SI	GENETIC TEST	TECHNIQUE	TEST	TAT
NO	GENETIC 1E31	TECHNIQUE	CHARGE	IAI
110	CYTOGENETIC TESTS			
1	KARYOTYPE FROM PERIPHERAL BLOOD		RS 2000	3 WEEKS
	SAMPLE			
2	FISH		RS 5000	1 WEEK
	MOLECULAR TESTS			
3	Y MICRODELETION ANALYSIS	PCR	RS 4600	2 WEEKS
4	SRY MUTATION ANALYSIS	PCR	RS 1400	1 WEEK
5	BETA THALASSEMIA - 7 MUTATION PANEL	SANGER SEQUENCING	RS 3000	2 WEEKS
	[HBS, IVS 1-5(G>C), IVS 1-1 (G>T),CODON			
	15 G>A,CODON 30 G>A,FS 8/9 +G,FS			
	41/42 -CTTT]			
6	BETA THALASSEMIA - COMPLETE GENE	SANGER SEQUENCING	RS 4000	3 WEEKS
	SEQUENCING	ncp.	DC 2000	1 /4/55/
7	BETA THALASSEMIA - 619bp DELETION	PCR SANGER SEQUENCING	RS 2000 RS 2000	1 WEEK 2 WEEKS
8	BETA THALASSEMIA - ANY SINGLE MUTATION	SANGER SEQUENCING	N3 2000	2 WLLKS
9	SICKLE CELL ANEMIA	SANGER SEQUENCING	RS 2000	2 WEEKS
10	HEMOGLOBIN E DISEASE	SANGER SEQUENCING	RS 2000	1 WEEK
11	HEMOGLOBIN D DISEASE	SANGER SEQUENCING	RS 2000	1 WEEK
12	G6PD DEFICIENCY	SANGER SEQUENCING	RS 3500	2 WEEKS
13	DUCHENNE MUSCULAR DYSTROPHY	MLPA [KIT BASED]	RS 4000	2 WEEKS
	[DMD] 79			
1.4	EXONS	MI DA [KIT DACED]	DC 4000	2 ///[[/(
14	DUCHENNE MUSCULAR DYSTROPHY [DMD] CARRIER ANALYSIS	MLPA [KIT BASED]	RS 4000	2 WEEKS
15	SPINAL MUSCULAR ATROPHY [SMA]	MLPA [KIT BASED]	RS 3300	2 WEEKS
13	MUTATION ANALYSIS	WELA [KIT DASED]	113 3300	Z WLLKS
16	SPINAL MUSCULAR ATROPHY [SMA]	MLPA [KIT BASED]	RS 3300	2 WEEKS
	CARRIER ANALYSIS			
17	MICRODELETION/ MICRODUPLICATION	MLPA [KIT BASED]	RS 3700	2 WEEKS
	ANALYSIS			
18	SPINOCEREBELLAR ATAXIA [SCA]	FRAGMENT ANALYSIS	RS 5500	2 WEEKS
	TYPES 1,2 & 3	EDA CAMENIT ANIALIYOIS	DC 7500	2 14/55/6
19	SPINOCEREBELLAR ATAXIA [SCA] TYPES 1,2, 3, 6	FRAGMENT ANALYSIS	RS 7500	2 WEEKS
20	SPINOCEREBELLAR ATAXIA [SCA]	FRAGMENT ANALYSIS	RS 9500	2 WEEKS
20	TYPES 1,2, 3, 6 & 12	TRAGINENT ANALISIS	K3 9300	2 WLLK3
21	SPINOCEREBELLAR ATAXIA [SCA] - ANY	FRAGMENT ANALYSIS	RS 2500	2 WEEKS
	ONE TYPE			
22	SPINOBULBAR MUSCULAR ATROPHY	FRAGMENT ANALYSIS	RS 2500	2 WEEKS
	[SBMA]			
23	HUNTINGTON DISEASE [HD]	FRAGMENT ANALYSIS	RS 2500	2 WEEKS
24	DENTATORUBROPALLIDOLUYSIAN	FRAGMENT ANALYSIS	RS 2500	2 WEEKS
25	ATROPHY [DRPLA] MUTATION ANALYSIS	TP-PCR [KIT BASED]	BS 6000	2 MEERS
25	FRAGILE X SYNDROME [FRX] MUTATION ANALYSIS	IT FOR [KIT DASED]	RS 6000	2 WEEKS
26	FRAGILE X CARRIER ANALYSIS	TP-PCR [KIT BASED]	RS 6000	2 WEEKS
27	MYOTONIC DYSTROPHY [MD]	TP-PCR	RS 2500	2 WEEKS
28	FRIEDREICH ATAXIA [FA]	TP-PCR	RS 2500	2 WEEKS
29	CYSTIC FIBROSIS - 36 MUTATION PANEL	ALLELE SPECIFIC PCR [KIT BASED]	RS 7500	1 WEEK
30	CYSTIC FIBROSIS [CF] - 4 MUTATION	SANGER SEQUENCING	RS 2800	2 WEEKS
50	PANEL	S. INGER SEQUENCING	113 2000	Z VVLLNJ
	[dF508, G542X, G551D, R553X]			
31	CYSTIC FIBROSIS [CF] - SINGLE	SANGER SEQUENCING	RS 2000	2 WEEKS
	MUTATION			
32	CYSTIC FIBROSIS [CF]- 5T/7T/9T	SANGER SEQUENCING	RS 2000	1 WEEK
	POLYMORPHISM			

SI NO	GENETIC TEST	TECHNIQUE	TEST CHARGE	TAT
33	CHRONIC PANCREATITIS [CP] SPINK1 N34S MUTATION ANALYSIS	SANGER SEQUENCING	RS 2000	2 WEEKS
34	CHRONIC PANCREATITIS [CP] SPINK1 [N34S] & PRSS1 [N29I, R122H]	SANGER SEQUENCING	RS 3000	2 WEEKS
35	LEBER HEREDITARY OPTIC NEUROPATHY [LHON] 3 MUTATION [G3460A, G11778A, T14484C]	SANGER SEQUENCING	RS 3500	2 WEEKS
36	LEBER HEREDITARY OPTIC NEUROPATHY [LHON] SINGLE MUTATION ANALYSIS	SANGER SEQUENCING	RS 2000	1 WEEK
37	LEIGH SYNDROME - 3 MUTATION PANEL (T12706C, A13084T, G13513A)	SANGER SEQUENCING	RS 3500	2 WEEKS
38	LEIGH SYNDROME - SINGLE MUTATION ANALYSIS	SANGER SEQUENCING	RS 2000	1 WEEK
39	MELAS - A3243G MUTATION ANALYSIS	SANGER SEQUENCING		
40	HEREDITARY HEMOCHROMATOSIS H63D & C282Y MUTATION ANALYSIS	SANGER SEQUENCING	RS 2500	2 WEEKS
41	MTHFR - 677C>T and 1298A>C POLYMORPHISMS	SANGER SEQUENCING	RS 2500	2 WEEKS
42	PROTHROMBIN [FACTOR II] G20210A MUTATION ANALYSIS	SANGER SEQUENCING	RS 2000	1 WEEK
43	FACTOR V LEIDEN MUTATION ANALYSIS	SANGER SEQUENCING	RS 2000	1 WEEK
44	NON SYNDROMIC HEARING LOSS GJB2 MUTATION ANALYSIS [AW24X & AW77X]	SANGER SEQUENCING	RS 2500	2 WEEKS
45	JAK2 V617F MUTATION ANALYSIS	SANGER SEQUENCING	RS 2500	2 WEEKS
46	ACHONDROPLASIA - FGFR3 MUTATION ANALYSIS [c.1138G>A, c.1138G>C]	SANGER SEQUENCING	RS 2500	2 WEEKS
47	UGT1A1 PROMOTER MUTATION ANALYSIS	SANGER SEQUENCING	RS 2500	2 WEEKS
48	NOTCH3 - c.421C>T MUTATION ANALYSIS	SANGER SEQUENCING	RS 2500	2 WEEKS
49	LGMD 2A - CAPN3 SINGLE MUTATION ANALYSIS [C.2051-1G>T & C.2051-1C>T]	SANGER SEQUENCING	RS 2500	2 WEEKS
50	PRENATAL DIAGNOSIS ON CVS/AF SAMPLE WITH MCC	BASED ON PROBAND'S DISEASE	RS 8500	1 WEEK
51	PRENATAL DIAGNOSIS ON CVS/AF SAMPLE ONLY	BASED ON PROBAND'S DISEASE	RS 5500	1 WEEK
52	MATERNAL CELL CONTAMINATION [MCC] ONLY	FRAGMENT ANALYSIS	RS 2500	1 WEEK
53	EXOME SEQUENCING ANALYSIS	NEXT GENERATION SEQUENCING	RS 16500	5 WEEKS
54	EXOME SEQUENCING RAW DATA ONLY	NEXT GENERATION SEQUENCING	RS 15000	3 WEEKS
55	GENOME SEQUENCING ANALYSIS	NEXT GENERATION SEQUENCING	RS 70000	6 WEEKS
56	GENOME SEQUENCING RAW DATA ONLY	NEXT GENERATION SEQUENCING SANGER SEQUENCING	RS 60000 RS 5500	4 WEEKS 3 WEEKS
57	TARGETED MUTATION ANALYSIS [TMA] BY PRIMER DESIGN [1 VARIANT IN 1 PERSON]			
58	TARGETED MUTATION ANALYSIS [TMA] BY PRIMER DESIGN [1 VARIANT IN EVERY ADDITIONAL PERSON]	SANGER SEQUENCING	RS 2500	3 WEEKS
59	DNA EXTRACTION	KIT BASED	RS 1000	NA
60	D4Z4 ARRAY CONTRACTION TESTING FOR FSHD-1	OPTICAL GENOME MAPPING	RS 46500	4 WEEKS
61	STRUCTURAL VARIATION ANALYSIS FOR CONSTITUTIONAL CHROMOSOMAL ABNORMALITIES	OPTICAL GENOME MAPPING	RS 53000	4 WEEKS