



## Group A Strep results in an actionable timeframe. No culture confirmation required.

Today's Strep A diagnostic landscape consists of rapid tests that are fast and easy, but lack sensitivity; or culture methods that are sensitive, but complex and slow. Solana delivers results you can trust in an actionable timeframe, without the need for culture back up.

# Prompt initiation of patient management and antimicrobial therapy

- Results in just 30 minutes
- High volume throughput capabilities
   Batch up to 12 tests per run

#### No Culture Back Up

- Gold standard sensitivity
- Confidence in your prescription decision

#### Ease and Affordability

- Reduced hands-on time vs. culture
- Simple procedure
- Competitive reagent pricing
- Flexible instrument placement options

The Accurate. Sustainable. Molecular **Strep** Solution.

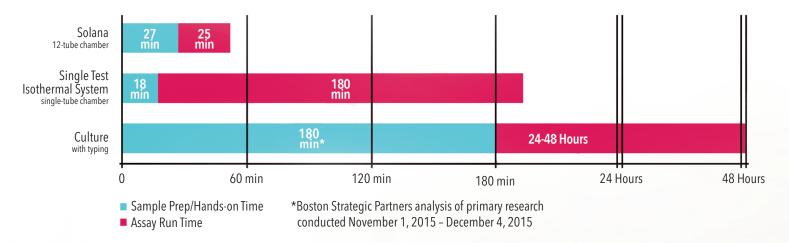
Solana

Solana Strep Complete Assay run with the Solana bench top instrument a mere 9.4" x 9.4" x 5.9"

### Procedure

#### **Amplification** Sample Preparation **Detection** 50 µL 50 µL 95°C 5 Min. Review results. Optional print or export Dilution Reaction Close lid tightly and Insert into Solana, input sample ID and Tube Tube proceed to next step. initiate program

## Time to 12 results



## Clinical performance

#### Performance Results of Solana GAS Assay for Group A β-hemolytic Streptococcus

Composite Culture						
		POS	NEG	Total	Sensitivity	98.2%
Solana GAS	POS	220	24*	244	Specificity	97.2%
	NEG	4**	833	837		4144/10
	Total	224	857	1081		10

<sup>\*</sup>Of the twenty-four (24) discordant specimens, sixteen (16) of these specimens were positive for GAS when tested with an additional FDA-cleared molecular device, eight (8) were negative.



SSM301302EN00 (03/17)



Solana GAS Assay – 48 Test Kit: Cat. #23043100 Quidel Molecular Strep A+G Control Set – 20-Reaction Kit: Cat. #23043150 Solana GAS Startup Kit: Cat. #23043101

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#### In the United States:

For customer service, call 1-800-640-0640 To fax an order, use 1-800-290-0290 To order online: www.fisherhealthcare.com



<sup>\*\*</sup>Of the four (4) discordant specimens, three (3) were negative when tested with an additional FDA-cleared molecular device.

Refer to Package Insert for additional performance claims.