

K812485 DAKO PAP KIT K504Feb 10, 1982
163 days to decisionK812485 · 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

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Device record: <https://www.510kdatabase.net/k812485/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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