NOTIFICATION

The following notification is being circulated in accordance with Article 10.6

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| **1.** | **Notifying Member:**Egypt**If applicable, name of local government involved (Article 3.2 and 7.2):** |
| **2.** | **Agency responsible:**Egyptian Organization for Standardization and Quality16 Tadreeb El-Modarrebeen St., Ameriya, Cairo - EgyptE-mail: eos@idsc.net.eg/eos.tbt@eos.org.egWebsite: <http://www.eos.org.eg>Tel.: + (202) 22845528Fax: + (202) 22845504**Name and address (including telephone and fax numbers, email and website addresses, if available) of agency or authority designated to handle comments regarding the notification shall be indicated if different from above:** |
| **3.** | **Notified under Article 2.9.2 [****X],** **2.10.1[****],** **5.6.2 [****],** **5.7.1 [****],** **other****:** |
| **4.** | **Products covered (HS or CCCN where applicable, otherwise national tariff heading. ICS numbers may be provided in addition, where applicable):** Syringes, needles and catheters (ICS 11.040.25), Other medical equipment (ICS 11.040.99) |
| **5.** | **Title, number of pages and language(s) of the notified document:** Draft of Egyptian standard for "Sharps injury protection — requirements and test methods — sharps protection features for single-use hypodermic needles, introducers for catheters and needles used for blood sampling" (18 page(s), in Arabic) |
| **6.** | **Description of content:** This draft standard gives requirements and test methods for evaluating the performance parameters of sharps injury protection features, whether active or passive in design, for medical devices  containing  (sharp) hypodermic needles for single use, introducers for catheters used in blood sampling. The sharps injury protection devices it covers may be provided integral to the device or and lancets, and   other needles used  in containing combined with the device prior to use to achieve the sharps injury protection.  It does not give requirements for the storage and handling of the sharps protection before its intended use, or for the medical device itself.Worth mentioning is that this draft standards is technically identical with  ISO  23908/2011 |
| **7.** | **Objective and rationale, including the nature of urgent problems where applicable:** Health and Safety requirements |
| **8.** | **Relevant documents:**ISO  23908/2011 |
| **9.** | **Proposed date of adoption:**To be determined**Proposed date of entry into force:**To be determined |
| **10.** | **Final date for comments:** 60 days from notification |
| **11.** | **Texts available from:National enquiry point[****X]** **or address, telephone and fax numbers and email and website addresses, if available, of other body:**Egyptian Organization for Standardization and QualityAddress: 16 Tadreeb El-Modarrebeen St.Ameriya, Cairo - EgyptE-mail: eos@idsc.net.eg/eos.tbt@eos.org.egWebsite: <http://www.eos.org.eg>Tel.: + (202) 22845528Fax: + (202) 22845504 |