Ο ΠΕΡΙ ΤΥΠΟΠΟΙΗΣΗΣ, ΔΙΑΠΙΣΤΕΥΣΗΣ ΚΑΙ ΤΕΧΝΙΚΗΣ ΠΛΗΡΟΦΟΡΗΣΗΣ ΝΟΜΟΣ ΤΟΥ 2002

Γνωστοποίηση βάσει του Άρθρου 5(γ) του Νόμου 156(I)/2002

Με το παρόν γνωστοποιείται ότι με βάσει το Άρθρο 5(γ) του Νόμου 156(I) του 2002 που ρυθμίζει τις δραστηριότητες της Τυποποίησης, Διαπίστευσης και Τεχνικής Πληροφόρησης, ο Κυπριακός Οργανισμός Τυποποίησης δημοσιεύει ως Κυπριακά πρότυπα τα πιο κάτω Ευρωπαϊκά πρότυπα:

STANDARDS FROM EUROPEAN COMMITTEE FOR STANDARDIZATION (CEN)

01	GENERALITIES. TERMINOLOGY. STANDARDIZATION. DOCUMENTATION

CYSEN Document:	Title	ICS1	Withdrawn
CYS EN 13460:2009	Maintenance - Documentation for maintenance	01.110	S/S CYS EN 13460:2002 =W/D
CYS EN 13888:2009	Grout for tiles - Requirements, evaluation of conformity, classification and designation	01.040.91	S/S CYS EN 13888:2002 =W/D
CYS EN 15663:2009	Railway applications - Definition of vehicle reference masses	01.040.45	
CYS EN 934-2:2009	Admixtures for concrete, mortar and grout - Part 2: Concrete admixtures - Definitions, requirements, conformity, marking and labelling	01.040.91	S/S CYS EN 934-2:2001-iss2=W/D
CYS EN 1330-9:2009	Non-destructive testing - Terminology - Part 9: Terms used in acoustic emission testing	01.040.19	S/S CYS EN 1330-9:2000 =W/D

03	SERVICES. COMPANY ORGANIZATION, MANAGEMENT AND QUALITY. ADMINISTRATION. TRANSPORT. SOCIOLOGY

CYSEN Document:	Title	ICS1	Withdrawn
CYS EN 9131:2009	Aerospace series - Quality management systems - Nonconformance documentation	03.120.10	
CYS EN ISO/IEC 19796-1:2009	Information technology - Learning, education and training -Quality management, assurance and metrics - Part 1: General approach	03.100.30	
CYS EN ISO/IEC 17030:2009	Conformity assessment - General requirements for third-party marks of conformity	03.120.20	

07	MATHEMATICS. NATURAL SCIENCES

CYSEN Document:	Title	ICS1	Withdrawn
CYS EN 14166:2009	Foodstuffs - Determination of vitamin B6 by microbiological assay	07.100.30	S/S CYS ENV 14166:2001= W/D

11	HEALTH CARE TECHNOLOGY

CYSEN Document:	Title	ICS1	Withdrawn
CYS EN 794-1:1997 +A2:2009	Lung ventilators - Part 1: Particular requirements for critical care ventilators	11.040.10	S/S CYS EN 794-1:1997-iss1=W/D
CYS EN ISO 12870:2009	Ophthalmic optics - Spectacle frames - Requirements and test methods	11.040.70	S/S CYS EN ISO 12870:2004 =W/D
CYS EN ISO 15004-1:2009	Ophthalmic instruments - Fundamental requirements and test methods - Part 1: General requirements applicable to all ophthalmic instruments	11.040.70	S/S CYS EN ISO 15004-1:2006=W/D
CYS EN ISO 11979-8:2009	Ophthalmic implants - Intraocular lenses - Part 8: Fundamental requirements	11.040.70	S/S CYS EN ISO 11979-8:2006=W/D
CYS EN ISO 9919:2009	Medical electrical equipment - Particular requirements for the basic safety and essential performance of pulse oximeter equipment for medical use	11.040.10	S/S CYS EN ISO 9919:2005 =W/D
CYS EN ISO 8835-4:2009	Inhalational anaesthesia systems - Part 4: Anaesthetic vapour delivery devices	11.040.10	S/S CYS EN ISO 8835-4:2004-iss1=W/D
CYS EN ISO 8185:2009	Respiratory tract humidifiers for medical use – Particular requirements for respiratory humidification systems	11.040.10	S/S CYS EN ISO 8185:2007 =W/D
CYS EN ISO 8359:2009	Oxygen concentrators for medical use - Safety requirements	11.040.10	S/S CYS EN ISO 8359:1997 =W/D
CYS EN ISO 10651-4:2009	Lung ventilators - Part 4: Particular requirements for operator-powered resuscitators	11.040.10	S/S CYS EN ISO 10651-4:2002=W/D
CYS EN ISO 5366-1:2009	Anaesthetic and respiratory equipment - Tracheostomy tubes - Part 1: Tubes and connectors for use in adults	11.040.10	S/S CYS EN ISO 5366-1:2004=W/D
CYS EN ISO 8835-2:2009	Inhalational anaesthesia systems - Part 2: Anaesthetic breathing systems	11.040.10	S/S CYS EN ISO 8835-2:2007=W/D
CYS EN ISO 10651-2:2009	Lung ventilators for medical use - Particular requirements for basic safety and essential performance - Part 2: Home care ventilators for ventilator-dependent patients	11.040.10	S/S CYS EN ISO 10651-2:2004=W/D
CYS EN ISO 10651-6:2009	Lung ventilators for medical use - Particular requirements for basic safety and essential performance - Part 6: Home-care ventilatory support devices	11.040.10	S/S CYS EN ISO 10651-6:2004=W/D
CYS EN ISO 7376:2009	Anaesthetic and respiratory equipment - Laryngoscopes for tracheal intubation	11.040.55	S/S CYS EN ISO 7376:2003-iss1=W/D
CYS EN ISO 11810-1:2009	Lasers and laser-related equipment - Test method and classification for the laser resistance of surgical drapes and/or patient protective covers - Part 1: Primary ignition and penetration	11.040.30	S/S CYS EN ISO 11810-1:2005=W/D
CYS EN ISO 11197:2009	Medical supply units	11.040.10	S/S CYS EN ISO 11197:2004 =W/D

CYS EN ISO 5360:2009	Anaesthetic vaporizers - Agent-specific filling systems	11.040.10	S/S CYS EN ISO 5360:2007 =W/D
CYS EN ISO 18778:2009	Respiratory equipment - Infant monitors – Particular requirements	11.040.10	S/S CYS EN ISO 18778:2005 =W/D
CYS EN ISO 14408:2009	Tracheal tubes designed for laser surgery - Requirements for marking and accompanying information	11.040.10	S/S CYS EN ISO 14408:2005 =W/D
CYS EN ISO 21647:2009	Medical electrical equipment - Particular requirements for the basic safety and essential performance of respiratory gas monitors	11.040.10	S/S CYS EN ISO 21647:2004-iss1=W/D
CYS EN ISO 24415-1:2009	Tips for assistive products for walking - Requirements and test methods - Part 1: Friction of tips	11.180.10	
CYS CEN/TR 15831:2009	Method for testing compression in medical hosiery	11.120.20	S/S CYS ENV 12718:2001, CYS ENV 12719:2001 =W/D
CYS EN ISO 9360-2:2009	Anaesthetic and respiratory equipment - Heat and moisture exchangers (HMEs) for humidifying respired gases in humans - Part 2: HMEs for use with tracheostomized patients having minimum tidal volumes of 250 ml	11.040.10	S/S CYS EN ISO 9360-2:2002=W/D
CYS EN ISO 9360-1:2009	Anaesthetic and respiratory equipment - Heat and moisture exchangers (HMEs) for humidifying respired gases in humans - Part 1: HMEs for use with minimum tidal volumes of 250 ml	11.040.10	S/S CYS EN ISO 9360-1:2000=W/D
CYS EN ISO 8836:2009	Suction catheters for use in the respiratory tract	11.040.25	S/S CYS EN ISO 8836:2008 =W/D
CYS EN ISO 11138-2:2009	Sterilization of health care products - Biological indicators - Part2: Biological indicators for ethylene oxide sterilization processes	11.080.01	S/S CYS EN ISO 11138-2:2006=W/D
CYS EN ISO 11138-3:2009	Sterilization of health care products - Biological indicators - Part3: Biological indicators for moist heat sterilization processes	11.080.01	S/S CYS EN ISO 11138-3:2006=W/D
CYS EN ISO 11140-1:2009	Sterilization of health care products - Chemical indicators - Part1: General requirements	11.080.01	S/S CYS EN ISO 11140-1:2005=W/D
CYS EN ISO 11140-3:2009	Sterilization of health care products - Chemical indicators - Part3: Class 2 indicator systems for use in the Bowie and Dick-type steam penetration test	11.080.01	S/S CYS EN ISO 11140-3:2007-iss1=W/D
CYS EN 285:2006 +A2:2009	Sterilization - Steam sterilizers - Large sterilizers	11.080.10	S/S CYS EN 285:2006+ A1:2008=W/D
CYS EN ISO 15193:2009	In vitro diagnostic medical devices - Measurement of quantities in samples of biological origin - Requirements for content and presentation of reference measurement procedures	11.100.10	
CYS EN ISO 15194:2009	n vitro diagnostic medical devices - Measurement of quantities in samples of biological origin - Requirements for certified reference materials and the content of supporting documentation	11.100.10	
CYS EN ISO 21534:2009	Non-active surgical implants - Joint replacement implants - Particular requirements	11.040.40	S/S CYS EN ISO 21534:2007 =W/D
CYS EN ISO 21535:2009	Non-active surgical implants - Joint replacement implants - Specific requirements for hip-joint replacement implants	11.040.40	S/S CYS EN ISO 21535:2007 =W/D

CYS EN ISO 21536:2009	Non-active surgical implants - Joint replacement implants - Specific requirements for knee-joint replacement implants	11.040.40	S/S CYS EN ISO 21536:2007 =W/D
CYS EN ISO 25539-1:2009	Cardiovascular implants - Endovascular devices - Part 1:Endovascular prostheses	11.040.40	S/S CYS EN ISO 25539-1:2008=W/D
CYS EN ISO 25539-2:2009	Cardiovascular implants - Endovascular devices - Part 2:Vascular stents	11.040.40	S/S CYS EN ISO 25539-2:2008=W/D
CYS EN 12006-2:1998+A1:2009	Non active surgical implants - Particular requirements for cardiacand vascular implants - Part 2: Vascular prostheses including cardiac valve conduits	11.040.40	S/S CYS EN 12006-2:1998=W/D
CYS EN 12006-3:1998+A1:2009	Non active surgical implants - Particular requirements for cardiacand vascular implants - Part 3: Endovascular devices	11.040.40	S/S CYS EN 12006-3:1999=W/D
CYS CEN/TR 12401:2009	Dentistry - Guidance on the classification of dental devices and accessories	11.060.01	S/S CEN/TR 12401:2003 =W/D
CYS EN 1422:1997 +A1:2009	Sterilizers for medical purposes - Ethylene oxide sterilizers - Requirements and test methods	11.080.10	S/S CYS EN 1422:1988-iss1=W/D
CYS EN 13060:2004 +A:2009	Small steam sterilizers	11.080.10	S/S CYS EN 13060:2004 =W/D
CYS EN 14180:2003 +A1:2009	Sterilizers for medical purposes - Low temperature steam and formaldehyde sterilizers - Requirements and testing	11.080.10	S/S CYS EN 14180:2003 =W/D
CYS EN ISO 10555-1:2009	Sterile, single-use intravascular catheters - Part 1: General requirements	11.040.25	S/S CYS EN ISO 10555-1:1997-iss1=W/D
CYS EN ISO 7197:2009	Neurosurgical implants - Sterile, single-use hydrocephalus shunts and components	11.040.40	S/S CYS EN ISO 7197:2006 =W/D
CYS EN ISO 9713:2009	Neurosurgical implants - Self-closing intracranial aneurysm clips	11.040.40	S/S CYS EN ISO 9713:2004 =W/D
CYS EN ISO 14630:2009	Non-active surgical implants - General requirements	11.040.40	S/S CYS EN ISO 14630:2008 =W/D
CYS EN ISO 14602:2009	Non-active surgical implants - Implants for Osteosynthesis - Particular requirements	11.040.40	S/S CYS EN ISO 14602:1998 =W/D
CYS EN ISO 5840:2009	Cardiovascular implants - Cardiac valve prostheses	11.040.40	S/S CYS EN ISO 5840:2005 =W/D
CYS EN ISO 14607:2009	Non-active surgical implants - Mammary implants – Particular requirements	11.040.40	S/S CYS EN ISO 14607:2007 =W/D
CYS EN ISO 7439:2009	Copper-bearing intra-uterine contraceptive devices - Requirements, tests	11.200	S/S CYS EN ISO 7439:2002 =W/D
CYS EN ISO 3950:2009	Dentistry - Designation system for teeth and areas of the oral cavity	11.060.01	S/S CYS EN ISO 3950:1997 =W/D

CYS EN ISO 7711-1:1998-iss1 (Including A1:2009)	Dental rotary instruments - Diamond instruments - Part 1:Dimensions, requirements, marking and packaging	11.060.20	S/S CYS EN ISO 7711-1:1998=W/D
CYS EN 868-2:2009	Packaging for terminally sterilized medical devices - Part 2:Sterilization wrap - Requirements and test methods	11.080.30	S/S CYS EN 868-2:1999=W/D
CYS EN 868-3:2009	Packaging for terminally sterilized medical devices - Part 3:Paper for use in the manufacture of paper bags (specified in EN868-4) and in the manufacture of pouches and reels (specified in EN 868-5) - Requirements and test methods	11.080.30	S/S CYS EN 868-3:1999=W/D
CYS EN 868-4:2009	Packaging for terminally sterilized medical devices - Part 4:Paper bags - Requirements and test methods	11.080.30	S/S CYS EN 868-4:1999=W/D
CYS EN 868-5:2009	Packaging for terminally sterilized medical devices - Part 5:Sealable pouches and reels of porous materials and plastic film construction - Requirements and test methods	11.080.30	S/S CYS EN 868-5:1999-iss1=W/D
CYS EN 868-6:2009	Packaging for terminally sterilized medical devices - Part 6:Paper for low temperature sterilization processes -Requirements and test methods	11.080.30	S/S CYS EN 868-6:1999=W/D
CYS EN 868-7:2009	Packaging for terminally sterilized medical devices - Part 7:Adhesive coated paper for low temperature sterilization processes - Requirements and test methods	11.080.30	S/S CYS EN 868-7:1999=W/D
CYS EN 868-8:2009	Packaging for terminally sterilized medical devices - Part 8: Re-usable sterilization containers for steam sterilizers conforming to EN 285 - Requirements and test methods	11.080.30	S/S CYS EN868-8:1999=W/D
CYS EN 868-9:2009	Packaging for terminally sterilized medical devices - Part 9:Uncoated non woven materials of polyolefines – Requirements and test methods	11.080.30	S/S CYS EN 868-9:2000=W/D
CYS EN 868-10:2009	Packaging for terminally sterilized medical devices - Part 10:Adhesive coated non woven materials of polyolefines - Requirements and test methods	11.080.30	S/S CYS EN 868-10:2000=W/D
CYS EN ISO 10993-3:2009	Biological evaluation of medical devices - Part 3: Tests for genotoxicity, carcinogenicity and reproductive toxicity	11.100.20	S/S CYS EN ISO 10993-3:2003-iss1=W/D
CYS EN ISO 10993-4:2009	Biological evaluation of medical devices - Part 4: Selection of tests for interactions with blood	11.100.20	S/S CYS EN ISO 10993-4:2002-iss2=W/D
CYS EN ISO 10993-9:2009	Biological evaluation of medical devices - Part 9: Framework for identification and quantification of potential degradation products	11.100.20	S/S CYS EN ISO 10993-9:1999=W/D
CYS EN ISO 10993-6:2009	Biological evaluation of medical devices - Part 6: Tests for local effects after implantation	11.100.20	S/S CYS EN ISO 10993-6:2007=W/D

13	ENVIRONMENT. HEALTH PROTECTION. SAFETY

CYSEN Document:	Title	ICS1	Withdrawn
CYS EN 379:2003 +A1:2009	Personal eye-protection - Automatic welding filters	13.340.20	S/S CYS EN 379:2003 =W/D
CYS EN 1760-2:2001 +A1:2009	Safety of machinery - Pressure sensitive protective devices - Part 2: General principles for the design and testing of pressure sensitive edges and pressure sensitive bars	13.110	S/S CYS EN 1760-2:2001=W/D
CYS EN 1760-1:1997 +A1:2009	Safety of machinery - Pressure sensitive protective devices - Part 1: General principles for the design and testing of pressure sensitive mats and pressure sensitive floors	13.110	S/S CYS EN 1760-1:1998=W/D