

K101768 MIE SCINTRON GAMMA CAMERA SYSTEMSOct 5, 2010
104 days to decisionK101768 · Product code: **KPS** · Radiology
Source: <https://www.510kdatabase.net/k101768/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Jun 23, 2010
Decision date	Oct 5, 2010
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Miegmbh
Location	Elk Grove Village, IL, US
Contact	THOMAS KUHL
510(k) history	4 submissions · 4 cleared · 2010-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101768/> 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 June 18, 2026