

**K810767 RPMI 1640 MEDIUM**May 8, 1981  
46 days to decisionK810767 · Product code: **KIT** · Chemistry  
Source: <https://www.510kdatabase.net/k810767/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media And Components, Synthetic Cell And Tissue Culture (KIT)
Date received	Mar 23, 1981
Decision date	May 8, 1981
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Dutchland Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	75 submissions · 75 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k810767/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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