

Understanding Your VUS Cardiovascular Genetic Test Result Information for patients with a variant of unknown significance

| Result | VUS | Your test result shows you have a variant of unknown significance (VUS), a change in a gene that can cause an inherited cardiovascular disorder. In this case, we do not know if it is the source or not. |
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| Diagnosis | NO CHANGE | This test result does not change your cardiovascular diagnosis. If you were diagnosed with cardiomyopathy, arrhythmia, or another, that remains the same. |
| Possible Testing | FOR FAMILY MEMBERS | Your report will indicate if testing family members may help us learn more about your specific VUS. Please speak with your healthcare provider to determine if they might also benefit from a test to evaluate their personal risk of developing a disease. |
| Management Options | PATIENTS WITH ARRHYTHMIA OR CARDIOMYOPATHY | Treatment options include medications, surgery, pacemakers, implantable cardioverter defibrillators (ICDs), or avoiding certain athletic activities. Talk to your doctor about which may be right for you. |
| Screening Options | PATIENTS WITH A FAMILY HISTORY OF CARDIOMYOPATHY OR ARRHYTHMIA, BUT NO PERSONAL SIGNS | Options for screening and early detection include physical exams, echocardiograms, electrocardiograms (EKGs), or cardiac MRI. Talk to your doctor about whether these options are right for you. |
| Next Steps | DISCUSS | Please share this with family members so they can talk to their doctors and learn more. |
| Reach Out | RESOURCES | Ambry's Cardiology Site for Families ambrygen.com/patients/cardiology National Society of Genetic Counselors nsgc.org Hypertrophic Cardiomyopathy Association 4hcm.org Sudden Arrhythmia Death Syndromes (SADS) Foundation sads.org Children's Cardiomyopathy Foundation childrenscardiomyopathy.org Genetic Information Nondiscrimination Act (GINA) ginahelp.org |

Cardiomyopathy or Arrhythmia in the Family

Even though your genetic testing result was a VUS, some cardiomyopathies and arrhythmias can still run in families. All close family members of someone with an inherited cardiomyopathy or arrhythmia (like parents, brothers, sisters, children) should talk with their doctor about screening.

Please speak to your doctor or genetic counselor about this. The field of genetics is continuously changing, so updates related to your result, medical recommendations, and/or potential treatments may be available over time. This information is not meant to replace a discussion with a healthcare provider and should not be considered or taken as medical advice.

