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510(k) Premarket Notification Database

Device Classification Name	Stopcock, I.V. Set
510(K) Number	K032393
Device Name	INDUCTION AND SAMPLING MANIFOLD (OR STOPCOCK) ELCAM MEDICAL ACAL
Applicant	7600 North 15th Street Suite 217 Phoenix, AZ 85020
Contact	Bruce Ward
Regulation Number	880.5440
Classification Product Code	FMG
Date Received	08/08/2003
Decision Date	09/12/2003
Decision	Substantially Equivalent (SE)
Classification Advisory Committee	General Hospital
Review Advisory Committee	General Hospital
Statement/Summary/Purged Status	Summary Only
Summary	Summary
Type	Traditional
Reviewed By Third Party	No
Expedited Review	No

Database Updated 08/06/2008

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