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### 510(k) Premarket Notification Database

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

Database Updated 12/07/2004

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