

**K080122 SOPRO 670 ARTHROSCOPIC PUMP**Jul 11, 2008  
176 days to decisionK080122 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k080122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jan 17, 2008
Decision date	Jul 11, 2008
Days to decision	176 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sopro</b>
Location	Marseille, FR
Contact	STEVE SALESKY
Website	<a href="http://www.soprole.cl/">http://www.soprole.cl/</a>
510(k) history	25 submissions · 25 cleared · 1997-2023

Sopro specialized in surgical visualization and imaging devices for general and plastic surgery applications. The company operated a manufacturing facility in Marseille, France. Sopro received FDA 510(k) clearances from total submissions between 1997 and 2023. The company focused exclusively on General & Plastic Surgery devices, particularly endoscopy cameras, digital operating room cameras, and light source systems. All submissions resulted in clearance with no denials on record. The company's product portfolio included digital endoscopy cameras, laparoscopes, operating ...

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