April 1, 2005                      N.L: 05-0405
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TO: ALL COUNTY CALIFORNIA CHILDREN’S SERVICES (CCS)
ADMINISTRATORS AND MEDICAL CONSULTANTS, CHILD HEALTH
AND DISABILITY PREVENTION (CHDP) PROGRAM DIRECTORS AND
DEPUTY DIRECTORS, AND STATE CHILDREN’S MEDICAL SERVICES
(CMS) STAFF

SUBJECT: AUTHORIZATION OF DIAGNOSTIC SERVICES FOR INFANTS
REFERRED BY THE CALIFORNIA NEWBORN SCREENING PROGRAM

I.  Background

The California Newborn Screening Program currently screens more than 500,000
newborns annually for phenylketonuria (PKU), galactosemia, primary congenital
hypothyroidism, and hemoglobinopathies including sickle cell disease. All the
conditions for which the California Newborn Screening Program screens are CCS-
eligible. Over 99 percent of the babies born in California are tested. Through this
mandatory screening program, about 500 newborns with these disorders are identified
and treated every year. When these disorders are diagnosed late or left undiagnosed,
serious disabilities and even death can result.

Annually, we estimate that approximately 75 referrals are made to metabolic centers
(PKU & galactosemia), 175 to sickle cell disease/hematology centers, and 700 to
endocrine centers for diagnostic evaluations throughout the state.

II.  Policy

A. CCS shall issue an authorization to a CCS-approved Special Care Center
(SCC) to perform a diagnostic evaluation on ALL infants referred by the
California Newborn Screening Program. The California Newborn
Screening Program staff will identify the SCC to which the infant will be
referred. The SCC may be metabolic, endocrine, sickle cell or hematology
depending on the screening result.
B. These authorizations shall be issued within five working days of receipt of the referral.

C. Issuance of this authorization for diagnostic services shall be done upon receipt of a CCS Request for Service form and a signed CCS application. Authorizations shall be issued without a signed CCS application for infants who have full scope no share of cost Medi-Cal or who are Healthy Families subscribers.

D. The CCS $20 assessment fee is to be waived for these services.

III. Policy Implementation

A. Authorizations shall include the following information:

Claims for services provided to children with other third-party coverage must be submitted to the insurance carrier or health maintenance organization prior to billing either the CCS or Medi-Cal program for the services. A denial of payment from the third-party payer must accompany the claim.

B. For infants whose diagnostic evaluation confirms the presence of a metabolic, endocrine or hematologic disorder, the CCS program shall initiate the steps to determine eligibility for ongoing treatment services.

1. Authorizations for treatment services shall be issued to CCS approved Metabolic, Endocrine, Sickle Cell or Hematology SCCs for infants who have full scope, no share of cost Medi-Cal.

2. Families of other infants must complete program eligibility requirements prior to the issuance of treatment authorizations.

On or before August 1, 2005, the California Newborn Screening Program will be expanded to include congenital adrenal hyperplasia and a large number of inborn errors of metabolism. A subsequent numbered letter will address these conditions and provide direction for authorizations.
If you have any questions regarding the above policy, please contact your Regional Office Medical Consultant.

Original Signed by Marian Dalsey, M.D., M.P.H.

Marian Dalsey, M.D., M.P.H., Acting Chief
Children's Medical Services Branch

Enclosure
Overview of the Genetic Disease Branch Newborn Screening Program

The Newborn Screening Program begins with the education of prenatal care providers and hospital staff. The prenatal care providers distribute a copy of the informational booklet, *Important Information for Parents about the Newborn Screening Test*, to women during their pregnancy. This booklet describes mandatory newborn screening. Because some women do not receive prenatal care, the booklet is also provided to women upon admission to a licensed perinatal health facility for delivery. The actual sample of newborn blood is obtained after 12 hours of age and before the sixth day of life by a heel-stick. The blood sample is collected on special filter paper, and mailed to the pre-assigned regional state contract laboratory for testing. These laboratories process the specimen and enter demographic data and test results on terminals linked to a Genetic Disease Branch central computer in Richmond, California. A computer-generated written report of all tests, referred to as a "result mailer" is mailed to the hospital that collected the specimen. Another copy is mailed to the baby’s physician as listed by the hospital of birth on the Newborn Screening Specimen Collection Form (also referred to as the “test request form” or TRF).

Because of the urgency for treatment, all initial positive test results for PKU, galactosemia and primary congenital hypothyroidism are immediately reported by telephone by the NAPS lab to assigned newborn screening coordinators. These coordinators are located at one of the seven state-funded Area Service Centers (ASC) and are linked to the state central computer in Richmond, CA. The State Genetic Disease Laboratory (GDL) reviews and releases hemoglobin results. Potentially clinically significant hemoglobinopathies and other initial positive results are reported on a daily "Interesting Case Report" to the Centers. The coordinators from the Centers immediately telephone the newborn’s physician to provide interpretation of the test and explain necessary follow-up. Medical consultants and specialists are also available to provide additional information and consultation when necessary.

If the initial tests are not within normal limits, a second blood sample called a "recall specimen" is collected (see attached flow chart). The recall specimens for phenylketonuria are sent to GDL for testing. Recall specimens for galactosemia are sent to a state-contract galactosemia confirmation laboratory at Children’s Hospital of Los Angeles, while those for hemoglobin disorders are sent to the state hemoglobin reference laboratory at Children’s Hospital and Research Center at Oakland. In the past, filter paper recall specimens for TSH were also sent to GDL for testing. As of February 2005, the state laboratory will no longer conduct recall testing on filter paper blood specimens for primary congenital hypothyroidism, and providers will be required to have a venous blood specimen collected/tested at a private California licensed laboratory (including hospital laboratories). Test results will then be reported to the NBS Program via the assigned (ASC).

The NBS coordinators notify the newborn's physician of the recall test results. Because primary care physicians are often unfamiliar with these rare disorders, the coordinator will assist the provider in referring a family to a California Children's Services (CCS)-approved metabolic, endocrine or sickle cell disease
center/hematology center for specialized diagnosis and treatment. The comprehensive
team approach to care offered at the Centers is particularly important in treatment of
these disorders.

The coordinators also directly contact the parents of the newborn by telephone to
discuss the need for confirmatory testing or a diagnostic evaluation. After contacting
the parents, the coordinator faxes a referral form to the appropriate CCS office
requesting an authorization for diagnostic evaluation at a CCS approved center, or in
certain circumstances, a CCS paneled specialist (e.g. pediatric endocrine, pediatric
hematologist) not affiliated with a center. The coordinator schedules the appointment
at the center and will send the parents a letter to the mother containing information
about CCS and CCS application form, with instructions for submission and the date of
the appointment. It is anticipated that this process will expedite the referral and the
care of the newborn. Once care is authorized, the center will make the arrangements
for any further confirmatory/diagnostic testing and develop if needed the treatment
plan.