SaintFrancis
Center for Genetics

6161 South Yale Avenue, Tulsa, OK, 74136 | 918-502-1720

Preferred Client Price List **v. 1/1/2019**

Procedure Name	CPT codes	McKesson Z-code	
Achondroplasia {FGFR 3}	81401		
Albright Hereditary Osteodystrophy {GNAS1}	81479		
Amyotrophic Lateral Sclerosis {SOD1}	81404		
Androgen Insensitivity Syndrome {AR}	81173		
Androgen Insensitivity Syndrome {AR}; Familial variant	81174		
Angelman Syndrome {UBE3A - Methylation}/ PWS	81331		
Apert Syndrome - FGFR2 exon 8	81404	ZB7K1	
Blau Syndrome - NOD2/CARD15 Complete Gene Analysis	81479		
BRAF codon 600	81210		
Hereditary Breast and Gynecological Cancer Panel (25 genes)	81432, 81433		
BRCA1/2 Gene Sequence w/ Del/Dup Analysis	81162		
BRCA1 gene, familial known variant	81215		
BRCA2 gene, familial known variant	81217		
Hereditary Colon Cancer Panel (18 genes)	81435, 81436		
Hereditary Cancer Comprehensive Panel (33 genes)	81432, 81433, 81435, 81436		
Congenital Adrenal Hyperplasia {CAH}	81405		
Connexin 26 {CX26}	81252	ZB7LH	
Connexin 30 {CX30}	81254	ZB7JV	
Craniodysmorphology Screen {FGFR 1,2 & 3}	81400, 81401, 81404		
Crohn's Disease {NOD2 - Markers}	81401		
Crouzon Syndrome with Acanthosis Nigricans	81403		
Cystic Fibrosis - DNA Probe	81220		
Cystic Fibrosis, known familial variant	81221		
Epidermal Growth Factor Receptor {EGFR - Mutation Analysis}	81235		
FGFR 2	81479		
Factor V Leiden Mutation	81241		
Fragile X Syndrome {FRAX1}	81243		
Fragile X syndrome by Southern Blot (an add-on charge)	81243		
Friederich's Ataxia {FRDA}	81284		
Frontotemporal Dementia - Full Gene (MAPT)	81406		
Frontotemporal Dementia - Familial Variant (MAPT)	81403		
Hereditary Dentatorubral - Pallidolysian Atrophy {DRPLA}	81177		
Huntington's Disease {HD}	81271	ZB7JW	
Hypochondroplasia {FGFR 3}	81404		
JAK2 Mutation for Myeloproliferative Disorders	81270		
KRAS codons 12 and 13	81275		
Li-Fraumeni Syndrome {P53}	81405		
Li Fraumeni - known variant	81403		
Marfan Syndrome - known mutation	81403		
Marfan Syndrome Type 1 {TGFBR1}	81405		
Marfan Syndrome Type 2 {TGFBR2}	81405		
Marfan Syndrome {Fib1 - unknown mutation - DNA}	81408		
Maternal Cell Contamination	81265		
McCune - Albright Syndrome {targeted GNAS1}	81479		

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Methylenetetrahydrofolate Reductase Gene	81291		
Mitochondrial Mutation A1555G {mito 12S DNA}	81401		
Mitochondrial Serine tRNA Mutations	81403		
Myotonic Dystrophy by PCR reflexed to Southern	81234	ZB7JY	
Myotonic dystrophy - Southern blot (an add-on charge)	81239		
NOD2/CARD15 Complete Gene Analysis	81479	ZB75YN	
Nonsyndromic Craniosynostosis	81400		
PGRN	81406		
Pendred Syndrome {SLC26A4}	81406		
Prader-Willi Syndrome {SNRPN - Methylation}/ Angelman	81331		
Prothrombin 20210 Mutation	81240		
Saethre - Chotzen Syndrome {TWIST}	81404		
Spinal Muscular Atrophy, Types 1,2 & 3 {SMN1}	81400	ZB7JX	
Spinocerebellar Ataxia, Type 1 {SCA1}	81178	ZB5YK	
Spinocerebellar Ataxia, Type 2 {SCA2}	81179		
Spinocerebellar Ataxia, Type 3 {SCA3}	81180		
Spinocerebellar Ataxia, Type 6 {SCA6}	81184		
Spinocerebellar Ataxia, Type 7 {SCA7}	81181		
Waardenburg 1 {PAX3}	81479		
Waardenburg 2 {MITF}	81479		
Sequencing - Known Familial Variant by gene			
AR	81174		
CYP21A2 (qty = 2)	81403		
FGFR2	81403		
GJB2	81253		
GNAS1	81479		
MAPT	81403		
MITF	81479		
NOD2	81479		
PAX3	81479		
PGRN	81403		
SLC26A4 (qty = 2)	81403		
SOD1	81403		
TGFBR1	81403		
TGFBR2	81403		
TWIST	81403		
Chromosomal MicroArray, Postnatal			
TAT = 2 weeks	81229		

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Sample Min = 2ml EDTA and 2ml NaHep
Newborn Min = 1ml EDTA and 1ml NaHep

Discounted Panels

	Total		
Connexin Hearing Loss Panel (GJB2, GJ6)	81252, 81254	ZB7LI	
Craniodysmorphology Panel (FGFR1,2,3,TWIST)	81400, 81401		
	81404 x 2		
Dwarfism Panel (FGFR3 - ACH / HCH)	81401, 81404		
Marfan Syndrome - TGFBR Panel	81405 x 2		
	81405		
Neurological Panel (HD,DRPLA,FRDA,SCA1,2,3,6,7)	81271, 81284		
	81177, 81178		
	81179, 81180		
	81181, 81184		
Newborn Hypotonia Panel - (DM,SMA,PWS)	81400, 81401		
	81331	ZB7JZ	
Nonsyndromic Hearing Loss Panel(GJB2,GJB6,mt1555)	81252, 81254		
	81401		
Spinocerebellar ataxia Panel (SCA1,2,3,6,7)	81178, 81179		
	81180, 81181		
	81184	ZB5YL	

(Prices may be subject to change)