

K883369 L-15 MEDIUM, CATALOG NO. 210-4004Aug 29, 1988
20 days to decisionK883369 · Product code: **KIT** · Pathology
Source: <https://www.510kdatabase.net/k883369/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media And Components, Synthetic Cell And Tissue Culture (KIT)
Date received	Aug 9, 1988
Decision date	Aug 29, 1988
Days to decision	20 days
Third-party review	No

APPLICANT

Company	J R Scientific
Location	Woodland, CA, US
Contact	G KONISH,PHD
510(k) history	132 submissions · 132 cleared · 1987-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883369/>; Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026