

We would be delighted to consider issuing certification under the Supply of Machinery (Safety) Regulations for your machinery that has already been certificated for the Machinery Directive by a European Notified Body, but first we need to find out more information about your company, your machinery and your Notified Body. Please provide the following:

Company Name:		
Address and post code:		
Contact Person:		
Position:		
Tel. No:		
Email:		
Website:		
Number of certificate	es in this application	
Name of the machine	ery (per certificate)	
Please describe the machines):	machine (or range of	
macrimos).		
What is the Annex I\	/ catagon/2	
List all sites of manu machinery including		
Is this a single mode machines?	l or a range of	



If a range of machines, please give details on all the machines in the range and explain differences:	
Please give details of options and accessories, if any:	
Volume of machinery already on the market:	
If you are not the manufacturer of the equipment, please provide the name and address of the original manufacturer:	
address of the original mandracturer.	
Which harmonised standards have been used in the design process?	
Is there series production of the machinery	
If yes, how do you ensure quality assurance in this process?	
Do you operate a Quality Management System certified to ISO 9001?	
If yes, please state the current certificate number and expiry date.	
Give the business name and full address of the Notified Body that has issued your	
EC type-examination certificate for the machinery:	
That in ory.	
Notified Body number:	
EC type-examination certificate number: (please attache a copy to this application)	
Date of expiry:	



Contact for amtri veritas (Name, telephone and em		
	,	
Give the name and full ad Authorised Representative		
a written mandate to mee the duties of compliance i	t some or all of	
stated on the Declaration	of Conformity. Usi	be confused with the 'person authorised to compile the technical file' ng an Authorised Representative is optional – it is a way of liance to another legal person.
		ease ensure that the Authorised Representative is stated on the formity and in the Instructions.
Please provide any other would be helpful in descri		
Do you want amtri verita "person authorised to con	npile the technical	
file" established in the UK		
Do you want amtri verita translation services for yo		
	JK Declaration of C	the market in the UK must declare a "person authorised to compile Conformity. amtri veritas offers this as an extra service and will express an interest.
		mitted and parts of it need to be submitted in English. amtri veritas ce and will send you a separate quotation if you express an interest.
Please note:	certified under the	or certification will only be accepted for machinery that is already a Machinery Directive. No planned or actual modifications to the nges to its manufacture or manufacturer(s) will be considered.
Certification will be completed based on evaluation work undertaken by your Not Body.		
We will require your Notified Body to enter into an agreement with amtrive will require the applicant to enter into a <i>UK Certification Agreement</i> with amtriveritas .		



By submitting this application, I declare that:

- the information provided is true and accurate,
- the above machinery has not been modified since the issue of the certificate supplied, and the place(s) and methods of manufacture remain the same,
- the copy of the technical file to be submitted is identical to the copy that was evaluated for certification by the Notified Body,
- my EU Notified Body is authorised to submit technical file and evaluation information to amtri veritas,
- I am authorised to submit this application as the responsible person on my company's behalf, and
- the application has not been submitted to, or lodged with, another Approved Body.

Completed by (name):	
Position in Company:	
Date:	

IMPORTANT: For shared certification, **amtri veritas** require <u>the entire</u> technical file that has been retained by the Notified Body in compliance with the Machinery Directive, Annex IX, clause 4.

Please complete the Stage 1 and Stage 2 information requirements tables below and send all required documents to **amtri veritas** at **office@amtriveritas.com**.



Stage 1 information requirements

Ref.	Document	Language required	Check if supplied with your application
1	This completed application form	English	
2	Copy of the EC type-examination certificate for the machinery	English/EU language	
3	Specification of the machinery or link to website	Any	

Stage 2 information requirements

Ref.	Document	Language required	Check if supplied with your application
Technical file (Annex VII A.1)			
1.1	A general description of the machinery	English	
1.2	Overall drawings of the machinery and control circuits	Any	
1.3	Full detailed drawings required to check conformity with the EHSRs or C-type designated standards	Any	
1.4	Calculations, test results, certificates, etc. required to check conformity with the EHSRs or C-type designated standards	English	
1.5	A risk assessment	English	
1.6	A statement of compliance with the EHSRs of the Regulations or with the clauses of a relevant C-type standard	English	
1.7	A list of designated and other standards used	Any	
1.8	Any technical reports giving results of tests carried out by the manufacturer or their representative	English	
1.9	A copy of the instructions	English	
1.10	Declarations of Incorporation and assembly instructions for any partly completed machinery used	Any EU language	
1.11	Declarations of Conformity for other products used in the machinery.	Any EU language	
1.12	A copy of the UK Declaration of Conformity	English	
1.13	A picture or drawing of the machine marking showing the UKCA mark	English	
Serie	es manufacture (if applicable) (Annex VII A.2)		
2	Documentation of the internal measures to be taken to ensure that the machinery remains in conformity with the regulations.	English	
Eval	uation records (Annex IX 3)		
3.1	Reports and other documents describing the evaluation and testing carried out by the Notified Body.	English	
3.2	Calibration records for the equipment used during the evaluation and testing carried out by the Notified Body	English	
3.3	Reports and other documents describing any non-conformances found and how they have been resolved.	English	