

Declaration of Conformity

Company	GENOMICA, S.A.U. Parque Empresarial Alvento Edificio B.Vía de los Poblados, 1 -1ª planta. 28033 Madrid
Product:	CLART® CMA EGFR
Classification:	Other products
Conformity assessment route:	Annex III of Directive 98/79/EC (EC Declaration of Conformity).
GMDN Code	58280

GENOMICA S.A.U, declares that the diagnostic device *CLART® CMA EGFR*, conforms to the relevant provisions of the EC Council Directive 98/79/EC on in vitro diagnostic medical devices, transposed to the Spanish legislation in the Royal Decree 1662/2000 of 29th September 2000.

Standards applied	ISO 9001:2008, ISO13485:2012, ISO 14971:2013, ISO 18113-1:2011, ISO18113-2:2011, ISO18303:2011, EN 980:2008, EN 13612:2002/AC 2002, EN 23640:2015, EN 62304:2006, EN 61010-2-101:2011, EN 61326-2 6:2006.
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Notified Body	NA
EC Certificate	NA
Period of validity:	NA

Authorised Signatory

Name:	Dra. Rosario Cospedal
Position:	Managing Director
Date	Madrid, 13 th July 2017