



## **DECLARATION OF CONFORMITY**

LEGAL MANUFACTURER	Danville Materials		
	2875 Loker Avenue East		
	Carlsbad, CA 92010 USA		
AUTHORIZED REPRESENTATIVE	MDSS GmbH		
	Schiffgraben 41		
HE WEST IN SECTION THE THE	30175 Hannover, GERMANY		
RESPONSIBILITY	This declaration of conformity is issued under the sole responsibility of Danville Materials		
Product(s)	Retractor		
	(refer to attached product schedule)		
RISK CLASS	Class I		
CONFORMITY ASSESSMENT ROUTE	Annex VII		
DECLARATION	We, the manufacturer, Danville Materials, declare that the above mentioned product meets the		
	provision of Council Directive 93/42/EEC for Medical Devices		
NOTIFIED BODY	N/A – not applicable to Class I devices		
EC CERTIFICATE	N/A – not applicable to Class I devices		
STANDARDS APPLIED	EN ISO 13485:2012	Medical devices - Quality management systems - Requirements for	
		regulatory purposes	
	EN ISO 14971:2012	Medical devices - Application of risk management to medical devices	
	ISO 10993-1:2009	Biological evaluation of medical devices - Part 1: Evaluation and testing	
		within a risk management process	
	ISO 15223-1:2012	Medical devices Symbols to be used with medical device labels, labelling	
		and information to be supplied Part 1: General requirements	
	EN 1041:2008	Information supplied by the manufacturer of medical devices	
	EN 980:2008	Symbols for use in the labelling of medical devices	
PLACE OF ISSUE	Carlsbad, CA, USA		
Signed	Mark	Hay 25, 2017	
	Mark Stavro, Directo	or Regulatory Affairs Date	





Part Number	Description	GMDN Code	UMDNS Code
89445	RETRACT,STANDARD	13380	13-380
89448	RETRACT,LARGE	13380	13-380