

CORRIGENDA

Corrigendum to Commission communication in the framework of the implementation of Directive 98/79/EC of the European Parliament and of the Council on in vitro diagnostic medical devices*(Publication of titles and references of harmonised standards under Union harmonisation legislation)**(Official Journal of the European Union C 173 of 13 May 2016)**(2016/C 249/05)*

On page 139, standard EN ISO 15197:2015:

for:

'CEN	EN ISO 15197:2015 In vitro diagnostic test systems - Requirements for blood-glucose monitoring systems for self-testing in managing diabetes mellitus (ISO 15197:2013)	This is the first publication	EN ISO 15197:2013 Note 2.1	30.6.2017
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For blood glucose test strips and control solutions, the date of cessation of presumption of conformity of the superseded standard shall be 30.6.2017',

read:

'CEN	EN ISO 15197:2015 In vitro diagnostic test systems - Requirements for blood-glucose monitoring systems for self-testing in managing diabetes mellitus (ISO 15197:2013)	13.5.2016	EN ISO 15197:2003 Note 2.1	31.7.2016
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For blood glucose test strips and control solutions, the date of cessation of presumption of conformity of the superseded standard shall be 30.6.2017',

on page 140, standard EN ISO 23640:2015:

for:

'CEN	EN ISO 23640:2015 In vitro diagnostic medical devices - Evaluation of stability of in vitro diagnostic reagents (ISO 23640:2011)	This is the first publication	EN ISO 23640:2013 Note 2.1	30.6.2017',
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read:

'CEN	EN ISO 23640:2015 In vitro diagnostic medical devices – Evaluation of stability of in vitro diagnostic reagents (ISO 23640:2011)	13.5.2016	EN ISO 13640:2002 Note 2.1	30.6.2017'.
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