

Registration Number of Grouping: GE117.
 Date of Registration of Grouping: 18th March 1997.
 Place of Registration of Grouping: Companies House, Crown Way,
 Cardiff CF4 3UZ.
 Official Address of Grouping: 31 St. Georges Street, London W1R
 0HQ.
 Particulars of Amendment to the Contract for the Formation of the
 Grouping: EEIG4 amending the name of the grouping from

Global Chamber European Economic Interest Grouping to
 World Chamber European Economic Interest Grouping.

J. Holden, Chief Executive and Registrar of Companies for
 England and Wales

Companies House,
 Cardiff CF4 3UZ.

(8 SI)

DEPARTMENT OF HEALTH

THE MEDICAL DEVICES AGENCY

Notice is hereby given that the following Standards are deemed to be "relevant national standards" for the purposes of the Medical Device Regulations 1994 and the Active Implantable Medical Devices Regulations 1992.

<i>Standard and OJ Ref</i>	<i>Title</i>	<i>BS Adoption</i>
CEN EN 600:1996 OJ96/C245/02	Natural rubber latex condom.	BSEN600:1996
CEN EN1640:1996 OJ97/C149/03	Dentistry. Medical devices for dentistry. Equipment.	BSEN 1640:1996
CEN ENISO4135:1996 OJ97/C149/03	Anaesthesiology. Vocabulary.	BSENISO4135:1996
CEN ENISO10079-1:1996 OJ97/C149/03	Medical suction equipment. Part 1: Electrically powered suction equipment. Safety requirements.	BSENISO10079-1:1997
CEN ENISO10079-2:1996 OJ97/C149/03	Medical suction equipment. Part 2: Manually powered suction equipment.	BSENISO10079-2:1997
CEN ENISO10079-3:1996 OJ97/C149/03	Medical suction equipment. Part 3: Suction equipment powered vacuum or pressure source.	BSENISO10079-3:1997
CEN ENISO10555-1:1996 OJ97/C149/03	Sterile, single use intravascular catheters. Part 1 General requirements.	BSENISO 10555-1:1996
CEN ENISO10993-10:1995 OJ96/C245/02	Biological evaluations of medical devices. Part 10 Tests for irradiation and sensitisation.	BSENISO10993-10:1996
CENELEC EN60601-2-17:1996 OJ97/C149/04	Medical electrical equipment. Part 2: Particular requirements for the safety of remote controlled automatically driven gamma ray after-loading equipment.	BSEN60601-2-17:1996
CENELEC EN60601-2-21:1994 JO95/C307/15	Medical electrical equipment. Part 2: Particular requirements for the safety of infant radiant warmers.	BSEN60601-2-21:1995
CENELEC EN 60601-2-22:1995 JO97/C149/04	Medical electrical equipment. Part 2: Particular requirements for the safety of diagnostic and therapeutic laser equipment.	BSEN60602-2-22:1996
CENELEC EN60601-2-25:1995 JO97/C149/04	Medical electrical equipment. Part 2: Particular requirements for the safety of electrocardiographs.	BSEN60601-2-25:1996
CENELEC EN60601-2-30:1995 OJ97/C149/04	Medical electrical equipment. Particular requirements for the safety of automatic cycling indirect blood pressure monitoring equipment.	BSEN60601-2-30:1995
CENELEC EN60601-2-33:1994 JO97/C149/04	Medical electrical equipment. Part 2: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis.	BSEN60601-2-33:1996
CEN EN46002:1995 OJ96/C245/05	Quality systems, medical devices. Particular requirements for the application of EN29002.	BSEN46002:1997
CENELEC EN60601-1:1990 OJ96/C245/06	Medical electrical equipment. Part 1: General requirements for safety.	BS5724-1
CEN ISOEN10993-10:1995 OJ97/C149/05	Biological evaluation of medical devices. Part 10: Tests for irradiation and sensitization.	BSENISO10993-10:1996
CEN ISOEN10993-12:1996 OJ97/C149/05	Biological evaluation of medical devices. Part 12: Sample preparation and reference materials.	BSENISO10993-12:1997

The above-mentioned Standards are available from the British Standards Institution, Linford Wood, Milton Keynes MK14 6LE1, telephone 01901 22 11 66.

D. Batten

The Medical Devices Agency, Hannibal House,
 Elephant and Castle, London SE1 6TQ.

(2 SI)