### MEDICARE COVERAGE OF LABORATORY TESTING

Please remember when ordering laboratory tests that are billed to Medicare/Medicaid or other federally funded programs, the following requirements apply:

1. Only tests that are medically necessary for the diagnosis or treatment of the patient should be ordered. Medicare does not pay for screening tests except for certain specifically approved procedures and may not pay for non-FDA approved tests or those tests considered experimental.
2. If there is reason to believe that Medicare will not pay for a test, the patient should be informed. The patient should then sign an Advance Beneficiary Notice (ABN) to indicate that he or she is responsible for the cost of the test if Medicare denies payment.
3. The ordering physician must provide an ICD-10 diagnosis code or narrative description, if required by the fiscal intermediary or carrier.
4. Organ- or disease-related panels should be billed only if all components of the panel are medically necessary.
5. Both ARUP- and client-customized panels should be billed to Medicare only if every component of the customized panel is medically necessary.
6. Medicare National Limitation Amounts for CPT codes are available through the Centers for Medicare & Medicaid Services (CMS) or its intermediaries. Medicaid reimbursement will be equal to or less than the amount of Medicare reimbursement.

The CPT Code(s) for test(s) profiled in this bulletin are for informational purposes only. The codes reflect our interpretation of CPT coding requirements, based upon AMA guidelines published annually. CPT codes are provided only as guidance to assist you in billing. ARUP strongly recommends that clients reconfirm CPT code information with their local intermediary or carrier. CPT coding is the sole responsibility of the billing party.

The regulations described above are only guidelines. Additional procedures may be required by your fiscal intermediary or carrier.

### Summary of Changes by Test Name

<table>
<thead>
<tr>
<th>Hotline Page #</th>
<th>Test Number</th>
<th>Name Change</th>
<th>Methodology</th>
<th>Performed/Reported Schedule</th>
<th>Specimen Requirements</th>
<th>Reference Interval</th>
<th>Interpretive Data</th>
<th>Note</th>
<th>CPT Code</th>
<th>Component Change</th>
<th>Other Interface Change</th>
<th>New Test</th>
<th>Inactive</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0099174</td>
<td>Lipoprotein (a)</td>
<td></td>
<td></td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**0099174  Lipoprotein (a) LIPO A**

**Specimen Required:**
- **Collect:** Serum Separator Tube (SST) or Plasma Separator Tube (PST). Also acceptable: Green (Lithium Heparin), Lavender (EDTA), or Pink (K$_2$EDTA).
- **Specimen Preparation:** Allow specimen to clot completely at room temperature. Separate from cells ASAP or within 2 hours of collection. Transfer 1 mL serum or plasma to an ARUP Standard Transport Tube. (Min: 0.5 mL)
- **Storage/Transport Temperature:** Frozen.
- **Unacceptable Conditions:** Body Fluids.
- **Stability (collection to initiation of testing):** After separation from cells: Ambient: 8 hours; Refrigerated: 2 days; Frozen: 3 months