



NEWS RELEASE

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Arlington Scientific, Inc. (ASI)

Springville, Utah

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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/fda-cleared-asi-evolution%C2%AE.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.

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