Commission communication in the framework of the implementation of Council Directive 90/385/EEC of 20 June 1990 in relation to 'Active Implantable Medical Devices' [1]

(2004/C 42/08)

(Text with EEA relevance)

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

ESO [1]	Reference	Title of the harmonised standards
CEN	EN ISO 14971:2000/AC:2002	Medical devices — Application of risk management to medical devices (ISO 14971:2000)

[1] ESO (European standardisation organisations):
— CEN: rue de Stassart 36, B-1050 Brussels, tel. (32-2) 550 08 11, fax (32-2) 550 08 19 (http://www.cenorm.be)
— Cenelec: rue de Stassart 35, B-1050 Brussels, tel. (32-2) 519 68 71, fax (32-2) 519 69 19 (http://www.cenelec.org)
— ETSI: 650, route des Lucioles, F-06921 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 European Parliament and Council Directive 98/34/EC [2] amended by the Directive 98/48/EC [3].

— Publication of the references in the *Official Journal of the European Union* does not imply that the standards are available in all the Community languages.

— The Commission ensures the updating of this list.

[1] OJ L 189, 20.7.1990, p. 17.
[2] OJ L 204, 21.7.1998, p. 37.
[3] OJ L 217, 5.8.1998, p. 18.