Department for Business & Trade

Office for Product Safety & Standards

Appointment to act as a Conformity Assessment Body

The Non-automatic Weighing Instruments Regulations 2016 (SI 2016/1152), as amended.

Conformity assessment body details

Body Name	***Eidgenössisches Institut für Metrologie METAS - Konformitätsbewertungs-stelle METAS-Cert***
Appointment Type	Approved Body
Appointment Number	***1259***

Appointment details

Appointment	Having been successfully assessed the above Conformity Assessment Body has been appointed to assess the conformity of the product categories and for the modules identified below and has been appointed under the terms of the Mutual Recognition Agreement between the United Kingdom and Switzerland
Northern Ireland	The body has not been appointed to act in respect of Conformity Assessment activities undertaken for goods for supply in Northern Ireland.

Accreditation details

Accreditation Body	***Swiss Accreditation Service SAS***
Accreditation Standard	17065
	The scope of the accreditation covers the product categories and conformity assessment procedures concerned in this appointment.

Revision details

Version	*Date*	*Details*
1	31 October 2023	Initial issue

The current validity of this document should be confirmed through the CAB's current listing on the UK Market Conformity Assessment Database.

Details of appointment

Product categories	*Assessment Procedure as defined in Schedule 7 of the Regulations*
Non-automatic weighing instruments for the purpose of the: (a) determination of mass for commercial transactions; (b) determination of mass for the calculation of a toll, tariff, tax, bonus, penalty, remuneration, indemnity or similar type of payment; (c) determination of mass for the application of laws or regulations or for an expert opinion given in court proceedings; (d) determination of mass in the practice of medicine for weighing patients for the purposes of monitoring, diagnosis and medical treatment; (e) determination of mass for making up medicines on prescription in a pharmacy and determination of mass in analyses carried out in medical and pharmaceutical laboratories; (f) determination of price on the basis of mass for the purposes of direct sales to the public and the making-up of pre-packages;	- **Module B** (Paragraph 1) Type examination **Module D** (Paragraph 2) Conformity to type based on quality assurance of the production process **Module D1** (Paragraph 3) Quality assurance of the production process **Module F** (Paragraph 4) Conformity to type based on product verification **Module F1** (Paragraph 5) Conformity based on product verification **Module G** (Paragraph 6) Conformity based on unit verification

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