FDA 510(k) Premarket Notification

K041022 NEUROQ - PET DP

CLEARED

Jun 17, 2004
58 days to decision

K041022 · 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