

**THIS IS NOT A TEST REQUEST FORM.**  
**The information below is required to perform OTC deficiency testing.**  
**Please fill out this form and submit it with the test request form or electronic packing list.**

**PATIENT HISTORY FOR ORNITHINE TRANSCARBAMYLASE DEFICIENCY (OTC) TESTING**

**Patient Name** \_\_\_\_\_ **Date of Birth** \_\_\_\_/\_\_\_\_/\_\_\_\_ **Gender**  F  M  
**Physician** \_\_\_\_\_ **Physician Phone** (\_\_\_\_) \_\_\_\_\_ **Practice Specialty** \_\_\_\_\_  
**Genetic Counselor** \_\_\_\_\_ **Counselor Phone #** (\_\_\_\_) \_\_\_\_\_

**PATIENT ETHNICITY (check all that apply)**

- |   |   |  |                                      |
|---|---|--|--------------------------------------|
| <input type="checkbox"/> African American | <input type="checkbox"/> Ashkenazi Jewish | <input type="checkbox"/> Asian           | <input type="checkbox"/> Caucasian   |
| <input type="checkbox"/> Hispanic         | <input type="checkbox"/> Middle Eastern   | <input type="checkbox"/> Native American | <input type="checkbox"/> Other _____ |

**SYMPTOMS of OTC**  No  Yes  Unknown

If yes, check all that apply

- |   |  |   |
|---|--|---|
| <input type="checkbox"/> Hyperammonemia | <input type="checkbox"/> Cyclical vomiting | <input type="checkbox"/> Seizures           |
| <input type="checkbox"/> Lethargy       | <input type="checkbox"/> Coma              | <input type="checkbox"/> Reye-like syndrome |
| <input type="checkbox"/> Encephalopathy | <input type="checkbox"/> Protein aversion  | <input type="checkbox"/> Other _____        |

**LABORATORY FINDINGS**

Plasma amino acids  Normal  Abnormal (result: \_\_\_\_\_)  Not performed  Unknown  
Orotic Acid  Normal  Abnormal (result: \_\_\_\_\_)  Not performed  Unknown  
Ammonia Level  Normal  Abnormal (result: \_\_\_\_\_)  Not performed  Unknown

**FAMILY HISTORY OF OTC DEFICIENCY?**  No  Yes  Unknown

If yes, describe relationship(s) to the patient \_\_\_\_\_

Has DNA testing for OTC been performed for these family member(s)?  No  Yes  Unknown

If yes, please attach a copy of the laboratory result (REQUIRED for familial mutation testing)

**Circle the OTC test you intend to order.**

**2004896 Ornithine Transcarbamyase Deficiency (OTC) Sequencing and Deletion/Duplication:** Sequencing and deletion/duplication analysis of the *OTC* coding regions and intron/exon boundaries. Clinical sensitivity approaches 90%.

**2004901 Ornithine Transcarbamyase Deficiency (OTC) Sequencing:** sequencing of the *OTC* coding regions and intron/exon boundaries. Clinical sensitivity approximately 80%.

**2001961 Familial Mutation, Targeted Sequencing:** Targeted sequencing for an *OTC* mutation previously identified in a family member.

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

Master Label