FDA 510(k) Premarket Notification

K883369 L-15 MEDIUM, CATALOG NO. 210-4004

CLEARED

Aug 29, 1988
20 days to decision

K883369 · 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