

# DIN EN ISO 18113-2:2013-01 (E)

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

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<b>Contents</b>		<b>Page</b>
<b>1</b>	<b>Scope .....</b>	<b>5</b>
<b>2</b>	<b>Normative references .....</b>	<b>5</b>
<b>3</b>	<b>Terms and definitions .....</b>	<b>5</b>
<b>4</b>	<b>General .....</b>	<b>6</b>
<b>4.1</b>	<b>Essential requirements .....</b>	<b>6</b>
<b>4.2</b>	<b>Identification of kit components .....</b>	<b>6</b>
<b>5</b>	<b>Content of the outer container label .....</b>	<b>6</b>
<b>5.1</b>	<b>Manufacturer .....</b>	<b>6</b>
<b>5.2</b>	<b>Identification of the IVD reagent .....</b>	<b>6</b>
<b>5.3</b>	<b>Contents .....</b>	<b>6</b>
<b>5.4</b>	<b>Intended use .....</b>	<b>6</b>
<b>5.5</b>	<b>In vitro diagnostic use .....</b>	<b>7</b>
<b>5.6</b>	<b>Storage and handling conditions .....</b>	<b>7</b>
<b>5.7</b>	<b>Expiry date .....</b>	<b>7</b>
<b>5.8</b>	<b>Warnings and precautions .....</b>	<b>7</b>
<b>6</b>	<b>Content of the immediate container label .....</b>	<b>7</b>
<b>6.1</b>	<b>General provisions .....</b>	<b>7</b>
<b>6.2</b>	<b>Manufacturer .....</b>	<b>8</b>
<b>6.3</b>	<b>Identification of the IVD reagent .....</b>	<b>8</b>
<b>6.4</b>	<b>Contents .....</b>	<b>8</b>
<b>6.5</b>	<b>In vitro diagnostic use .....</b>	<b>8</b>
<b>6.6</b>	<b>Storage and handling conditions .....</b>	<b>8</b>
<b>6.7</b>	<b>Expiry date .....</b>	<b>8</b>
<b>6.8</b>	<b>Warnings and precautions .....</b>	<b>8</b>
<b>7</b>	<b>Content of the instructions for use .....</b>	<b>9</b>
<b>7.1</b>	<b>Manufacturer .....</b>	<b>9</b>
<b>7.2</b>	<b>Identification of the IVD reagent .....</b>	<b>9</b>
<b>7.3</b>	<b>Intended use .....</b>	<b>9</b>
<b>7.4</b>	<b>Principles of the examination method .....</b>	<b>9</b>
<b>7.5</b>	<b>Traceability of values assigned to calibrators and trueness-control materials .....</b>	<b>9</b>
<b>7.6</b>	<b>Components .....</b>	<b>10</b>
<b>7.7</b>	<b>Additional required equipment .....</b>	<b>10</b>
<b>7.8</b>	<b>Reagent preparation .....</b>	<b>10</b>
<b>7.9</b>	<b>Storage and shelf life after first opening .....</b>	<b>10</b>
<b>7.10</b>	<b>Warnings and precautions .....</b>	<b>10</b>
<b>7.11</b>	<b>Primary sample collection, handling and storage .....</b>	<b>11</b>
<b>7.12</b>	<b>Examination procedure .....</b>	<b>11</b>
<b>7.13</b>	<b>Control procedure .....</b>	<b>11</b>
<b>7.14</b>	<b>Calculation of examination results .....</b>	<b>11</b>
<b>7.15</b>	<b>Interpretation of results .....</b>	<b>11</b>
<b>7.16</b>	<b>Performance characteristics .....</b>	<b>11</b>
<b>7.17</b>	<b>Biological reference intervals .....</b>	<b>12</b>
<b>7.18</b>	<b>Limitations of the examination procedure .....</b>	<b>12</b>

<b>7.19</b>	<b>Literature references .....</b>	<b>12</b>
	<b>Bibliography .....</b>	<b>13</b>
	<b>Annex ZA (informative) Relationship between this European Standard and the Essential Requirements of the EU Directive 98/79/EC on "in vitro Diagnostic Medical Devices" .....</b>	<b>15</b>
	<b>Foreword .....</b>	<b>3</b>
	<b>Introduction .....</b>	<b>4</b>