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

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

Mission® U120 Smart Urine Analyzer

classified as others of 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 self-declaration is according to Annex III (excluding Section 6) of the Directive.

Authorized Representative: Medical Device Safety Service GmbH Schiffgraben 41 30175 Hannover, Germany

Signed this 16 day of June, 2017 in San Diego, CA USA

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

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

Insight® U120 Smart Urine Analyzer

classified as others of 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 self-declaration is according to Annex III (excluding Section 6) of the Directive.

Authorized Representative: Medical Device Safety Service GmbH Schiffgraben 41 30175 Hannover, Germany

Signed this 16th day of June, 2017 in San Diego, CA USA

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

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

Mission® U120 Smart Urine Analyzer
Mission® U120 Smart Urine Analyzer Data Transfer Kit
Mission® Barcode Reader
Mission® Printer Paper Rolls

classified as others of 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 self-declaration is according to Annex III (excluding Section 6) of the Directive.

Authorized Representative: Medical Device Safety Service GmbH Schiffgraben 41 30175 Hannover, Germany

Signed this 16th day of June, 2017 in San Diego, CA USA



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

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

Insight® U120 Smart Urine Analyzer
Insight® U120 Smart Urine Analyzer Data Transfer Kit
Insight® Barcode Reader
Insight® Printer Paper Rolls

classified as others of 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 self-declaration is according to Annex III (excluding Section 6) of the Directive.

Authorized Representative: Medical Device Safety Service GmbH Schiffgraben 41 30175 Hannover, Germany

Signed this 1677 day of True, 2017 in San Diego, CA USA

