

## *Medical Genetics*

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**TEST: Oncology - Solid Tumors/Fine Needle Aspirates (FNA)  
Chromosome Analysis**

**Purpose of Test:** To identify acquired chromosome abnormalities detectable by routine cytogenetic analysis. This includes: translocations, deletions, duplications, inversions, and numerical aberrations.

**Required Information:** Please provide indications (ICD-9) for testing on our referral form with each specimen. Accurate testing and interpretation may otherwise be compromised.

### **Solid Tumor/FNA**

- Aseptically place a 0.5 cm or larger tumor biopsy or the FNA in a sterile transport container with 10 mL of sterile medium.
- Alternatively Hank's balanced salt solution, Ringer's solution, RPMI medium or physiologically normal saline can be used.
- Label container with patient's name and specimen type.
- **SEND SPECIMEN REFRIGERATED, DO NOT FREEZE.**

**Adjunct Test:** FISH specific per physician's request  
No additional specimen required.

**Turn Around Time:** 12-18 days

### **CPT Code Information:**

88239	Other tissue culture (solid tumor)
88262	Count 15-20 cells, 2 karyotypes, GTG
88280	Additional karyotypes, each study
88285	Additional cells counted, each study