

K812738 BOUNDARY PROCESS CHANGENov 16, 1981
48 days to decisionK812738 · Product code: **KKX** · General Hospital
Source: <https://www.510kdatabase.net/k812738/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Sep 29, 1981
Decision date	Nov 16, 1981
Days to decision	48 days
Third-party review	No

APPLICANT

Company	The Buckeye Cellulose Corp.
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1980-1983

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Device record: <https://www.510kdatabase.net/k812738/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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