

**K911419 SENSITITRE ARIS MODULE**Jul 1, 1992  
457 days to decisionK911419 · Product code: **LRG** · Microbiology  
Source: <https://www.510kdatabase.net/k911419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument For Auto Reader & Interpretation Of Overnight Suscept. Systems (LRG)
Date received	Apr 1, 1991
Decision date	Jul 1, 1992
Days to decision	457 days
Third-party review	No

**APPLICANT**

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Company	<b>Sensititre , Ltd.</b>
Location	Salem, NH, US
Contact	LAURA BASISTA
510(k) history	3 submissions · 3 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911419/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026