BS EN ISO 18113-2:2011
Incorporating corrigendum December 2011

BSI Standards Publication

In vitro diagnostic medical devices — Information supplied by the manufacturer (labelling)

Part 2: In vitro diagnostic reagents for professional use (ISO 18113-2:2009)

National foreword

This British Standard is the UK implementation of EN ISO 18113-2:2011. It is identical to ISO 18113-2:2009. It supersedes BS EN ISO 18113-2:2009, which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee CH/212, IVDs.

A list of organizations represented on this committee can be obtained on request to its secretary.

This publication does not purport to include all the necessary provisions of a contract. Users are responsible for its correct application.

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Compliance with a British Standard cannot confer immunity from legal obligations.

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English Version

In vitro diagnostic medical devices - Information supplied by the manufacturer (labelling) - Part 2: In vitro diagnostic reagents for professional use (ISO 18113-2:2009)

Dispositifs médicaux de diagnostic in vitro - Informations fournies par le fabricant (étiquetage) - Partie 2: Réactifs de diagnostic in vitro à usage professionnel (ISO 18113-2:2009)

In-vitro-Diagnostika - Bereitstellung von Informationen durch den Hersteller - Teil 2: In-vitro-diagnostische Reagenzien für den Gebrauch durch Fachpersonal (ISO 18113-2:2009)

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