

K770305 CUTTER, P607 BONEMar 9, 1977
21 days to decisionK770305 · Product code: **FZT** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k770305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cutter, Surgical (FZT)
Date received	Feb 16, 1977
Decision date	Mar 9, 1977
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Dentronix, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1977-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k770305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 7, 2026