FDA 510(k) Premarket Notification

# K123818 URSA ANGIOGRAPHIC CATHETER

**CLEARED**

Mar 4, 2013
82 days to decision

K123818 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k123818/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Catheter, Intravascular, Diagnostic (DQO) |
| Date received | Dec 12, 2012 |
| Decision date | Mar 4, 2013 |
| Days to decision | 82 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Curative Medical Technology, Inc.** |
| Location | Lake Jackson, TX, US |
| Contact | AMY MCKINNEY |
| 510(k) history | 2 submissions · 2 cleared · 2010-2013 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k123818/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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