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## EC DECLARATION OF CONFORMITY

According Directive 98/79/EC on in vitro diagnostic medical devices, Annex III

**Manufacturer:** ACRO BIOTECH LLC 9500 7th Street Unit M Rancho Cucamonga,  
CA 91730

**Product:** HIV/HCV/HBSAG/Syphilis Combo Rapid Test Cassette (Whole Blood/Serum/Plasma)

**Product code:** IMID-445

**Classification:** Other device (all devices except Annex II and self-testing devices)

We herewith declare on our sole responsibility that all batches of the above mentioned product are conform with the Essential Requirements of the directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices.

ACRO BIOTECH INC

Dr, Mathias Volk  
Chemist, Safety officer

This document is valid until 31-12-2025