FDA 510(k) Premarket Notification

K031248 CONSULTIVA REPORT STATION, MODEL RS-1

CLEARED

Jun 20, 2003
63 days to decision

K031248 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k031248/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 18, 2003
Decision date	Jun 20, 2003
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Miramedica, Inc.**
Location	Los Gatos, CA, US
Contact	WIDO MENHARDT
510(k) history	1 submissions · 1 cleared · 2003-2003