



Fetal Care Center of Southern California Referral Request Form

Fetal Center Appointment: **714.509.3696**

Fax: **714.509.3697**

Fetal Center Information: **833.OC.FETAL**

Thank you for referring your patient to **The Fetal Care Center of Southern California!**

Patient Information

Patient Name: _____ Date of Birth: _____ / _____ / _____

EDD: _____ / _____ / _____ Gravida: _____ Para: _____ Gestational Age: _____

Patient Phone: _____ Insurance: _____

Reason for Referral (Fetal Diagnosis): _____

Please see page 2 for an alphabetical list of Fetal Diagnosis examples and ICD-10 Codes

Services Requested: Fetal Care Center Comprehensive Visit is recommended for all patients and Nurse Navigators are available to assist for a customized experience

Fetal Care Center Comprehensive Visit

- Obstetric Ultrasound
 - CPT code 76811
- Fetal Echocardiogram
 - CPT codes 99205, 93325, 76825, 76827
- MFM Consultation
 - CPT code 99205
- Neonatology Consultation
 - CPT code 99205
- Pediatric Subspecialty Consultation
 - Indicate condition-specific subspecialist in column to the right
 - CPT code 99205
- Fetal MRI
 - CPT code 74712

Condition-Specific Pediatric Subspecialists

- Pediatric Cardiology
- Pediatric Cardiovascular Surgery
- Pediatric Ear Nose & Throat (ENT)
- Pediatric General and Thoracic Surgery
- Pediatric Nephrology
- Pediatric Neurology
- Pediatric Neurosurgery
- Pediatric Orthopedic Surgery
- Pediatric Plastic Surgery
- Pediatric Urology
- Palliative Care
- Other: _____

Use CPT code 99205 for each consultation requested

To expedite appointment scheduling, please provide the following by fax to 714-509-3697

- Complete Prenatal Records
- Patient demographics
- Insurance information
- Authorization including all appropriate CPT codes

Referring Provider Name: _____ Phone: _____ Fax: _____

Provider Address: _____ City: _____ Zip: _____

Provider Signature: _____ **Date:** _____

Please contact our office if you have questions about authorizations

ICD-10 code O28.3: abnormal ultrasonic finding on antenatal screening of mother
ICD-10 code O35.8XX1 maternal care for other (suspected fetal abnormality and damage)

Agenesis of corpus callosum	Q04.0	Hydronephrosis/hydroureter	Q62.0
Amniotic band syndrome	PO2.8	Hydrops fetalis	P83.2
Anorectal malformations including imperforate anus	Q42.3	Hypoplastic left heart syndrome	Q23.4
Aortic and pulmonary valve stenosis	Q23.0/Q22.1	Isoimmunization and other hematologic disease	O36.1
Arachnoid cysts		Large chorangioma	O43.893
Atrioventricular canal defects	Q21.2	Laryngeal atresia	Q31.8
Bladder exstrophy	Q64.10	Limb-length discrepancies	M21.7
Brain and spinal vascular malformations	varies	Lissencephaly	Q04.3
Brain cysts	O35.0XX0	Lower urinary tract obstruction	
Brain tumors	O35.0XX0	Lymphatic malformations	Q89.9
Bronchial atresia, bronchopulmonary sequestration and congenital lobar emphysema	Q32.4/Q33.2	Malrotation and volvulus	Q43.3
Bronchogenic cyst	Q33.0	Megacystis and megacystis microcolon intestinal hypoperistalsis syndrome	Q64.79 Q43.8
Cardiac masses and tumors	O36.8390	Megalourethra	Q64.79
Cardiomyopathy (enlarged or thickened heart)		Micrognathia	M26.09
Cervical teratoma	D48.7	Multicystic dysplastic kidney/polycystic kidneys	Q61.4
Choledochal cyst	Q44.4	Meningomyelocele/myelomeningocele (spina bifida)	Q05.9
Chromosome anomalies including Trisomy 13, Trisomy 18 and Trisomy 21 (Down syndrome)	T13: Q91.4 T18: Q91.0 T21: Q90.0	Neural tube defects	Q05.9
Cleft lip and palate (unilateral or bilateral)	Q37.1/Q37.0	Obstructive epulis	Q38.5
Cloacal exstrophy	Q64.12	Omphalocele, including OEIS	Q79.2
Cloacal malformation	Q43.7	Pachygyria	Q04.3
Clubfoot	O66.0	Polydactyly of the hand	Q69.0
Coarctation of the aorta	Q25.1	Posterior urethral valves	Q64.2
Colonic atresia	Q42.9	Proximal focal femoral deficiency	Q68.8
Congenital cytomegalovirus infection	P35.1	Prune belly syndrome	Q79.4
Congenital diaphragmatic hernia (CDH)	Q79.0	Pulmonary atresia	Q25.5
Congenital goiter	E03.0	Renal duplication anomalies	Q63.8
Congenital hemangioma with airway compromise	D18.00	Renal agenesis (bilateral/unilateral)	Q60.1/60.0
Congenital high airway obstruction syndrome (CHAOS)	Q31.8	Sacroccygeal teratoma	D48.0
Congenital pulmonary airway malformation (CPAM)	Q33.0	Schizencephaly	Q04.6
Conjoined twins	Q89.4	Selective fetal growth restriction	O36.5990
Cortical dysplasia	Q04.6	Single ventricle type complex congenital heart disease	Q24.9
Craniosynostosis	Q75.0	Skeletal dysplasia	Q77.9
Dandy-Walker malformation	Q03.1	Small bowel atresia (duodenal, jejunal and ileal)	Q41.0
Double outlet right ventricle	Q20.1	Solitary kidney	Q60.0
Ebstein's anomaly	Q22.5	Spinal deformities	varies
Echogenic bowel	O28.3	Tethered cord	
Encephalocele	Q01.8	Tetralogy of Fallot	Q21.3
Esophageal atresia with or without tracheoesophageal fistula	Q39.0/Q39.1	TRAP sequence	O30.039
Fetal arrhythmias	O36.83	Twin-twin transfusion syndrome	O43.029
Gastroschisis	Q79.3	Ureterocele	Q64.6
Genetic syndromes including Beckwith-Wiedemann and Noonan's syndrome	Q87.3 Q87.19	Ureteropelvic junction obstruction	Q62.11
Head and neck vascular malformations	Q27.9	Vein of Galen malformations	Q28.2
Hirschsprung's disease	Q43.1	Ventricular septal defects	Q21.0
Hydrocephalus	Q03.8	Ventriculomegaly	Q04.8