Will my child always need to see a heart doctor?

Yes. Just like your child needs special heart care today, he/she will always need special heart care—even as a teenager and adult. It’s very important that people born with heart defects get regular heart checks. Heart problems can start with no symptoms. Finding and treating small problems early can prevent big problems later on.

Will my child need a special kind of heart doctor when he/she grows up?

Yes. Most adult cardiologists do not have the special training needed to take care of complex heart defect patients. Your child will always need to look for a doctor with special training in congenital heart disease. Many children’s hospitals now work with an adult congenital heart program. Ask your child’s heart care team what they recommend. The Adult Congenital Heart Association (www.achaheart.org) also has an online and print listing of adult congenital heart programs throughout the country.

Can’t my child just stay with a pediatric heart team his/her whole life?

No, but many young adults do continue to get care from their pediatric team. If your child chooses to go away to school or move for work, he/she will need to start with a new heart care team. As your child gets older, he/she may have new adult health issues, such as pregnancy. There are also heart problems that older adults with heart defects get that teens and young adults do not. Adult congenital heart programs treat congenital heart patients who “outgrow” pediatric care. ACHA has many resources, including webinars and FAQ sheets, to help you and your child learn more about lifelong CHD heart health.

Have more questions?

Your child’s heart care team can help you learn more about his or her lifelong heart needs. We encourage you to talk to them now. The more questions you ask, the better you can plan for your child’s future.

The mission of the Adult Congenital Heart Association (ACHA) is to empower the congenital heart disease community by advancing access to resources and specialized care that improve patient-centered outcomes.

ACHA offers resources to help heart kids thrive as adults. Call us toll-free at (888) 921-ACHA or visit www.achaheart.org

The information in this brochure was based on the ACC/AHA Joint Guidelines on Managing Adults with Congenital Heart Disease. Learn more at www.achaheart.org/guidelines.
Was your child born with a heart defect?

The good news is that children born with heart problems now live longer than ever before. But getting the right kind of heart care will help your child’s special heart stay as healthy as possible. When your child is an adult, he/she will still need to get his/her heart checked regularly. This is true even if he/she had successful surgery or other treatment. Children with congenital heart disease (CHD) will need lifelong care from congenital heart experts. This brochure lists congenital heart defects considered simple, moderately complex, and highly complex. If your child’s problem is listed as complex, this brochure is for you.

How can you lay the foundation to help your child thrive as a teen and adult with CHD? Your child’s heart team can guide you through what you and your child need to know. Ask them what you can do now to plan for the future. You and your child can learn together. Here are some questions and answers to get you started:

Should I tell my child about his/her heart defect?

Yes. You and your child should learn the name of the congenital heart defect before the teen years. At your child’s next visit ask your heart care team to write it down here:

Practice saying this at home. Sometimes it is easy to forget. Don’t be afraid to ask to review it at every heart check. Your child’s heart care team can also give you tips on how to talk with your child about her heart.

www.achaheart.org

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Your child’s heart care team are the experts on your child’s heart. Ask them to tell you more about your child’s long-term care needs. The chart below lists recommendations for adults with congenital heart disease. Learn more about lifelong care for congenital heart patients at www.achaheart.org.

**SIMPLE CONGENITAL HEART DISEASE**

Periodic heart checks should occur. General cardiologists can usually oversee care. One visit to an ACHD program to verify diagnosis and health status is recommended. ACHD care is usually not needed unless new problems occur.

**Unrepaired conditions:**
- Isolated small atrial septal defect (ASD)
- Isolated small ventricular septal defect (VSD)
- Mild pulmonic stenosis
- Isolated dextrocardia, no other heart problems

**Repaired conditions:**
- Patent ductus arteriosu (PDA)
- Secundum atrial septal defect (ASD)
- Isolated ventricular septal defect (VSD)

**Repair or unrepaired conditions:**
- Isolated aortic valve disease
- Isolated mitral valve disease
- Isolated patent foramen ovale (PFO)

These patients should be seen every year or more frequently at an adult congenital heart program.

**MILDLY COMPLEX CONGENITAL HEART DISEASE**

- Congenitally Corrected Transposition of the Great Arteries (ccTGA or L-TGA)
- Double outlet ventricle
- Mitral atresia
- Pulmonary atresia (all forms)
- Pulmonary arterial hypertension
- Shone’s syndrome
- Single ventricle – all forms (i.e., double-inlet ventricle; HLHS, HRHS, common/primitive ventricle)
- Transposition of the Great Arteries (d-TGA)

**Repaired conditions:**
- Coarctation of the aorta
- Ebstein’s anomaly
- Infundibular right ventricular outflow obstruction (moderate to severe)
- Pulmonary valve regurgitation (moderate to severe)
- Pulmonic valve stenosis (moderate to severe)
- Sinus of Valsalva fistula/aneurysm
- Subvalvar or supravalvar aortic stenosis
- Tetralogy of Fallot
- Ventricular septal defect (VSD) with any valve problems and/or obstructions

**HIGHLY COMPLEX CONGENITAL HEART DISEASE**

These patients should be seen every year or more frequently at an adult congenital heart program.

**Repaired or unrepaired conditions:**
- Tricuspid atresia
- Truncus arteriosus/hemitruncus
- Other abnormalities of AV connections (i.e., criss-cross heart, isomerism, heterotaxy syndromes)

All patients who have undergone any of the following procedures:
- Arterial switch procedure
- Blalock-Taussig shunt

All patients with Eisenmenger syndrome
All patients who are cyanotic (“blue”)