

Declaration of Conformity

Certificate Identification:

ARCH Sys Acc LC

IRIS V4

Legal Manufacturer's Name:

Abbott Laboratories Diagnostics Division

Legal Manufacturer's Address:

Supersedes:

Abbott Park, IL 60064 USA

List Numbers and Size Code of Devices	GMDN Code	N	ames and Description of Devices	Classification
4D18-03	56701	ARCHITECT	Septum	Self-declared
4D19-01	56701	ARCHITECT	Replacement Caps	Self-declared
7C14-01	56676	ARCHITECT	Sample Cups	Self-declared
7C15-02	56676	ARCHITECT	Reaction Vessels	Self-declared
7C15-03	56676	ARCHITECT	Reaction Vessels	Self-declared
Representative Max-Plance				
Storage site of technical Abbott Laboration Diagnostics I (Name and Address) Abbott Park,		MICH. 1992 (1997)		
Harmonized Standards Listed in the		Listed in the	echnical Documentation	

We, the undersigned, hereby declare that the in vitro diagnostic medical devices described above and bearing the CE marking. conform with the applicable provisions of the EC Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on In Vitro Diagnostic Medical Devices as they are transposed into the laws of the member states.

This declaration is made in accordance with Annex III of the IVD Directive and is issued under the sole responsibility of the manufacturer.

Signature: Full Name: Position: Date of Approval: Date of Approval: Abbott Laboratories, Diagnostics Place Issued: Date Issued: Division, Abbott Park, IL 60064 USA Effective (Date or 02 June 2015

Lot Number):