FDA 510(k) Premarket Notification

CLEARED

K142767 ThinkMed Intermittent Catheter with/without Hydrophilic Coating

Feb 3, 2015
131 days to decision

K142767 · Product code: **GBM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k142767/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urethral (GBM)
Date received	Sep 25, 2014
Decision date	Feb 3, 2015
Days to decision	131 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Thinkmed Medical Technology Co., Ltd.**
Location	Kunshan, CN
Contact	Garfield Wang
510(k) history	2 submissions · 2 cleared · 2015-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k142767/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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