EC Declaration of Conformity

TRIMERO Diagnostics, SL
c. València 558, 4t 2a - 08026 Barcelona (Spain)
+34 93 244 86 79 - www.3diag.com

CE IVD

Products for the determination of Soluble Transferrin Receptor (sTfR)

The Manufacturer,

TRIMERO Diagnostics SL - c. València, 558, 4t 2a - 08026 Barcelona (Spain)

declares under his responsibility that the products:

3diag - sTfR - 800 KIT	REF	TD-42680	Soluble Transferrin Receptor - KIT - for *Immage 800*
3diag - sTfR - 800	REF	TD-42681	Soluble Transferrin Receptor - for *Immage 800*
3diag - sTfR - TIA KIT	REF	TD-42694	Soluble Transferrin Receptor - for Turbidimetry
3diag - sTfR - TIA	REF	TD-42691	Soluble Transferrin Receptor - for Turbidimetry
3diag - sTfR - ADV	REF	TD-42695	Soluble Transferrin Receptor - for *Advia*
3diag - sTfR - ADV (5)	REF	TD-42696	Soluble Transferrin Receptor - for *Advia* (500 test)
3diag - sTfR - CAL SET	REF	TD-42682	Soluble Transferrin Receptor - Calibrators (6 lev.)
3diag - sTfR - CAL	REF	TD-42692	Soluble Transferrin Receptor - Calibrator
3diag - sTfR - CONTROL	REF	TD-42683	Soluble Transferrin Receptor - Controls (2 lev.)

comply with the essential requirements of Annex I of the 98/79/CE Directive on *In Vitro* Diagnostic Medical Devices, that are applicable to them.

Product Classification: **IVDMD, neither of Annex II, nor of Self-diagnosis.**

Assessment Conformity: **In accordance with Annex III of 98/79/CE Directive.**

TRIMERO Diagnostics SL
Barcelona, March 8, 2021

Joan Carles Gómez
General Manager