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Accelerate Diagnostics Completes Clinical Data Collection, Will Present Data at the William Blair 36th Annual Growth Stock Conference

TUCSON, Ariz., June 14, 2016 (GLOBE NEWSWIRE) -- Accelerate Diagnostics, Inc. ("Accelerate") announced today the completion of clinical data collection intended for submission to the FDA in conjunction with its *de novo* 510k request for premarket clearance of the Accelerate Pheno™ system and Accelerate Pheno™ BC kit for positive blood cultures. An overview of the data collected will be included in the previously announced presentation at the William Blair 36th Annual Growth Stock Conference. The Accelerate Pheno™ system and Accelerate PhenoTest™ BC kit were formerly referred to in generic terms as the Accelerate ID/AST System and Blood Culture Assay Kit.

The above-mentioned overview will be accessible on the company's website at ir.axdx.com concurrently with the scheduled conference presentation.

About Accelerate Diagnostics, Inc.

Accelerate Diagnostics, Inc. ("Accelerate Diagnostics,") (Nasdaq:AXDX), is an *in vitro* diagnostics company dedicated to providing solutions for the global challenge of antibiotic resistance and hospital acquired infections. The company's fully automated ID/AST system, Accelerate Pheno™, and direct from positive blood culture test, Accelerate PhenoTest™ BC, utilize proprietary molecular and phenotypic detection technologies which have the potential to substantially reduce the time to antimicrobial susceptibility results while achieving high sensitivity and specificity. For more information about Accelerate Diagnostics, visit www.acceleratediagnostics.com.

The "ACCELERATE DIAGNOSTICS" and "ACCELERATE PHENO" and "ACCELERATE PHENOTEST" logos and marks are trademarks or registered trademarks of Accelerate Diagnostics, Inc.

Forward-Looking Statements

Certain of the statements made in this press release are forward looking, such as those, among others, about our projections as to when certain key business milestones may be achieved, including completion of the trial intended to support marketing authorization by the FDA of the Accelerate Pheno™ system and Accelerate PhenoTest™ BC kit for positive blood cultures, the commercial launch of the Accelerate Pheno™ system and Accelerate PhenoTest™ BC kit for positive blood cultures, the potential of our technology, the growth of the market, our estimates as to the size of our market opportunity and potential pricing, our competitive position and estimates of time reduction to results, and our future development plans and growth strategy. Actual results or developments may differ materially from those projected or implied in these forward-looking statements. Information about the risks and uncertainties faced by Accelerate Diagnostics is contained in the section captioned "Risk Factors" in the company's most recent Annual Report on Form 10-K, filed with the Securities and Exchange Commission on March 9, 2016. In addition, the company's forward-looking statements could be affected by general industry and market conditions. Except as required by federal securities laws, the company undertakes no obligation to update or revise these forward-looking statements to reflect new events, uncertainties or other contingencies.

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