



Declaration of Conformity

FORM-QA-031.0
Level 4
Effective Date: 2/1/11

Product Designation: AdenoPlus™

EDMA- Code: 15-70-90-90-00

Model Number: RPS-AD

We herewith declare that the products listed above are in compliance with the requirements set out in the Council Directive on the harmonization of the Laws of the Member States concerning In-Vitro-Diagnostic Directive 98/79/EC.

Conformity Assessment Procedure:

In Vitro Diagnostic Directive 98/79/EC Annex III other than List A, List B or Self Testing

Applied Harmonized Standards:

EN/ISO 11737-2:2009

EN/ISO 14971:2012

EN 62366:2008

EN 13612:2002

EN 13612/AC:2002

EN ISO15223-1:2012

EN/ISO13485:2012

EN/ISO18113-1:2011

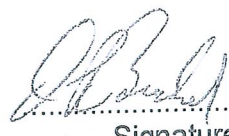
EN/ISO18113-2:2011

EU Authorized Representative: MT Promedt Consulting GmbH
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Company Name: Rapid Pathogen Screening, Inc.

Address: 7227 Delainey Court
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September 19, 2012
Date


Signature
Douglas Bueschel, VP QA/RA

STATE OF FLORIDA
COUNTY OF SARASOTA

On this 13th day of January, 2014, I attest that the preceding document is a true, exact, complete, and unaltered photocopy made by me of the RPS Declaration of Conformity for AdenoPlus™ presented to me by the document's custodian, Karen O'Toole, and, to the best of my knowledge, that the photocopied document is neither a public record nor a publicly recordable document, certified copies of which are available from an official source other than a notary public.



Lauren Milford