Proc Code	Test ID	Test Name	LCD	LCD A56199	LCD A56867	LCD A52378
81120	65661A	IDH1 COMMON VARIANTS	ICD10 LIMITATIONS A56199			
81120	92361A	IDH1 COMMON VARIANTS	ICD10 LIMITATIONS A56199			
81120	IDH12	IDH1/IDH2 MUTATION ANALYSIS, TUMOR	ICD10 LIMITATIONS A56199			
81120	NGAMT	AML 4 Gene Panel, Therapeutic	ICD10 LIMITATIONS A56199			
81121	65661B	IDH2 COMMON VARIANTS	ICD10 LIMITATIONS A56199			
81121	92361B	IDH2 COMMON VARIANTS	ICD10 LIMITATIONS A56199			
81121	IDH12	IDH1/IDH2 MUTATION ANALYSIS, TUMOR	ICD10 LIMITATIONS A56199			
81121	NGAMT	AML 4 Gene Panel, Therapeutic	ICD10 LIMITATIONS A56199			
81161	DBMD	DMD/BMD Deletion/Duplication	NON COVERED A56199			
81162	614318F	BRCA1&2 GEN FULL SEQ DUP/DEL	ICD10 LIMITATIONS A56199			
81162	614319C	BRCA1&2 GEN FULL SEQ DUP/DEL	ICD10 LIMITATIONS A56199			
81162	614574D	BRCA1&2 GEN FULL SEQ DUP/DEL	ICD10 LIMITATIONS A56199			
81162	614576B	BRCA1&2 GEN FULL SEQ DUP/DEL	ICD10 LIMITATIONS A56199			
81162	COMCP	Hereditary Common Cancer Panel	ICD10 LIMITATIONS A56199			
81162	HBOCZ	BRCA1/2 Full Gene Analysis	ICD10 LIMITATIONS A56199			
81162	PANCP	Hereditary Pancreatic Cancer Panel	ICD10 LIMITATIONS A56199			
81162	PRS8P	Hereditary Prostate Cancer Panel	ICD10 LIMITATIONS A56199			
81162	XCP	Hereditary Expanded Cancer Panel	ICD10 LIMITATIONS A56199			
81170	BAKDM	BCR/ABL1 MUTATION, SEQUENCING	ICD10 LIMITATIONS A56199			
81177	DRPL	DRPLA Gene Analysis	INDIVIDUAL REVIEW A56199			
81178	609505A	ATXN1 GENE DETC ABNOR ALLELE	INDIVIDUAL REVIEW A56199			
81178	G204	ATXN1 (SCA 1) Gene Analysis	INDIVIDUAL REVIEW A56199			
81178	SCAP	Spinocerebellar Ataxia Panel	INDIVIDUAL REVIEW A56199			
81179	609505B	ATXN2 GENE DETC ABNOR ALLELE	INDIVIDUAL REVIEW A56199			
81179	G205	ATXN2 (SCA 2) Gene Analysis	INDIVIDUAL REVIEW A56199			
81179	SCAP	Spinocerebellar Ataxia Panel	INDIVIDUAL REVIEW A56199			
81180	609505C	ATXN3 GENE DETC ABNOR ALLELE	INDIVIDUAL REVIEW A56199			
81180	G206	ATXN3 (SCA 3) Gene Analysis	INDIVIDUAL REVIEW A56199			
81180	SCAP	Spinocerebellar Ataxia Panel	INDIVIDUAL REVIEW A56199			
81181	609505E	ATXN7 GENE DETC ABNOR ALLELE	INDIVIDUAL REVIEW A56199			
81181	G207	ATXN7 (SCA 7) Gene Analysis	INDIVIDUAL REVIEW A56199			
81181	SCAP	Spinocerebellar Ataxia Panel	INDIVIDUAL REVIEW A56199			
81184	609505D	CACNA1A GEN DETC ABNOR ALLEL	INDIVIDUAL REVIEW A56199			
81184	G208	CACNA1A (SCA 6) Gene Analysis	INDIVIDUAL REVIEW A56199			
81184	SCAP	Spinocerebellar Ataxia Panel	INDIVIDUAL REVIEW A56199			
81185	616524A	CACNA1A GENE FULL GENE SEQ	INDIVIDUAL REVIEW A56199			
81185	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81186	G169	Gene CACNA1A	INDIVIDUAL REVIEW A56199			
81188	CSTB	CSTB, repeat expansion analysis	INDIVIDUAL REVIEW A56199			
81190	G168	Gene CSTB	INDIVIDUAL REVIEW A56199			
81194	NTRK	NTRK Gene Fusion Panel	INDIVIDUAL REVIEW A56199			
81201	614318G	APC GENE FULL SEQUENCE	NON COVERED A56199			
81201	614319A	APC GENE FULL SEQUENCE	NON COVERED A56199			
81201	614580A	APC GENE FULL SEQUENCE	NON COVERED A56199			
81201	COMCP	Hereditary Common Cancer Panel	NON COVERED A56199			

81201	THYRP	Hereditary Thyroid Cancer Panel	NON COVERED A56199			
81201	XCP	Hereditary Expanded Cancer Panel	NON COVERED A56199			
81202	_G050	Gene APC	NON COVERED A56199			
81204	SBULB	Spinobulbar Musc Atrophy, Kennedy's	NON COVERED A56199			
81206	65248C	BCR/ABL1 GENE MAJOR BP	ICD10 LIMITATIONS A56199			
81206	89006Z1	BCR/ABL1 GENE MAJOR BP	ICD10 LIMITATIONS A56199			
81206	B210R	BCR/ABL1, P210, QUANT, REFLEX	ICD10 LIMITATIONS A56199			
81206	BADX	BCR/ABL, RNA-Qual, Diagnostic	ICD10 LIMITATIONS A56199			
81206	BCRAB	BCR/ABL, p210, Quant, Monitor	ICD10 LIMITATIONS A56199			
81206	BCRFX	BCR/ABL1 REFLEX, QUAL/QUANT	ICD10 LIMITATIONS A56199			
81207	65248B	BCR/ABL1 GENE MINOR BP	ICD10 LIMITATIONS A56199			
81207	89006Z2	BCR/ABL1 GENE MINOR BP	ICD10 LIMITATIONS A56199			
81207	B190R	BCR/ABL1, P190, QUANT, REFLEX	ICD10 LIMITATIONS A56199			
81207	BA190	BCR/ABL, p190, Quant, Monitor	ICD10 LIMITATIONS A56199			
81207	BADX	BCR/ABL, RNA-Qual, Diagnostic	ICD10 LIMITATIONS A56199			
81207	BCRFX	BCR/ABL1 REFLEX, QUAL/QUANT	ICD10 LIMITATIONS A56199			
81208	65248A	BCR/ABL1 GENE OTHER BP	ICD10 LIMITATIONS A56199			
81208	89006Z3	BCR/ABL1 GENE OTHER BP	ICD10 LIMITATIONS A56199			
81208	BADX	BCR/ABL, RNA-Qual, Diagnostic	ICD10 LIMITATIONS A56199			
81208	BCRFX	BCR/ABL1 REFLEX, QUAL/QUANT	ICD10 LIMITATIONS A56199			
81210	36517A	BRAF GENE	ICD10 LIMITATIONS A56199			
81210	BBRAF	BRAF Analysis	ICD10 LIMITATIONS A56199			
81210	BRAFB	CFDNA BRAF V600 TEST, BLOOD	ICD10 LIMITATIONS A56199			
81210	BRAFD	BRAF V600 Somatic Mutation Analysis, Tumor	ICD10 LIMITATIONS A56199			
81210	RASFP	RAS/RAF Panel, Tumor	ICD10 LIMITATIONS A56199			
81215	_G085	GENE BRCA1	ICD10 LIMITATIONS A56199			
81217	_G086	GENE BRCA 2	ICD10 LIMITATIONS A56199			
81218	CEBPA	CEBPA Mutations, Sequencing	ICD10 LIMITATIONS A56199			
81219	CALR	MPN, CALR Gene Mutation, Exon 9	ICD10 LIMITATIONS A56199			
81219	CALX	CALR, Gene Mutation, Exon 9, Reflex	ICD10 LIMITATIONS A56199			
81219	MPNCM	MPN (CALR, MPL) REFLEX	ICD10 LIMITATIONS A56199			
81220	CFP	Cystic Fibrosis Mutation Panel	ICD10 LIMITATIONS A56199			
81221	_G003	Gene CFTR_SEQ	ICD10 LIMITATIONS A56199			
81222	_G004	Gene CFTR_MLPA	ICD10 LIMITATIONS A56199			
81222	35388A	CFTR GENE DUP/DELET VARIANTS	ICD10 LIMITATIONS A56199			
81222	35640D	CFTR GENE DUP/DELET VARIANTS	ICD10 LIMITATIONS A56199			
81222	CFTRZ	CFTR Gene, Full Gene Analysis	ICD10 LIMITATIONS A56199			
81222	HPPAN	Hereditary Pancreatitis Panel	ICD10 LIMITATIONS A56199			
81223	35388B	CFTR GENE FULL SEQUENCE	ICD10 LIMITATIONS A56199			
81223	35640C	CFTR GENE FULL SEQUENCE	ICD10 LIMITATIONS A56199			
81223	CFTRZ	CFTR Gene, Full Gene Analysis	ICD10 LIMITATIONS A56199			
81223	HPPAN	Hereditary Pancreatitis Panel	ICD10 LIMITATIONS A56199			
81225	2C19R	CYP2C19 Genotype, V	ICD10 LIMITATIONS A56199			
81225	610060C	CYP2C19 GENE COM VARIANTS	ICD10 LIMITATIONS A56199			
81225	PSYQP	Psychotropic PGx Panel, V	ICD10 LIMITATIONS A56199			
81226	610060B	CYP2D6 GENE COM VARIANTS	ICD10 LIMITATIONS A56199			

81226	PSYQP	Psychotropic PGx Panel, V	ICD10 LIMITATIONS A56199			
81227	2C9QT	CYP2C9 Genotype, V	ICD10 LIMITATIONS A56199			
81227	610060D	CYP2C9 GENE COM VARIANTS	ICD10 LIMITATIONS A56199			
81227	PSYQP	Psychotropic PGx Panel, V	ICD10 LIMITATIONS A56199			
81228	_G075	SINGLE-GENE LARGE DEL/DUP	NON COVERED A56199			
81228	COLAB	Hereditary Colon Cancer CGH Array	NON COVERED A56199			
81229	CMACB	Chromosomal Microarray, Blood	NON COVERED A56199			
81229	CMAMT	CHROMOSOMAL MICROARRAY, POC, FFPE	NON COVERED A56199			
81229	CMAP	Chromosomal Microarray, Prenatal	NON COVERED A56199			
81229	CMAPC	Chromosomal Microarray, POC	NON COVERED A56199			
81229	CMAPC_GC	Chromosomal Microarray, POC	NON COVERED A56199			
81229	CMAPC_PC	PROF COMP Chromosomal Microarray, POC	NON COVERED A56199			
81229	CMAPC_TC	TECH COMP Chromosomal Microarray, POC	NON COVERED A56199			
81229	CMTFT	CMA, TECH-ONLY, FRESH, CONGENITAL	NON COVERED A56199			
81229	CMTPE	CMA, TECH-ONLY, FFPE, CONGENITAL	NON COVERED A56199			
81230	3A4Q	CYP3A4 Genotype, V	NON COVERED A56199			
81230	610060E	CYP3A4 GENE COMMON VARIANTS	NON COVERED A56199			
81230	PSYQP	Psychotropic PGx Panel, V	NON COVERED A56199			
81231	3A5Q	CYP3A5 Genotype, V	NON COVERED A56199			
81231	610060F	CYP3A5 GENE COMMON VARIANTS	NON COVERED A56199			
81231	PSYQP	Psychotropic PGx Panel, V	NON COVERED A56199			
81232	DPYDG	DPYD FULL GENE SEQUENCING	NON COVERED A56199			
81232	DPYDQ	DPYD Genotype, V	NON COVERED A56199			
81235	EGFRD	CFDNA EGFR MUTATION ANALYSIS, BLOOD	NON COVERED A56199			
81235	EGFRS	EGFR Gene, Mutation Analysis, Tumor	NON COVERED A56199			
81235	EGFRW	EGFR with ALK Reflex, Tumor	NON COVERED A56199			
81235	T790M	CFDNA EGFR T790M TEST, BLOOD	NON COVERED A56199			
81238	NGSF9	F9 Gene, Full Gene NGS	NON COVERED A56199			
81240	AATHR	Thrombophilia Prof	NON COVERED A56199			
81240	PR360	Venous Thrombosis Panel	NON COVERED A56199			
81240	PR520	Coagulation Profile	NON COVERED A56199			
81240	PTNT	PROTHROMBIN G20210A MUTATION, B	NON COVERED A56199			
81241	F5DNA	FACTOR V LEIDEN (R506Q) MUTATION, B	NON COVERED A56199			
81241	PR360	Venous Thrombosis Panel	NON COVERED A56199			
81241	PR520	Coagulation Profile	NON COVERED A56199			
81243	FXS	Fragile X Synd,Mol Analys	NON COVERED A56199			
81244	FUFXS	Fragile X, Follow Up Analysis	NON COVERED A56199			
81245	19739Z	FLT3 GENE	ICD10 LIMITATIONS A56199			
81245	65661C	FLT3 GENE	ICD10 LIMITATIONS A56199			
81245	FLT	FLT3 Mutation Analysis, V	ICD10 LIMITATIONS A56199			
81245	NGAMT	AML 4 Gene Panel, Therapeutic	ICD10 LIMITATIONS A56199			
81246	19739D	FLT3 GENE ANALYSIS	ICD10 LIMITATIONS A56199			
81246	65661D	FLT3 GENE ANALYSIS	ICD10 LIMITATIONS A56199			
81246	FLT	FLT3 Mutation Analysis, V	ICD10 LIMITATIONS A56199			
81246	NGAMT	AML 4 Gene Panel, Therapeutic	ICD10 LIMITATIONS A56199			
81248	_G126	Gene G6PD_SEQ	NON COVERED A56199			

81249	G6PDB	G6PD FULL GENE SEQUENCING	NON COVERED A56199			
81253	_G127	Gene GJB2_SEQ	NON COVERED A56199			
81256	HFE	Hemochromatosis HFE Gene Analysis,B	ICD10 LIMITATIONS A56199			
81258	_G128	Gene HBA1/HBA2_SEQ	NON COVERED A56199			
81259	WASEQ	ALPHA GLOBIN GENE SEQUENCING, B	NON COVERED A56199			
81259	WASQR	ALPHA GLOBIN GENE SEQUENCING, B	NON COVERED A56199			
81261	31141Z3	IGH GENE REARRANGE AMP METH	ICD10 LIMITATIONS A56199			
81261	31142Z3	IGH GENE REARRANGE AMP METH	ICD10 LIMITATIONS A56199			
81261	83123Z3	IGH GENE REARRANGE AMP METH	ICD10 LIMITATIONS A56199			
81261	BCGBM	Immunoglobulin Gene Rearrange, BM	ICD10 LIMITATIONS A56199			
81261	BCGR	Immunoglobulin Gene Rearrange, B	ICD10 LIMITATIONS A56199			
81261	BCGRV	Immunoglobulin Gene Rearrange, V	ICD10 LIMITATIONS A56199			
81263	BCLL	IGH Somatic Hypermutation in B-CLL	ICD10 LIMITATIONS A56199			
81264	31141Z2	IGK REARRANGEABN CLONAL POP	ICD10 LIMITATIONS A56199			
81264	31142Z2	IGK REARRANGEABN CLONAL POP	ICD10 LIMITATIONS A56199			
81264	83123Z2	IGK REARRANGEABN CLONAL POP	ICD10 LIMITATIONS A56199			
81264	BCGBM	Immunoglobulin Gene Rearrange, BM	ICD10 LIMITATIONS A56199			
81264	BCGR	Immunoglobulin Gene Rearrange, B	ICD10 LIMITATIONS A56199			
81264	BCGRV	Immunoglobulin Gene Rearrange, V	ICD10 LIMITATIONS A56199			
81265	_STR1	Comp analysis using STR (Bill only)	ICD10 LIMITATIONS A56199			
81265	CHRGB	Chimerism-Recipient Germline	ICD10 LIMITATIONS A56199			
81265	MULT	Zygosity Testing	ICD10 LIMITATIONS A56199			
81265	SPECI	Specimen Source Identification	ICD10 LIMITATIONS A56199			
81266	_STR2	Add'l comp analysis w/STR (Bill Only)	ICD10 LIMITATIONS A56199			
81266	ADONO	ADDITIONAL CHIMERISM DONOR	ICD10 LIMITATIONS A56199			
81267	CHIMU	CHIMERISM TRANSPLANT NO CELL SORT	ICD10 LIMITATIONS A56199			
81268	SORT1	CHIMERISM CELL SORT 1	ICD10 LIMITATIONS A56199			
81268	SORT2	CHIMERISM CELL SORT 2	ICD10 LIMITATIONS A56199			
81269	ATHAL	ALPHA-GLOBIN GENE ANALYSIS	NON COVERED A56199			
81269	ATHL	ALPHA-GLOBIN GENE ANALYSIS (ATHL)	NON COVERED A56199			
81270	88715A	JAK2 GENE	ICD10 LIMITATIONS A56199			
81270	JAK2B	JAK2 V617F Mutation Detection, B	ICD10 LIMITATIONS A56199			
81270	JAK2M	JAK2 V617F MUTATION DETECTION, BM	ICD10 LIMITATIONS A56199			
81270	JAK2V	JAK2 V617F MUTATION DETECTION, V	ICD10 LIMITATIONS A56199			
81270	MPNR	MPN (JAK2 V617F, CALR, MPL) Reflex	ICD10 LIMITATIONS A56199			
81270	PVJAK	JAK2 EXON 12-15 SEQUENCING, REFLEX	ICD10 LIMITATIONS A56199			
81271	HAD	Huntington Disease Analysis	NON COVERED A56199			
81272	35342B	KIT GENE TARGETED SEQ ANALYS	ICD10 LIMITATIONS A56199			
81272	GISTP	GIST Panel, Tumor	ICD10 LIMITATIONS A56199			
81272	KITE	KIT MUTATION, HEMATOLOGIC NEOPLASM	ICD10 LIMITATIONS A56199			
81273	KITVS	KIT Asp816Val Mutation Analysis, V	ICD10 LIMITATIONS A56199			
81275	36517B	KRAS GENE	ICD10 LIMITATIONS A56199			
81275	610679A	KRAS Gene Analysis, variants in Exon 2	ICD10 LIMITATIONS A56199			
81275	616452A	KRAS GENE	ICD10 LIMITATIONS A56199			
81275	68003A	KRAS GENE	ICD10 LIMITATIONS A56199			
81275	KRASD	CFDNA KRAS 12, 13, 61, 146 BLOOD	ICD10 LIMITATIONS A56199			

81275	KRASP	KRAS Somatic Mutation Analysis, Tumor	ICD10 LIMITATIONS A56199			
81275	KRASW	KRAS Mutation Analysis, Peritoneal	ICD10 LIMITATIONS A56199			
81275	RASFP	RAS/RAF Panel, Tumor	ICD10 LIMITATIONS A56199			
81276	610679B	KRAS Gene Analysis, addl variants	ICD10 LIMITATIONS A56199			
81276	616452B	KRAS GENE ADDL VARIANTS	ICD10 LIMITATIONS A56199			
81276	68003B	KRAS GENE ADDL VARIANTS	ICD10 LIMITATIONS A56199			
81276	KRASD	CFDNA KRAS 12, 13, 61, 146 BLOOD	ICD10 LIMITATIONS A56199			
81276	KRASP	KRAS Somatic Mutation Analysis, Tumor	ICD10 LIMITATIONS A56199			
81276	KRASW	KRAS Mutation Analysis, Peritoneal	ICD10 LIMITATIONS A56199			
81277	CMAH	Chromosomal Microarray, Hematologic	INDIVIDUAL REVIEW A56199			
81277	CMAPT	Chromosomal Microarray, Tumor, FFPE	INDIVIDUAL REVIEW A56199			
81277	CMAPT_GC	Chromosomal Microarray, Tumor, FFPE	INDIVIDUAL REVIEW A56199			
81277	CMAPT_PC	PROF COMP Chromosomal Microarray, Tumor, FFPE	INDIVIDUAL REVIEW A56199			
81277	CMAPT_TC	TECH COMP Chromosomal Microarray, Tumor, FFPE	INDIVIDUAL REVIEW A56199			
81277	CMAT	Chromosomal Microarray, Tumor	INDIVIDUAL REVIEW A56199			
81283	IL28Q	IL28B Genotype, V	NON COVERED A56199			
81284	AFXN	FXN, Repeat Expansion Analysis	NON COVERED A56199			
81287	MGMT	MGMT Promoter Methylation, Tumor	ICD10 LIMITATIONS A56199			
81288	BMLHH	MLH1 Hypermethylation Analysis	INDIVIDUAL REVIEW A56199			
81288	ML1HM	MLH1 Hypermethylation	INDIVIDUAL REVIEW A56199			
81288	MLHPB	MLH1 Hypermethylation Analysis, B	INDIVIDUAL REVIEW A56199			
81291	610060G	MTHFR GENE	NON COVERED A56199			
81291	PSYQP	Psychotropic PGx Panel, V	NON COVERED A56199			
81292	614318C	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	614319H	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	614572C	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	614574F	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	614576D	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	64333A	MLH1 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81292	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81292	LYNCP	Lynch Syndrome Panel	INDIVIDUAL REVIEW A56199			
81292	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81292	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81292	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81293	_G005	Gene MLH1	INDIVIDUAL REVIEW A56199			
81295	614318D	MSH2 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81295	614319J	MSH2 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81295	614572D	MSH2 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81295	614574G	MSH2 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81295	614576E	MSH2 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81295	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81295	LYNCP	Lynch Syndrome Panel	INDIVIDUAL REVIEW A56199			
81295	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81295	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81295	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81296	_G006	Gene MSH2	INDIVIDUAL REVIEW A56199			

81298	614318E	MSH6 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81298	614319K	MSH6 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81298	614572E	MSH6 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81298	614574H	MSH6 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81298	614576F	MSH6 GENE FULL SEQ	INDIVIDUAL REVIEW A56199			
81298	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81298	LYNCP	Lynch Syndrome Panel	INDIVIDUAL REVIEW A56199			
81298	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81298	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81298	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81299	_G007	Gene MSH6	INDIVIDUAL REVIEW A56199			
81301	TMSI	Tumor, Microsatellite Instability	ICD10 LIMITATIONS A56199			
81302	35484A	MECP2 GENE FULL SEQ	NON COVERED A56199			
81302	MCP2Z	MECP2 Gene, Full Gene Analysis	NON COVERED A56199			
81302	MECPZ	MECP2 Gene, Full Gene Analysis	NON COVERED A56199			
81303	_G008	Gene MECP2_SEQ	NON COVERED A56199			
81304	_G076	Gene MECP2_MLPA	NON COVERED A56199			
81304	35484B	MECP2 GENE DUP/DELET VARIANT	NON COVERED A56199			
81304	MECPZ	MECP2 Gene, Full Gene Analysis	NON COVERED A56199			
81305	LPLFX	REFLEX TESTING OF MYD88 AND CXCR4	ICD10 LIMITATIONS A56199			
81305	MYD88	MYD88 L265P Gene Mutation Analysis	ICD10 LIMITATIONS A56199			
81307	614318H	PALB2 GENE FULL GENE SEQ	NON COVERED A56199			
81307	614319L	PALB2 GENE FULL GENE SEQ	NON COVERED A56199			
81307	614574J	PALB2 GENE FULL GENE SEQ	NON COVERED A56199			
81307	614576G	PALB2 GENE FULL GENE SEQ	NON COVERED A56199			
81307	COMCP	Hereditary Common Cancer Panel	NON COVERED A56199			
81307	PANCP	Hereditary Pancreatic Cancer Panel	NON COVERED A56199			
81307	PRS8P	Hereditary Prostate Cancer Panel	NON COVERED A56199			
81307	XCP	Hereditary Expanded Cancer Panel	NON COVERED A56199			
81310	NPM1Q	NPM1 Mutation Analysis, V	ICD10 LIMITATIONS A56199			
81311	36517E	NRAS GENE VARIANTS EXON 2&3	ICD10 LIMITATIONS A56199			
81311	63161E	NRAS GENE VARIANTS EXON 2&3	ICD10 LIMITATIONS A56199			
81311	65562C	NRAS GENE VARIANTS EXON 2&3	ICD10 LIMITATIONS A56199			
81311	NSRGP	Noonan Syndrome and Related Panel,B	ICD10 LIMITATIONS A56199			
81311	PMNSR	POSTMORTEM NOONAN AND RELATED PANEL	ICD10 LIMITATIONS A56199			
81311	RASFP	RAS/RAF Panel, Tumor	ICD10 LIMITATIONS A56199			
81314	35342C	PDGFRA GENE	ICD10 LIMITATIONS A56199			
81314	GISTP	GIST Panel, Tumor	ICD10 LIMITATIONS A56199			
81315	PMLR	PML/RARA QUANTITATIVE,PCR	ICD10 LIMITATIONS A56199			
81317	_PMS2	PGL_PMS2C (BILL ONLY)	INDIVIDUAL REVIEW A56199			
81317	65791A	PMS2 GENE FULL SEQ ANALYSIS	INDIVIDUAL REVIEW A56199			
81318	_G068	GENE PMS2_LR AND PMS2_SEQ	INDIVIDUAL REVIEW A56199			
81319	_G069	GENE PMS2_MLPA	INDIVIDUAL REVIEW A56199			
81319	_PMS2	PGL_PMS2C (BILL ONLY)	INDIVIDUAL REVIEW A56199			
81319	614318A	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	614319M	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			

81319	614320B	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	614572A	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	614574A	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	614576H	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	65791B	PMS2 GENE DUP/DELET VARIANTS	INDIVIDUAL REVIEW A56199			
81319	BRGYP	Hereditary Breast/Gyn Cancer Panel	INDIVIDUAL REVIEW A56199			
81319	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81319	LYNCP	Lynch Syndrome Panel	INDIVIDUAL REVIEW A56199			
81319	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81319	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81319	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81321	614318J	PTEN GENE FULL SEQUENCE	INDIVIDUAL REVIEW A56199			
81321	614319N	PTEN GENE FULL SEQUENCE	INDIVIDUAL REVIEW A56199			
81321	614575B	PTEN Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81321	614580B	PTEN GENE FULL SEQUENCE	INDIVIDUAL REVIEW A56199			
81321	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81321	PTNZ	PTEN Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81321	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81321	THYRP	Hereditary Thyroid Cancer Panel	INDIVIDUAL REVIEW A56199			
81321	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81322	_G014	Gene PTEN	INDIVIDUAL REVIEW A56199			
81324	PMPDD	PMP22 GENE, DELETION/DUPLICATION	NON COVERED A56199			
81326	_G088	GENE PMP22_MLPA	NON COVERED A56199			
81326	_G125	Gene PMP22_SEQ	NON COVERED A56199			
81328	SLC1Q	SLCO1B1 Genotype, V	NON COVERED A56199			
81329	SMNCS	SMA CARRIER BY DEL/DUP	NON COVERED A56199			
81329	SMNDX	SMA DIAGNOSTIC BY DEL/DUP	NON COVERED A56199			
81331	PWAS	Prader-Willi/Angelman Syn	NON COVERED A56199			
81336	SMN1Z	SMN1 Full Gene Analysis	NON COVERED A56199			
81337	_G115	Gene SMN1	NON COVERED A56199			
81339	MPLR	MPL Exon 10 Mutation Detection, R	ICD10 LIMITATIONS A56199			
81339	MPLVS	MPL Exon 10 Mutation Detection, V	ICD10 LIMITATIONS A56199			
81339	MPNML	MPL EXON 10 SEQUENCING, REFLEX	ICD10 LIMITATIONS A56199			
81340	31139Z1	TRB@ GENE REARRANGE AMPLIFY	ICD10 LIMITATIONS A56199			
81340	31140Z1	tTRB@ GENE REARRANGE AMPLIFY	ICD10 LIMITATIONS A56199			
81340	83122Z1	TRB@ GENE REARRANGE AMPLIFY	ICD10 LIMITATIONS A56199			
81340	TCGBM	T Cell Receptor Gene Rearrange, BM	ICD10 LIMITATIONS A56199			
81340	TCGR	T Cell Receptor Gene Rearrange, B	ICD10 LIMITATIONS A56199			
81340	TCGRV	T Cell Receptor Gene Rearrange, V	ICD10 LIMITATIONS A56199			
81340	TCRB	TCRVB Spectratyping, B	ICD10 LIMITATIONS A56199			
81340	TCRVB	TCR Spectratyping, B	ICD10 LIMITATIONS A56199			
81342	31139Z2	TRG GENE REARRANGEMENT ANAL	ICD10 LIMITATIONS A56199			
81342	31140Z2	TRG GENE REARRANGEMENT ANAL	ICD10 LIMITATIONS A56199			
81342	83122Z2	TRG GENE REARRANGEMENT ANAL	ICD10 LIMITATIONS A56199			
81342	TCGBM	T Cell Receptor Gene Rearrange, BM	ICD10 LIMITATIONS A56199			
81342	TCGR	T Cell Receptor Gene Rearrange, B	ICD10 LIMITATIONS A56199			
81342	TCGRV	T Cell Receptor Gene Rearrange, V	ICD10 LIMITATIONS A56199			

81345	TERT	TERT PROMOTER ANALYSIS, TUMOR	ICD10 LIMITATIONS A56199			
81350	60343Z	UGT1A1 GENE	NON COVERED A56199			
81350	83949Z	UGT1A1 GENE	NON COVERED A56199			
81350	U1A1Q	UGT1A1 TA Repeat Genotype, V	NON COVERED A56199			
81351	614318K	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614319O	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614574K	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614575E	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614576J	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614577A	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	614580D	TP53 GENE FULL GENE SEQUENCE	ICD10 LIMITATIONS A56199			
81351	COMCP	Hereditary Common Cancer Panel	ICD10 LIMITATIONS A56199			
81351	PANCP	Hereditary Pancreatic Cancer Panel	ICD10 LIMITATIONS A56199			
81351	PRS8P	Hereditary Prostate Cancer Panel	ICD10 LIMITATIONS A56199			
81351	RENCP	Hereditary Renal Cancer Panel	ICD10 LIMITATIONS A56199			
81351	THYRP	Hereditary Thyroid Cancer Panel	ICD10 LIMITATIONS A56199			
81351	WILMP	Hereditary Wilms Tumor Panel	ICD10 LIMITATIONS A56199			
81351	XCP	Hereditary Expanded Cancer Panel	ICD10 LIMITATIONS A56199			
81352	65661F	TP53 GENE ANALYSIS TARGETED SEQUENCE ANALYSIS	ICD10 LIMITATIONS A56199			
81352	NGAMT	AML 4 Gene Panel, Therapeutic	ICD10 LIMITATIONS A56199			
81352	P53CA	TP53 gene somatic mutation analysis	ICD10 LIMITATIONS A56199			
81353	_G017	Gene TP53	INDIVIDUAL REVIEW A56199			
81362	_G129	Gene HBB_SEQ	NON COVERED A56199			
81363	BTHAL	Beta-Globin Gene, Large Del/Dup	NON COVERED A56199			
81363	WBDD	BETA GLOBIN CLUSTER LOCUS DEL/DUP	NON COVERED A56199			
81363	WBDDR	BETA GLOBIN CLUSTER LOCUS DEL/DUP,B	NON COVERED A56199			
81364	64924D	HBB FULL GENE SEQUENCE	NON COVERED A56199			
81364	64938E	HBB FULL GENE SEQUENCE	NON COVERED A56199			
81364	NGCDA	CDA SEQUENCING, V	NON COVERED A56199			
81364	NGMEM	RBC MEMBRANE SEQUENCING, V	NON COVERED A56199			
81364	WBSEQ	BETA GLOBIN GENE SEQUENCING, B	NON COVERED A56199			
81364	WBSQR	BETA GLOBIN GENE SEQUENCING, B	NON COVERED A56199			
81371	68300C	HLA I & II TYPE VERIFY LR	ICD10 LIMITATIONS A56199			
81371	68301C	HLA I & II TYPE VERIFY LR	ICD10 LIMITATIONS A56199			
81371	CFABR	HLA A,B,DR Mol. Phenotype, Confirm	ICD10 LIMITATIONS A56199			
81371	LRBUD	LR ABDR, Related Donor, Swab	ICD10 LIMITATIONS A56199			
81371	LRBUR	Confirm Typing, Recip, Swab	ICD10 LIMITATIONS A56199			
81371	LRDNR	LR ABDR, Related Donor	ICD10 LIMITATIONS A56199			
81371	LRREC	Confirmatory Typing, Recipient	ICD10 LIMITATIONS A56199			
81372	1DIS	HLA-A-B-C DisAssoc Typing LowRes,B	ICD10 LIMITATIONS A56199			
81372	609354A	HLA I TYPING COMPLETE LR	ICD10 LIMITATIONS A56199			
81375	2DIS	HLA-DR-DQ DisAssoc Typing LowRes,B	ICD10 LIMITATIONS A56199			
81375	609356B	HLA II TYPING AG EQUIV LR	ICD10 LIMITATIONS A56199			
81376	2DIS	HLA-DR-DQ DisAssoc Typing LowRes,B	ICD10 LIMITATIONS A56199			
81376	31928A	HLA II TYPING 1 LOCUS LR	ICD10 LIMITATIONS A56199			
81376	609356A	HLA II TYPING 1 LOCUS LR	ICD10 LIMITATIONS A56199			

81376	88906A	HLA II TYPING 1 LOCUS LR	ICD10 LIMITATIONS A56199			
81376	CDCOM	Celiac Disease Comprehensive Cascade	ICD10 LIMITATIONS A56199			
81376	CDGF	Celiac Disease Gluten-Free Cascade	ICD10 LIMITATIONS A56199			
81376	CELI	Celiac Associated HLA-DQ Typing	ICD10 LIMITATIONS A56199			
81376	CELI2	HLA-DQ Typing	ICD10 LIMITATIONS A56199			
81376	NARC	NARCOLEPSY ASSOCIATED AG	ICD10 LIMITATIONS A56199			
81379	68295A	HLA I TYPING COMPLETE HR	ICD10 LIMITATIONS A56199			
81379	68296A	HLA I TYPING COMPLETE HR	ICD10 LIMITATIONS A56199			
81379	68297B	HLA I TYPING COMPLETE HR	ICD10 LIMITATIONS A56199			
81379	HRDNR	HR Class I and II, Related Donor	ICD10 LIMITATIONS A56199			
81379	HRMDB	HR Class I and II, NMDP Donor	ICD10 LIMITATIONS A56199			
81379	HRREC	HR Class I and II, Recipient	ICD10 LIMITATIONS A56199			
81379	HRSI	HIGH RESOLUTION CLASS I PHENOTYPE	ICD10 LIMITATIONS A56199			
81381	610048A	HLA I TYPING 1 ALLELE HR	ICD10 LIMITATIONS A56199			
81381	610060H	HLA I TYPING 1 ALLELE HR	ICD10 LIMITATIONS A56199			
81381	64369A	HLA I TYPING 1 ALLELE HR	ICD10 LIMITATIONS A56199			
81381	CARBR	Carbamazepine PGx Panel, V	ICD10 LIMITATIONS A56199			
81381	HL57R	HLA-B 5701 Genotype, V	ICD10 LIMITATIONS A56199			
81381	HL58R	HLA-B 5801 Genotype, V	ICD10 LIMITATIONS A56199			
81381	PSYQP	Psychotropic PGx Panel, V	ICD10 LIMITATIONS A56199			
81382	68295B	HLA II TYPING 1 LOC HR	ICD10 LIMITATIONS A56199			
81382	68296B	HLA II TYPING 1 LOC HR	ICD10 LIMITATIONS A56199			
81382	68297A	HLA II TYPING 1 LOC HR	ICD10 LIMITATIONS A56199			
81382	DPHRS	High Resolution DP Phenotype, B	ICD10 LIMITATIONS A56199			
81382	HRDNR	HR Class I and II, Related Donor	ICD10 LIMITATIONS A56199			
81382	HRMDB	HR Class I and II, NMDP Donor	ICD10 LIMITATIONS A56199			
81382	HRREC	HR Class I and II, Recipient	ICD10 LIMITATIONS A56199			
81382	HRSL2	HR Single Locus, Class II	ICD10 LIMITATIONS A56199			
81401	35376B	H19 METHYLATION ANALYSIS	INDIVIDUAL REVIEW A56199			
81401	35376C	KCNQ1OT1 METHYLATION ANALYSIS	INDIVIDUAL REVIEW A56199			
81401	65332B	MOPATH PROCEDURE LEVEL 2	INDIVIDUAL REVIEW A56199			
81401	AIHL	Aminoglycoside-Induced Hearing Loss	INDIVIDUAL REVIEW A56199			
81401	APOEG	Apolipoprotein E Geno	INDIVIDUAL REVIEW A56199			
81401	BWRS	BWS/RSS Molecular Analysis	INDIVIDUAL REVIEW A56199			
81401	GAL14	Galactosemia Gene Analysi	INDIVIDUAL REVIEW A56199			
81401	IN16Q	Inv(16); CBFB-MYH11, Quant, V	INDIVIDUAL REVIEW A56199			
81401	KD2T	KRABBE DISEASE 2ND TIER NBS, BS	INDIVIDUAL REVIEW A56199			
81401	T821Q	RUNX1/RUNX1T1, t(8;21), Quant, V	INDIVIDUAL REVIEW A56199			
81402	37445B	MOPATH PROCEDURE LEVEL 3	INDIVIDUAL REVIEW A56199			
81402	CYPZ	CYP21A2 GENE, FULL GENE ANALYSIS	INDIVIDUAL REVIEW A56199			
81402	UNIPD	Uniparental Disomy	INDIVIDUAL REVIEW A56199			
81403	_G001	Gene GRHPR	INDIVIDUAL REVIEW A56199			
81403	_G002	Gene PPOX	INDIVIDUAL REVIEW A56199			
81403	_G009	Gene MLH3	INDIVIDUAL REVIEW A56199			
81403	_G010	Gene CHEK2	INDIVIDUAL REVIEW A56199			
81403	_G011	Gene IDUA	INDIVIDUAL REVIEW A56199			
81403	_G012	Gene AXIN2	INDIVIDUAL REVIEW A56199			

81403	_G013	Gene BMPR1A	INDIVIDUAL REVIEW A56199			
81403	_G015	Gene SMAD4	INDIVIDUAL REVIEW A56199			
81403	_G016	Gene STK11	INDIVIDUAL REVIEW A56199			
81403	_G018	Gene IDS	INDIVIDUAL REVIEW A56199			
81403	_G019	Gene FLCN	INDIVIDUAL REVIEW A56199			
81403	_G020	Gene SPINK1	INDIVIDUAL REVIEW A56199			
81403	_G021	Gene PRSS1	INDIVIDUAL REVIEW A56199			
81403	_G022	Gene CTRC	INDIVIDUAL REVIEW A56199			
81403	_G023	Gene MEFV	INDIVIDUAL REVIEW A56199			
81403	_G024	Gene TNFRSF1A	INDIVIDUAL REVIEW A56199			
81403	_G025	Gene ABCD1	INDIVIDUAL REVIEW A56199			
81403	_G026	Gene CDH1	INDIVIDUAL REVIEW A56199			
81403	_G027	Gene NAGLU	INDIVIDUAL REVIEW A56199			
81403	_G028	Gene SGSH	INDIVIDUAL REVIEW A56199			
81403	_G029	Gene ARSB	INDIVIDUAL REVIEW A56199			
81403	_G030	Gene GNPTAB	INDIVIDUAL REVIEW A56199			
81403	_G032	Gene ACADVL	INDIVIDUAL REVIEW A56199			
81403	_G033	Gene ACADM	INDIVIDUAL REVIEW A56199			
81403	_G034	Gene ACADS	INDIVIDUAL REVIEW A56199			
81403	_G035	Gene FECH	INDIVIDUAL REVIEW A56199			
81403	_G036	Gene MAPT	INDIVIDUAL REVIEW A56199			
81403	_G037	Gene PKHD1	INDIVIDUAL REVIEW A56199			
81403	_G038	Gene GRN	INDIVIDUAL REVIEW A56199			
81403	_G039	Gene FTCD	INDIVIDUAL REVIEW A56199			
81403	_G040	Gene CDNK1C	INDIVIDUAL REVIEW A56199			
81403	_G041	Gene CPOX	INDIVIDUAL REVIEW A56199			
81403	_G042	Gene ATP7B	INDIVIDUAL REVIEW A56199			
81403	_G043	Gene GAA	INDIVIDUAL REVIEW A56199			
81403	_G044	Gene HMBS	INDIVIDUAL REVIEW A56199			
81403	_G045	Gene GALT	INDIVIDUAL REVIEW A56199			
81403	_G046	Gene GLA	INDIVIDUAL REVIEW A56199			
81403	_G047	Gene BTD	INDIVIDUAL REVIEW A56199			
81403	_G048	Gene HEXA	INDIVIDUAL REVIEW A56199			
81403	_G049	Gene AGXT	INDIVIDUAL REVIEW A56199			
81403	_G051	Gene MLYCD	INDIVIDUAL REVIEW A56199			
81403	_G052	Gene MMACHC	INDIVIDUAL REVIEW A56199			
81403	_G053	Gene GBA	INDIVIDUAL REVIEW A56199			
81403	_G054	Gene SMPD1	INDIVIDUAL REVIEW A56199			
81403	_G055	Gene CPT2	INDIVIDUAL REVIEW A56199			
81403	_G056	Gene TTR	INDIVIDUAL REVIEW A56199			
81403	_G057	Gene UBE3A	INDIVIDUAL REVIEW A56199			
81403	_G058	Gene GALC	INDIVIDUAL REVIEW A56199			
81403	_G059	Gene GSN	INDIVIDUAL REVIEW A56199			
81403	_G060	Gene LYZ	INDIVIDUAL REVIEW A56199			
81403	_G061	Gene FGA	INDIVIDUAL REVIEW A56199			
81403	_G062	Gene APOA1	INDIVIDUAL REVIEW A56199			
81403	_G063	Gene APOA2	INDIVIDUAL REVIEW A56199			

81403	_G064	Gene MMADHC	INDIVIDUAL REVIEW A56199			
81403	_G065	Gene SLC25A20	INDIVIDUAL REVIEW A56199			
81403	_G066	Gene ARSA	INDIVIDUAL REVIEW A56199			
81403	_G067	Gene NPC1/2_SEQ and NPC1/2_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G070	Gene RAI1	INDIVIDUAL REVIEW A56199			
81403	_G071	Gene MYH	INDIVIDUAL REVIEW A56199			
81403	_G072	Gene HGSNAT	INDIVIDUAL REVIEW A56199			
81403	_G073	Gene GNS and GRHPR_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G074	Gene PSAP	INDIVIDUAL REVIEW A56199			
81403	_G077	Gene RET	INDIVIDUAL REVIEW A56199			
81403	_G078	Gene SUMF1	INDIVIDUAL REVIEW A56199			
81403	_G079	Gene CASR_Seq	INDIVIDUAL REVIEW A56199			
81403	_G080	Gene VHL_SEQ	INDIVIDUAL REVIEW A56199			
81403	_G081	Gene VHL_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G082	Gene SHDP_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G083	Gene SDHB, SDHC, and SDHD_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G084	Gene SDHC, SDHC, SDHD_Seq	INDIVIDUAL REVIEW A56199			
81403	_G087	GENE DMD_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G089	GENE MPZ_MLPA	INDIVIDUAL REVIEW A56199			
81403	_G098	SKELETAL MUSCLE CHANNELOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81403	_G102	Gene SERPINA1, Bill Only	INDIVIDUAL REVIEW A56199			
81403	_G112	Gene SDHAF2, Bill Only	INDIVIDUAL REVIEW A56199			
81403	_G113	Gene TMEM127	INDIVIDUAL REVIEW A56199			
81403	_G114	Gene MAX	INDIVIDUAL REVIEW A56199			
81403	_G130	Known Familial Variant,Other	INDIVIDUAL REVIEW A56199			
81403	36517C	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	37510A	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614318B	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614319F	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614320C	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614572B	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614574B	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	614576C	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	63162B	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	65589C	MOPATH PROCEDURE LEVEL 4	INDIVIDUAL REVIEW A56199			
81403	BRGYP	Hereditary Breast/Gyn Cancer Panel	INDIVIDUAL REVIEW A56199			
81403	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81403	CTNNB	B-CATENIN MUTATION ANALYSIS, TUMOR	INDIVIDUAL REVIEW A56199			
81403	F81B	HA F8 Intron 1 Inversion KM, B	INDIVIDUAL REVIEW A56199			
81403	F81P	HA F8 Intron 1 Inversion, AF or CVS	INDIVIDUAL REVIEW A56199			
81403	F822B	HA F8 Intron 22 Inversion KM, B	INDIVIDUAL REVIEW A56199			
81403	F822P	HA F8 Int22 Inversion KM, AF or CVS	INDIVIDUAL REVIEW A56199			
81403	F8INP	HA F8 Int 1/22 Inversion, AF or CVS	INDIVIDUAL REVIEW A56199			
81403	F8INV	HA F8 Intron 1/22 Inversion, B	INDIVIDUAL REVIEW A56199			
81403	F9KMP	F9 GENE KNOWN MUTATION, AF OR CVS	INDIVIDUAL REVIEW A56199			
81403	FIXKM	F9 GENE KNOWN MUTATION, B	INDIVIDUAL REVIEW A56199			

81403	LQTGP	LONG QT SYNDROME MULTI-GENE PANEL,B	INDIVIDUAL REVIEW A56199			
81403	LYNCP	Lynch Syndrome Panel	INDIVIDUAL REVIEW A56199			
81403	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81403	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81403	RASFP	RAS/RAF Panel, Tumor	INDIVIDUAL REVIEW A56199			
81403	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81403	YMCRO	Y Microdeletion	INDIVIDUAL REVIEW A56199			
81404	_G092	MYOFIBRILLAR MYOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81404	_G096	EMERY-DREIFUSS PANEL	INDIVIDUAL REVIEW A56199			
81404	_G123	Febrile Seizure Panel	INDIVIDUAL REVIEW A56199			
81404	35640A	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	603783A	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	605191A	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	614318L	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	614319E	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	614574E	CDKN2A gene	INDIVIDUAL REVIEW A56199			
81404	614574M	STK11 dup/del	INDIVIDUAL REVIEW A56199			
81404	614575D	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	63161B	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	63162C	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	63163B	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	65562B	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	65582A	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	65586A	MOPATH PROCEDURE LEVEL 5	INDIVIDUAL REVIEW A56199			
81404	ATTRZ	Familial Amyloidosis, DNA Sequence	INDIVIDUAL REVIEW A56199			
81404	BRGGP	Brugada Syndrome Multi-Gene Panel,B	INDIVIDUAL REVIEW A56199			
81404	BTDZ	BTD Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81404	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81404	HPPAN	Hereditary Pancreatitis Panel	INDIVIDUAL REVIEW A56199			
81404	LQTGP	LONG QT SYNDROME MULTI-GENE PANEL,B	INDIVIDUAL REVIEW A56199			
81404	NSRGP	Noonan Syndrome and Related Panel,B	INDIVIDUAL REVIEW A56199			
81404	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81404	PMNSR	POSTMORTEM NOONAN AND RELATED PANEL	INDIVIDUAL REVIEW A56199			
81404	PRSSZ	PRSS1 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81404	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81404	UGTFG	UGT1A1 FULL GENE SEQUENCING	INDIVIDUAL REVIEW A56199			
81404	VHLE	VHL Gene Erythrocytosis Mutations	INDIVIDUAL REVIEW A56199			
81404	VHLZZ	VHL Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81404	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81405	_G092	MYOFIBRILLAR MYOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81405	_G096	EMERY-DREIFUSS PANEL	INDIVIDUAL REVIEW A56199			
81405	_G123	Febrile Seizure Panel	INDIVIDUAL REVIEW A56199			
81405	_G124	Epilepsy with Migraine Panel	INDIVIDUAL REVIEW A56199			
81405	35640E	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	37445A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	603783B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			

81405	603784A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	605191B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	608015A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	608023A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	608031A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	608032A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	614318M	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	614319G	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	614574L	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	614575A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	614577B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	616524B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	63161C	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	63161F	Title MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	64938B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	65143A	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	65562D	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	65582C	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	65586B	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	APGP	Acute Porphyria Gene Panel	INDIVIDUAL REVIEW A56199			
81405	CASRZ	CASR Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81405	CSRSM	CASR Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	CYPZ	CYP21A2 GENE, FULL GENE ANALYSIS	INDIVIDUAL REVIEW A56199			
81405	DHCRZ	DHCR7 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	FABRZ	Fabry Dis Full Gene Anal	INDIVIDUAL REVIEW A56199			
81405	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81405	HPPAN	Hereditary Pancreatitis Panel	INDIVIDUAL REVIEW A56199			
81405	LRCCZ	FH Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	LUNGR	LUNG REARRANGEMENT TESTING, TUMOR	INDIVIDUAL REVIEW A56199			
81405	MPS2Z	Hunter Syndrome, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	MSUDP	MSUD Gene Panel	INDIVIDUAL REVIEW A56199			
81405	NGMEM	RBC MEMBRANE SEQUENCING, V	INDIVIDUAL REVIEW A56199			
81405	NSRGP	Noonan Syndrome and Related Panel,B	INDIVIDUAL REVIEW A56199			
81405	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81405	PCGP	Porphyria Comprehensive Gene Panel	INDIVIDUAL REVIEW A56199			
81405	PHEGP	Phenylalanine Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81405	PKLRG	PKLR FULL GENE AND DELETION	INDIVIDUAL REVIEW A56199			
81405	PMNSR	POSTMORTEM NOONAN AND RELATED PANEL	INDIVIDUAL REVIEW A56199			
81405	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81405	STK1Z	MOPATH PROCEDURE LEVEL 6	INDIVIDUAL REVIEW A56199			
81405	WILMP	Hereditary Wilms Tumor Panel	INDIVIDUAL REVIEW A56199			
81405	XALDZ	X-ALD, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81405	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81406	_G092	MYOFIBRILLAR MYOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81406	_G096	EMERY-DREIFUSS PANEL	INDIVIDUAL REVIEW A56199			

81406	_G098	SKELETAL MUSCLE CHANNELOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81406	_G124	Epilepsy with Migraine Panel	INDIVIDUAL REVIEW A56199			
81406	_G131	Tuberous Sclerosis Panel	INDIVIDUAL REVIEW A56199			
81406	603784B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	603785A	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	605191C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	605192A	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	608015B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	608023B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	608031B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	608032B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	608033A	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	614318N	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	614319D	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	614575C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	614580C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	616524C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	616537A	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	63161D	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	63162D	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	63162G	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	63163C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	65562E	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	65582B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	65586C	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	65589B	MOPATH PROCEDURE LEVEL 7	INDIVIDUAL REVIEW A56199			
81406	65747B	MOLECULAR PATHOLOGY LEVEL 7 INIT, PANEL	INDIVIDUAL REVIEW A56199			
81406	65747C	MOLECULAR PATHOLOGY LEVEL 7 ADDTL, PANEL A3	INDIVIDUAL REVIEW A56199			
81406	65748B	MOLECULAR PATHOLOGY LEVEL 7 INIT, PANEL	INDIVIDUAL REVIEW A56199			
81406	65748C	MOLECULAR PATHOLOGY LEVEL 7 ADDL, PANEL	INDIVIDUAL REVIEW A56199			
81406	APGP	Acute Porphyria Gene Panel	INDIVIDUAL REVIEW A56199			
81406	BRGGP	Brugada Syndrome Multi-Gene Panel,B	INDIVIDUAL REVIEW A56199			
81406	BTKFP	BTK Full-Gene Panel, B	INDIVIDUAL REVIEW A56199			
81406	BTKS	BTK, Full Gene Sequence	INDIVIDUAL REVIEW A56199			
81406	BTKSP	BTK, Full Gene Sequence	INDIVIDUAL REVIEW A56199			
81406	CDHZ	CDH1 Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81406	FHRGP	HYPERCHOLESTEROLEMIA GENE PANEL	INDIVIDUAL REVIEW A56199			
81406	GAAZ	Pompe Disease, Full Gene Sequencing	INDIVIDUAL REVIEW A56199			
81406	GALZ	Galactosemia, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	GRNZ	Progranulin Gene Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	HEXAZ	HEXA Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	HHTGP	HEREDITARY HEMORRHAGIC TELAN PANEL	INDIVIDUAL REVIEW A56199			
81406	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81406	KRABZ	Krabbe Disease, Full Gene Analysis	INDIVIDUAL REVIEW A56199			

81406	LQTGP	LONG QT SYNDROME MULTI-GENE PANEL,B	INDIVIDUAL REVIEW A56199			
81406	MAPTZ	MAPT Screening Sequence Analysis	INDIVIDUAL REVIEW A56199			
81406	MPS1Z	Hurler Syndrome, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	MSUDP	MSUD Gene Panel	INDIVIDUAL REVIEW A56199			
81406	NSRGP	Noonan Syndrome and Related Panel,B	INDIVIDUAL REVIEW A56199			
81406	PCGP	Porphyria Comprehensive Gene Panel	INDIVIDUAL REVIEW A56199			
81406	PHEGP	Phenylalanine Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81406	PMNSR	POSTMORTEM NOONAN AND RELATED PANEL	INDIVIDUAL REVIEW A56199			
81406	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81406	RETZZ	RET Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	THYRP	Hereditary Thyroid Cancer Panel	INDIVIDUAL REVIEW A56199			
81406	TSCP	Tuberous Sclerosis Gene Panel	INDIVIDUAL REVIEW A56199			
81406	TYRGP	Tyrosine Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81406	UBE3Z	UBE3A Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81406	VLCZ	VLCAD Deficiency,Full Gene	INDIVIDUAL REVIEW A56199			
81406	WDZ	Wilson Disease Mutat Scrn	INDIVIDUAL REVIEW A56199			
81406	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81407	_G123	Febrile Seizure Panel	INDIVIDUAL REVIEW A56199			
81407	_G124	Epilepsy with Migraine Panel	INDIVIDUAL REVIEW A56199			
81407	_G131	Tuberous Sclerosis Panel	INDIVIDUAL REVIEW A56199			
81407	603783C	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	603784C	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	603785B	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	614318P	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	614575F	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	616524D	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	616537B	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	63162E	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	63163D	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	65748D	MOPATH PROCEDURE LEVEL 8	INDIVIDUAL REVIEW A56199			
81407	BRGGP	Brugada Syndrome Multi-Gene Panel,B	INDIVIDUAL REVIEW A56199			
81407	F8NGS	F8 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81407	FHRGP	HYPERCHOLESTEROLEMIA GENE PANEL	INDIVIDUAL REVIEW A56199			
81407	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81407	LQTGP	LONG QT SYNDROME MULTI-GENE PANEL,B	INDIVIDUAL REVIEW A56199			
81407	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81407	TSCP	Tuberous Sclerosis Gene Panel	INDIVIDUAL REVIEW A56199			
81407	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81408	614318Q	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	614319B	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	614574C	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	614576A	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	616524E	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	65749B	MOPATH PROCEDURE LEVEL 9	INDIVIDUAL REVIEW A56199			
81408	ARPKZ	ARPKD Mutation Screen	INDIVIDUAL REVIEW A56199			
81408	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			

81408	EDSGP	Ehlers-Danlos Syndrome Gene Panel	INDIVIDUAL REVIEW A56199			
81408	FBN1B	FBN1 Full Gene Sequence	INDIVIDUAL REVIEW A56199			
81408	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81408	NF1Z	NF1 Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81408	PANCP	Hereditary Pancreatic Cancer Panel	INDIVIDUAL REVIEW A56199			
81408	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81408	VWFNG	VWF Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81408	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81410	MFRGP	Marfan and Related Genetic Panel	NON COVERED A56199			
81410	PMMFR	POSTMORTEM MARFAN AND RELATED PANEL	NON COVERED A56199			
81415	WESDX	Whole Exome Sequencing	NON COVERED A56199			
81416	G226	Number of Comparators for WESDX	NON COVERED A56199			
81419	_G116	Epilepsy Expanded Panel	INDIVIDUAL REVIEW A56199			
81419	603776K	EPILEPSY GENOMIC SEQUENCE ANALYSIS PANEL	INDIVIDUAL REVIEW A56199			
81419	EPPAN	Comprehensive Epilepsy Gene Panel	INDIVIDUAL REVIEW A56199			
81430	606144A	HEARING LOSS SEQUENCE ANALYS	NON COVERED A56199			
81430	HHLP	Hereditary Hearing Loss Panel	NON COVERED A56199			
81431	606144B	HEARING LOSS DUP/DEL ANALYS	NON COVERED A56199			
81431	HHLP	Hereditary Hearing Loss Panel	NON COVERED A56199			
81432	614320A	HRDTRY BRST CA-RLATD DSORDRS	NON COVERED A56199			
81432	BRGYP	Hereditary Breast/Gyn Cancer Panel	NON COVERED A56199			
81435	CRCGP	Hereditary GI Cancer Panel	NON COVERED A56199			
81437	ENDCP	Hereditary Endocrine Cancer Panel	NON COVERED A56199			
81437	HPGLP	Hereditary PGL/PCC Panel	NON COVERED A56199			
81439	ARVGP	ARRHYTHMOGENIC CARDIOMYOPATHY, B	NON COVERED A56199			
81439	CCMGP	CARDIOMYOPATHY GENETIC PANEL, B	NON COVERED A56199			
81439	DCMGP	DILATED CARDIOMYOPATHY PANEL, B	NON COVERED A56199			
81439	HCMGP	HYPERTROPHIC CARDIOMYOPATHY PANEL,B	NON COVERED A56199			
81439	PMCMP	POSTMORTEM CARDIOMYOPATHY PANEL	NON COVERED A56199			
81440	65212B	MITOCHONDRIAL GENE	NON COVERED A56199			
81440	MITON	MITOCHONDRIAL NUCLEAR GENE PANEL	NON COVERED A56199			
81440	MITOT	COMBINED MITOCHONDRIAL ANALYSIS	NON COVERED A56199			
81443	_G090	MOTOR NEURON DISEASE PANEL	NON COVERED A56199			
81443	_G091	MUSCULAR DYSTROPHY PANEL	NON COVERED A56199			
81443	_G093	CONGENITAL MYOPATHY PANEL	NON COVERED A56199			
81443	_G094	CONGENITAL MYASTHENIC SYNDROMES	NON COVERED A56199			
81443	_G095	METABOLIC MYOPATHY PANEL	NON COVERED A56199			
81443	_G097	DISTAL MYOPATHY PANEL	NON COVERED A56199			
81443	_G099	MYOPATHY EXPANDED PANEL	NON COVERED A56199			
81443	_G100	DISTAL WEAKNESS EXPANDED PANEL	NON COVERED A56199			
81443	_G101	RHABDOMYOLYSIS AND MYOPATHY PANEL	NON COVERED A56199			
81443	_G109	Comprehensive CDG Panel	NON COVERED A56199			
81443	_G110	Normal Transferrin CDG Panel	NON COVERED A56199			
81443	_G111	Abnormal Transferrin CDG Panel	NON COVERED A56199			
81443	_G117	Encephalopathy with Seizures Panel	NON COVERED A56199			

81443	_G118	Early Epileptic Encephalopathy Panel	NON COVERED A56199			
81443	_G119	Neuronal Migration Disorders Panel	NON COVERED A56199			
81443	_G120	Progressive Myoclonic Epilepsy Panel	NON COVERED A56199			
81443	_G121	Infantile Spasms Panel	NON COVERED A56199			
81443	_G122	Focal Epilepsy Panel	NON COVERED A56199			
81443	3MGAP	3-Methylglutaconic Aciduria Panel	NON COVERED A56199			
81443	603777H	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	603778H	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	603779G	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	603780E	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	603781G	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	603782F	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	608019A	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	62190F	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	63685E	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	63686D	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	64937G	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	64939H	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65559F	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65578D	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65579F	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65580G	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65581L	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65583H	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65584F	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65585G	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65587J	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65588H	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65664D	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65666C	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65669D	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	65824E	GENETIC TSTG SEVERE INH COND	NON COVERED A56199			
81443	AUTOP	AUTOINFLAMMATORY PID GENE PANE	NON COVERED A56199			
81443	BCLGP	B-CELL DEFICIENCY PID GENE PANEL	NON COVERED A56199			
81443	CDGGP	CDG Gene Panel	NON COVERED A56199			
81443	CHLGP	Cholestasis Gene Panel	NON COVERED A56199			
81443	CLADP	Congenital Lactic Acidosis Panel	NON COVERED A56199			
81443	G150	Hereditary Custom Gene Panel Tier 6	NON COVERED A56199			
81443	G158	Lysosomal Storage Disease Panel	NON COVERED A56199			
81443	G159	NCL (Batten Disease) Panel	NON COVERED A56199			
81443	G160	Peroxisomal Disorder Panel	NON COVERED A56199			
81443	G161	Glycogen Storage Disease Panel	NON COVERED A56199			
81443	GSDGP	Glycogen Storage Disease Gene Panel	NON COVERED A56199			
81443	HFAOP	Fatty Acid Oxidation Gene Panel	NON COVERED A56199			
81443	IBDGP	IBD PID Gene Panel	NON COVERED A56199			
81443	LSDGP	LSD Gene Panel	NON COVERED A56199			
81443	MMAGP	MMAGP	NON COVERED A56199			

81443	MPAGP	MMA PA Combined Gene Panel	NON COVERED A56199			
81443	NCLGP	NCL (Batten Disease) Gene Panel	NON COVERED A56199			
81443	NGENZ	RBC ENZYME SEQUENCING, V	NON COVERED A56199			
81443	NGHHA	HEREDITARY HEMOLYTIC ANEMIA SEQ, V	NON COVERED A56199			
81443	PDGP	Peroxisomal Disorder Gene Panel	NON COVERED A56199			
81443	PMARP	POSTMORTEM ARRHYTHMIA PANEL	NON COVERED A56199			
81443	SCDGP	SCID PID GENE PANEL	NON COVERED A56199			
81443	SCNGP	CONGENITAL NEUTROPENIA PID PANEL	NON COVERED A56199			
81443	UCDP	Urea Cycle Disorders Gene Panel	NON COVERED A56199			
81445	CAPN	Solid Tumor Targeted Cancer Panel	ICD10 LIMITATIONS A56867			
81445	G179	Oncology Custom Gene Panel Tier 2	ICD10 LIMITATIONS A56867			
81445	LNGPR	LUNG PANEL WITH REARRANGEMENT TUMOR	ICD10 LIMITATIONS A56867			
81445	LUNGP	LUNG CANCER PANEL, TUMOR	ICD10 LIMITATIONS A56867			
81445	MELP	Melanoma Panel, Tumor	ICD10 LIMITATIONS A56867			
81448	_G103	Custom Gene Panel(CPT 81448) Tier 2	NON COVERED A56199			
81448	_G104	Motor and Sensory Neuropathy Panel	NON COVERED A56199			
81448	_G105	Hereditary Sensory Neuropathy Panel	NON COVERED A56199			
81448	_G106	Hereditary Motor Neuropathy Panel	NON COVERED A56199			
81448	_G107	Spastic Paraplegia Neuropathy Panel	NON COVERED A56199			
81448	_G108	Metabolic or Syndromic Neuropathies	NON COVERED A56199			
81448	G151	Custom Gene Panel(CPT 81448) Tier 2	NON COVERED A56199			
81448	G152	Custom Gene Panel(CPT 81448) Tier 3	NON COVERED A56199			
81448	G153	Custom Gene Panel(CPT 81448) Tier 4	NON COVERED A56199			
81448	G154	Custom Gene Panel(CPT 81448) Tier 5	NON COVERED A56199			
81450	NGAML	Next Gen Sequencing, AML, 11 Gene				
81450	NGSFX	Reflex Analysis, NGSHM				
81450	NGSHM	NGS for Myeloid Neoplasms (NGSHM)				
81455	G181	Oncology Custom Gene Panel Tier 3	ICD10 LIMITATIONS A56867			
81455	MCSTP	MayoComplete Solid Tumor Panel	ICD10 LIMITATIONS A56867			
81455	NGSMM	NGS MULTIPLE MYELOMA	ICD10 LIMITATIONS A56867			
81455	NONCP	NEURO-ONC EXPANDED PANEL	ICD10 LIMITATIONS A56867			
81455	SARCP	Sarcoma Targeted Gene Fusion Panel	ICD10 LIMITATIONS A56867			
81460	608019B	WHOLE MITOCHONDRIAL GENOME	NON COVERED A56199			
81460	62510A	WHOLE MITOCHONDRIAL	NON COVERED A56199			
81460	65212A	WHOLE MITOCHONDRIAL GENOME	NON COVERED A56199			
81460	CLADP	Congenital Lactic Acidosis Panel	NON COVERED A56199			
81460	MITOP	Mitochondrial Full Genome Analysis	NON COVERED A56199			
81460	MITOT	COMBINED MITOCHONDRIAL ANALYSIS	NON COVERED A56199			
81465	608019C	WHOLE MITOCHONDRIAL GENOME	NON COVERED A56199			
81465	62510B	WHOLE MITOCHONDRIAL GENOME	NON COVERED A56199			
81465	65212C	WHOLE MITOCHONDRIAL GENOME	NON COVERED A56199			
81465	CLADP	Congenital Lactic Acidosis Panel	NON COVERED A56199			
81465	MITOP	Mitochondrial Full Genome Analysis	NON COVERED A56199			
81465	MITOT	COMBINED MITOCHONDRIAL ANALYSIS	NON COVERED A56199			
81479	_G031	Gene SEPT9	INDIVIDUAL REVIEW A56199			

81479	_G092	MYOFIBRILLAR MYOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81479	_G096	EMERY-DREIFUSS PANEL	INDIVIDUAL REVIEW A56199			
81479	_G098	SKELETAL MUSCLE CHANNELOPATHY PANEL	INDIVIDUAL REVIEW A56199			
81479	_G123	Febrile Seizure Panel	INDIVIDUAL REVIEW A56199			
81479	_G124	Epilepsy with Migraine Panel	INDIVIDUAL REVIEW A56199			
81479	_G132	SEPT9 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	2B6Q	CYP2B6 Genotype, V	INDIVIDUAL REVIEW A56199			
81479	2OHGP	2-OH Glutaric Aciduria Gene Panel	INDIVIDUAL REVIEW A56199			
81479	603783D	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	603784D	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	605191D	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	605192B	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	608015C	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	608023C	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	608031C	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	608032C	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	608033B	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	610060A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614318R	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614319P	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614575G	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614576K	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614577C	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	614580E	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	616524F	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	63161A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	63162A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	63163A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	64924B	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	64938A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65143B	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65562A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65582D	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65586D	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65589A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	65747A	UNLISTED MOLECULAR PATHOLOGY, A6, PANEL	INDIVIDUAL REVIEW A56199			
81479	65748A	UNLISTED MOLECULAR PATHOLOGY, A5, PANEL	INDIVIDUAL REVIEW A56199			
81479	65749A	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	83301Z	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	83303Z	UNLISTED MOLECULAR PATHOLOGY	INDIVIDUAL REVIEW A56199			
81479	AGXTZ	AGXT Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	AHUSP	COMPLEMENT AHUS/TMA GENE PANEL	INDIVIDUAL REVIEW A56199			
81479	APGP	Acute Porphyria Gene Panel	INDIVIDUAL REVIEW A56199			
81479	APO1Z	APOA1 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	APO2Z	APOA2 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	APOL1	APOL1 Genotype, V	INDIVIDUAL REVIEW A56199			

81479	ATNGS	SERPINC1 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	BAP1Z	BAP1 Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	BHDZ	FLCN Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	BPGMM	BPGM FULL GENE SEQUENCING	INDIVIDUAL REVIEW A56199			
81479	BRGGP	Brugada Syndrome Multi-Gene Panel,B	INDIVIDUAL REVIEW A56199			
81479	C9ORF	C9orf72, Molecular Analysis	INDIVIDUAL REVIEW A56199			
81479	CDKZ	CDKN1C Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	COMCP	Hereditary Common Cancer Panel	INDIVIDUAL REVIEW A56199			
81479	CSF3R	CSF3R EXON 14/17 MUTATION DETECTION	INDIVIDUAL REVIEW A56199			
81479	CXCFX	CXCR4, GENE MUTATION, REFLEX	INDIVIDUAL REVIEW A56199			
81479	CXLPL	CXCR4 MUTATION IN B-CELL LYMPHOMA	INDIVIDUAL REVIEW A56199			
81479	CYSGP	Cystinuria Gene Panel	INDIVIDUAL REVIEW A56199			
81479	EDSGP	Ehlers-Danlos Syndrome Gene Panel	INDIVIDUAL REVIEW A56199			
81479	EPOR	EPOR GENE, MUTATION ANALYSIS, B	INDIVIDUAL REVIEW A56199			
81479	ESR1	ESR1 MUTATION ANALYSIS, TUMOR	INDIVIDUAL REVIEW A56199			
81479	F10NG	F10 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F11NG	F11 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F12NG	F12 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F13NG	F13A1 and B Genes, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F2NGS	F2 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F5NGS	F5 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	F7NGS	F7 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	FGAZ	FGA Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	FHRGP	HYPERCHOLESTEROLEMIA GENE PANEL	INDIVIDUAL REVIEW A56199			
81479	FIBNG	FGA/B/G Genes, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	GA2P	Glutaric Aciduria Type 2 Gene Panel	INDIVIDUAL REVIEW A56199			
81479	GATA2	GATA2 COMPREHENSIVE GENE SEQUENCING	INDIVIDUAL REVIEW A56199			
81479	GBAZ	Gaucher Disease, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	GRHPZ	GRHPR Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	GSNZ	GSN Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	HEMP	HEREDITARY ERYTHROCYTOSIS MUT, B	INDIVIDUAL REVIEW A56199			
81479	HEXBZ	HEXB Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	HHTGP	HEREDITARY HEMORRHAGIC TELAN PANEL	INDIVIDUAL REVIEW A56199			
81479	HIF2A	HIF2A GENE, MUTATION ANALYSIS, B	INDIVIDUAL REVIEW A56199			
81479	HMEP	Hemiplegic Migraine Gene Panel	INDIVIDUAL REVIEW A56199			
81479	KCNN4	KCNN4 Full Gene Sequencing, V	INDIVIDUAL REVIEW A56199			
81479	KETGP	Ketone Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81479	LQTGP	LONG QT SYNDROME MULTI-GENE PANEL,B	INDIVIDUAL REVIEW A56199			
81479	LUNGR	LUNG REARRANGEMENT TESTING, TUMOR	INDIVIDUAL REVIEW A56199			
81479	LYZZ	LYZ Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	MCADZ	MCAD Mutation Screen	INDIVIDUAL REVIEW A56199			
81479	MINT	MOLECULAR INTERPRETATION	INDIVIDUAL REVIEW A56199			
81479	MSUDP	MSUD Gene Panel	INDIVIDUAL REVIEW A56199			
81479	NAT2	NAT2, Full Gene Sequence	INDIVIDUAL REVIEW A56199			
81479	NAT2F	NAT2, FULL GENE SEQUENCE	INDIVIDUAL REVIEW A56199			
81479	NAT2Q	NAT2 Genotype, V	INDIVIDUAL REVIEW A56199			

81479	NGCDA	CDA SEQUENCING, V	INDIVIDUAL REVIEW A56199			
81479	NGMEM	RBC MEMBRANE SEQUENCING, V	INDIVIDUAL REVIEW A56199			
81479	NSRGP	Noonan Syndrome and Related Panel,B	INDIVIDUAL REVIEW A56199			
81479	NTC3Z	NOTCH3 Gene, Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	PCGP	Porphyria Comprehensive Gene Panel	INDIVIDUAL REVIEW A56199			
81479	PCNGS	PROC Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	PHAGP	PHAGOCYTIC PID GENE PANEL	INDIVIDUAL REVIEW A56199			
81479	PHD2	PHD2 GENE, MUTATION ANALYSIS, B	INDIVIDUAL REVIEW A56199			
81479	PHEGP	Phenylalanine Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81479	PMNSR	POSTMORTEM NOONAN AND RELATED PANEL	INDIVIDUAL REVIEW A56199			
81479	PRCNG	PROCR Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	PRKSD	PRKAR1A Full Gene Analysis	INDIVIDUAL REVIEW A56199			
81479	PRS8P	Hereditary Prostate Cancer Panel	INDIVIDUAL REVIEW A56199			
81479	PRSNG	PROS1 Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	PSYQP	Psychotropic PGx Panel, V	INDIVIDUAL REVIEW A56199			
81479	RENCP	Hereditary Renal Cancer Panel	INDIVIDUAL REVIEW A56199			
81479	SERPZ	SERPINA1 GENE, FULL GENE ANALYSIS	INDIVIDUAL REVIEW A56199			
81479	TBNGS	Susceptibility, Mtb Complex, NGS	INDIVIDUAL REVIEW A56199			
81479	TELGP	TELOMERE DEFECTS GENE PANEL	INDIVIDUAL REVIEW A56199			
81479	THBNG	THBD Gene, Full Gene NGS	INDIVIDUAL REVIEW A56199			
81479	THYRP	Hereditary Thyroid Cancer Panel	INDIVIDUAL REVIEW A56199			
81479	TREC	TREC, IMMUNE RECONSTITUT	INDIVIDUAL REVIEW A56199			
81479	TRECS	TREC Analysis, B	INDIVIDUAL REVIEW A56199			
81479	TYRGP	Tyrosine Disorders Gene Panel	INDIVIDUAL REVIEW A56199			
81479	WGSEQ	GAMMA GLOBIN FULL GENE SEQUENCING	INDIVIDUAL REVIEW A56199			
81479	WGSQR	GAMMA GLOBIN FULL GENE SEQUENCING	INDIVIDUAL REVIEW A56199			
81479	WILMP	Hereditary Wilms Tumor Panel	INDIVIDUAL REVIEW A56199			
81479	XCP	Hereditary Expanded Cancer Panel	INDIVIDUAL REVIEW A56199			
81479	XL2	FOXL2 MUTATION ANALYSIS, TUMOR	INDIVIDUAL REVIEW A56199			
81596	FIBRO	FIBROTEST-ACTITEST, S				
0027U	JAKXB	JAK2 EXON 12 MUTATION DETECTION, B	ICD10 LIMITATIONS A56199			
0027U	JAKXM	JAK2 EXON 12 MUTATION DETECTION, BM	ICD10 LIMITATIONS A56199			
0027U	JAKXR	JAK2 EXON 12-15 SEQUENCING, REFLEX	ICD10 LIMITATIONS A56199			
0029U	PGXQP	Focused Pharmacogenomics Panel, V				
0030U	WARSQ	Warfarin Response Genotype, V				
0031U	1A2Q	CYP1A2 Genotype, V				
0032U	COMTQ	COMT Genotype, V				
0034U	65160D	TPMT (THIOPURINE S-METHYLTRANSFERASE), NUDT15 (NUDIX HYDROXY				
0034U	TPNUQ	TPMT and NUDT15 Genotype, V				
0070U	2D6Q	CYP2D6 Genotype Cascade, V	ICD10 LIMITATIONS A56199			
0120U	PM3CX	Lymph3Cx, Lg Bcell Lymphoma,mRNA,Ts				
0154U	TFGFR	FGFR Mutation/Fusion Analysis Tumor				
0219U	HIVDR	HIV-1 Genotypic Drug Resistance, P				
0239U	75469P	Targeted genomic sequence analysis panel				