BIOMÉRIEUX

SCANRDI®

When you Need to Know Now
Bring your Sterility Testing to the Next Level

Your Ally in Advancing Quality

PIONEERING DIAGNOSTICS

Help ensure sterility and patient safety.

SCANRDI® enables same-shift detection of microbial contaminants in filterable drug products. The US Food and Drug Administration (FDA), European Medicines Agency (EMA), and other regulatory bodies support the use of validated Rapid Microbiological Methods (RMM) for sterile product release - especially in the case of shelf-life products, where a rapid result is more suitable than a compendial 14-day test.

4 Steps in 4 Hours

ULTRA-RAPID

- Same-shift results
- Release your product faster

ACCURATE

- Tested & proven technology
- Make decisions with confidence

COMPLIANT

- Full validation support: URS completion guide, IOQ guide, primary validation guide, suitability guide and Drug Master File (DMF)
- 21 CFR part 11 enabled software

Improve productivity and compliance.

The right Solution for your Lab & Business

PROVEN TECHNOLOGY

- Long history of use for final product sterility, in-process bioburden control & water testing
- Down to 1 microbe/filtrable volume
- >200 compatible products

BROAD SPECIFICITY

- Universal fluorescent labeling
- Bacteria, yeast, and mold detection

>200 compatible products*	
	Antibiotics
	Isotonic solutions
	Chemotherapy drugs
	Hormones
	Anti-inflammatories
	Anti-coagulants
	Analgesics
	Diuretics
	Vitamins

*non-exhaustive list

INCREASE YOUR MARKET COMPETITIVENESS

- Safely release your short shelf-life & generic products faster
- Optimize your distribution logistics with just-in-time inventory management of stock and choose your most affordable shipping option

CONTROL YOUR MANUFACTURING PROCESS

- Rapid in-process, bioburden and sterility results give you confidence that your production is operating properly
- Facilitate your investigation process to quickly return to normal operations
- Eliminate lab bottlenecks

BE CONFIDENT IN YOUR COMPLIANCE

- Ensure your electronic compliance with 21 CFR part 11 enabled software, barcode scanning traceability, several approval levels and a full audit trail
- Feel reassured in your validation process with support from bioMérieux including URS completion guide, IOQ guide, primary validation guide, suitability guide and Drug Master File (DMF)

Services: Feasibility, Validation, and Instrument Support

With bioMérieux services, your productivity and product quality is our priority. Our mission is to provide the pharmaceutical industry with rapid, accurate, and cost-effective solutions to ensure smooth implementation and maximizing your return on investment.

Feasibility Studies

Quickly confirm the compatibility of your product and method

Validation Services

Expedite implementation of your investment by reducing validation time and expense

Instrument Service Plans and Support

Maximize the uptime of your instruments and ensure continuous operations

Smart Remote Support Solutions

VILINK® ensures the continuity of your operations

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