510(k) Premarket Notification Database

Device Classification Name	Catheter, Percutaneous
510(K) Number	K041151
Regulation Number	870.1250
Device Name	KERBEROS OCCLUDING GUIDE CATHETER AND ACCESSORIES
Applicant	KERBEROS PROXIMAL SOLUTIONS, INC. 1400 Terra Bella Ave, Suite K Mountain View, CA 94043
Contact	Michael A Daniel
Product Code	DQY
Date Received	05/03/2004
Decision Date	07/22/2004
Decision	Substantially Equivalent (SE)
Classification Advisory Committee	Cardiovascular
Review Advisory Committee	Cardiovascular
Statement/Summary/Purged Status	Summary Only
Summary	Summary
Type	Traditional
Reviewed By Third Party	No
Expedited Review	No

Database Updated 12/07/2004

Center for Devices and Radiological Health / CDRH