FDA 510(k) Premarket Notification

K173023 Symbia T16, Symbia Intevo 16 and Symbia Intevo Bold

CLEARED

Nov 17, 2017
50 days to decision

K173023 · Product code: **KPS** · Radiology
Source: https://www.510kdatabase.net/k173023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Sep 28, 2017
Decision date	Nov 17, 2017
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	Tabitha Estes
510(k) history	778 submissions · 778 cleared · 1980-2026