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### 510(k) Premarket Notification Database

<b>Device Classification Name</b>	<a href="#">Suture, Nonabsorbable, Synthetic, Polyamide</a>
<b>510(K) Number</b>	K994176
<b>Regulation Number</b>	<a href="#">878.5020</a>
<b>Device Name</b>	SUTRALON NYLON SUTURES <a href="#">VISIONARY MEDICAL SUPPLIES, INC.</a>
<b>Applicant</b>	4740 Connecticut Ave. N.W. Suite 708 Washington, DC 20008
<b>Contact</b>	Jonathan Green
<b>Classification Product Code</b>	<a href="#">GAR</a>
<b>Date Received</b>	12/10/1999
<b>Decision Date</b>	02/08/2000
<b>Decision</b>	Substantially Equivalent (SE)
<b>Classification Advisory Committee</b>	General & Plastic Surgery
<b>Review Advisory Committee</b>	General & Plastic Surgery
<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 9/06/2005