FDA 510(k) Premarket Notification

K820117 AUTOMATIC TRAY (ACT I & ACT II

CLEARED

Feb 18, 1982
31 days to decision

K820117 · 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