

K173023 Symbia T16, Symbia Intevo 16 and Symbia Intevo BoldNov 17, 2017
50 days to decisionK173023 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173023/> 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 15, 2026