FDA 510(k) Premarket Notification

CLEARED

K831026 INDIRECT FLUORESCENT ANTIBODY REAGENTS

Jun 22, 1983
84 days to decision

K831026 · Product code: **LHL** · Microbiology
Source: https://www.510kdatabase.net/k831026/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagents, Antibody, Legionella, Direct & Indirect Fluorescent (LHL)
Date received	Mar 30, 1983
Decision date	Jun 22, 1983
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Zeus Technologies**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1984