TEST CHANGE

# Alpha-Iduronidase Enzyme Activity in Leukocytes

2011415, A-I LEUK

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| Specimen Requirements: | |
| Patient Preparation: | |
| Collect: | Yellow (ACD Solution B), lLavender (K2EDTA), or lLavender (K3EDTA). |
| Specimen Preparation: | Transport 3 mL whole blood. (Min: 1 mL) |
| Transport Temperature: | Room temperature (preferredTemperature (Preferred) or rRefrigerated |
| Unacceptable Conditions: | Grossly hemolyzed or heparinized specimens. |
| Remarks: | Additional information is required: Clinical Indication for testing. |
| Stability: | Ambient: 3 days; Refrigerated: 3 days; Frozen: Unacceptable |
| Methodology: | Quantitative Fluorometry |
| Performed: | MonVaries |
| Reported: | 3-10 days |
| Note: | |
| CPT Codes: | 82657 |
| New York DOH Approval Status: | This test is New York DOH approved. |
| Interpretive Data: | |
| Refer to report. Refer to report. | |
| This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes. | |
| Reference Interval: | |
| 12- -65 nmol hydrolyzed/hr/mg protein | |

HOTLINE NOTE: There is a unit of measure change associated with this test. Refer to the Hotline Test Mix for interface build information.