St. Number	Assigned disease/trait
1- 16701958	Ehler-Danlos Syndrome
2- 149261	Beta-thalassemia
3- 16700623	Sickle-cell Anemia
4- 17701925	Phenylketonuria
5- 16701178	Huntington's Disease
6- 16330793	Cystic Fibrosis
7- 17330559	Color Blindness
8- 16700647	Muscular Dystrophy
9- 16701811	Polydactyly
10-16701454	Early-onset Familial Alzheimer's Disease
11-17000261	Friedreich's Ataxia
12-17701065	Osteogenesis Imperfecta
13-15702305	Fragile X Syndrome
14-16330821	Ehler-Danlos Syndrome
15-17701327	Beta-thalassemia
16-16700560	Sickle-cell Anemia
17-16330801	Phenylketonuria
18-19500771	Huntington's Disease
19-16450182	Cystic Fibrosis
20-15000342	Color Blindness
21-15702206	Muscular Dystrophy
22-17700367	Polydactyly
23-15701978	Early-onset Familial Alzheimer's Disease
24-15330341	Friedreich's Ataxia
25-15701968	Osteogenesis Imperfecta
26-17700209	Fragile X Syndrome
27-17701179	Ehler-Danlos Syndrome
28-16701739	Beta-thalassemia
29-17701206	Sickle-cell Anemia
30-16300006	Phenylketonuria
31-17700320	Huntington's Disease
32-133580	Cystic Fibrosis
33-148034	Color Blindness
34- 17330581	Muscular Dystrophy
35- 16330804	Polydactyly
36- 17330523	Early-onset Familial Alzheimer's Disease
37- 17701447	Friedreich's Ataxia