ARUP TEST REQUEST FORM

IRB Registration Required to Order Test

ARUP Client Number		ARUP Client Name
Patient Name (Last) (First) (M.I.)		
Patient I.D. Number		Lab I.D. Number
Date of Birth _____ mo / day / yr	Sex	Physician Name (Last) (First) (M.I.) Physician Contact Number
Specimen Collection Date: _____ Time: _____ [ ] AM [ ] PM mo / day / year		

0081284 [ ] Soluble Mesothelin Related Peptides (MESOMARK®)

Soluble Mesothelin Related Proteins MESOMARK is performed exclusively at ARUP Laboratories. It is an enzyme-linked immunosorbent assay for the quantitative measurement of soluble mesothelin related peptides (SMRP) in human serum. Measurement of SMRP may aid in the monitoring of patients diagnosed with epithelioid or biphasic mesothelioma. Epidemiologic studies have established exposure to asbestos fibers as the primary cause of malignant mesothelioma.

Because MESOMARK is a test approved by the FDA as a Humanitarian Use Device, testing must be ordered using the following procedure(s):

1. **The ordering physician must register with the Internal Review Board (IRB) for MESOMARK testing. Go to fdi.com/mesomark to obtain IRB certification online.**

2. **The test should be ordered using the ARUP Test Request Form. The full name of the ordering physician must be included on the ARUP form to ensure timely testing of the sample. Samples submitted with incomplete information may delay specimen testing.**

3. **ARUP does not accept samples directly from physician offices. ARUP only accepts samples from established clients. To send a sample to ARUP, contact your local hospital/reference laboratory to determine if they are an ARUP client and can send the sample. If they cannot send the sample to ARUP, contact ARUP Client Services at (800) 522-2787 to find an ARUP client in your area.**

4. **Specimen Collection Instructions:**
   Collect: One 4 mL SST or red top tube.
   Transport: 0.2 mL serum, frozen (Min: 0.1 mL) Submit specimen in an ARUP Standard Transport Tube.
   Remarks: CRITICAL FROZEN.

5. **Forms and information about MESOMARK testing, and IRB registration, may be accessed at fdi.com/mesomark.**

Humanitarian Device. Authorized by Federal Law for use as an aid in the monitoring of patients diagnosed with biphasic or epithelioid mesothelioma. The effectiveness of this device for this use has not been demonstrated.

THIS BOX FOR ARUP USE ONLY

Qty_____ RT R F ID# _____

Master Label