

List of the analysed genes and investigated genetic diseases

Table 1: CardioScreen-Prevention sudden cardiac arrest

	DISEASE NAME	PhenoMIM	GENE
1	Atrial fibrillation, familial, 12	614050	<b>ABCC9</b>
2	Atrial septal defect 5	612794	<b>ACTC1</b>
3	Dilated cardiomyopathy 1AA	612158	<b>ACTN2</b>
4	Cardiomyopathy, hypertrophic/Cardiomyopathy, dilated	609599	<b>ANKRD1</b>
5	Ventricular tachycardia, catecholaminergic polymorphic, 2	611938	<b>CASQ2</b>
6	Cardiomyopathy, familial hypertrophic	192600	<b>CAV3</b>
7	Cardiomyopathy, dilated, 1II	615184	<b>CRYAB</b>
8	Cardiomyopathy, dilated, 1M	607482	<b>CSRP3</b>
9	Cardiomyopathy, dilated	600435	<b>CTF1</b>
10	Cardiomyopathy, dilated, 1I	604765	<b>DES</b>
11	Arrhythmogenic right ventricular dysplasia 11	610476	<b>DSC2</b>
12	Arrhythmogenic right ventricular dysplasia 10	610193	<b>DSG2</b>
13	Arrhythmogenic right ventricular dysplasia 8	607450	<b>DSP</b>
14	Left ventricular noncompaction 1, with or without congenital heart defects	604169	<b>DTNA</b>
15	Emery-Dreifuss muscular dystrophy 1, X-linked	310300	<b>EMD</b>
16	Cardiomyopathy, dilated	602633	<b>FHL2</b>
17	Fabry disease, cardiac variant	301500	<b>GLA</b>
18	Arrhythmogenic right ventricular dysplasia 12	611528	<b>JUP</b>
19	Cardiomyopathy, dilated, 1JJ	615235	<b>LAMA4</b>
20	Danon disease	300257	<b>LAMP2</b>
21	Cardiomyopathy, dilated, 1C, with or without LVNC	601493	<b>LDB3</b>
22	Cardiomyopathy, dilated, 1A	115200	<b>LMNA</b>
23	Cardiomyopathy, dilated, 1MM	615396	<b>MYBPC3</b>
24	Atrial septal defect 3	614089	<b>MYH6</b>
25	Cardiomyopathy, dilated, 1S	613426	<b>MYH7</b>
26	Cardiomyopathy, hypertrophic, 10	608758	<b>MYL2</b>
27	Cardiomyopathy, hypertrophic, 8	608751	<b>MYL3</b>
28	Cardiomyopathy, hypertrophic, 1, digenic	192600	<b>MYLK2</b>
29	Cardiomyopathy, hypertrophic, 16	613838	<b>MYOZ2</b>
30	Cardiomyopathy, dilated, 1CC	613122	<b>NEXN</b>
31	Arrhythmogenic right ventricular dysplasia 9	609040	<b>PKP2</b>
32	Cardiomyopathy, dilated, 1P	609909	<b>PLN</b>
33	Cardiomyopathy, hypertrophic 6	600858	<b>PRKAG2</b>
34	Cardiomyopathy, dilated, 1DD	613172	<b>RBM20</b>
35	Arrhythmogenic right ventricular dysplasia 2	600996	<b>RYR2</b>

36	Cardiomyopathy, dilated, 1L	606685	<b>SGCD</b>
37	Barth syndrome	302060	<b>TAZ</b>
38	Cardiomyopathy, hypertrophic, 25	607487	<b>TCAP</b>
39	Arrhythmogenic right ventricular dysplasia 5	604400	<b>TMEM43</b>
40	Cardiomyopathy, dilated, 1Z	611879	<b>TNNC1</b>
41	Cardiomyopathy, dilated, 2A	611880	<b>TNNI3</b>
42	Cardiomyopathy, dilated, 1D	601494	<b>TNNT2</b>
43	Cardiomyopathy, dilated, 1Y	611878	<b>TPM1</b>
44	Cardiomyopathy, dilated, 1G	604145	<b>TTN</b>
45	Amyloidosis, hereditary, transthyretin-related	105210	<b>TTR</b>
46	Cardiomyopathy, dilated, 1W	611407	<b>VCL</b>