FDA 510(k) Premarket Notification

K811721 H-101 KAPPA,H-102 LAMBDA,H-107 LYSOZME

CLEARED

Jul 16, 1981
28 days to decision

K811721 · Product code: **DFH** · Pathology
Source: https://www.510kdatabase.net/k811721/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kappa, Antigen, Antiserum, Control (DFH)
Date received	Jun 18, 1981
Decision date	Jul 16, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Immulok, Inc.**
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1981-1983