4 DECLARATION OF CONFORMITY

CF Declaration of Conformity CF					
according to Directive 98/79/EC, on in vitro diagnostic medical devices					
Maker: (Name, Address)	Biopanda Reagents Ltd				
Production address	Unit 14 Carrowreagh Business Park, Carrowreagh Road, Belfast BT16 1QQ United Kingdom.				
Medical device	Description	on : COVID-19 Rapid Test Kit (RAPG-COV-019)			
	Classification of products according to directive			•	Others
	Batch/serial No. type, production term (if applicable)			••	
Applicable coordination standards:	EN ISO 14971:2012 ISO 13485:2016 EN ISO 18113:2011 EN 13612:2002 EN 23640:2015 EN ISO 15223-1:2016				
Signatory representative declares herein the above mentioned device meets the basic				ned device meets the basic	
requirements of the European Parliament and the Council's in vitro diagnostic medical					
devices directive: 98/79/EC Annex III.					
This declaration of conformity is based on European Parliament and the Council's					
98/79/EC directive Annex III.					
Managing Director:					
24/02/2020		Han Yan			
(place and date of issue)		(name and signature or equivalent marking of authorised person)			