FDA 510(k) Premarket Notification

K791354 CODE QAS 103 ANTI-DNA CONTROLS

CLEARED

Sep 17, 1979
61 days to decision

K791354 · Product code: **KTL** · Immunology
Source: https://www.510kdatabase.net/k791354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anti-dna Indirect Immunofluorescent Solid Phase (KTL)
Date received	Jul 18, 1979
Decision date	Sep 17, 1979
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Amersham Corp.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1976-1992