



UCSF Benioff Children's Hospitals

For more information or to refer a patient, please contact us:





UCSF Fetal Treatment Center

HyDROPS Team

1-800-RX-FETUS (1-800-793-3887)

hydrops@ucsf.edu

Teresa Sparks, MD, MAS
Mary Norton, MD
Billie Lianoglou, LCGC
Tippi MacKenzie, MD
Stephan Sanders, MD, PhD
Renata Gallagher, MD

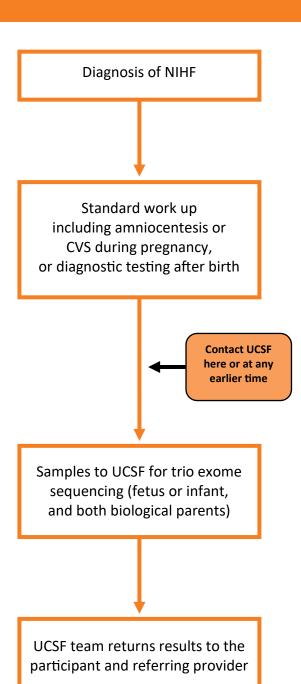


- **Hy**drops:
- **D**iagnosing &
- **R**edefining
- Outcomes with
- Precision
- <u>S</u>tudy



Our research contributes novel information about genetic disorders that cause Non-Immune Hydrops Fetalis.

This enables more accurate **counseling**, **individualized antenatal care**, and **anticipation of neonatal needs**.



Enrollment

The UCSF team can consent and enroll patients locally at UCSF or remotely via video or telephone.

Inclusion Criteria

Living or demised fetus or infant with nondiagnostic karyotype and/or microarray, and one or more of the following:

- Non-immune hydrops fetalis
- Single abnormal fetal fluid compartment (such as isolated ascites)
- Cystic hygroma
- Nuchal translucency of ≥ 3.5 mm

Exclusion Criteria (any)

- Alloimmunization
- Twin-twin transfusion syndrome

Approach to Testing

- Trio exome sequencing on fetus/infant and (ideally) both biological parents
- If patient declines diagnostic testing during pregnancy, infant or other tissue sample may be sent to UCSF
- Exome sequencing at no cost to families
- Multidisciplinary panel review and classification of genetic variants
- Turnaround time is 2 4 weeks for ongoing pregnancies and live births
- Our CLIA-approved lab issues a formal report to the patient and ordering provider

Clinical Data

Our team will request a limited amount of clinical information, such as results of karyotype and/or microarray and ultrasound findings.

How to Refer a Patient

hydrops@ucsf.edu or 1-800-RX-FETUS
See insert for recommended work up of NIHF.

Send Us a Specimen

- Our team can assist with sample transfer
- Saliva kits shipped directly to biological parents with pre-paid return to UCSF
- Only one sample type needed for the fetus/ infant as well as each biological parent

Fetal: Cultured amniocytes, chorionic

villi, cord blood, products of conception, extracted DNA, or

other tissue

Neonatal: Blood, buccal/saliva, skin, or

other tissue

Parental: Blood or saliva





Work up of Non-Immune **Hydrops Fetalis (NIHF)**

Adapted from SMFM Clinical Guideline, 2015

Ultrasound including fetal echocardiogram & MCA Dopplers; type and screen; MCV*; KB Fetal anemia NO YES Amniocentesis or fetal **Amniocentesis** blood sample if IUT

Send for:

- Karvotype and/or microarray;
- +/-PCR for CMV. toxoplasmosis, parvovirus

If karyotype and/or microarray are nondiagnostic, consider:

- RASopathy panel
- Other targeted testing if high clinical suspicion
- Exome sequencing

rev. 2-2020

Send for:

- Karvotype and/or microarray;
- PCR for CMV. toxoplasmosis, parvovirus

If evaluations are nondiagnostic, consider:

- Targeted testing for hereditary anemias if clinical suspicion
- Exome sequencing

* Low MCV: test for alpha thalassemia

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