

**K831680 VERO CELLS**Jul 6, 1983  
43 days to decisionK831680 · Product code: **KIR** · Pathology  
Source: <https://www.510kdatabase.net/k831680/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cells, Animal And Human, Cultured (KIR)
Date received	May 24, 1983
Decision date	Jul 6, 1983
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Viomed Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	25 submissions · 25 cleared · 1983-1996

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

Device record: <https://www.510kdatabase.net/k831680/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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