

**K810888 BOUNDARY TUBING HOLDERS**Apr 23, 1981  
21 days to decisionK810888 · Product code: **FZX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k810888/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Surgical, Instrument (FZX)
Date received	Apr 2, 1981
Decision date	Apr 23, 1981
Days to decision	21 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810888/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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