FDA 510(k) Premarket Notification

K812485 DAKO PAP KIT K504

CLEARED

Feb 10, 1982
163 days to decision

K812485 · Product code: **DFH** · Pathology
Source: https://www.510kdatabase.net/k812485/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kappa, Antigen, Antiserum, Control (DFH)
Date received	Aug 31, 1981
Decision date	Feb 10, 1982
Days to decision	163 days
Third-party review	No

APPLICANT

Company	**Dako Corp.**
Location	Mchenry, IL, US
510(k) history	54 submissions · 54 cleared · 1981-2002