

K843250 BHK-21Sep 6, 1984
17 days to decisionK843250 · Product code: **KIR** · Pathology
Source: <https://www.510kdatabase.net/k843250/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cells, Animal And Human, Cultured (KIR)
Date received	Aug 20, 1984
Decision date	Sep 6, 1984
Days to decision	17 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k843250/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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