Administrative Resolution No. (62) of 2020
Approving the Charges for Laboratory Tests Conducted at the Dubai Health Authority Laboratories[1]

The Director General of the Dubai Health Authority,

After perusal of:

Law No. (14) of 2009 Concerning the Pricing of Government Services in the Emirate of Dubai and its amendments;

Law No. (1) of 2016 Concerning the Financial Regulations of the Government of Dubai;

Law No. (6) of 2018 Concerning the Dubai Health Authority (the "**DHA**");

Decree No. (9) of 2012 Approving a Pricing Method for Dubai Health Authority Services;

Decree No. (17) of 2018 Establishing the Corporations Affiliated to the Dubai Health Authority and Determining their Functions;

Decree No. (18) of 2018 Appointing the Director General of the Dubai Health Authority;

Executive Council Resolution No. (18) of 2018 Approving the Organisational Structure of the Dubai Health Authority;

Administrative Resolution No. (148) of 2014 Concerning Health Service Fees; and

The Department of Finance approval of the charges for new laboratory tests, dated 28 June 2020, Ref. DOF/OUT/2020/0001504,

Does hereby issue this Resolution.

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[1] *Every effort has been made to produce an accurate and complete English version of this legislation. However, for the purpose of its interpretation and application, reference must be made to the original Arabic text. In case of conflict, the Arabic text will prevail.*

Approval of Charges
Article (1)

Pursuant to this Resolution, the charges for laboratory tests conducted at the DHA laboratories, as stated in the Schedule attached hereto, are approved.

Implementation
Article (2)

All Organisational Units of the DHA, each within its own powers, must take the necessary action to implement this Resolution.

Repeals
Article (3)

Any provision in any other administrative resolution will be repealed to the extent that it contradicts the provisions of this Resolution.

Commencement and Publication
Article (4)

This Resolution comes into force on the day on which it is issued, and will be published in the Official Gazette.

Humaid Al Qatami

Director General

Dubai Health Authority

Issued in Dubai on 6 July 2020

Corresponding to 15 Thu al-Qidah 1441 A.H.

Charges for Laboratory Tests Conducted at the Dubai Health Authority Laboratories

SN	Service Code	Service Description	Proposed Price in Dirhams
1	81214	BRCA1 (breast cancer 1) (e.g. hereditary breast and ovarian cancer) gene analysis; full sequence analysis and common duplication/deletion variants (i.e. exon 13 del 3.835kb, exon 13 dup 6kb, exon 14-20 del 26kb, exon 22 del 510bp, exon 8-9 del 7.1kb)	6,490.00
2	81408	Molecular pathology procedure, Level 9 (e.g. analysis of > 50 exons in a single gene by DNA sequence analysis) FBN1 (fibrillin 1) (e.g. Marfan syndrome), full gene sequence NF1 (neurofibromin 1) (e.g. neurofibromatosis, type 1), full gene sequence RYR1 (ryanodine receptor 1, skeletal) (e.g. malignant hyperthermia), full gene sequence VWF (von Willebrand factor) (e.g. von Willebrand disease types 1 and 3), full gene sequence	6,511.00
3	81407	Molecular pathology procedure, Level 8 (e.g. analysis of 26-50 exons by DNA sequence analysis, mutation scanning or duplication/deletion variants of > 50 exons, sequence analysis of multiple genes on 1 platform) SCN1A (sodium channel, voltage-gated, type 1, alpha subunit) (e.g. generalized epilepsy with febrile seizures), full gene sequence	8,545.00
4	2008127	ANTI SACCHAROMYCES CEREVI. ABS	174.00
5	2090018	Aicardi-Goutières syndrome (NGS)	6,830.00
6	2090019	Aldosterone-Sensitive Distal Nephron (NGS)	6,830.00
7	2090020	Alport syndrome (NGS)	6,830.00
8	2090021	Amyotrophic lateral sclerosis (ALS) (NGS)	10,430.00

9	2090022	Angelman -like syndrome (NGS)	8,630.00
10	2090023	Aortopathy (NGS)	10,430.00
11	2090024	Cardiomyopathy (ARVD/ARVC)/Arrhythmogenic right ventricular dysplasia (NGS)	8,630.00
12	2090025	Ataxia (NGS)	10,430.00
13	2090026	Ataxic polyneuropathies (NGS)	10,430.00
14	2090027	Auditory neuropathy (NGS)	6,830.00
15	2090028	Burkitt lymphoma (NGS)	6,830.00
16	2090029	C3-Glomerulopathies (C3G) (NGS)	8,630.00
17	2090030	Cardiofaciocutaneous syndrome (NGS)	8,630.00
18	2090031	Cardiomyopathy, dilated (NGS)	10,430.00
19	2090032	CFHR5-Nephropathy (NGS)	8,630.00
20	2090033	Chondrodysplasia punctate (NGS)	6,830.00
21	2090034	Congenital disorders of glycosylation (CDG) (NGS)	10,430.00
22	2090035	Congenital heart defects (NGS)	10,430.00
23	2090036	Corneal dystrophies (NGS)	8,630.00
24	2090038	Craniosynostosis syndromes (NGS)	10430.00
25	2090039	Cutaneous T-cell lymphoma (NGS)	6,830.00
26	2090040	Cystinuria (NGS)	6,830.00
27	2090041	Dent syndrome (NGS)	6,830.00
28	2090042	Diabetes insipidus (NGS)	6,830.00
29	2090043	Diabetes mellitus,monogenic (NGS)	10,430.00
30	2090044	Diffuse large B-cell lymphoma (NGS)	8,630.00
31	2090045	Disorders of sex development (DSD (NGS)	10,430.00
32	2090046	Dyskeratosis congenita (NGS)	8,630.00
33	2090047	Dystonie (NGS)	10,430.00
34	2090048	Ectodermal dysplasia (NGS)	10,430.00
35	2090049	Ellis-van-Crefeld syndrome (NGS)	7,280.00
36	2090050	Epidermolysis bullosa (NGS)	8,630.00
37	2090051	Epilepsy (NGS)	10,430.00
38	2090052	Epilepsy, metabolic (NGS)	10430.00
39	2090053	Fanconi anemia (NGS)	8,630.00
40	2090054	Fatty acid oxidation disorders (NGS)	8,630.00
41	2090055	Follicular lymphoma (NGS)	6,830.00
42	2090056	Frontotemporal dementia (NGS)	8,630.00
43	2090057	Glucocorticoid Deficiency (NGS)	6,830.00

44	2090058	Familial Gluconeogenesis (NGS)	6,830.00
45	2090059	Glycogen Storage Disease (NGS)	7,223.00
46	2090060	Growth Hormone Deficiency (NGS)	10,430.00
47	2090062	Hereditary sensory and autonomic neuropathy (HSAN) (NGS)	8,630.00
48	2090063	Hermansky-Pudlak syndrome (NGS)	6,830.00
49	2090064	Holoprosencephaly Hyperekplexia (NGS)	6,830.00
50	2090065	Hyperoxaluria Hypertriglyceridemia (NGS)	6,830.00
51	2090066	Familial Hypoglycemia (NGS)	10,430.00
52	2090067	Hypogonadotropic hypogonadism (NGS)	8,630.00
53	2090068	Hypophosphatemic rickets/Phosphate diabetes (NGS)	8,630.00
54	2090069	Ichthyosis and related disorders of cornification (NGS)	10,430.00
55	2090070	Cholestasis, progressive familial intrahepatic (NGS)	6,830.00
56	2090071	Kartagener syndrome (NGS)	10,430.00
57	2090072	Ketogenesis disorder (NGS)	6,830.00
58	2090073	Ketolysis disorder (NGS)	6,830.00
59	2090074	Left ventricular noncompaction (NGS)	6,830.00
60	2090075	Leigh syndrome (NGS)	10,430.00
61	2090076	Acute Myeloid Leukemia (NGS)	8,630.00
62	2090077	Atypical Chronic Myeloid Leukemia (NGS)	6,830.00
63	2090078	Chronic Myelomonocytic Leukemia (NGS)	6,830.00
64	2090079	Chronic Neuthrophilic Leukemia (NGS)	6,830.00
65	2090080	Leukodystrophy (NGS)	10,430.00
66	2090081	T-cell acute lymphoblastic Leukemia (NGS)	8,630.00
67	2090082	Limb-girdle muscular dystrophy (LGMD) (NGS)	10,430.00
68	2090083	Lissencephaly (NGS)	10,430.00
69	2090084	Loeys-Dietz syndrome (NGS)	6,830.00
70	2090085	Lysosomal disorders (NGS)	10,430.00
71	2090086	Macrocephaly (NGS)	10,430.00
72	2090087	Malignant Hyperthermia (NGS)	6,830.00
73	2090088	Mantle Cell Lymphoma (NGS)	6,830.00

74	2090089	Medullary Cystic Kidney Disease (MCKD) (NGS)	6,830.00
75	2090090	Micromelic Dysplasia (NGS)	10,430.00
76	2090091	Metaphyseal dysplasia (NGS)	6,830.00
77	2090092	Mitochondrial encephalopathy (NGS)	10,430.00
78	2090093	Morbus Waldenström (NGS)	6,830.00
79	2090094	mtDNA Depletion/Integrity panel (NGS)	6,830.00
80	2090095	Mucopolysaccharidosis (NGS)	8,630.00
81	2090096	Multiple Epiphyseal Dysplasia and Pseudoachondroplasia (NGS)	6,830.00
82	2090097	Multiple Myeloma (NGS)	6,830.00
83	2090098	Muscular Dystrophies,Congenital (NGS)	10,430.00
84	2090099	Myasthenic Syndrome, Congenital (NGS)	8,630.00
85	2090100	Myelodysplastic syndrome (NGS)	10,430.00
86	2090101	Myopathy, Distal (NGS)	10,430.00
87	2090103	Myotonia (NGS)	6,830.00
88	2090104	Nephrocalcinosis (NGS)	10,430.00
89	2090105	Neurodegeneration with Brain Iron Accumulation (NBIA) (NGS)	8,630.00
90	2090106	Neurofibromatosis (NF) (NGS)	6,830.00
91	2090107	Neuronal Migration Disorder (NGS)	10,430.00
92	2090108	NK/T-Cell Lymphoma (NGS)	6,830.00
93	2090109	Nystagmus (NGS)	6,830.00
94	2090110	Obesity (NGS)	10,430.00
95	2090112	Pancreas Carcinoma (NGS)	8,630.00
96	2090113	Pancreatitis, Chronic (NGS)	6,830.00
97	2090114	Parkinson Disease (NGS)	13,966.00
98	2090115	Pena-Shokeir Syndrome (NGS)	8,630.00
99	2090116	Pendred Syndrome (NGS)	8,630.00
100	2090117	Periodic Fever Syndromes/Autoinflammation (NGS)	10,430.00
101	2090118	Peripheral T-Cell Lymphoma (NGS)	6,830.00
102	2090119	Perrault Syndrome (NGS)	6,830.00
103	2090120	Polymicrogyria (NGS)	8,630.00
104	2090121	Pontocerebellar Hypoplasia (NGS)	6,830.00
105	2090122	Porphyria (NGS)	6,830.00
106	2090123	Progeria Syndromes (NGS)	8,630.00
107	2090124	Progressive External Ophthalmoplegia (PEO) (NGS)	8,630.00

108	2090125	Prostate Cancer (NGS)	10,430.00
109	2090126	Pulmonary Hypertension (NGS)	6,830.00
110	2090127	Pyruvate Dehydrogenase Deficiency (NGS)	6,830.00
111	2090128	Rasopathies (NGS)	8,630.00
112	2090129	Refsum Syndrome (NGS)	8,630.00
113	2090131	Renal Carcinoma (NGS)	8,630.00
114	2090132	Retinitis Pigmentosa (NGS)	10,430.00
115	2090133	Retinoblastoma (NGS)	6,830.00
116	2090134	SANDD Syndrome (NGS)	6,830.00
117	2090135	Schizencephaly (NGS)	6,830.00
118	2090136	Short Stature (NGS)	8,630.00
119	2090137	Skeletal Dyplasia with Abnormal Bone Density/Mineralisation (NGS)	8,630.00
120	2090138	Spastic Paraplegia (NGS)	10,430.00
121	2090139	Spinal Muscular Atrophy, Distal (NGS)	10,430.00
122	2090140	Splenic Marginal Zone Lymphoma (NGS)	6,830.00
123	2090141	Spondylometaphyseal Dysplasia and Spondyloepiphyseal Dysplasia (NGS)	10,430.00
124	2090142	Stargardt Disease (NGS)	6,830.00
125	2090143	Stickler Syndrome (NGS)	6,830.00
126	2090145	Thrombotic Microangiopathy (TMA) (NGS)	8,630.00
127	2090146	Thrombotic Thrombocytopenic Purpura (TTP) (NGS)	8,630.00
128	2090147	Treacher Collins Syndrome (NGS)	6,830.00
129	2090148	Walker-Warburg Syndrome (NGS)	8,630.00
130	2090149	Warburg-Micro Syndrome (NGS)	10,430.00
131	2090150	Xanthinuria (NGS)	6,830.00
132	2090151	Xeroderma Pigmentosum (NGS)	6,830.00
133	2090152	X-linked mental retardation (NGS)	10,430.00
134	2090154	Albinism, Oculocutaneous (MC1R)	1,655.00
135	2090155	Alpha-1-Antitrypsin Deficiency (SERPINA1)	980.00
136	2090156	Alpha-Thalassemia (HBA, HBA2, sequencing + MLPA)	2,780.00
137	2090157	Pulmonary Alveolar Microlithiasis (SLC34A2)	3,680.00

138	2090158	Amyloidosis, Hereditary, Transthyretin-related (TTR, stage 1)	980.00
139	2090159	Amyloidosis, Hereditary, Transthyretin-related (TTR, stage 2)	1,295.00
140	2090160	Arthrogryposis (TNNI2)	2,105.00
141	2090161	Arts Syndrome (PRPS1)	2,330.00
142	2090162	Ataxia Teleangiectasia (ATM)	6,830.00
143	2090163	Ataxia Teleangiectasia (ATM, MLPA)	1880.00
144	2090164	Beta-Propeller Protein-Associated Neurodegeneration (WDR45)	2,555.00
145	2090166	Blepharophimosis (FOXL2)	980.00
146	2090167	Hypertension and Brachydactyly Syndrome (PDE3A)	5,399.00
147	2090169	Coenzyme Q10 Deficiency (NGS)	8,630.00
148	2090170	Cohesinopathies (e.g. Cornelia de Lange Syndrome) (NGS)	6,830.00
149	2090171	Cowden Syndrome 3 (NGS)	6,830.00
150	2090172	CTNNB1-Associated Diseases (CTNNB1)	3,905.00
151	2090173	Donnai-Barrow Syndrome (LRP2)	6,830.00
152	2090174	Dravet Syndrome (GABRG2)	3455.00
153	2090176	Ellis-Van-Creveld Syndrome (EVC, EVC2, MLPA)	1,880.00
154	2090177	Encephalopathy Syndrome, Lethal Neonatal Spasticity-Epileptic (BRAT1)	5,030.00
155	2090178	Epileptic Encephalopathy (WWOX)	3,005.00
156	2090180	Fleck Retina, Familial Benign (PLA2G5)	1,565.00
157	2090181	Exudative Vitreoretinopathy (LRP5)	6,830.00
158	2090182	Exudative Vitreoretinopathy Type 1 (FZD4)	1,115.00
159	2090183	Familial Isolated Arrhythmogenic Ventricular Dysplasia (PKP2)	5,317.00
160	2090184	X-Linked Intellectual Disability (OPHN1)	4,580.00
161	2090185	Limb-Girdle Muscular Dystrophy Type 2D (SGCA)	3,157.00
162	2090186	Glioma (POT1) (NGS)	4,580.00
163	2090187	Hemolytic-Uremic Syndrome, Atypical (AHUS3/CFI)	3,110.00
164	2090188	Hemolytic-Uremic Syndrome, Atypical (AHUS5/C3)	5,930.00

165	2090189	Hemolytic-Uremic Syndrome, Atypical (AHUS7/DGKE)	3,005.00
166	2090190	Hermansky-Pudlak syndrome 1 (HPS1)	5,030.00
167	2090191	Hermansky-Pudlak Syndrome 3 (HPS3)	5,030.00
168	2090192	Hermansky Pudlak Syndrome 5 (HPS5)	5,030.00
169	2090193	Hermansky-Pudlak Syndrome 6 (HPS6)	2,371.00
170	2090194	Hermansky-Pudlak Syndrome 8 (HPS8/BLOC1S3)	980.00
171	2090195	HIV Infection, Susceptibility/Resistance to (CCR5)	980.00
172	2090196	Corneal Dystrophy (SLC4A11)	5,480.00
173	2090197	Hyper-IgE Syndrome-NGS (NGS)	6,830.00
174	2090198	Hyperlipoproteinemia (APOA5) (NGS)	6,830.00
175	2090199	Hypophosphatemia with Hypercalciuria (SLC34A3)	2,555.00
176	2090200	Hypophosphatemia with Nephrolithiasis or Osteoporosis (SLC34A1)	4,008.00
177	2090201	Hypophosphatemic Rickets, Autosomal Dominant (FGF23, MLPA)	1,913.00
178	2090202	Imerslund-Grasbeck Syndrome (AMN)	3,095.00
179	2090203	Jalili Syndrome (cone-rod retinal dystrophy and amelogenesis imperfecta) (CNNM4)	3,026.00
180	2090204	Jervell- and Lange-Nielsen Syndrome (NGS)	6,830.00
181	2090205	Joubert Syndrome (POC1B)	3,905.00
182	2090206	Campomelic Dysplasia (SOX9, MLPA)	1,880.00
183	2090207	Cataract Syndrome (CRYAA)	1,880.00
184	2090208	Cataract 14, Multiple Types (GJA3)	980.00
185	2090210	Lathosterolosis (SC5DL)	3,680.00
186	2090211	Legius syndrome (SPRED1)	2,465.00
187	2090213	Lesch-Nyhan syndrome (HPRT1, MLPA)	2,371.00
188	2090215	Lujan-Fryns Syndrome (MED12) (NGS)	6,830.00
189	2090216	Mayer-Rokitansky-Küster-Hauser Syndrome (WNT4)	2,015.00
190	2090217	Megalencephaly-Capillary Malformation-Polymicrogyria Syndrome, Somatic (PIK3CA)	5,930.00
191	2090218	Microcephaly (ASPM)	6,830.00

192	2090219	Microcephaly (KIF11)	4,130.00
193	2090220	Microcephaly (PCNT)	10,430.00
194	2090221	Microcephaly (RBBP8)	5,930.00
195	2090222	Microcephaly (SLC25A19)	2,555.00
196	2090223	Mitochondrial DNA Depletion Syndrome (DGUOK)	2,830.00
197	2090224	Wilson Disease (ATP7B, Stage 3, MLPA)	1,880.00
198	2090226	Myotonia Congenita Type Becker/Thomsen (CLCN1, MLPA)	2,155.00
199	2090227	Nephronophthisis (NPHP4)	5,930.00
200	2090228	Nephronophthisis-like Nephropathy (NPHPL1/XPNPEP3)	3,353.00
201	2090230	Nephrotic Syndrome (WDR73)	2,465.00
202	2090231	Retinal Dystrophy (RGS9BP)	980.00
203	2090232	Hereditary Sensory and Autonomic Neuropathy Type 5 (NGF)	1,295.00
204	2090233	Noonan-like Syndrome with Loose Anagen Hair (SHOC2)	3,095.00
205	2090234	Osteopathia Striata, Cranial Sclerosis (AMER1/WTX)	2,465.00
206	2090235	Paralysis, Thyrotoxic Periodic (KCNJ18)	1,295.00
207	2090236	Perrault Syndrome (CLPP)	1,880.00
208	2090237	Perrault Syndrome (LARS2)	4,662.00
209	2090238	Polycystic Liver Disease (GANAB)	6,155.00
210	2090240	Retinol Dystrophy, Iris coloboma, and Comedogenic Acne Syndrome (RBP4)	1,655.00
211	2090241	Retinitis Pigmentosa (IMPG2)	6,830.00
212	2090242	Retinitis Pigmentosa (LRAT)	1,115.00
213	2090243	Retinitis Pigmentosa (NR2E3)	3,095.00
214	2090244	Rett-like syndrome (NTNG1, MLPA)	1,880.00
215	2090245	Rett-like syndrome (NTNG1, Seq.)	2,195.00
216	2090246	Robinow Syndrome (WNT5A)	4,355.00
217	2090247	Deafness (GPSM2)	4,580.00
218	2090248	Very Long Chain Acyl-CoA Dehydrogenase Deficiency (ACADVL)	6,380.00
219	2090249	Simpson-Golabi-Behmel Syndrome (GPC3) (Testicular Feminization Syndrome)	3,353.00
220	2090251	TARP Syndrome (RBM10)	5,930.00

221	2090254	Hyperphosphatemic Tumoral Calcinosis, Familial (GALNT3)	5,534.00
222	2090255	Vasculopathy, ADA2 Deficiency (CECR1)	3,005.00
223	2090256	Wolman Disease (LIPA)	3,005.00
224	2090257	Cone-Rod Dystrophy (CTNNA1)	4,355.00
225	2090258	Congenital Adrenal Hyperplasia (NGS)	8,630.00
226	2090259	Lysosomal Perioxisomal Disorders NGS	10,430.00
227	2090260	CYP21A2 Deletion & Duplication (MLPA)	1,880.00
228	2090261	Microarray CGH 180 K Amniotic Fluid/CVS	6,750.00
229	2090262	PTEN Gene Amniotic fluid/CVS	2,925.00
230	2090263	Skeletal Dysplasia and Pena Shokeir Syndrome	8,630.00
231	2090264	Methylation Test Russel Siver Syndrome	1,880.00
232	2090265	Ichthyoses and Related Disorder panels	10,430.00
233	2090266	Heteroxy and Situs Inversion Panel	10,430.00
234	2090267	Multi Gene Panel- Mitochondrial	10,430.00
235	2090268	Sister Chromatid Exchange for Bloom Syndrome.	1,871.00
236	2090269	Breast Cancer Mutigene Panel-NGS	8,630.00
237	2090270	Medium Chain Acyl CoA Deficiecy	3,905.00
238	2090271	Gastrointestinal Atresia Panel	6,626.00
239	2090272	KGB Syndrome	3,680.00
240	2090273	Neurofascins Abs. (140 & 155)	1,954.00
241	2090274	Contactin- 1 (CASPR1)	1,138.00
242	2090275	Amlodipine	418.00
243	2090276	Beta HCG(CSF)	454.00
244	2090277	Alpha Feto protein (AFP) (CSF)	205.00
245	2090278	HIV Viral load(CSF)	945.00
246	2090279	5 Methyl Tetra Hydro folate(CSF)	1,221.00
247	2090280	Pregnanacy Associated Plasma Protein(PAPP-A)	305.00
248	2090281	Rituximab Neutralizing Abs	2,905.00
249	2090282	Omega 6 Fatty Acids Screen	306.00
250	2090283	Chlamydia Trachomatis IgA & IgG ABS	251.00
251	2090284	Chlamydia Pneumonia IgM & IgG ABS	251.00
252	2090285	Zika Virus IgG/IgM ABS	305.00
253	2090286	Entero Virus RNA PCR	741.00
254	2090287	Legionella Culture Specimen	162.00

255	2090288	Porphyrin	251.00
256	2090289	Porphobilinogen	251.00
257	2090290	Pseudomonas Aeruginosa Abs.	445.00
258	2090291	Oncotype DX (21 GENES)	22,080.00
259	2090292	Iodine Urine	113.00
260	2090293	Cytomegalovirus PCR CSF	233.00
261	2090294	Chickengunya Virus RNA PCR	395.00
262	2090295	Dihydropteridine Reductase (DHPR) Activity	305.00
263	2090296	Fibroblast Growth Factor 23 Level	180.00