neuroquip

CE

5 061008 370002

DECLARATION OF CONFORMITY

Neuroquip Ltd
St John's Innovation Centre
Cowley Road
Cambridge
CB4 0WS
United Kingdom

We hereby declare that the following device has been classified according to the classification rules and conforms to the essential principles for safety and performance specified in the Medical Devices Regulations EU 2017/745.

Manufacturing Site

The Workshop
Violet House
Chapel Lane
Elm
Wisbech
Cambridgeshire
PE14 0DJ

Medical Device

Neuroquip 'Action Research Arm Test Kit' – Model# ARAT001 / UDI: 5061008370002

Risk Classification

Class 1 (Non-Sterile) Medical Device as classified in MDR Annex VIII, Chapter 3, Rule 4.1

Conformity Assessment

Conformity assessment procedure: Class 1 Annex I and III of MDR (EU) 2017/745

This declaration of conformity is valid from 1st June 2018

Andrew D Teager

Andrew D Teager (Managing Director)
Authorised Signatory

Dated: 1st June 2018