

K082667 VISIONSENSE STEREOSCOPIC VISION SYSTEM (VS SYSTEM)Oct 30, 2008
48 days to decisionK082667 · Product code: **EOB** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k082667/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nasopharyngoscope (flexible Or Rigid) (EOB)
Date received	Sep 12, 2008
Decision date	Oct 30, 2008
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Visionsense, Ltd.
Location	Washington, DC, US
Contact	GERARD J PRUD'HOME
510(k) history	13 submissions · 13 cleared · 2008-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082667/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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