

Directive 98/79/EC of the European Parliament and of the  
Council of 27 October 1998 on in vitro diagnostic medical devices

*F1* Article 7

1 The Commission shall be assisted by the Committee set up by Article 6(2) of Directive 90/385/EEC.

2 Where reference is made to this paragraph, Articles 5 and 7 of Council Decision 1999/468/EC<sup>(1)</sup> shall apply, having regard to the provisions of Article 8 thereof.

The period laid down in Article 5(6) of Decision 1999/468/EC shall be set at three months.

3 Where reference is made to this paragraph, Article 5a(1) to (4) and Article 7 of Decision 1999/468/EC shall apply, having regard to the provisions of Article 8 thereof.

4 Where reference is made to this paragraph, Article 5a(1), (2), (4) and (6) and Article 7 of Decision 1999/468/EC shall apply, having regard to the provisions of Article 8 thereof.]

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**Textual Amendments**

**F1** Substituted by [Regulation \(EC\) No 596/2009 of the European Parliament and of the Council of 18 June 2009 adapting a number of instruments subject to the procedure referred to in Article 251 of the Treaty to Council Decision 1999/468/EC with regard to the regulatory procedure with scrutiny](#) Adaptation to the regulatory procedure with scrutiny — Part Four.

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*Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.*

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(1) [<sup>F1</sup>OJ L 184, 17.7.1999, p. 23.]

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