

Company Information Form for Own-Brand/Private Label Under the Medical Devices Directives.

Please complete all relevant sections & sign in the space provided

COMPANY	INFORMAT	ION							
Company Name			Viamed Ltd.						
Address	Address		15,Station Road, Crosshills.						
						-			
City:	City: Keighley Post C		Post Code:	BD207DT		Country:	W.Yorkshire		
Phone:				Fax:	+4	4 1535 636			
Contact:		D.I.Lamb		Positio	n: .	Managing Directo	or		
Alternati	Alternative Contact:			Positio	n:				
E-Mail:	E-Mail: DIL@viame		ed.co.uk		te:	www.viamed.co.uk			
<u> </u>									
	DUCT INFOR		-				···		
Details of certificat	of product(s) to e	be cove	ered by OBL						
Name of	Name of Original Equipment Supplier (OES)								
Address of OES					•				
						,-,-,,,			
OES Product Name(s) & Variants			<u> </u>						
	·								
Is the product bought fully finished from the OES? (e.g. packaged, sterilized etc).			YES						
OLOT (8:9. packaged, atermized olor.				Not Sterile					
What further processing (if any) do you carry out?			Basic Functional Test QA.						
			Add Tracking nos. & Bar Codes						



DOCUMENTATION

Are the following documents available for review?	
OES & OBL Product Labels & Instructions For Use	YES
Copies of OES CE Certificates	YES
OBL Draft Declaration of Conformity	YES
OES Declaration of Conformity (Signed)	YES
Vigitance, PMS & Recall Procedures (if OBL is responsible)	YES
Contract / Agreement between OBL & OES	YES

OES / OBL AGREEMENT

	Page Reference		
Description of product & variants	Section 3.0		
Design Drawings	Section 4.0		
Methods of manufacture	Section 4.0		
Risk Management	Section 9.0		
Evidence of compliance with Essential Requirements	Section 6.0		
If sterile, details of protocol & validation -	Not Applicable		
Test reports, clinical data, etc	Section 6.0		
Artwork for labels & instructions for use	Section 4.0.		
Access to documentation by BSI & Competent Authorities	Section 6.0		
Documentation retention period	Section 2.0		
Outline of arrangement to exchange PMS data	Section 10.0		
Outline of arrangement & responsibility for vigilance reporting (including recalls & advisory notices)	Section 9.0		
How each party & the Notified Body will be advised of any changes to the product or process	Section 10.0 Section 4.0		
OES & OBL approval signatures	Section 11		

Signature:	Si	g,	٦a	tu	re:	
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Date:

Please return this Company Information Form to:-

BSI Product Services
Maylands Avenue, Hemel Hempstead, Hertfordshire, HP2 4SQ
Telephone: +44 (0) 1442 278607 Fax: +44 (0) 1442 278575



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Contact		D.I.La	D.I.Lamb		n:	Managing Director		
Alternati	ive Contact:			Positio	n:			
E-Mail:	DIL@vian	ned.co.ul	<	Web s	ite:	www.viamed.co.uk		
	DUCT INFOF of product(s) to te							
Name o	Name of Original Equipment Supplier (OES)							
Address	of OES							
OES Pr	OES Product Name(s) & Variants							
Is the p OES?	Is the product bought fully finished from the OES? (e.g. packaged, sterilized etc).			Yes				
What fu	What further processing (if any) do you carry out?				Basic Functional Test & QA Addition of tracking Bar codes			