Lloyd's Register
LRQA

APPROVAL OF CONFORMITY CERTIFICATE

In accordance with the requirements of the In Vitro Diagnostic Medical Devices Directive 98/79/EC and the Medical Devices Regulations 2002, UK Statutory Instrument 2002 No. 618

This is to certify that the Quality Management System of:

Microgen Bioproducts Ltd
1 Admiralty Way,
Camberley, Surrey GU15 3DT
United Kingdom

has been assessed against the requirements of Annex IV of the In Vitro Diagnostic Medical Devices Directive 98/79/EC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown on the attached certificate schedule.

Approval is subject to the maintenance of the quality system in accordance with the requirements of the above Directive and Regulations.

Authorisation is hereby given to use the LRQA Notified Body Registration Number in accordance with the requirements of the specified Directives/Regulations in relation to the products as identified above.

Certificate No:	LRQ 0942206/C
Original Approval:	25 June 2003
Current Certificate:	31 October 2016
Certificate Expiry:	30 October 2019

LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited

1 Trinity Park, Bickenhill Lane, Birmingham, B37 7ES, United Kingdom

APPROVAL OF CONFORMITY CERTIFICATE
CERTIFICATE LRQ 0942206/C SCHEDULE

has been assessed against the requirements of Annex IV of the In Vitro Diagnostic Medical Devices Directive 98/79/EC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown below:

Microgen Bioproducts Ltd
1 Admiralty Way,
Camberley, Surrey GU15 3DT
United Kingdom

Annex II List B Products
Mercia Rubella-G (M506 & M5066)

Schedule Issue: 01

Date of Schedule Issue: 31 October 2016

LRQA Notified Body Number 0088

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