22q11.2 Deletion Syndrome

(Included conditions: DiGeorge syndrome, Velocardiofacial syndrome, Conotruncal anomaly face syndrome, Autosomal dominant Opitz G/BBB syndrome, Sedlackova syndrome, Cayler cardiofacial syndrome)

If you have just found out through prenatal testing that your baby has or *may* have 22q11.2 deletion syndrome and are looking for more information, the Genetic Support Foundation is a good starting point.

There are many other important resources out there, including your doctor and genetic counselor. We have also compiled a list of resources that can provide you with additional information and support. After reviewing the information below, if you have a question that you are having difficulty finding an answer for, please feel free to contact us.