



# EC Declaration of Conformity

Name and address of the company

Authorised Representative

QIAGEN Sciences LLC (QIAGEN)  
19300 Germantown Road  
Germantown, MD 20874  
USA

QIAGEN GmbH  
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40724 Hilden  
Germany

We herewith declare under our sole responsibility that the product

***digene*<sup>®</sup> HC2 DNA Collection Device**

**REF 619234**

Classification

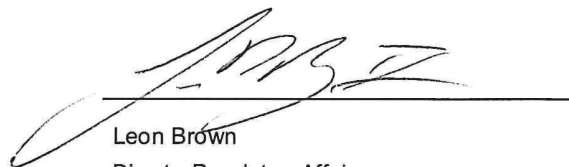
Class I (Rule 5)

Conformity Assessment Route

Annex VII

and meets all applicable requirements of the following European Directive:  
Medical Device Directive (MDD) 93/42/EEC

QIAGEN GmbH, Hilden  
03 November 2021



Leon Brown

Director Regulatory Affairs  
QIAGEN

