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Declaration of Conformity

Identification of the Legal Manufacturer:	Natera, Inc. 201 Industrial Rd., Suite 410 San Carlos, CA 94070, U.S.A Tel: +1-(650) 249-9090
Identification of Authorized Representative:	QARAD B.V. 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™ Prenatal Screen Collection Kit
Part numbers covered by this Declaration of Conformance	NAT-801141
Product Class	The Panorama Prenatal Screen Collection Kit is classified as "General IVD" under Article 1(2)(b) and Article 9 Section 1 of the In-Vitro Diagnostics Medical Device Directive 98/79/EC.
Date of first CE Marking:	October 5, 2015
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 product meets the provisions of Council Directive 98/79/EC for *In vitro* Diagnostics Medical Devices as transposed in the national laws of the member states.	
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: October 5, 2015