Commission communication in the framework of the implementation of Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in-vitro diagnostic medical devices [1]

(2001/C 319/04)

(Text with EEA relevance)

(Publication of titles and references of European harmonised standards under the Directive)

ESO [1]	Reference	Title of the harmonised standards	Year of ratification
CEN	EN 980/A1	Graphical symbols for use in the labelling of medical devices	1996 1999
CEN	EN 12286/A1	In-vitro diagnostic medical devices — Measurement of quantities in samples of biological origin — Presentation of reference measurement procedures	1998 2000

[1] ESO (European standardisation organisation):
— CEN: rue de Stassart/Stassartstraat 36, B-1050 Brussels; tel. (32-2) 550 08 11, fax (32-2) 550 08 19 (http://www.cenorm.be);
— Cenelec: rue de Stassart/Stassartstraat 35, B-1050 Brussels; tel. (32-2) 519 68 71, fax (32-2) 519 69 19 (http://www.cenelec.be);
— ETSI: F-06561 Sophia Antipolis Cedex, tel. (33-4) 92 94 42 00, fax (33-4) 93 65 47 16 (http://www.etsi.org).

NOTE:

— any information concerning the availability of the standards can be obtained either from the European standardisation organisations or from the national standardisation bodies of which the list is annexed to the Directive 98/34/EC [2] of the European Parliament and of the Council of 22 June 1998,

— publication of the references in the *Official Journal of the European Communities* does not imply that the standards are available in all the Community languages,

— the Commission ensures the updating of this list [3].

[1] OJ L 331, 7.12.1998, p. 1.

[2] OJ L 204, 21.7.1998, p. 37.

[3] OJ C 227, 10.8.1999, p. 16, OJ C 288, 9.10.1999, p. 41, OJ C 293, 14.10.2000, p. 11.