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 QIAGEN Strasse 1 40724 Hilden Germany

We herewith declare under our sole responsibility that the product

***digene*® HC2 DNA Collection Device** **REF 619234**

Classification Class I (Rule 5)

Conformity Asesement 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

CE