon 2 & 3
81312 - Oculopharyngeal Dys

81313 - PROSTATE CANCER BIOMARKER
81314 - PDGFRA Gene Ana Targ Seq
81315 - PML/RARalpha translocate
81317 - OVANEXT ANALYSES OF 24 GENES
81319 - GASTROINTESTINAL HEREDITARY CA
81321 - PTEN RELATED DISORDERS
81322 - SPECIFIC SITE ANALYSIS OF PTEN
81323 - PTEN RELATED DISORDERS
81324 - COMPLETE HNPP EVALUATION
81325 - COMPLETE HNPP EVALUATION
81327 - EPI PROCOLON
81329 - Sp Musc Atrophy CNA
81330 - HORIZON PANEL 27
81331 - Prader-Willi Syndrome
81332 - A1AT DEFICIENCY PROFILE
81335 - TPMT Gene Analy Comm Vari
81340 - TCR BETA GENE REARRANGEMENT
81342 - TRG gene rearrangement
81350 - UGT1A1
81355 - WARFARIN SENSITIVITY GENOTYPE
81361 - BETA GLOBIN (HBB) HBS, HBC
81364 - ERYTHROCYTOSIS EVALUATION
81370 - Low res HLA type
81371 - HLA Typing, A,B,DR Loci
81372 - HLA CLASS I MOLECULAR TYPING
81373 - HLA Typing, B Locus
81374 - HLA B27
81375 - DRDQDRp Loci Low Res
81376 - HLA CELIAC
81378 - HLA Cl I&II (A,B,C,DR)

TRICORE REFERENCE LABORATORIES

CPT CODES GENETIC/MOLECULAR TESTS 2019, continued

81380 - A Locus High Res
81381 - HLA B5701
81382 - HLA Cl II (DQ)
81383 - NARCOLEPSY GENOTYPING
81400 - ASHKENAZI JEWISH CARRIER TEST (2)
81400 - HORIZON PANEL 3
81401 - Spinal Musc. Atrophy
81401 - TPMT Genetics
81402 - KIT Mut. AML Masto
81403 - JAK2, exon 12
81404 - KIT Mutation Melanoma
81404 - HORIZON PANEL 3
81405 - MEN2 Mutation Screen
81406 - POLG DNA SEQUENCING TEST (ALPS
81407 - TSC FAMILIAL MUTATION EVAL
81408 - COMPLETE DMD EVALUATION
81410 - AORTOPATHY SEQUENCING, 21 GENS
81411 - AORTOPATHY PANEL, SEQUENCING
81413 - ARHYTHMIA SEQ/DEL/DUP PANEL
81414 - ARHYTHMIA SEQ/DEL/DUP PANEL
81414 - LONG QT SYNDROME SEQUENCING
81415 - EXOME SEQUENCING SYMPTOM GUIDD
81416 - EXOME SEQUENCING SYMPTOM GUIDD (2)
81420 - NON INVASIVE PRENATAL TESTING
81422 - Fetal Chrom Microdeletion
81432 - CANCER PANEL, HEREDITARY
81433 - CANCER PANEL, HEREDITARY
81434 - RETINITIS PIGMENTOSA
81435 - CANCER PANEL, HEREDITARY
81436 - CANCER PANEL, HEREDITARY

81437 - CANCER PANEL, HEREDITARY
81438 - CANCER PANEL, HEREDITARY
81440 - MITOCHONDRIAL DISORDERS PANEL
81442 - NOONAN SPECTRUM DISORDERS PANL
81445 - Targ Solid Organ Neoplasm
81448 - CMT ADVANCED EVALUATION
81450 - Targ Hematolymphoid Neo
81455 - MYELOID MALIGNANCIES MUTATION
81460 - MITOCHONDRIAL DISORDERS PANEL
81465 - MITOCHONDRIAL DISORDERS PANEL
81479 - MGMT Methylation,unlisted
81503 - OVA 1
81507 - PRENATAL TESTING FETAL ANEUPLY
81507 - PANORAMA PRENATAL & HORIZON 4
81508 - MATERNAL SERUM SCREEN
81512 - PENTA SCREEN
81539 - PROSTATE SPECIFIC KALLIKREIN
81599 - SEQUENTIAL MATERNAL SCREEN
83520 - Kappa Lambda/Ratio *consided by some insurance guidelines to be a genetic test that requires a Prior Authorization be obtained before testing is ordered
END