

K101861 TRUEVISION 3D VISUALIZATION AND GUIDANCE SYSTEMDec 22, 2010
173 days to decisionK101861 · Product code: **HKI** · Ophthalmic
Source: <https://www.510kdatabase.net/k101861/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Jul 2, 2010
Decision date	Dec 22, 2010
Days to decision	173 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Truevision Systems, Incorporated
Location	Santa Barbara, CA, US
Contact	BURTON TRIPATHI
510(k) history	1 submissions · 1 cleared · 2010-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101861/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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