FDA 510(k) Premarket Notification

CLEARED

K844164 REACTIFS, IBF ULTROGELS ACA & ULTROGELS A

Dec 10, 1984
46 days to decision

K844164 · Product code: **DII** · Chemistry
Source: https://www.510kdatabase.net/k844164/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Columns, Glc (DII)
Date received	Oct 25, 1984
Decision date	Dec 10, 1984
Days to decision	46 days
Third-party review	No

APPLICANT

Company	**Lkb Instruments, Inc.**
Location	Mchenry, IL, US
Contact	ED ZIOR
510(k) history	52 submissions · 52 cleared · 1976-1987