FDA 510(k) Premarket Notification

K822347 LIDOCAINE FLUORESCENT IMMUNOASSAY

CLEARED

Aug 25, 1982
19 days to decision

K822347 · Product code: **KLR** · Toxicology
Source: https://www.510kdatabase.net/k822347/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Lidocaine (KLR)
Date received	Aug 6, 1982
Decision date	Aug 25, 1982
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**American Diagnostic Corp.**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1980-2017