FDA 510(k) Premarket Notification

K810767 RPMI 1640 MEDIUM

CLEARED

May 8, 1981
46 days to decision

K810767 · Product code: **KIT** · Chemistry
Source: https://www.510kdatabase.net/k810767/

SUBMISSION DETAILS

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

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