

Products liability questionnaire

1.4.1.

NOTE

If sufficient space is not available for any information required, please provide it on a separate sheet of paper containing your official letterhead and date and sign each sheet.

GENE	GENERAL				
1.1.	Insured (list subsidiaries included as well).				
1.2.	Situation of business premises.				
1.3.	Description of business i.e. whether you manufacture, service, repair or distribute the products and year established.				
1.4.	Description and Annual Turnover of all products. (Please list your main products and include a miscellaneous item for the balance. Do not include exports to USA and Canada as these are not insured by the policy.				

Products manufactured by you.

	Annual Turnover				
	Local Use	Exports to			
Product and Intended Use	Local Ose	Europe	Australia	Other	Total
Miscellaneous					
Total					



1.4.2. Products supplied by you.

	Annual Turnover				
	Local Use	Exports to			
Product and Intended Use	Local Ose	Europe	Australia	Other	Total
Miscellaneous					
Total					

1.4.3. Products serviced or repaired by you.

	Annual Turnover				
	l a sal lila a	Exports to			
Product and Intended Use	Local Use	Europe	Australia	Other	Total
Miscellaneous					
Total					

1.5. Special Industrie	I Industries
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Are any of your products used in the following industries?				
1.5.1.	Aviation	R		
1.5.2.	Aerospace	R		
1.5.3.	Nuclear energy	R		
1.5.4.	Shipping	R		



2. COMPLAINTS/CLAIMS HISTORY

3.

Describe the complaints you have received in the last 5 years about your products or the claims that have been made against you for injury or damage allegedly caused by your products in the past 5 years. Please provide this information in the following table

С	ate	Product	Type of Accident	Cause of Accident	Amount Paid or Outstanding		
What s	teps have	you taken to prevent a	a recurrence of the abo	ve incidents?			
QUALI	TY CONT	ROL					
			pelow the steps taken by pels or inadequate war	y you to prevent the supply o	f defective products or		
·			oolo of madoquato wan	migo orranom.			
3.1.		Products VEC/NO					
	3.1.1.		Do all your products carry the SABS mark?				
		If not, please give re	easons on a separate s	neet.			
	3.1.2.	inclusion into your		ality control system in use t ingredients/ components r			
	3.1.3.		eparate sheet the quality of defective products to	y control system in force per your customers.	item / batch to		
	3.1.4.	Who audits the conf	trol systems to ensure	that theyare maintained at a	all times?		
3.2.	2. Incorrect Labels						
	What precautions are taken to prevent incorrect labelling?						



	3.3.	Inadequa	te Warnings / Instructions on Labels	
		3.3.1.	Do the labels on your products carry adequate warnings about the possible dange of the products or their incorrect use?	
		3.3.2.	Do the labels / instructions contained in or on the product carry clear instructions at to the manner in which they should be used?	
		3.3.3.	Are the warnings / instructions / labels referred for legal opinion before being used?	.YES/NO
		3.3.4.	Who proof reads the labels after printing?	
4.			DUCTS/INGREDIENTS	
			contain any components or ingredients that could be harmful to anyone using	YES/NO
	If so, p	lease deta	il on a separate sheet:	
	4.1.	The harm	oful ingredient / component and its possible harmful effects.	
	4.2.	The steps	s taken to eliminate or reduce the possibility of such harm.	
5.	DESIG	N OF THE	PRODUCT	
	5.1.		ent to which goods are manufactured to your own design formula or specification with full details.	
		•••••		
	5.2.	Do you ha	ave a separate design section or laboratory?	YES/NO
	5.3.	specificat	e, experience and qualifications of each employee concerned with design, formula or ion of goods.	
	5.4.	Detail ste	ps in design or formula process e.g. drawing board model, prototype.	
	5.5.	Detail ext	ent and manner of tests and checks on new product before production.	



56	. To what extent do you introduce new products? Please detail.
57.	. Give the period during which existing products have been in production and on sale.
58	. To what extent do you manufacture to customer's design, formula or specification?
	leas e detail.
59	. What checks and tests are made by customer before production run actually commenced?
PO.	To what extent are goods designed by third parties, on your behalf? Detail a tending and
50	. To what extent are goods des igned by third parties on your behalf? Detail s tanding and experience of third parties and details of conditions of agreement under which des ign work is undertaken.
5 11.	. State extent to which you design 'one off jobs' as opposed to general production runs.
OI.	. Otale extent to which you design one on jobs as opposed to general production rans.
512	. Is design, formula or specification risk confined to improvements and modifications of existing range of goods?



6.	TERM	IS OF PURCHASE / SALE
	6.1.	When you purchase your products for resale or the components / ingredients used in your products, do you retain full rights of recourse against the seller?YES/NO
		If not, please provide details?
	6.2.	Do you disclaim liability for any injury or damage caused by your products?
	6.3.	Have you in the past had any claim made against you which was not paid by you or your insurers because of the disclaimer included in your terms of sale?
		In terms of the E.C. Directive it should be noted that your disclaimer will no longer be enforceable for goods exported to these countries
7.	ACTIC SUPP	ON ON DISCOVERY OF A DEFECT IN PRODUCTS SUPPLIED / RECORDS OF PRODUCTS LIED
	7.1.	If it is necessary to recall products, can you identify to whom they were sold?YES/NO
	7.2.	Give details of your product recall contingency programme.
	7.3.	Has your programme been approved by attorneys and will future amendments also be vetted by them?YES/NO
	7.4.	Do you keep adequate records for up to 10 years to clearly identify by batch number, serial number, date stamp or similar, the date the actual product was put into circulation?YES/NO
		N.B. These records (as mentioned in 7.4) are required as a defence against any claim brought against you in terms of the E.C. / E.F.T.A. Directives issued in Europe and it is a policy condition that you maintain such a record. It is not required to keep the record for raw materials.
8.	INDE	MNITY LIMIT
	What	indemnity limit is required for any one 12-month period of insurance for your products R?
DE	CLARA	TION
info		nat the particulars in this questionnaire are true and complete and that I have not withheld any material in nor am I aware (after due enquiry) of any incidents that could lead to a claim other than those listed in 2
DA	TE	AUTHORISED SIGNATORY
		DESIGNATION