

K913230 UNIDIRECTIONAL OCCLUDEROct 2, 1991
72 days to decisionK913230 · Product code: **HOY** · Ophthalmic
Source: <https://www.510kdatabase.net/k913230/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shield, Eye, Ophthalmic (including Sunlamp Protective Eyewear And Post-mydratic Eyewear) (HOY)
Date received	Jul 22, 1991
Decision date	Oct 2, 1991
Days to decision	72 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Welch Allyn, Inc.
Location	Mchenry, IL, US
Contact	CRAIG MULLIN
Website	http://www.welchallyn.com/
510(k) history	111 submissions · 111 cleared · 1977-2025

Welch Allyn, Inc. is a medical device manufacturer based in McHenry, US. The company specializes in patient monitoring and diagnostic equipment for healthcare settings. Welch Allyn has maintained a strong FDA 510(k) regulatory record since 1977. The company has received FDA 510(k) clearances from total submissions. Cardiovascular monitoring devices represent the dominant category in recent clearances, including the Connex vital signs monitor series and central station systems. The company's latest clearance in 2025 demonstrates continued regulatory activity and product in...
