

TEST CHANGE

N-Telopeptide, Cross-Linked, Urine

0070062, NTX

Specimen Requirements:

Patient Preparation: For monitoring therapy, a baseline specimen should be

collected prior to initiation of therapy. Subsequent specimens for comparison should be collected at the same time of day as

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the baseline specimen.

Collect: Second-morning void or 24-hour urine. Refrigerate during

collection. Collect without preservative.

Specimen Preparation: Transfer a 1 mL aliquot of urine from a well-mixed, second-

morning void or 24-hour collection to an ARUP Standard

Transport Tube. (Min: 0.5 mL)

Transport Temperature: Frozen.

Unacceptable Conditions: Specimens contaminated with blood or extensive hemolysis.

Remarks:

Stability: Ambient: 24 hours; Refrigerated: 1 week; Frozen: 4 weeks2

years

Methodology: Quantitative Chemiluminescent Immunoassay

Performed: Tue-Sat

Reported: 1-4 days

Note:

CPT Codes: 82523

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

NTx Units = nM BCE/mM creatinine

A decrease of 30-40% from the NTx baseline after three months of therapy is a typical response to anti-resorptive therapy.

NTx = Cross-linked N-telopeptide of Type I Collagen

BCE = Bone Collagen Equivalent

Reference Interval:

BCE/mM creatinine



Age Male Female 7-9 years 167-578 nM 201-626 nM BCE/mM BCE/mM creatinine creatinine 10-12 years 152-505 nM 173-728 nM BCE/mM BCE/mM creatinine creatinine 13-15 years 103-776 nM 38-515 nM BCE/mM BCE/mM creatinine creatinine 20-144 nM 16-17 years 34-313 nM BCE/mM BCE/mM creatinine creatinine 18 years and 21-83 nM older BCE/mM creatinine 17-94 nM Premenopausal BCE/mM creatinine 26-124 nM Postmenopausal

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