

**MANUFACTURER'S DECLARATION OF CONFORMITY**  
*AUSTRALIAN THERAPEUTIC GOODS (MEDICAL DEVICES) REGULATIONS 2002*  
**DECLARATION OF CONFORMITY PROCEDURES**

SAP DIR No.: 80020236                      Version: A

This is a declaration of conformity made under clause 6.6 of Schedule 3 to the Therapeutic Goods (Medical Devices) Regulations 2002.

Manufacturer's name: Welch Allyn, Inc.

Business address: 4341 State Street Road  
Skaneateles Falls, NY 13153-0220  
U.S.A.

Product name: GS Lights



DEVICES:  
44416, 44456, 44606, 44616, 44906, 44900-C, 44900-W, 48816

ACCESSORIES:  
44215, 48200, 48605, 48805, 48850, 48950, 48955, 48960, 52630, 52640-B,  
405966

Classification: I

GMDN code and term: 12276 – Light, examination

Scope of application: All

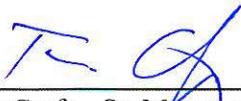
Each kind of medical device to which the technical documentation applies complies with the applicable provisions of the essential principles and the classification rules before being supplied.

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Standards applied:	EN 14971	2007	Medical Devices- Application of Risk Assessment to medical devices
	EN 60601-1 (incl. Amendments)	1990	Medical Electrical Equipment- part 1: General requirements for basic safety and essential requirements
	EN 60601-1-1	2000	Medical Electrical Equipment- part 1-1-General requirements for safety- Collateral Standard: Safety requirements for Medical Electrical Equipment
	EN 60601-1-2	2004	Medical Electrical Equipment- part 1-2- General requirements for safety- Collateral Standard: Electromagnetic Compatibility- Requirements and Test
	EN 60601-1-4	1997	Medical Electrical Equipment- part 1-4- General requirements for safety- Collateral Standard: General requirements for programmable electrical medical systems
	EN 60601-1-6	2004	Medical Electrical Equipment- part 1-6- General requirements for safety- Collateral Standard: Usability
	EN 1041	2008	Information supplied by the manufacturer with Medical devices
	EN 980	2008	Graphical symbols for use in the labeling of medical devices
	EN 62366	2008	Medical devices- Application of usability engineering to medical devices
	ISO 14155	2003	Clinical investigation of medical devices for human subjects
	ISO 10993-1	2003	Biological evaluation of medical devices -- Part 1: Evaluation and testing

Authorised Signatory:



Tim Croft    Sr. Manager, Regulatory Affairs - JAPAC

2015-09-21

Date

Rydalmere, NSW

Place of Issue

*This authorisation is given in the signatory's capacity as representative of the "Manufacturer" (as recorded on page 1 of this declaration)*