

**K901108 FIBERLUME**Mar 28, 1990  
19 days to decisionK901108 · Product code: **HBI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k901108/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Illuminator, Fiberoptic, Surgical Field (HBI)
Date received	Mar 9, 1990
Decision date	Mar 28, 1990
Days to decision	19 days
Third-party review	No

**APPLICANT**

---

Company	<b>Burton</b>
Location	Van Nuys, CA, US
Contact	GRETEL ANDERSON
Website	<a href="http://www.burtonalbionfc.co.uk/">http://www.burtonalbionfc.co.uk/</a>
510(k) history	1 submissions · 1 cleared · 1990-1990

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901108/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 7, 2026