FDA 510(k) Premarket Notification

K030870 WBR, MODEL HR

CLEARED

Apr 9, 2003
21 days to decision

K030870 · Product code: **KPS** · Radiology
Source: https://www.510kdatabase.net/k030870/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Mar 19, 2003
Decision date	Apr 9, 2003
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**U.C.G. Technologies , Ltd.**
Location	Haifa, IL
Contact	DAN LAOR
510(k) history	2 submissions · 2 cleared · 2003-2003