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510(k) Premarket Notification Database

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|--|--|
| Device Classification Name | Stopcock, I.V. Set |
| 510(K) Number | K022895 |
| Device Name | ELCAM STOPCOCKS AND MANIFOLDS |
| Applicant | MEDICAL DEVICE CONSULTANTS, INC. 49 Plain St. North Attleboro, MA 02760 4153 |
| Contact | Daniel J Dillon |
| Regulation Number | 880.5440 |
| Classification Product Code | FMG |
| Date Received | 09/03/2002 |
| Decision Date | 10/18/2002 |
| 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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