

TGA

THERAPEUTIC  
GOODS  
ADMINISTRATION

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ABN 40 939 406 804



Commonwealth Department of  
Health and  
Ageing

## **CERTIFICATE FOR INCLUSION OF A MEDICAL DEVICE**

**ARTG Number** 150061

**ARTG Labelname** IQ Medical - Eye dye

**Sponsor** IQ Medical

**Commencement  
Date** 13/02/2008

**Manufacturer** Fluoron GmbH Germany

**Device Class** Class IIb  
**GMDN Code** 45180 Eye dye

**ARTG Product Number and Name**  
238776 Eye dye

**The above Medical Device is Included in the Australian Register of Therapeutic Goods subject to the following conditions**

### ***Standard Conditons***

*The automatic conditions applicable to the inclusion of all kinds of medical devices in the Register are as specified in section 41FN of the Therapeutic Goods Act 1989.*

*The standard conditions that are imposed under section 41FO of the Therapeutic Goods Act 1989 when kinds of medical devices are included in the Register are as set out in the following paragraphs.*

*For a medical device included in the Register under Chapter 4 and imported into Australia, the Sponsor must ensure that information about the Sponsor is provided in such a way as to allow the sponsor to be identified.*

*Each sponsor shall retain records of the distribution of all of the sponsor's medical*

**CERTIFIED ORIGINAL  
CERTIFICATE  
WHEN RED**

## **CERTIFICATE FOR INCLUSION OF A MEDICAL DEVICE**

**ARTG Number** 152235

**ARTG LabelName** IQ Medical - Retinal tamponade medium, intraoperative

**Sponsor** IQ Medical

**Commencement Date** 09/05/2008

**Manufacturer** Fluoron GmbH Germany

**Device Class** Class IIa

**GMDN Code** 45283 Retinal tamponade medium, intraoperative

**ARTG Product Number and Name**  
241990 Retinal tamponade medium, intraoperative

**The above Medical Device is Included in the Australian Register of Therapeutic Goods subject to the following conditions**

**Standard Conditions**

*The automatic conditions applicable to the inclusion of all kinds of medical devices in the Register are as specified in section 41FN of the Therapeutic Goods Act 1989.*

*The standard conditions that are imposed under section 41FO of the Therapeutic Goods Act 1989 when kinds of medical devices are included in the Register are as set out in the following paragraphs.*

*For a medical device included in the Register under Chapter 4 and imported into Australia, the Sponsor must ensure that information about the Sponsor is provided in such a way as to allow the sponsor to be identified.*

*Each sponsor shall retain records of the distribution of all of the sponsor's medical*

## **CERTIFICATE FOR INCLUSION OF A MEDICAL DEVICE**

**ARTG Number** 152234

**ARTG Labelname** IQ Medical - Retinal tamponade medium, postoperative

**Sponsor** IQ Medical

**Commencement  
Date** 09/05/2008

**Manufacturer** Fluoron GmbH Germany

**Device Class** Class IIb  
**GMDN Code** 45125 Retinal tamponade medium, postoperative

### **ARTG Product Number and Name**

241989 Retinal tamponade medium, postoperative

**The above Medical Device is Included in the Australian Register of Therapeutic Goods subject to the following conditions**

#### **Standard Conditions**

*The automatic conditions applicable to the inclusion of all kinds of medical devices in the Register are as specified in section 41FN of the Therapeutic Goods Act 1989.*

*The standard conditions that are imposed under section 41FO of the Therapeutic Goods Act 1989 when kinds of medical devices are included in the Register are as set out in the following paragraphs.*

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