

K761023 SIMULATOR, 10 LEAD ECGJan 31, 1977
81 days to decisionK761023 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k761023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 11, 1976
Decision date	Jan 31, 1977
Days to decision	81 days
Third-party review	No

APPLICANT

Company	Birtcher Corp.
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1988

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k761023/> 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 May 22, 2026