

THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

PATIENT HISTORY FOR MULTIPLE ENDOCRINE NEOPLASIA TYPE 2 (MEN2) TESTING

Patient Name: _____ **Date of Birth:** _____ **Sex:** Female Male
Ordering Provider: _____ **Provider's Phone:** _____
Practice Specialty: _____ **Provider's Fax:** _____
Genetic Counselor: _____ **Counselor's Phone:** _____

Patient's Ethnicity/Ancestry (check all that apply)

African American/Black Asian Hispanic White Other: _____

List country of origin (if known): _____

Does the patient have symptoms? No Yes (check all that apply and describe)

- Medullary thyroid carcinoma (MTC) (age of onset: _____)
 - Bilateral / Unilateral
 - Monoclonal / Multifocal
- Pheochromocytoma (age of onset: _____)
 - Bilateral / Unilateral
- Skeletal abnormalities (describe): _____
- Eye abnormalities (describe): _____
- Neuromas (describe): _____
- Other symptom(s): _____

Laboratory Findings

- Calcitonin..... Abnormal Normal Unknown
- Calcium..... Abnormal Normal Unknown
- Parathyroid hormone..... Abnormal Normal Unknown
- Catecholamines..... Abnormal Normal Unknown

Has the patient had an allogeneic bone marrow or umbilical cord blood transplant?..... No Yes Unknown

Has the patient undergone previous DNA testing?..... No Yes Unknown

If yes, describe the genes, disorder, methodology, and results: _____

Is there any relevant family history?..... No Yes Unknown

If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset:

Has DNA testing been performed for the family member(s)?..... No Yes Unknown

If yes, attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing).

Check the test you intend to order.

- 3004572 Multiple Endocrine Neoplasia Type 2 (MEN2), RET Sequencing:** Full sequencing of the *RET* gene with >95%, >98% and >88-95%, clinical sensitivity for MEN2A, MEN2B, and familial MTC, respectively.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a sequence variant previously identified in a family member; a copy of a relative's lab result is REQUIRED.



For questions, contact an ARUP genetic counselor at 800-242-2787 ext. 2141.