

**K812484 DAKO PAP KIT K505**Feb 10, 1982  
163 days to decisionK812484 · Product code: **DEW** · Pathology  
Source: <https://www.510kdatabase.net/k812484/>**SUBMISSION DETAILS**

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

**APPLICANT**

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

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Device record: <https://www.510kdatabase.net/k812484/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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