North Carolina Women’s Hospital
Cleft Lip and Cleft Palate

What is a cleft lip?
A cleft lip is an opening of the lip. The lips form during the 4th to 7th week of pregnancy. If the tissue in the lip does not form completely, a cleft lip will occur. The opening can be a small slit or a large opening that goes through the lip into the nose. It can be on one side of the mouth (unilateral), on both sides (bilateral) or in the center of the lip.

What is a cleft palate?
An opening of the roof of the mouth is called a cleft palate. The roof of the mouth forms during the 6th to 9th weeks of pregnancy. A cleft palate occurs when the tissue in the roof of the mouth does not join correctly. A cleft palate can occur alone or with a cleft lip. It can sometimes be difficult to see if there is a cleft plate on ultrasound, so it may not be known if the baby has a cleft lip alone or a cleft lip with a cleft palate until after delivery. A cleft palate without a cleft lip is usually not diagnosed during pregnancy.

What causes cleft lip and cleft palate?
Most babies born with a cleft lip and/or cleft palate have no additional health problems. In these cases, it is thought that the cleft lip could have been caused by both genetic and environmental factors. In most cases, the exact cause cannot be determined. Certain medications or other exposures during early pregnancy may have led to the cleft lip or palate. Other times cleft lip and cleft palate are a part of a genetic condition in the baby.

It is important to remember that anyone can have a baby with a birth defect. It is estimated that 1 in 1000 babies is born with a cleft lip (with or without cleft palate) and 1 in 2500 babies is born with just a cleft palate. Although it is normal for parents to worry about what caused the problem, birth defects are rarely caused by something that the parents did or did not do during pregnancy.

Can cleft lip and cleft palate be repaired?
A cleft lip and/or cleft palate can be repaired with surgery after your baby is born. Surgery to repair a cleft lip is recommended before 12 months and is usually done in the first few months of life. Surgery for cleft palate is usually done within the first 18 months. Some children will require more surgeries as they get older. A pediatric surgical specialist will talk with you about the options for surgery and help you decide what is best for your child.

Your child will likely be followed by the UNC Craniofacial Center, which is a team of healthcare professionals who care for children and families with cleft lip and cleft palate. The team includes many different specialists who will be involved in the care of your child.

Will my baby have other health problems?
Approved by UNC Women’s Hospital Patient Education Steering Committee on June 2007, Reviewed & Revised on August 2010, April 2011, January 2016
Most babies with a cleft lip and/or cleft palate do not have additional birth defects or health problems. Some babies born with a cleft lip have speech or dental problems later in childhood. Some babies born with cleft palate may have frequent ear infections.

Some babies with cleft lip and/or cleft palate are found to have additional birth defects. When a cleft lip is diagnosed on routine ultrasound, a more detailed ultrasound is usually recommended to look for other birth defects. If other problems are found on the ultrasound, your genetic counselor or healthcare provider will discuss the findings with you in detail. Additional testing during pregnancy may be offered to you to help determine the cause of the birth defects.

Your baby will have more evaluations after delivery because not all health problems can be diagnosed during pregnancy. You may meet with a pediatric geneticist (doctor who studies genetic conditions) after your baby is born. The geneticist may help you understand why the cleft lip and cleft palate happened and the chance of having another child with a similar condition.

**How will my baby be delivered?**
Having a baby with a cleft lip with or without cleft palate and no other known health problems should not change your plans for delivery. Because some of these babies may have breathing problems at birth, a hospital delivery is recommended. You and your healthcare provider will discuss the safest choice for you and your baby.

**What can I expect after my baby is born?**
If your baby does not have any other health problems, he or she will be cared for like a typical newborn. Your baby may need to go to the Newborn Critical Care Center (NCCC) if he or she has any problems, such as problems breathing, after delivery. A pediatrician will examine your baby before you leave the hospital.

A pediatric surgical specialist may also examine your baby.

**Will my baby have feeding problems?**
Most babies with cleft lip have few problems with breast or bottle-feeding. Babies with cleft palates, with or without the cleft lip, often have problems with feeding. Special bottles and nipples are available to help your baby. Mother’s milk is important for these babies to lower the risk of frequent infections. Lactation consultants (specialists who help mothers breast feed), nurses, and other hospital staff will help you develop a plan for feeding your baby.

**Where can I get additional information?**
We expect that you will have many more questions about the diagnosis of a cleft lip and/or palate for your baby. We encourage you to speak with your genetic counselor and other healthcare providers about any other questions you may have.

Additional information and resources
- Cleft Palate Foundation: [www.cleftline.org](http://www.cleftline.org) 1-800-24-CLEFT.
- March of Dime Birth Defects Foundation: [www.marchofdimes.com](http://www.marchofdimes.com) 1-888-MODIMES

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