VSY Biotechnology BV
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DECLARATION OF CONFORMITY

We

Name of Manufacturer **VSY Biotechnology BV**

Address of Manufacturer: **Strawinskylaan 1143 1077XX**
Amsterdam-The Netherlands

declare on our own responsibility that the medical device

Name of Medical Device: **Intraocular Lens**

Acriva UD 613, Acriva UDB625, Acriva UDC625, Acriva HAF

Acriva UDM 611, Acriva Reviol MF 613, Acriva Reviol MFB 625

Acriva Reviol MFM 611, Acriva Reviol Tri-ED TF613

Acriva Reviol Tri-ED TFB625, Acriva Reviol Tri-ED TFM611,

Ocuva A625, Ocuva 625, Ocuva AB625, Ocuva B625,

Acriva BB UD 613, Acriva BB UDM611, Acriva Reviol BB MF613,

Acriva Reviol BB MFM611, Acriva Reviol Tri-ED 613,

Acriva Reviol Tri-ED 611, Acriva BB T UDM 611,

Acriva Reviol BB T MFM 611, Acriva Reviol Tri-ED T 611

GMDN-Code:**35658**

Duration of validity :(see below)

Class: **IIb**

According to annex IX of direct 93/42/EEC

Meets all applicable requirements of the Directive 93/42/EEC

Name, Address and Identification **ECM, Bismarckstr. 106 52066 Aachen Germany**

Number of Notified Body **0481**
Conformity Assessment Procedure:**93/42/EEC, Annex II(excluding 4)**
CE Certificate Number: **Z/15/03635E**

18.11.2018
Validity
Amsterdam, 01.10.2015
Place, Date

Director
VSY Biotechnology B.V.
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