DECLARATION OF CONFORMITY

The undersigned legal representative of the company Faster S.r.l. hereby declares that the follow products:

GLOVE FAST CYTO PHARMA

are in compliance with the following directives:

2006/42/EC	Directive of the European Parliament and of the Council on machinery
2014/30/UE	Directive of the European Parliament and of the Council on the approximation of the laws of the Member States relating to electromagnetic compatibility
2014/35/UE	Directive of the European Parliament and of the Council on the harmonisation of the laws of Member States relating to electrical equipment designed for use within certain voltage limits

and with the following standards:

ISO 14644-1	Clean rooms and associated controlled environments: Part 1: Classification of air cleanliness.
ISO 14644-7	Clean rooms and associated controlled environments: Part 7:Separative devices (clean air hoods, gloveboxes, isolator and mini-environments).
EN 61010-1	Safety requirements for electrical equipment for measurement, control and laboratory use Part 1: general requirements
EN 61326-1	Electrical equipment for measurement, control and laboratory use EMC requirements
DIN 12980:2017	Laboratory installations – Safety cabinets and glove boxes for cytotoxic substances and other CMR drugs

and, according to the above-mentioned directives, the CE IIA mark has been applied.

The undersigned also declares that the person who is authorised to compile the relevant technical documentation is Ing. Pietro Bascapè

Cornaredo, May 16, 2018

Faster S.r.l.

Maria Giulia Turzi

Maria Giulia Turzi

Chairman of the board