| N -Telopeptide, Cross-Linked, 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 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, secondmorning 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 weeksz 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 $=\mathrm{nM}$ BCE/ $/ \mathrm{mM}$ creatini |  |
| A decrease of $30-40 \%$ from the $N$ anti-resorptive therapy. | baseline after three months of therapy is a typical response to |
| NTx = Cross-linked N -telopeptide <br> $B C E=$ Bone Collagen Equivalent | f Type I Collagen |
| Reference Interval: |  |


| Age |  |  |
| :---: | :---: | :---: |
| 7-9 years | $167-578 \mathrm{nM}$ <br> BCE/mM creatinine | 201-626 nM <br> BCE/mM creatinine |
| 10-12 years | $152-505 \mathrm{nM}$ BCE/mM creatinine | 173-728 nM BCE/mM creatinine |
| 13-15 years | 103-776 nM BCE/mM creatinine | $38-515 \mathrm{nM}$ <br> BCE/mM creatinine |
| 16-17 years | 34-313 nM BCE/mM creatinine | 20-144 nM <br> BCE/mM creatinine |
| 18 years and older | 21-83 nM BCE/mM creatinine |  |
| Premenopausal |  | 17-94 nM BCE/mM creatinine |
| Postmenopausal |  | 26-124 nM <br> BCE/mM creatinine |

