Sonora Quest Test Code	Sonora Quest Test Name	CPT* Codes
903398	ABL KINASE DOMAIN MUTATION IN CML CELL-BASED	81170
904449	ABL Kinase Domain Mutin CML Plasma-based Leumeta(R)	81170
904071	ABL T315I Mutation IN CML C-Based	81401
904468	ACCUTYPE CP CLOPIDOGREL CYP2C19 GENOTYPE	81225
904520	ACCUTYPE(R) WARFARIN	81227,81355
901995	Alpha-1 Antitrypsin (AAT) Mutation Analysis	81332
902148	Alpha-Globin Common Mutations Analysis	81257
903923	Alpha-Globin Gene Deletion or Duplication	81269
902350	ALPHA-GLOBIN GENE SEQUENCING	81259
906877	AML Molecular Profile	81455
901928	AML1-ETO t(8 21)	81401
906698	APO E GENOTYPING CARDIOVASCULAR RISK	81401
906357	B CELL RECEPTOR IGH REARRANGEMENT PCR	81261
906359	B-CELL CLONALITY PNL (IGH IGK) PCR	81261,81264
906356	B-CELL RECP IGK GENE REARRANGEMENT	81264
906564	BCR-ABL 1 p190 (Minor) Quantitative	81206,81207
905842	BCR-ABL 1 p190 (Minor), Quantitative	81206,81207
905013	BCR-ABL1 KINASE DOMAIN MUTATION 35-NUCLEOTIDE INSERTION	81170
901424	BETA-GLOBIN COMPLETE	81364
905045	Biotinidase Mutation	81404
700811	Bloom Syndrome DNA Mutation Analysis	81209
905639	BRAD V600 Mutation Cobas(R)	81210
904908	BRAF Mutation Analysis	81210
906367	BRCA vantage(TM) Rearrangements	81164
906366	BRCAVANTAGE (TM) ASHKENAZI JEWISH SCREEN	81212
906474	BRCAVANTAGE(TM) ASHKENAZI JEWISH SCREEN W RFLX COMPREHENSIVE	81212
906369	BRCAVANTAGE(TM) COMPREHENSIVE	81162
903278	C KIT MUTATION ANALPLASMA BASED LEUMETA	81272
901422	CAH(21-HYDROXYLASE DEFICIENCY)	81402
906471	Calreticulin (CALR) Mutation Analysis	81219
900810	CANAVAN DISEASE DNA MUTATION	81200
906136	CATECHOL-O-METHYLTRANSFERASE GENO	81479
905098	CCL IgVH MUTATSTATUS LEU ETA (R)	81263
705554	CEBPA Mutation Analysis	81218
901590	CFTR Intron 8 Poly-T Analysis	81224
906672	CFVANTAGE (R) CYSTIC FIBROSIS EXPANDED SCREEN	81220
905362	CHIMERISM DONOR	81267
905001	CHIMERISM POST-TRANSPLANT	81267
905000	CHIMERISM POST-TRANSPLANT SORTED T-CELLS	81268
905099	CHIMERISM RECIPIENT PRE-TRANSPLANT	81479
905439	CHIMERISM SORTED B-CELLS	81268
905441	CHIMERISM SORTED GRANULOCYTES	81268
905440	CHIMERISM SORTED-OTHER-CELLS	81268
905645	Chromosomal Microarray POC ClariSure(R) Oligo-SNP	81229
904806	Chromosomal Microarray Postnatal ClariSure (R) Oligo-SNP	81229
705646	Chromosomal Microarray Prenatal ClariSure (R) Oligo-SNP	81229
906372	Chromosome Microarray Congenital Blood	81229
904941	C-Kit Mutation Analysis Cell Based	81272
901734	CLL IgVH MUTATION CELL	81263
904413	COLOVANTAGE(R) (METHYLATED SEPTIN9)	81327
901625	Cystic Fibrosis Comp Rare Mutation	81223
905587	Cystic Fibrosis D1152H Mutation Analysis	81221
900830	Cystic Fibrosis DNA Analysis Fetus	81220
902291	Cystic Fibrosis Gene Deletion or Duplication	81222
901182	Cystic Fibrosis Rare Mutation Analysis One Exon	81221
900616	CYSTIC FIBROSIS SCREEN	81220
902746	CYTOCHROME P450 2D6	81226
901695	DPD Gene Mutation	81232
906741	EGFR mutation	81235
904556	EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) MUTATION ANALYSIS	81235
21315	FACTOR V LEIDEN MUTATION	81241
905789	FACTOR XIII V34L MUTATION ANALYSIS	81400
900884	Familial Dysautonomia Mutation Analysis	81260
906126	FAMILIAL MEDITERRANEAN FEVER MUTATION	81402
900812	Fanconi Anemia DNA Mutation Analysis	81242
904495	FRAGILE X DNA ANALYSIS FETUS	81243
101548	Gaucher Disease DNA Mutation Analysis	81251
906840	GIVANTAGE HEREDITARY COLORECTAL CANCER PANEL	81435,81436
905683	GSDIa MUTATION ANALY(ASH JEW)	81250
906878	Hematology Molecular Profile	81455
109626	HEREDITARY HEMOCHROMATOSIS DNA MUTATION ANALYSIS	81256

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905598	Hereditary Persistence of Fetal Hemoglobin(HPFH)8 Mutation	81479
901709	HLA (A B DR) IR (SSP)	81371
906710	HLA (A B DR) IR (LUMINEX)	81371
904513	HLA (A B) IR for Patient Platelet Transfusion	81373 002
101050	HLA Class 1 Phenotype (ABC)	81372
901071	HLA CLASS I DNA HR	81379
101513	HLA CLASS II DNA HR	81382 003
101049	HLA CLASS II DNA LR	81375
901192	HLA ONE LOCUS IR	81373
701843	HLA TYPING FOR CELIAC DISEASE	81376 002,81382 002
905388	HLA-A LOW RESOLUTION	81373
905873	HLA-A29 DNA TYPING	81374
902652	HLA-B 5701 TYPING	81381
906404	HLA-B HIGH RESOLUTION	81380
905384	HLA-B LOW RESOLUTION	81373
904791	HLA-B27 DNA TYPING	81374
904362	HLA-B51 DNA TYPING	81374
903570	HLA-DQA 1HR	81383
903571	HLA-DQB1 Supplemental HR	81383
906406	HLA-DQB1 HIGH RESOLUTION	81382
906405	HLA-DRB1 HIGH RESOLUTION	81382
906125	HLA-DRB3 4 5 DNA TYPING LOW RESOLUTION	81376
15716	HUNTINGTONS DISEASE DNA	81271
903430	JAK2 V617F MUTATION QL PCR LEUMETA (R) W RFLX EX 12 13	81270
902434	JAK2 V617F MUTATION DETECTION	81270
906423	JAK2 V617F Mutation Detection Qualitative	81270
906100	JAK2 V617F QL Leumeta (R) w Rflx to Exons 12 13 and MPL	81270
902002	KELL ANTIGEN TESTING PARENTAL	81403
903929	KRAS MUTATION ANALYSIS	81275,81276
905609	Long Chain Acyl-CoA Dehydrog Mutation Analysis	81479
906541	LYNCH SYNDROME PANEL	81292,81294,81295,81297,81298,81300,81317,81319,81403
902294	Maple Syrup Disease (MSUD) Mut Analysis (Ashkenazi Jewish)	81205
900831	Maternal Cell Contamination Study STR Analysis	81265
900773	MED CHAIN ACYL Coa MOL ANAL	81400
901893	METHYLENETETRAHYDROFOLATE REDUCTASE (MTHFR) DNA MUTATION	81291
906664	METHYLENETETRAHYDROFOLATE REDUCTASE (MTHFR) DNA MUTATION	81291
906879	MPN Molecular Profile	81450
906692	MUCOLIPIDOSIS TYPE IV MUTATION	81290
906745	MYVANTAGE HEREDITARY COMPREHENSIVE CANCER PANEL	81432,81433,81435,81436
90071	NEO ALLOIMMTHROMBOCYTOPENIA	81400,86022
906805	NGS Colon Target Gene Panel Colon Cancer	81445
906808	NGS Full Target Gene Panel with fusions	81445
906807	NGS Lung Target Gene Panel Lung Cancer with fusions	81445
906806	NGS Melanoma Target Gene Panel Melanoma	81445
900832	Niemann-Pick (Type A) Mutation Analysis	81330
906109	NPM (EXON 12) MUT PLASMA LEUMETA	81310
904055	NPM (Exon 12) MutaAnalCell-Based	81310
904931	PDGFRA MUTATION ANALYSIS	81314
906382	PDM CYP450 3A4 Genotoye Qual	81230
906383	PDM CYP450 3A5 Genotype Qual	81231
905588	Phenylketonuria (PKU) Mutation Analysis	81406
904738	PLASMINOGEN ACTIVATOR INHIB-1 4G 5G	81400
905626	PLATELET ANTIGEN GENOTYPING PANEL	81105,81106,81107,81108,81109,81110,81111,81112, 86022
90055	PML-RARA t(15 17)Quantitative RT-PCR	81315
901373	PRADER-WILLI ANGELMAN SYNDROME	81331
906646	PRENATAL CARRIER SCREEN (CF,FRAGILE X,SMA)	81220,81401,81243
903177	Prothrom(Factor II)20210GA Mutation Analysis	81240
11887	PROTHROMBIN (FACTOR II) 20210GA MUTATION	81240
906529	PTEN SEQUENCING AND DELETION DUPLICATION	81321,81323
906553	QNATAL (TM) ADVANCED	81420
901683	Rett Syndrome Mutation	81302
904986	Rett Syndrome Rearrangement Del Dup	81304
593430	Rflx JAK2 Exons 12 and 13 Mutations QL Leumeta (TM)	81403
906127	Rh C c AG GENOTYPING (PARENTAL)	81479
903184	Rh C c ANTIGEN GENOTYPING PRENATAL	81479
903147	Rh E e ANTIGEN GENOTYPING PREN	81479
903049	Rh E e ANTIGEN GENOTYPING-PARE	81479
903164	RHD ZYGOSITY	81403
905492	SMA Carrier Screen	81329
906028	SMA Diagnostic Test	81329

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905729	TAC1 FULL GENE ANALYSIS	81479
711526	Tay Sachs Mutation	81255
906358	T-CELL CLONALITY PNL(TCRB TCRG)PCR	81340,81342
906354	T-CELL RECEPTOR BETA REARRANGEMENT	81340
801610	THROMBOPHILIA PANEL	81240,81241,81291
803744	THROMBOPHILIA PANEL	81240,81241,81291
902136	TPMT Genotype	81335
906104	UGT1A1 GENE POLYMORPHISM	81350
905447	von Willebrand Disease Mutation Analysis	81408
10300	XSENSE FRAGILE X WITH REFLEX	81243
901247	Y CHROMOSOME MICRODELETION DNA	81403

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