

검사명 변경 항목 list (총 69항목)

※ 적용일: 03월 02일 접수분

No.	GC Labs 코드	변경 전	변경 후
1	B661	Achondroplasia FGFR3	FGFR3 gene Major mutation (Achondroplasia) [Sequencing]
2	Z020	AML1/ETO 유전자 재배열	AML1/ETO (RUNX1/RUNX1T1) 유전자 재배열 [RT-PCR]
3	S875	Apo E genotype	APOE genotype [Real-time PCR]
4	X779	Avellino corneal dystrophy mutation	TGFBI gene R124H mutation (Corneal dystrophy, Avellino type) [Real-time PCR]
5	M345	BRAF Mutation	BRAF gene mutation [Real-time PCR]
6	L446	BRAF Mutation (Sequencing)	BRAF gene mutation [Sequencing]
7	Z226	CADASIL	NOTCH3 gene mutation (CADASIL) [Sequencing]
8	P538	CADASIL family test	NOTCH3 gene, Familial mutation (CADASIL) [Sequencing]
9	L535	CALR gene mutation	CALR gene mutation [Fragment analysis & Sequencing]
10	N915	CBFB-MYH11 유전자 재배열	CBFB/MYH11 유전자 재배열 [RT-PCR]
11	M347	CEBPA Gene Mutation	CEBPA gene mutation [Sequencing]
12	N040	c-KIT gene mutation (BM)	c-KIT gene mutation (BM) [Sequencing]
13	X348	c-KIT gene mutation (Tissue)	c-KIT gene mutation (Tissue) [Sequencing]
14	X757	c-KIT gene mutation (WB)	c-KIT gene mutation (WB) [Sequencing]
15	X031	E2A-PBX1 유전자 재배열	E2A/PBX1 (TCF3/PBX1) 유전자 재배열 [RT-PCR]
16	N537	EGFR (cell-free DNA, Real-time PCR)	EGFR (cell-free DNA) [Real-time PCR]
17	L694	EGFR (PNA Clamp)	EGFR (Tissue) [PNA clamping Real-time PCR]
18	M040	EGFR (Pyrosequencing)	EGFR (Tissue) [Pyrosequencing]
19	M683	Factor V Leiden mutation	Factor V Leiden R534Q mutation
20	M745	FLT3-ITD	FLT3-ITD mutation [Fragment analysis & Sequencing]
21	M746	FLT3-TKD	FLT3-TKD mutation [Sequencing]
22	M019	GLA Gene (Fabry)	GLA gene mutation (Fabry disease) [Sequencing]
23	K599	HBV 약제내성돌연변이 24종(sequencing)	HBV 약제내성변이 24종 [Sequencing]
24	L265	HCV 1b NS5A L31/Y93 약제내성검사	HCV 1b NS5A L31/Y93 약제내성검사 [Sequencing]
25	N340	HCV genotype	HCV genotype [Real-time PCR]
26	M053	Hemavision 28종	Hemavision, Acute leukemia gene rearrangement profile [RT-PCR]
27	X157	HPRT1 Gene Mutation	HPRT1 gene mutation (Lesch-Nyhan syndrome) [Sequencing]
28	L020	IGH 유전자 재배열	IGH gene rearrangement (IGH 유전자 재배열)
29	M049	JAK2 gene Exon 12 Mutation	JAK2 gene Exon 12 mutation [Sequencing]
30	Z041	JAK2 V617F mutation	JAK2 V617F mutation [Real-time PCR]
31	M562	KRAS 전체 돌연변이	KRAS 전체 돌연변이 [Sequencing]
32	M563	KRAS 주요 돌연변이	KRAS 주요 돌연변이 [Pyrosequencing]
33	M015	MEN1 유전자검사	MEN1 gene mutation (Multiple Endocrine Neoplasia Type 1) [Sequencing]
34	L665	MPL gene W515 mutation	MPL gene W515 mutation [Sequencing]
35	M014	NF2 (Neurofibromatosis 2)	NF2 gene mutation (Neurofibromatosis 2) [Sequencing]

No.	GC Labs 코드	변경 전	변경 후
36	M346	NPM1 gene Mutation	NPM1 gene mutation [Sequencing]
37	M029	NRAS 전체 돌연변이	NRAS 전체 돌연변이 [Sequencing]
38	M030	NRAS 주요 돌연변이	NRAS 주요 돌연변이 [Pyrosequencing]
39	M555	PCR of Major bcr/abl	Major BCR/ABL1 [RT-PCR]
40	M556	PCR of Minor bcr/abl	Minor BCR/ABL1 [RT-PCR]
41	L025	PDGFRA gene mutation	PDGFRA gene mutation [Sequencing]
42	H188	PML/RARA Bone Marrow	PML/RARA 정성 (BM) [RT-PCR]
43	S884	PML/RARA Whole blood	PML/RARA 정성 (WB) [RT-PCR]
44	G229	PML/RARA 정량(BM)	PML/RARA 정량 (BM) [Real-time PCR]
45	M747	PML/RARA 정량(WB)	PML/RARA 정량 (WB) [Real-time PCR]
46	M678	Post-BMT	Post-BMT engraftment analysis
47	M684	Prothrombin G20210A Mutation	Prothrombin G20210A mutation
48	M007	PTEN	PTEN gene mutation (PTEN Hamartoma Tumor syndrome) [Sequencing]
49	Z166	PTPN11 gene mutation (Noonan syndrome)	PTPN11 gene mutation (Noonan syndrome) [Sequencing]
50	K157	Real time major BCR-ABL (IS) 정량	Major BCR/ABL1 (b2a2, b3a2) 정량 (IS) [Real-time PCR]
51	L002	RNF213 gene R4810K mutation	RNF213 gene R4810K mutation (Moyamoya disease) [Sequencing]
52	Z906	SCA 1	SCA 1 gene mutation (Spinocerebellar ataxia 1)
53	Z907	SCA 2	SCA 2 gene mutation (Spinocerebellar ataxia 2)
54	Z908	SCA 3	SCA 3 gene mutation (Spinocerebellar ataxia 3)
55	Z909	SCA 6	SCA 6 gene mutation (Spinocerebellar ataxia 6)
56	Z910	SCA 7	SCA 7 gene mutation (Spinocerebellar ataxia 7)
57	S160	SRY gene	SRY gene (Sex determining Region Y) [PCR]
58	S893	TEL/AML1 유전자 재배열	TEL/AML1 (ETV6/RUNX1) 유전자 재배열 [RT-PCR]
59	B341	Thalassemia mutation	HBB gene mutation (β -Thalassemia) [Sequencing]
60	N131	TTR gene mutation	TTR gene mutation (Hereditary Transthyretin Amyloidosis) [Sequencing]
61	L023	T세포 수용체 감마 유전자 재배열	TCR gamma gene rearrangement (T세포 수용체 감마 유전자 재배열)
62	N476	UGT1A1 gene mutation (Crigler-Najjar syndrome)	UGT1A1 gene mutation (Crigler-Najjar syndrome) [Sequencing]
63	P535	UGT1A1 genotype, major variants (irinotecan sensitivity)	UGT1A1 genotype, Major variants (Irinotecan sensitivity) [Sequencing]
64	M342	VKORC1 Genotype	VKORC1 genotype
65	Z963	Wilson Disease	ATP7B gene mutation (Wilson disease) [Sequencing]
66	L493	Wilson disease family test	ATP7B gene, Familial mutation (Wilson disease) [Sequencing]
67	M031	글리벡내성유전자(Major bcr/abl)	Imatinib 약제내성 (Major BCR/ABL1) [Sequencing]
68	M338	항결핵약제내성(INH,Sequencing)	항결핵약제내성 (INH) [Sequencing]
69	M337	항결핵약제내성(RFP,Sequencing)	항결핵약제내성 (RFP) [Sequencing]