2. DECLARATION OF CONFORMITY

Identification of the Legal Manufacturer:	Natera Corporation 201 Industrial Rd., Suite 410, San Carlos, CA 94070 Tel: (650) 249-9090 raqa@natera.com www.natera.com
Identification of Authorized Representative:	Qarad BV Flight Forum 40 5657 DB Eindhoven The Netherlands www.qarad.com
This Declaration of Conformity is issued under the sole responsibility of the Legal Manufacturer.	
Product Name:	Natera™ Multiplex PCR Reagents
Part numbers covered by this Declaration of Conformance and CE marked	gDNA Multiplex PCR Reagents (Part No. 121100, 121144) cfDNA Multiplex PCR Reagents (Part No. 111100, 111144) Library Preparation Reagents (Part No. 131100, 131144)
Product Class and Rule:	Natera Multiplex PCR reagents are classified as "General IVD" under Article 9 Sections 2 and 3 and Annex III of the Medical Device Directive 98/79/EC.
Date of first CE Marking:	May 19, 2015
The undersigned hereby declares, on behalf of Natera, Inc., that the above mentioned devices comply with the Essential Requirements laid down in Annex I and the above mentioned products meet the provisions of Council Directive 98/79/EC for *In vitro* Diagnostics Medical Devices as transposed in the national laws of the member states, California State and US Federal Clinical Laboratory Improvement Amendments (CLIA) and The College of American Pathologists Accreditation (CAP).	
Signed for and on behalf of Natera by:	Name: Jonathan Sheena Signature: _____ Title: Chief Technology Officer Place of Issue: Natera, Inc., San Carlos, CA 94070 Date: May 29, 2015