



251658240 **healthcare**

## Supplier Quality Questionnaire

<b>1. Company Details</b>	
Company Name: Viamed Ltd	
Company Address: 15 Station Road Cross Hills, Keighley West Yorkshire BD20 7DT	
Tel No: 01535 634542	Fax No: 01535 635582

Customer Service Contact: Tel No: 01535 634542	Email: enquiries@viamed.co.uk
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<b>2. Person responsible for Quality Assurance:</b>	
Name: Derek Lamb	Position: Managing Director
Tel No: 01535 634542	Email: Derek.lamb@viamed.co.uk
To whom is he/she responsible: Name:	Position:

<b>3. Person Responsible for Product Complaints:</b>	
Name: Steve Hardaker	Position: UK Sales Manager
Tel No: 01535 634542	Email: steve.hardaker@viamed.co.uk

<b>4. Do you have an ISO Accredited quality system?</b>		<b>Yes</b>	
If <b>YES</b> , please complete the following section and ignore section 5			
Name of system(s)	Certification Body	Certificate Number	Date of Registration
ISO9001:2008	BSI	FS 28344	15/06/94
ISO13485:2003	BSI	MD78787	27/01/04
CE Certification			23/08/96
ISO 14001:2004			
WDL			
<b>Please attach a copy of the certificate(s) and scope to this form</b>			

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Do you have a Quality Manual?	YES	NO	N/A
Do you have a Company Quality Policy?	YES	NO	N/A

## Do you have written procedures for the following?

Supplier Approval	YES	NO	N/A
Purchasing	YES	NO	N/A
Inspection of delivered materials	YES	NO	N/A
Design	YES	NO	N/A
Process Planning & Development	YES	NO	N/A
Cleanliness & contamination control	YES	NO	N/A
Production Control	YES	NO	N/A
Inspection & testing	YES	NO	N/A
Batch release	YES	NO	N/A
Equipment calibration & preventive maintenance	YES	NO	N/A
Risk Analysis	YES	NO	N/A
Retained samples	YES	NO	N/A
Complaints Handling	YES	NO	N/A
Batch Identification & traceability of product (recall)	YES	NO	N/A
Product Recall			
Change Control	YES	NO	N/A
Product packaging and labelling			
Sales	YES	NO	N/A
Storage & Distribution	YES	NO	N/A
Document and Record controls	YES	NO	N/A
Non-compliance	YES	NO	N/A
CAPA	YES	NO	N/A
Pest control	YES	NO	N/A
Continuous improvement	YES	NO	N/A
Training	YES	NO	N/A
Counterfeit awareness	YES	NO	N/A
<b>Quality Audits:</b>			
Internal	YES	NO	N/A
External	YES	NO	N/A
Management Review	YES	NO	N/A
Training	YES	NO	N/A

## 6.

If required, are you willing to supply a copy of your quality manual to us?	No
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If required, are you willing to allow a quality audit by us?	Yes
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7. Product Supplied

BUNZL Product Code	Supplier Product Code	Description	Technical Data Sheet Available? *	MSDS/ COSH Available? *	TSE Free Statement Available? *	ETO Compatible ?	Gamma Compatible	Contains Latex? *	Contains Phthalates?	Is the Device CE Marked ?	Medical Device Product Class	PPE Product Class	CE Number	Shelf Life	Instructions for Use	
			Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	I/IIa/IIb/III			Years	Required? Y/N	Supplied? * Y/N
	0021013	Posey Sensor Wraps (Box of 12)	N	N	N	N	N	N	N	Y	N/A		N/a		N	Y

\* Where the response is yes, please supply certification



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8. Form completed by:	
Name: Catrin Hollings	Position: Sales & Marketing Co-Ordinator
Signature: C Hollings	Date: 15/05/13