

# Calcium Sensing Receptor (CASR) Gene Testing Patient Information

The accurate interpretation and reporting of genetic results is contingent upon the reason for referral, clinical information, and family history. To help provide the best possible service, supply the information requested below and send paperwork with the specimen.

## Patient Information

Patient Name <i>(Last, First, Middle)</i>	Birth Date <i>(Month DD, YYYY)</i>	Gender <input type="checkbox"/> Male <input type="checkbox"/> Female
Requesting Physician <i>(Last, First)</i>	Phone	Fax
Genetic Counselor	Phone	Fax

*\*Fax number given must be from a fax machine that complies with applicable HIPAA regulation.*

## Reason for Testing

What is the suspected diagnosis? <input type="checkbox"/> Familial hypocalciuric hypercalcemia (FHH) <input type="checkbox"/> Neonatal severe primary hyperparathyroidism (NSPHPT) <input type="checkbox"/> Autosomal dominant hypocalcemia (ADH) <input type="checkbox"/> Bartter syndrome type V	
Patient presentation of hypercalcemia or hypocalcemia? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	If yes, <input type="checkbox"/> hypercalcemia or <input type="checkbox"/> hypocalcemia
Family history of hypercalcemia or hypocalcemia? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	If yes, indicate hyper or hypo and list affected family members:
Family member with known CASR mutation: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	If yes, indicate exon or nucleotide and list affected family members: