

**K821659 RESPIRACULT-STREP DEEP BED CULTURE PAD**Aug 5, 1982  
62 days to decisionK821659 · Product code: **JSH** · Microbiology  
Source: <https://www.510kdatabase.net/k821659/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culture Media, Non-selective And Differential (JSH)
Date received	Jun 4, 1982
Decision date	Aug 5, 1982
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Orion Diagnostica, Inc.</b>
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1980-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k821659/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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