

K820117 AUTOMATIC TRAY (ACT I & ACT II)Feb 18, 1982
31 days to decisionK820117 · Product code: **KPX** · Radiology
Source: <https://www.510kdatabase.net/k820117/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Changer, Radiographic Film/cassette (KPX)
Date received	Jan 18, 1982
Decision date	Feb 18, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	Burbury & Cutter Ind., Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820117/>; 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 21, 2026