

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)

bsi.

...making excellence a habit.™

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 It supersedes BS EN ISO 18113-2:2009, which is withdrawn.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.

© The British Standards Institution 2012

ISBN 978 0 580 77328 0

ICS 11.100.10

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

This British Standard was published under the authority of the Standards Policy and Strategy Committee on 30 November 2011.

Amendments/corrigenda issued since publication

Date	Text affected
31 January 2012	Implementation of CEN correction notice 9 November 2011: Corrected date of withdrawal in EN foreword