

K810836 DECALCIFER IIApr 23, 1981
28 days to decisionK810836 · Product code: **KDX** · Pathology
Source: <https://www.510kdatabase.net/k810836/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solution, Decalcifier, Acid Containing (KDX)
Date received	Mar 26, 1981
Decision date	Apr 23, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Surgipath
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k810836/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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