FDA 510(k) Premarket Notification

K761332 HI-SENSITIVITY MODE OPTION FOR S60-1A

CLEARED

Dec 30, 1976
3 days to decision

K761332 · Product code: **JAK** · Radiology
Source: https://www.510kdatabase.net/k761332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Dec 27, 1976
Decision date	Dec 30, 1976
Days to decision	3 days
Third-party review	No

APPLICANT

Company	**Syntex Corp.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1976-1992