

Pediatric Medical Genetics

PE 235.99

General Information

COURSE CHAIR: Pedro Sanchez, MD

STUDENT COORDINATOR: Calvin Lee PHONE: 310-423-4780 (option 1) or 310-423-8455 E-MAIL: <u>Calvin.Lee2@cshs.org</u>

STUDENTS / PERIOD: max 1

DURATION: 3 weeks Please contact the Student Program Coordinator for other arrangements

Electives at Cedars-Sinai start in May. Please visit our <u>VSLO</u> catalog for dates and to submit your application.

Description

The student will obtain a broad exposure to the clinical aspects of medical genetics, especially in regard to the diagnostic approach to genetic disease, dysmorphology, deep-phenotyping genetic counseling, prenatal diagnosis and targeted treatment of genetic diseases. Students will be involved in the work up and management of inpatient genetic consultations and will see out-patients in genetic clinic. Exposure to different multidisciplinary clinics can be tailored to student's needs and interests.

(craniofacial, skeletal dysplasia, neurofibromatosis, neurogenetics, geneticendocrine eXemplarY kids (45X, 47XXY, 47XYY,), Angelman Syndrome and others)

Course Objectives

- 1. Navigating the difficult conversation: Counseling techniques and the approach to the patient and family with hereditary diseases
- Deeper dive into data: Historical, physical, and laboratory evaluation of the patient with congenital anomalies, dysmorphic features, and developmental concerns.
- Understanding the spectrum, strength and limitations of genetic and genomic tests: from pre-conception (carrier, PGT-A, PGT-M), Prenatal diagnosis, (NIPT, amniocentesis and chorionic villus sampling) and postnatal options.
- 4. Identify helpful resources and tools to re-interpret laboratory reports
- 5. Integration of genetics into other subspecialties: Neurogenetics, cardiogenetics, endocrine and the ICU setting

Student Experiences

COMMON PROBLEMS/DISEASES

- Syndrome delineation and dysmorphology
- Approach to congenital anomalies
- Whole exome & Genome sequencing
- Chromosomal microarray analysis
- Prenatal diagnosis
- Carrier detection/screening
- Cardiovascular genetics
- Genetic counseling

INPATIENT: 10% PRIMARY CARE: 90%

CONSULTATION: 100%

CLOSE CONTACT WITH

- Full-Time Faculty
- Clinical Faculty
- Fellows
 Other: Genetic Counselors



Pediatric Medical Genetics

Typical Schedule

Approx. # of Patients Evaluated/Week by Student	8
Approx. # of Patients Eval./Each Week by Service	31
Typical Monday	9:00 am - 11:00 am - Genetics Training Program Lecture Series 1:00 pm - 3:00 pm - Chart Review for Clinics 3:00 pm - 4:00 pm – Lecture/ Case Review/ Prenatal Conference
Typical Tuesday	9:00 am – 9:30am case conference 9:00 am - 2:00 pm - Medical Genetics Clinic 2:00pm - 4:00pm- Plagiocephaly Clinic
Typical Wednesday	9:00 am-10am Medical Genetics Clinic 11:00am-12 perinatal/MFM conference 4 th floor 1 pm- 3:00 pm - Medical Genetics Clinic
Typical Thursday	8:00 am - 2:00 pm - Medical Genetics Clinic
Typical Friday	8:00am- 12:00pm - OCC Friday Results Clinic
On-Call Schedule	NONE
Weekend Activities	NONE
Additional Comments/Special Requirements	NONE