FDA 510(k) Premarket Notification

K770767 IN VITRO DIAG. FOR URIC ACID

CLEARED

Jul 5, 1977
68 days to decision

K770767 · Product code: **CDO** · Chemistry
Source: https://www.510kdatabase.net/k770767/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid, Uric, Uricase (u.v.) (CDO)
Date received	Apr 28, 1977
Decision date	Jul 5, 1977
Days to decision	68 days
Third-party review	No

APPLICANT

Company	**Smithkline Diagnostics, Inc.**
Location	Mchenry, IL, US
510(k) history	42 submissions · 42 cleared · 1976-1996