



VARIAN MEDICAL SYSTEMS UK LTD  
DECLARATION OF CONFORMITY WITH DIRECTIVE  
93/42/EEC

This declaration is hereby made under Annex II of the Council Directive concerning Medical Devices; 93/42/EEC of 14<sup>th</sup> June 1993.

Medical Devices covered by this declaration, manufactured by Varian Medical Systems UK Ltd, Gatwick Road, Crawley, West Sussex, RH10 9RG, UK; comply with the provisions of Council Directive 93/42/EEC which apply to them.

This declaration covers the devices listed in the description of product range in the Technical File referred to below.

The British Standards Institution have been appointed to undertake activities pursuant to Annex II in respect of all devices except those Class I devices supplied non-sterile and which do not have a measuring function.

<b>Product/Product Group:</b>	ACUITY Radiotherapy Simulator	<b>Technical File Ref. No.:</b>	AC/TF
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**Revision Control & Change History**

Issue	Change	ECO	Changed By
1	NEW TECHNICAL FILE (LIMITED SCOPE)	10019	VACE
2	FULL PRODUCTION RELEASE	10338	VACE

Authorised:

*S. Turner* 28/12/02  
*P. P. Ch Brown* 20<sup>th</sup> December 2002,  
Quality Assurance Manager  
VMS UK Ltd

Date:

20<sup>th</sup> December 2002,

