FDA 510(k) Premarket Notification

K140951 YEESCOPE LARYNGOSCOPES

CLEARED

Nov 10, 2014
210 days to decision

K140951 · Product code: **CCW** · Anesthesiology
Source: https://www.510kdatabase.net/k140951/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laryngoscope, Rigid (CCW)
Date received	Apr 14, 2014
Decision date	Nov 10, 2014
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Yeescope Pty. , Ltd.**
Location	Kings Park, Nsw, AU
Contact	PAUL DRYDEN
510(k) history	1 submissions · 1 cleared · 2014-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k140951/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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