

K041022 NEUROQ - PET DPJun 17, 2004
58 days to decisionK041022 · Product code: **KPS** · Radiology
Source: <https://www.510kdatabase.net/k041022/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Apr 20, 2004
Decision date	Jun 17, 2004
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Syntermed, Inc.
Location	Atlanta, GA, US
Contact	KENNETH F VAN TRAIN
510(k) history	12 submissions · 12 cleared · 2000-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041022/> 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 17, 2026