Declaration of Conformity

Identification of the Legal Manufacturer:	Natera Corporation 201 Industrial Rd., Suite 410, San Carlos, CA 94070 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:	Panorama™ Precision Genomics SaaS Platform
Model names covered by this Declaration of Conformance and CE marked	Panorama Software Module Version 2.0
Product Class	Panorama™ Precision Genomic SaaS Platform is classified under "General IVD" under Article 9 Section 1 and Annex III of the In vitro Diagnostic Medical Device Directive 98/79/EC.
Name and address of Notified Body:	Not Applicable
Date of first CE Marking:	June 26, 2014
The undersigned hereby declares, on behalf of Natera, Inc., that the above mentioned device(s) 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) The College of American Pathologists Accreditation.	
Signed for and on behalf of Natera by:	Name: Michelle Roeding Signature: Michelle Roeding Title: Director, Regulatory Affairs Place of Issue: Natera, Inc., San Carlos, CA 94070 Date: August 5, 2015