NEWS RELEASE

Date: 02 January 2018
Arlington Scientific, Inc. (ASI)
Springville, Utah
Contact: Mike LaDow, Director Sales and Marketing
Tel: 801-489-8911
Email: mladow@arlingtonscientific.com

For Immediate Release

Arlington Scientific, Inc. (ASI) Receives US Food and Drug Administration CBER Clearance for the ASI Automated RPR test for Syphilis; for use on the ASI Evolution. FDA 510(k) BK170114

The ASI Automated RPR (rapid plasma reagin) Test for syphilis is a qualitative nontreponemal flocculation test for the detection of reagin antibodies in human serum and plasma as a screening test for serological evidence of syphilis for use on the ASI Evolution analyzer. The test is intended for use in screening blood donors.

- The World's ***first and only*** FDA CBER Cleared Fully Automated Nontreponemal Syphilis screening system for screening human serum and plasma.
- The ability to do automated syphilis testing using the Centers for Disease Control (CDC) recommended algorithm for syphilis screening.
- ***Automation rules*** – Full walk-away platform provides lowest automated cost per test and dramatic labor savings.

FDA CBER Link:

https://www.fda.gov/BiologicsBloodVaccines/BloodBloodProducts/ApprovedProducts/SubstantiallyEquivalent510kDeviceInformation/ucm590922.htm

ASI Evolution Link:
http://www.arlingtonscientific.com/asi-evolution.html

About Arlington Scientific, Inc.

Arlington Scientific, Inc., is a leading global medical technology company that develops, manufactures and sells *in vitro* diagnostics, medical devices, diagnostic analyzers and blood donor lounges. Throughout its 30-year history, ASI remains focused on enhancing the quality and speed of diagnosing infectious diseases. As the "Syphilis Authority", ASI has a particular emphasis and expertise in syphilis diagnostic reagents and innovative analyzers.

CONTACT:
Mike LaDow, Director of Sales and Marketing
mladow@arlingtonscientific.com
http://www.arlingtonscientific.com