

Declaration of Conformity

ACON Laboratories, Incorporated
10125 Mesa Rim Road
San Diego, CA 92121, USA

We declare under our sole responsibility that the *in vitro* diagnostic device:

Mission® PT Coagulation Monitoring System (CCM-121)
Mission® PT Coagulation Meter (CCM-121)
Mission® PT Coagulation Test Strips (CCS-121)
Mission® PT Coagulation Control Solution

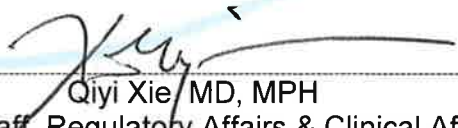
classified as Self Test in the directive 98/79/EC,

meets all the provisions of the directive 98/79/EC on *in vitro* diagnostic medical devices which apply to it

**This declaration is according to Annex IV of the Directive
and thus is based on approval by the notified body
TÜV SÜD Product Service GmbH,
Ridlerstraße 65,
80339 MÜNCHEN, Germany,
notified under No. 0123 to the EC Commission.**

Authorized Representative:
MDSS
Schiffgraben 41
30175 Hannover, Germany

Signed this 25 day of Feb, 2016
in San Diego, CA, USA


Qi Yi Xie MD, MPH
Senior Staff, Regulatory Affairs & Clinical Affairs
Acon Laboratories, Inc.

