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Contact: Nancy Winstanley, NIAAA Library, c/o CSR Incorporated 2107 Wilson Blvd., Suite 1000, Arlington, VA 22201; phone: 703-741-7147; e-mail: nwinstanley@csrincorporated.com

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Contact: Chevy Chase; MD; http://www.aacom.org/InfoFor/educators/Pages/thesaurus.aspx

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CCS2005 - Clinical Classifications Software (CCS). Release Date: April 2005. Phone: 301-594-1364.; Agency for Healthcare Research and Quality (AHRQ); Rockville,MD; http://www.hcup-us.ahrq.gov/toolssoftware/ccs/ccs.jsp; ENG

Contact: Anne Elixhauser, Ph.D.; Senior Research Scientist; Agency for Healthcare Research and Quality; 540 Gaither Road; Rockville; MD; United States; 20850; phone: (301) 427-1411, phone: 1-800-358-9295; fax: (301) 594-1430; email: <u>AElixhau@AHRQ.gov</u>

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COSTART has been superseded by the Medical Dictionary for Regulatory Activities (MedDRA) Terminology.

Contact: National Technical Information Service. http://www.ntis.gov/fcpc/cpn5580.htm

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DDB00 - Diseases Database 2000. May, 2000. London (England): Medical Object Oriented Software Enterprises Ltd., 2000. Contact: Malcolm Duncan < mhduncan@compuserve.com; http://www.diseasesdatabase.com/.

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GO2011_06_01 - The Gene Ontology; The Gene Ontology Consortium; June 1, 2011

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(HCPCS); Baltimore, MD; Centers for Medicare & Medicaid Services; 2012

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Contact: Cynthia Hake; Centers for Medicare & Medicaid Services (CMS); 7500 Security Blvd.; Baltimore; MD; United States; 21244; 1-410-786-3404; Cynthia. Hake@cms.hhs.gov; http://www.cms.gov/

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Contact: Virginia K. Saba, EdD, RN, FAAN, FACMI, LL, Distinguished Scholar, Adjunct, Georgetown University, Washington, DC; Professor, Adjunct, USUHS, Bethedsa, MD 2332 South Queen Street, Arlington VA 22202; Tel: 703-521-6132; Fax: 703-521-3866; e-mail: vsaba@worldnet.att.net

Read more information about this source

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Contact: Health Level Seven, 3300 Washtenaw Avenue, Suite 227, Ann Arbor MI 48104-4250; phone: (734)677-7777; fax: (734)677-6622; e-mail: HQ@HL7,ORG

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Contact: Health Level Seven; 3300 Washtenaw Avenue, Suite 227; Ann Arbor; MI; USA; 48104-4250; (734) 677-7777; (734) 677-6622; HQ@HL7.ORG

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Contact: Henk Lamberts (HLREL), University of Amsterdam; email: H.Lamberts@AMC.UVA.NL.

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Contact: Dr. Elspeth Bruford; Project Coordinator; Gene Nomenclature Committee (HGNC); EMBL Outstation; European Bioinformatics Institute Wellcome Trust Genome Campus; Cambridge; United Kingdom; CB10 1SD; +44 (0) 01223 492 624; +44 (0) 1223 494 468; elspeth@ebi.ac.uk; http://www.genenames.org/

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CATEGORY 3 RESTRICTIONS APPLY

Contact: Office of Publications, World Health Organization, 1211 Geneva 27, Switzerland

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CATEGORY 3 RESTRICTIONS APPLY

Contact: National Centre for Classification in Health, University of Sydney, Faculty of Health Sciences; PO Box 170 Lidcombe, NSW, Australia 1825; Phone: +61 2 9351 9461

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CATEGORY 3 RESTRICTIONS APPLY

Contact: National Centre for Classification in Health University of Sydney Faculty of Health Sciences PO Box 170 Lidcombe, NSW Australia 1825; phone: +61 2 9351 9461

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CATEGORY 4 RESTRICTIONS APPLY to U.S. UMLS USERS

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Contact: National Center for Health Statistics; 3311 Toledo Road; Hyattsville; MD; United States; 20782; 1-800-232-4636; cdcinfo@cdc.gov; http://www.cdc.gov/nchs

Read more information about this source

ICD10DUT_200403 - Hirs, W., H.W. Becker, C. van Boven, S.K. Oskam, I.M. Okkes, H. Lamberts. ICD-10, Dutch Translation, 200403. Amsterdam: Department of General Practice, Academic Medical Center/University of Amsterdam, Dutch College of General Practitioners (NHG), March 2004.

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Contact: UMLS Support; National Library of Medicine; custserv@nlm.nih.gov

Read more information about this source

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Contact: Pat Brooks; Center for Medicare and Medicaid Services; 7500 Security Blvd. C4-08-06; Baltimore; MD; United States; 21244; patricia.brooks2@cms.hhs.gov; http://www.cms.gov/

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NLM has generated fully specified titles for ICD-9-CM codes in cases in which the official ICD- 9-CM titles consist of extensions to higher levels in the ICD-9-CM hierarchy. The fully specified names were produced with reasonable care, but have not yet been reviewed and approved by the producers of ICD-9-CM.

Contact: Patricia Brooks; Contact for Procedures; Health Care Financing Administration; Centers for Medicare & Medicaid Services; 7500 Security Boulevard; Baltimore; MD; United States; 21244; pbrooks@hcfa.gov; http://www.cms.gov/

Read more information about this source

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Contact: Contact for Diseases: Donna Pickett, National Center for Health Statistics; e-mail: dfp4@cdc.gov; Contact for Procedures: Patricia Brooks, Health Care Financing Administration; e-mail: pbrooks@hcfa.gov

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Contact: Contact for Diseases: Donna Pickett, National Center for Health Statistics; e-mail: dfp4@cdc.gov; Contact for Procedures: Patricia Brooks, Health Care Financing Administration; e-mail: pbrooks@hcfa.gov

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Contact: Nenad Kostanjsek; Technical Officer, Classification, Terminology and Standards (CTS), Department of Health Statistics and Informatics (HSI); World Health Organization; 20, Avenue Appia; CH-1211 Geneva 27; Switzerland; +41.22.791.3242; +41.22.791.4894; kostanjsekn@who.int; http://www.who.int/classification/icf

Read more information about this source

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Contact: Amy Coenen; ICNP Programme Director; University of Wisconsin - Milwaukee, College of Nursing; P.O. Box 413; Milwaukee; WI; United States; 53201-0413; 1-414-229-6474; coenena@uwm.edu

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ICPC93 - The International Classification of Primary Care (ICPC). Denmark: World Organisation of Family Doctors, 1993.

This year, the Metathesaurus has also included translations of ICPC93 in the following languages:

- Basque (ICPCBAQ 1993),
- Danish (ICPCDAN_1993),
- Dutch (ICPCDUT_1993),
- Finnish (ICPCFIN_1993),
- French (ICPCFRE_1993),
- German (ICPCGER_1993),
- Hebrew (ICPCHEB_1993),
- Hungarian (ICPCHUN_1993),
- Italian (ICPCITA_1993),
- Norwegian (ICPCNOR_1993),
- Portuguese (ICPCPOR_1993),
- Spanish (ICPCSPA_1993), and
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 $Contact: \ UMLS \ Support; National \ Library \ of \ Medicine; \underline{custserv@nlm.nih.gov}$

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Contact: A/Prof Helena Britt, Director, Family Medicine Research Centre, Acacia House, Westmead Hospital, PO Box 533, Wentworthville NSW 2145, Australia. Phone: +61 2 9845 8150; Fax: +612 9845 8155; email: fmrc@fmrc.org.au; http://www.fmrc.org.au; http://www.fmrc.org.au/

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Contact: Bo-Young Jung; Korea Centers for Disease Control and Prevention; Korea; +82-2-3481-6791; +82-2-532-3820; happiness630@hanmail.net

Read more information about this source

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There are later editions of this source that are not reflected in the UMLS Metathesaurus. This source has considerable non-biomedical content and will never be included in the Metathesaurus in its entirety.

Contact: http://www.lcweb.loc.gov/

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Contact: LOINC c/o Medical Informatics; The Regenstrief Institute, Inc; Health Information and Translational Sciences Bldg (HITS), 410 West 10th Street; Suite 2000; Indianapolis; Indiana; United States; 46202; 1-317-423-5558; 1-317-423-5695; loinc@regenstrief.org; http://loinc.org/

Read more information about this source

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Contact: kmercer@regenstrief.org

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Contact: Prevention Plus; Jball79686@aol.com; http://www.bradenscale.com/copyright.asp

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Contact: Sharon K. Inouye, M.D., MPH; Professor of Medicine, Beth Israel Deaconess Medical Center, Harvard Medical School, Milton and Shirley F. Levy Family, Chair Director; Aging Brain Center; Institute of Aging Research, Hebrew SeniorLife; 1200 Centre Street; Boston; MA 02131; 02131; (617) 363-8020; (617) 363-

8901; sharoninouye@hrca.harvard.edu

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Contact: University of Michigan Medical Center; University of Michigan Health System; Ann Arbor; MI; United States; 48109; http://www.med.umich.edu/pain/pediatric.htm

Read more information about this source

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LNC238_MDS30 (updated) - Institute of Gerontology; Minimum Data Set; 3.0; University of Michigan; Ann Arbor, Michigan

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Contact: David Cella, PhD; Principal Investigator; Northwestern University Feinberg School of Medicine; d-cella@northwestern.edu; http://www.mss.northwestern.edu/

Read more information about this source

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Contact: Andrew Kramer; MD; andy.kramer@uchsc.edu

Read more information about this source

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Contact: American Physical Therapy Association; 1111 North Fairfax Street; Alexandria; VA; United States; 22314-1488; (800)-999-2782; (703)-683-6748; publications@apta.org; http://www.apta.org/OPTIMAL/

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Contact: 212-573-2273; requests for permission@pfizer.com

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Contact: Tim Benson; Routine Health Outcomes Ltd.; 14 Pinewood Crescent; Hermitage, Thatcham; Berkshire; UK; RG18 9WL; +44 1635 203 162, +44 7855 682 037; timbenson@routinehealthoutcomes.com; www.routinehealthoutcomes.com

Read more information about this source

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Contact: HIV/AIDS Department; World Health Organization; Geneva; Switzerland; 41-22-791-1497; 41-22-791-1580; http://www.who.int/hiv/pub/en

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MCM92 - Glossary of Methodologic Terms for Clinical Epidemiologic Studies of Human Disorders. Canada: McMaster University, 1992.

Contact: R. Brian Haynes, M.D., Ph.D.; e-mail: bhaynes@mcmaster.ca; Clinical Epidemiology & Biostatistics and Medicine, Faculty of Health Sciences, McMaster University, Room 2C10B, 1200 Main Street, West Hamilton Ontario, Canada L8N 3Z5; phone (905) 525-9140

Read more information about this source

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Contact: Karen Eckert, RPh; Medi-Span; 8425 Woodfield Crossing Blvd., Suite 490; Indianapolis; IN; United States; 46240; phone (800) 388-8884; fax (317) 735-5390; email: Keckert@drugfacts.com, ms-support@drugfacts.com; http://www.medi-span.com/products/product_mddb.asp

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Contact: MedDRA MSSO; Northrop Grumman Corporation; 3975 Virginia Mallory Drive; Chantilly; Virginia; United States; 20151; 1-877-258-8280(AT&T Worldwide); 1-703-272-5505(USA); mssosubscribe@ngc.com; http://www.meddramsso.com/

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Contact: MedDRA MSSO; Northrop Grumman Corporation; 3975 Virginia Mallory Drive; Chantilly; Virginia; United States; 20151; 1-877-258-8280(AT&T Worldwide); 1-703-272-5505(USA); mssosubscribe@ngc.com; http://www.meddramsso.com/

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MED12 (updated) - MEDLINE Current Files (2007-2012). Bethesda (MD): National Library of Medicine. Contact: http://www.nlm.nih.gov/.

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Contact: David Lareau; Chief Operating Officer; Medicomp Systems; 14500 Avion Parkway, Suite 175; Chantilly; VA; United States; 20151; 703-803-8080; 703-803-8035; david.lareau@medicomp.com; http://www.medicomp.com/

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MEDLINEPLUS_20111228 (updated) - MedlinePlus Health Topics; National Library of Medicine; December 28, 2011; Bethesda, MD

Contact: Ms. Naomi Miller; Manager, Consumer Health Information, PSD; National Library of Medicine; 8600 Rockville Pike; Bldg. 38, 1W22B; Bethesda; MD; United States; 20894; (301)-496-0240; (301)-402-1384; millern@mail.nlm.nih.gov; http://www.nlm.nih.gov/medlineplus/xmldescription.html

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MSHCZE2012 (updated) - Dept. of Bibliography, National Library of Medicine; Czech translation of the Medical Subject Headings; Prague, Czech Republic; 2012

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MSHDUT2005 - Nederlandse vertaling van MeSH [Dutch translation of MeSH); Nederlands Tijdschrift voor Geneeskunde [Dutch Journal of Medicine]; 2005; Amsterdam, The Netherlands; DUT

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MSHFIN2008 - Finnish Translations of Medical Subject Headings (MeSH); Finnish Medical Society Duodecim; 2008; Helsinki, Finland; FIN

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Contact: Dr.Yannick Pilatte; INSERM, DISC-Information Scientifique et Technique; Hopital de Bicetre Batiment Claude Bernard; 84, Rue du general Leclerc; Le Kremlin-Bicetre; France; 94276 Cedex; (33)-1-49-59-53-92; (33)-1-49-59-56-03; yannick.pilatte@inserm.fr; yannick.pilatte@inserm.fr; <a

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Contact: Yosuke Seyama; Japan Medical Abstracts Society; 2-5-18, Tokaido-Higashi, Suginami; Tokyo; Japan; 168-0072; seyama@jamas.or.jp

MSHLAV2012 (updated) - Latvian translation of the Medical Subject Headings; Medical Library of Latvia; 2012; Riga (Latvia)

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MTHCH2012 (updated) - National Library of Medicine; Metathesaurus CPT Hierarchical Terms; National Library of Medicine; 2012; Bethesda, MD

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Read more information about this source

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NCI 2011_02D - National Cancer Institute, National Institutes of Health; NCI Thesaurus; National Cancer Institute; February 2011, Protege version; Rockville, MD

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Contact: Sherri de Coronado; Center for Bioinformatics, National Cancer Institute; 2115 E. Jefferson St.; 6th Floor; Rockville; MD; United States of America; 20892-8335; 925-377-5960; decorons@osp.nci.nih.gov

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Contact: National Cancer Institute Bethesda, MD; phone: 301-496-8510

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NDFRT_2012_03_05 (updated) - National Drug File - Reference Terminology; 2012_03_05; Washington, DC; U.S. Department Of Veterans Affairs, Veterans Health Administration; June 2010

Contact: Michael J. Lincoln MD, FACMI; Chief Terminologist; VA Chief Health Information Office (VA CHIO); 550 Foothill Boulevard; Salt Lake City; UT; United States; 84113; phone: 801-588-5200; e-mail: michael.lincoln@va.gov

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Contact: Douglas M. Bowden, M.D., Regional Primate Research Center, University of Washington, Box 357330, Seattle, WA 98195; e-mail: dmbowden@u.washington.edu

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NIC2005 - Nursing Interventions Classification (NIC)

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Read more information about this source

PCDS97 - Ozbolt, Judy Grace. Patient Data Care Set (PCDS), Version 4.0, 1998.Contact: judy.ozbolt@mcmail.vanderbilt.edu; Vanderbilt University School of Nursing; 400-C Godchaux Hall; Nashville, TN 37240-0008; Telephone 615-343-3291

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Contact: Margaret Haber, NCI, Building 6116 - 6116 Exec Blvd, Room 3124, Rockville, MD; telephone: 301-594-9185; Fax: 301-480-8105; Email: mhaber@mail.nih.gov

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Contact: Sharon Giarrizzo-Wilson; info@aorn.org

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Contact: Scott Antall, American Pharmaceutical Association - Academy of Pharmaceutical Research and Science; email: ssa@mail.aphanet.org; Academy of Pharmaceutical Research and Science, 2215 Constitution Avenue NW, Washington DC 20037-2985

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Contact: Lisa A. Gallagher, American Psychological Association, 750 First Stree, NE, Washingtion, DC 20002-4242; phone: 202-336-5726; email" LGallagher@apa.org

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QMR96 - Quick Medical Reference (QMR). San Bruno (CA): First DataBank, 1997.

Contact: Quick Medical Reference, First Databank, 1111 Bayhill Drive San Bruno, CA 94066

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RAM99 - QMR clinically related terms from Randolf A. Miller, 1999.

Contact: Dr. Randolph A. Miller (email: randolph.a.miller@vanderbilt.edu), Chair, Dept. of Biomedical Informatics, Vanderbilt University, 436 Eskind Biomedical Library, 2209 Garland Ave., Nashville TN 37232-8340

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RCD99 - Clinical Terms Version 3 (CTV3) (Read Codes) (Q199): National Health Service National Coding and Classification Centre; March, 1999.

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http://umlsinfo.nlm.nih.gov/RxNorm.html

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SPN2003 - Standard Product Nomenclature (SPN). Rockville, (MD); U.S. Food and Drug Administration, 2003

Contact: <u>custserv@nlm.nih.gov</u>

Read more information about this source

SRC - UMLS Metathesaurus Source Terminologies. Bethesda, MD: National Library of Medicine.

Contact: Jan Willis, National Library of Medicine, UMLS Support, 38A-4th fl, 8600 Rockville Pike, Bethesda MD 20894; phone: 301 496-7715; e-mail: jwillis@nlm.nih.gov

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TKMT2011 - Traditional Korean Medical Terms: 2011: Korea: KOR

Contact: Jinhyun Kim; Researcher/O.M.D; Information Research Center, TKM Information Research Division, Korea Institute of Oriental Medicine; 483 Expo-ro, Yuseong-gu; Daejeon; Korea; 305-811; +82-42-868-9565, +82-10-2237-2378; +82-42-861-9421; <a href="https://kiphyrozoa.gov/

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ULT93 - Bell, Douglas. Ultrasound Structured Attribute Reporting (UltraSTAR). Boston (MA): Brigham & Womens Hospital, 1993.

CATEGORY 3 RESTRICTIONS APPLY

Contact: Robert Greenes, M.D., Ph.D., Brigham & Womens Hospital; Department of Radiology, 75 Francis Street, Boston MA 02115 e-mail: greenes@harvard.edu phone: (617) 732-6281

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UMD2012 (updated) - The Universal Medical Device Nomenclature System; ECRI; Plymouth Meeting; 2012

CATEGORY 1 RESTRICTIONS APPLY

Contact: Elizabeth Richardson; Director of Database and Nomenclature Systems; ECRI; 5200 Butler Pike; Plymouth Meeting; Pennsylvania; United States; 19462-1298; 1-610-825-6000; 1-610-834-1275; erichard@ecri.org; http://www.ecri.org/Products/Pages/UMDNS.aspx

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USPMG_2004 - United States Pharmacopeia (USP). Medicare Prescription Drug Benefit Model Guidelines: Drug Categories and Classes in Part D, 2004. http://www.usp.org/pdf/EN/mmg/comprehensiveDrugListing2004-12-31.pdf

Contact: http://www.usp.org/healthcareInfo/mmg/finalGuidelines.html

Read more information about this source

UWDA173 - University of Washington Digital Anatomist, (UWDA). Seattle (WA): University of Washinton, Version 1.7.3, March, 2003. Jose Mejino, M.D.; email: onard@biostr.washington.edu

Contact: Jose Mejino, M.D.; e-mail: onard@biostr.washington.edu; University of Washington Digital Anatomist Symbolic Knowledge Base, University of Washington Digital Anatomist Information System, Structural Informatics Group, Department of Biological Structure, University of Washington, Seattle WA 98195

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VANDF_2012_01_23 (updated) - Veterans Health Administration National Drug File; January 23, 2012; Washington, DC; U.S. Department of Veterans Affairs

*NOTE: Now a CATEGORY 0.

Contact: Michael Lincoln, M.D.; U.S. Department of Veterans Affairs, Veterans Health Administration; Washington; DC; United States; email: michael.lincoln@med.va.gov; URL: http://www.pbm.va.gov/default.aspx

WHO97 - WHO Adverse Drug Reaction Terminology (WHOART). Uppsala (Sweden): WHO Collaborating Centre for International Drug Monitoring, 1997.

CATEGORY 2 RESTRICTIONS APPLY

The Metathesaurus includes translations of WHO97 in:

- French (WHOFRE_1997),
- German (WHOGER_1997),
- · Portuguese (WHOPOR_1997), and
- Spanish (WHOSPA_1997).

Contact: WHO Collaborating Centre for International Drug Monitoring, Stora Target 3, S-753 20 Uppsala, Sweden; fax: +46-18-656080

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WHOFRE_1997 - WHO Adverse Drug Reaction Terminology (WHOART). French Translation. Uppsala (Sweden): WHO Collaborating Centre for International Drug Monitoring, 1997.

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Contact: WHO Collaborating Centre for International Drug Monitoring, Stora Target 3, S-753 20 Uppsala, Sweden; fax: +46-18-656080

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WHOGER_1997 - WHO Adverse Drug Reaction Terminology (WHOART). German Translation. Uppsala (Sweden): WHO Collaborating Centre for International Drug Monitoring, 1997.

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WHOPOR_1997 - WHO Adverse Drug Reaction Terminology (WHOART). Portuguese Translation. Uppsala (Sweden): WHO Collaborating Centre for International Drug Monitoring, 1997.

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WHOSPA_1997 - WHO Adverse Drug Reaction Terminology (WHOART). Spanish Translation. Uppsala (Sweden): WHO Collaborating Centre for International Drug Monitoring, 1997.

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APPENDIX 2

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In this Licence Agreement, terms defined in Appendix A (Defined Terms) have the meanings set out in that Appendix.

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- 2.1 The Licensor grants the Licensee, subject to the terms of this Licence Agreement, a perpetual (subject to revocation in accordance with clause 5), worldwide, non-exclusive, non-transferable licence for the term of this Licence Agreement to:
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 - 2.2.2 in the development and operation of the Licensee's information systems;
 - 2.2.3 for the Licensee's research purposes; and/or
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- 2.5 Each sub-licence granted by the Licensee under clause 2.1.5 must:
 - 2.5.1 not grant the End User any greater rights in respect of the International Release than the Licensee itself has under this Licence Agreement;
 - 2.5.2 not permit the End User to do any act or thing in respect of the International Release that the Licensee is prohibited from doing under this Licensee Agreement:

- 2.5.3 not permit the End User to sub-license or transfer any of its rights under the sub-licence (unless the End User is also an Affiliate, in which case that Affiliate shall be entitled to sub-license further its rights under the sub-licence with the Licensee, subject to the same restrictions as apply to sub-licensing the International Release under the Affiliate's licence agreement with the Licensor);
- 2.5.4 terminate automatically upon termination of this Licence Agreement;
- 2.5.5 provide that the End User may apply directly to the Licensor upon receiving notice that the sub-licence will terminate in accordance with **clause 2.5.4**, and that the Licensor may in such circumstances (but shall not be obliged to):
- (a) grant the End User a licence in respect of the International Release for a limited period in order to enable the End User to continue to use the Licensee Products that are subject to the sub-licence during that period; or
- (b) give the End User an assurance or undertaking that for a limited period the Licensor will not seek to prevent the End User from using the Licensee Products: and
- 2.5.6 permit the Licensee to disclose the terms of the sub-licence to the Licensor in accordance with clause 8.
- 2.6 If the Licensee becomes aware of any material error or change or correction needed in the International Release, the Licensee agrees to advise the Licensor promptly of such error, change or correction by following the Licensor's procedures for change notification that the Licensor prescribes by regulations and notifies to the Licensee from time to time.
- 2.7 The Licensee shall comply with the Internet security measures that the Licensor prescribes by regulations and notifies to the Licensee from time to time.

3. EXTENSIONS AND DERIVATIVES

- 3.1 The Licensee may not create any Standards-Based Extension or any Standards Based Derivative unless it has first been issued with a Namespace Identifier by or on behalf of the Licensor.
- 3.2 The Licensee may request that the Licensor issue it with a Namespace Identifier, and the Licensor shall not unreasonably refuse to do so taking into account amongst other things quality assurance, governance processes, Standards and Regulations.
- 3.3 The Licensee shall ensure that all Standards-Based Extensions and Standards Based Derivatives that the Licensee creates under this Licence Agreement are created in accordance with, and comply with, all applicable Standards (including, without limitation, as to the use of Namespace Identifiers).
- 3.4 Subject to **clauses 3.5** and **3.6**, the Licensee shall own all Intellectual Property Rights in all Extensions and Derivatives that the Licensee creates under this Licence Agreement. The Licensee may not assign or otherwise transfer those Intellectual Property Rights to any other person unless (i) that person is an Affiliate and, in the case of Standards-Based Extensions or Standards-Based Derivatives, has a Namespace Identifier; and (ii) the transfer is notified in writing to the Licensor within thirty (30) days after the transfer.
- 3.5 The Licensee shall, if requested by the Licensor, transfer to the Licensor or a Member nominated by the Licensor all of its Intellectual Property Rights in such Standards-Based Extensions (or parts thereof) as the Licensor may specify.
- 3.6 The Licensee shall, if requested by the Licensor and agreed by the Licensee in the Licensee's sole discretion, transfer to the Licensor or a Member nominated by the Licensor all of its Intellectual Property Rights in such Standards-Based Derivatives as the Licensor may specify.
- 3.7 Upon the transfer to the Licensor, or to a Member, of the Intellectual Property Rights in any Standards-Based Extension (or part thereof) or Standards-Based Derivative in accordance with clauses 3.5 or 3.6:
 - 3.7.1 responsibility for the maintenance and distribution of that Extension (or part thereof) or Derivative shall also transfer from the Licensee to the Licensor or the Member (as the case may be); and
 - 3.7.2 the Licensor hereby grants a licence back to the Licensee from the Licensor or may procure from the Member a licence back to the Licensee (as the case may be) of that Extension (or part thereof) or Derivative, on the same terms as apply to the International Release under clause 2 of this Licence Agreement, until that Extension (or part thereof) or Derivative becomes part of the International Release or the Member's National Release (as the case may be).

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- 4.3 The Licensee may, by written notice, request the Licensor to modify the SNOMED CT Core. Upon receipt of such written notice, the Licensor shall consult with the Licensee and shall give due consideration as to whether the proposed modification should be made based on the Licensor's editorial guidelines and policies. Following due consideration of the matter, including consideration of any information presented by the Licensee, the Licensor shall inform the Licensee whether the proposed modification shall be made and if the Licensor agrees that the proposed modification should be made, the Licensor shall give a non-binding indication of when, reasonably and in good faith, it anticipates that the proposed modification will be made. If the Licensee would like the content of the proposed modification to be developed more quickly than the Licensor has indicated, the Licensee may itself undertake or procure the undertaking of the development of the content of the proposed modification (outside of any existing Licensor's support services contract). On receipt of the developed content of the proposed modification, the Licensor will then give due consideration as to whether the developed content meets the Licensor's quality assurance, other governance processes, Standards and Regulations then the Licensor shall incorporate the modification into the SNOMED CT Core according to its schedule which will give due consideration as to when the proposed modification shall be incorporated into the SNOMED CT Core, taking into account other proposals for the modification of the SNOMED CT Core and the work required to include the proposed modification in the SNOMED CT Core.

5. TERM AND TERMINATION

- 5.1 This Licence Agreement shall commence on the date on which it comes into effect in accordance with the notice at the beginning of this Licence Agreement, and shall continue until terminated in accordance with this clause 5.
- 5.2 Either party may terminate this Licence Agreement if the other party commits a material breach of any of its obligations under this Licence Agreement in accordance with the following procedure:

- 5.2.1 the party seeking to terminate the License Agreement (the "**Terminating Party**") shall serve an escalation notice (the "**Escalation Notice**") on the other party (the "**Defaulting Party**") requiring the Defaulting Party to nominate a member of its senior management team to meet with a member of the Terminating Party's senior management team to seek to resolve in good faith the matter giving rise to the service of the escalation notice;
- 5.2.2 The representatives of the parties identified in accordance with clause 5.2.1 shall meet in good faith to seek to resolve the matter. If they are unable to resolve the matter within 45 days of the date of the Escalation Notice the Terminating Party may serve a formal breach notice (the "Breach Notice") on the Defaulting Party requiring it to remedy the breach within 90 days.
- 5.2.3 If the Defaulting Party does not remedy the breach within 90 days of the date of the Breach Notice the Terminating Party may terminate the License Agreement by giving 180 days written notice to the Defaulting Party (the "Termination Notice").
- 5.3 The Licensor may not terminate this Licence Agreement except in accordance with clause 5.2.
- 5.4 The Licensee may terminate this License Agreement by giving up to twelve (12) months' prior written notice to the Licensor at any time between the Licensor giving notice of a variation under clause 6.3 and that variation becoming effective in accordance with clause 6.3.
- 5.5 Upon termination of this Licence Agreement in accordance with this **clause 5**, all licences granted under this Licence Agreement shall automatically and immediately be revoked.
- 5.6 The Licensee shall, by no later than thirty (30) days after termination of this Licence Agreement for any reason, remove all copies of the International Release from its computer systems and destroy all copies of electronic, paper copy and other media containing or representing any part of the International Release. The Licensee shall, if requested by the Licensor, certify in writing to the Licensor that the Licensee has complied with its obligations under this clause 5.6.
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- 5.9 The Licensor shall be entitled to publicise the termination of this Licence Agreement to such persons (including Members, other Affiliates of the Licensor and End Users) and in such manner as it sees fit.
- 5.10 Clauses 5.6, 5.7, 5.8, 5.9, 7, 8 and 10 to 14 inclusive shall survive termination of this Licence Agreement.

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- 6.3 The Licensor may vary the terms of this Licence Agreement by giving written notice to the Licensee. Any such variation shall take effect not less than ninety (90) days after the notice is given, as specified in the notice. If the Licensee does not wish this Licence Agreement to continue subject to the variation, the Licensee may terminate this Licence Agreement in accordance with **clause 5.4**, and if the Licensee does so then the variation shall not take effect.
- 6.4 The College of American Pathologists, as originator of Intellectual Property Rights in the International Release, shall as a licensee have a specific exception to the Licensor's rights in Clause 6.3 in specific circumstances and for a specific fixed term period to be agreed with the Licensor, and the terms of such special exemption shall be deemed part of such licensee's Affiliate Licence Terms. The Licensor will publish the terms of the special exemption with the Articles.

7. LICENCE FEES

- 7.1 The Licensee shall pay the Licence Fees to the Licensor in respect of the Licensee's activities in Non-Member Territories. The Licence Fees shall be payable sixmonthly in arrear, and each annual fee set out in Appendix B shall be payable in two equal six-monthly instalments.
- 7.2 All Licence Fees and other amounts payable to the Licensor under this Agreement are exclusive of value added tax and any other tax of a similar nature, which shall be payable by the Licensee at the prevailing rate in addition to those amounts.
- 7.3 The Licensee shall, by no later than fourteen (14) days after 1st January and 1st July in each calendar year, submit a statement of account to the Licensor in such manner and form as the Licensor may prescribe, setting out the Licensee's activities in Non-Member Territories in the preceding six-month period, and the Licensee's calculation of the Licence Fees and other amounts payable to the Licensor in respect of that period. If the Licence Fees for any period are less than \$1,000 (one thousand United States dollars), a statement shall nevertheless be due but no payment shall be due until the period in which the accumulated Licence Fees of \$1,000 or greater are due, at which time the Licensee shall be liable to pay the accumulated Licence Fees.
- 7.4 The Licensee shall provide the Licensor with such information as the Licensor may reasonably request for the purpose of verifying any statement of account submitted to the Licensor under clause 7.3.
- 7.5 The Licensor shall, following receipt of a statement of account from the Licensee under **clause 7.3**, submit an invoice to the Licensee setting out the Licensee Fees and other amounts payable by the Licensee in respect of the period to which the statement of account relates. The Licensee shall pay to the Licensor all amounts set out on each invoice submitted under this **clause 7.5** within thirty (30) days of receipt of that invoice. The Licensee shall make payment under this **clause 7.5** by wire transfer or by such other means as the Licensor may make available to the Licensee for time to time.
- 7.6 Interest shall accrue on any outstanding Licence Fees and other amounts at the rate of the lesser of (a) 500 basis points above the European Inter-Bank Offer Rate (EURIBOR), calculated daily from the date on which payment was due and compounding at the end of each calendar month or (b) the maximum amount allowed under applicable law.

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- 8.2.3 abbreviate the marks SNOMED or SNOMED CT; or
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- 8.7 If any use of the International Release (including without limitation use through a Licensee Product) is reasonably determined by the Licensor to be below the standards of quality required under this Licence Agreement, the Licensor shall notify the Licensee of such deficiency in writing. Upon receipt of such notice, the Licensee shall take all necessary steps to correct such deficiency (including such steps as the Licensor may reasonably specify).
- 8.8 The Licensee shall maintain a complete, accurate and up-to-date register of all sub-licences granted by the Licensee under clause 2.1.5, and shall make that register available for inspection during normal business hours by the Licensor and its representatives upon the Licensor giving not less than fourteen (14) days' prior written notice. The register maintained by the Licensee under this clause 8.8 shall at a minimum contain the following information in respect of each sub-licence: the name and registered office of the sub-licensee; the Licensee Product subject to the sub-licence; and the version of the International Release included in that Licensee Product. In the absence of circumstances giving the Licensor reasonable grounds to suspect a breach of this Licence Agreement, the Licensor may not give notice under this clause 8.8 more frequently than once per year.

9. COMPLIANCE WITH LOCAL REQUIREMENTS IN MEMBER TERRITORIES

- 9.1 The Licensee may only exercise its rights under this Licence Agreement in a Member Territory in accordance with such conditions as the Member for that territory may prescribe from time to time.
- 9.2 Conditions prescribed by a Member under clause 9.1 may:
 - 9.2.1 include, without limitation, a requirement that the Licensee notify the Member before exercising its rights under this Licensee Agreement in that Member's territory and a requirement that the Licensee enter into a license agreement with the Member in respect of that Member's National Release; and 9.2.2 relate to the International Release, the Member's National Release or any part of either of them.

10. AFFILIATE STATUS

- 10.1 During the term of this Licence Agreement the Licensee shall be an Affiliate.
- 10.2 As an Affiliate, the Licensee shall be entitled to participate in the Licensor's Affiliates Forum, which is a forum in which the Licensee and other Affiliates may communicate with the Licenser and with each other. The Licensor may make regulations from time to time governing the Licensee's participation in the Affiliates Forum. New regulations that the Licensor shall make from time to time governing participation in the Affiliates Forum shall not remove the Licensee's right to participate in that forum.

11. REPRESENTATIONS AND WARRANTIES

- 11.1 To the extent permitted by law, the Licensor excludes all representations, warranties and conditions that would otherwise be implied by law in this Licence Agreement (including, without limitation, all implied warranties of merchantability or fitness for a particular purpose).
- 11.2 Without limiting clause 11.1, the Licensor does not represent or warrant that the International Release or any part of it will satisfy any of the Licensee's requirements, operate in combinations selected by the Licensee or be free from defects or errors.

12. LIMITATION OF LIABILITY

- 12.1 The Licensor shall not be liable to the Licensee or to any other person, whether in contract, tort (including negligence), misrepresentation, breach of statutory duty or otherwise, for any of the following arising under or in connection with this Licence Agreement (including, without limitation, in respect of the Licensee's use of or inability to use the International Release or any part of it):
 - 12.1.1 indirect or consequential loss;
 - 12.1.2 special or punitive damages;
 - 12.1.3 loss of profits, loss of savings and loss of revenue;
 - 12.1.4 loss of business, loss of reputation and loss of goodwill; and
 - 12.1.5 loss of data.
- 12.2 Neither the Licensor nor any Member shall be liable to the Licensee or any other person for any failure by the Licensor or the Member (as the case may be) to maintain or distribute any Extension (or part thereof) or Derivative transferred to the Licensor or the Member (as the case may be) in accordance with clauses 3.4 or 3.5.
- 12.3 The liability of the Licensor arising in any year under or in connection with this Licence Agreement, whether in contract, tort (including negligence), misrepresentation, breach of statutory duty or otherwise, shall not in any event exceed the Licence Fees paid by the Licensee in respect of that year.
- 12.4 Nothing in this Licence Agreement excludes or limits the liability of either party for:
 - 12.4.1 fraud (including fraudulent misrepresentation);
 - 12.4.2 death or personal injury caused by the negligence of that party;
 - 12.4.3 any breach of its obligations implied by section 12 of the Sale of Goods Act 1979; or
 - 12.4.4 any other liability that by law cannot validly be excluded or limited (but only to the extent that the liability cannot validly be excluded or limited).

13. ASSIGNMENT

- 13.1 The Licensee may not assign, novate or otherwise transfer any of its rights or obligations under this Licence Agreement to any person without the prior written consent of the Licensor not to be unreasonably withheld.
- 13.2 The Licensor may transfer all of its rights and obligations under this Licence Agreement to any person to whom the Licensor transfers the Intellectual Property Rights in respect of which the licences under this Licence Agreement are granted.

14. GENERAL PROVISIONS

- 14.1 This Licence Agreement contains the entire agreement between the parties relating to the subject matter of this Licence Agreement, supersedes all previous agreements between the Parties relating to that subject matter and sets out the entirety of the Licensee's rights in respect of the International Release.
- 14.2 Each party acknowledges that, in entering into this Licence Agreement, it has not relied on any representation, warranty, collateral contract or other assurance made by on behalf of the other party before the date of this Licence Agreement.
- 14.3 Except as provided in **clause 6.3**, this Licence Agreement may not be varied except in writing signed by both parties and expressed to vary this Licence Agreement.
- 14.4 Nothing in this Licence Agreement shall give either party the ability to act or incur obligations or liability on behalf of the other party or constitutes a joint venture, agency, partnership or employment relationship between the parties.
- 14.5 If any term of this Licence Agreement is or becomes illegal, invalid or unenforceable in any jurisdiction, that shall not affect the legality, validity or enforceability in that jurisdiction of any other term of this Licence Agreement, or the legality, validity or enforceability in any other jurisdiction of that or any other term of this Licence Agreement.

15. GOVERNING LAW AND JURISDICTION

- 15.1 This Licence Agreement shall be governed by, and construed in accordance with, English law.
- 15.2 The English courts shall have exclusive jurisdiction to settle any dispute arising out of or in connection with this Licence Agreement (including a dispute regarding its existence, validity or termination).
- 15.3 Clause 15.2 is for the benefit of the Licensor only. As a result, the Licensor shall not be prevented from taking proceedings relating to any dispute in any other courts with jurisdiction. To the extent permitted by law, the Licensor may take concurrent proceedings in any number of jurisdictions.

Appendix A

Defined Terms

In this Licence Agreement, the following defined terms have the following meanings:

Term Associated Definition in the Affiliate License agreement

Affiliate	an affiliate of the Licensor in accordance with the Articles;
Articles	the Licensor's Articles of Association (vedtægter);
Cross-Map	a work consisting of (i) SNOMED CT Content and (ii) content of another nomenclature, classification or knowledge structure, together with a set of relationships between (i) and (ii);
Data Analysis System	a computer system that is used to analyse records or other data that is encoded using SNOMED CT, but not if that system is also a Data Creation System;

Data Creation System	a computer system that is used to create records or other data that is encoded using SNOMED CT;		
Derivative	a work consisting of (a) SNOMED CT Content, from the SNOMED CT CORE or an Extension; together with (b) either (i) additional properties and/or information about such SNOMED CT content; and/or (ii) any set of relationships between that SNOMED CT Content and Content of other nomenclature, classification or knowledge structure, and includes a Cross-Map and a Sub-Set;		
End User	a third party user of a Licensee Product;		
Extension	A work consisting of SNOMED CT Content alone that is supplementary to the SNOMED CT Core and that depends on the SNOMED CT Core;		
Intellectual Property Rights	patents, trade marks, service marks, copyright (including rights in computer software), moral rights, database rights, rights in designs, trade secrets, know-how and other intellectual property rights, in each case whether registered or unregistered and including applications for registration, and all rights or forms of protection having equivalent or similar effect in any jurisdiction;		
Hospital	a health care facility consisting of multiple Practices and providing in-patient care;		
International Release	the release produced and distributed by or on behalf of the Licensor, consisting of the SNOMED CT Core, the Specifications and the Licensor's Derivatives and other documents and software;		
Licence Fees	the licence fees set out in Appendix B (Licence Fees in Non-Member Territories);		
Licensee Products	products distributed or licensed by the Licensee that include the International Release (or any part of it) and/or any Extensions or Derivatives created by the Licensee under this Licence Agreement;		
Member	a member of the Licensor;		
Member Territory	a territory that is represented by a Member (as published by the Licensor from time to time);		
Namespace Identifier	a code or that part of a code that identifies the organisation responsible for creating and maintaining a Standards-Based Extension or a Standards-Based Derivative and is used as an element of SNOMED CT Identifiers;		
National Release	in respect of each Member, the release produced and distributed by the Member, consisting of the International Release, the Member's Extensions, the Member's Derivatives and other documents and software;		
Non-Member Territory	a territory that is not a Member Territory;		
Practice	a single department of a Hospital, or any health care facility that provides principally out-patient care, including without limitation a pharmacy, an optician's facility, a physiotherapy centre, a general medical practice or a family medical practice;		
Qualifying Research Project	a discrete research project that meets all of the following criteria: (a) it is supported by a formal proposal that has been peer reviewed; (b) it has been ethically approved in accordance with the prevailing legislation, regulations and guidelines in effect in the relevant territory; (c) it is conducted within a definite timeframe; and (d) the results of the research are offered for publication in peer-reviewed public journals and are provided to the Licensor free of charge;		
Regulations	regulations of the Licensor made in accordance with the Articles;		
Relationship	a relationship, of a kind defined by the Licensor in Specifications, between concepts (which may be, without limitation, a hierarchical or an associative relationship) or between a concept and a description;		
SNOMED CT	the concept-based work of clinical nomenclature and classification with multiple hierarchies and semantic definitions known as SNOMED Clinical Terms (SNOMED CT);		
SNOMED CT Content	terminological content, consisting of concepts, descriptions and Relationships, each of which is identified using a SNOMED CT Identifier;		
SNOMED CT Core	the SNOMED CT Content that is controlled, maintained and distributed by the Licensor from time to time;		
SNOMED CT Identifier	a code, of a kind defined by the Licensor in Specifications, for identifying concepts, descriptions and Relationships;		
Specification	specifications promulgated by the Licensor for products and processing relating to SNOMED CT, including specifications of the internal logic of SNOMED CT, editorial policies, guidelines and characteristics;		
Sponsored Territory	a Non-Member Territory that has been recognised and designated by the Licensor as a sponsored territory (as published on the Licensor's web site);		
Standard	a Specification that is formally adopted by the Licensor; and		
Standards- Based	in respect of an Extension or a Derivative, an Extension or Derivative the creation of which is the subject of one or more Standards;		
Sub-Set	a sub-set of SNOMED CT Content that is grouped together for one or more purposes.		

Appendix B

Licence Fees in Non-Member Territories

1. Introduction

- 1.1 This Appendix B sets out the licence fees payable by the Licensee in respect of its activities in Non-Member Territories.
- 1.2 The licence fees set out in this Appendix B do not apply in respect of the Licensee's activities in any Non-Member Territory if that Non-Member Territory is a Sponsored Territory or was a Sponsored Territory at the time when the Licensee's activities in that Non-Member Territory were carried out.
- 1.3 The Licensor may, in its sole discretion, waive the Licensee's obligation to pay any or all of the licence fees set out in this Appendix B if the Licensor considers that the Licensee's activities in any Non-Member Territory are in support of charitable or humanitarian causes in that Non-Member Territory. Any waiver by the Licensor under this paragraph 1.3 may be revoked by the Licensor at any time, shall be without prejudice to any of the Licensor's other rights and remedies under this Licence Agreement and shall not relieve the Licensee of any of its other obligations under this Licence Agreement.

2. Data Creation Systems

- 2.1 The Licensee shall pay the following fees in respect of each hospital or Practice in a Non-Member Territory in or to which the Licensee:
- (a) deploys the International Release or any Licensee Product that contains the International Release (or any part of it) in a Data Creation System, unless that Data Creation System is used exclusively in connection with a Qualifying Research Project; or
- (b) distributes or licenses a Licensee Product that is or includes a Data Creation System, unless that Licensee Product is used exclusively in connection with a Qualifying Research Project.

Non-Member Territory (fee bands)	Associated License Fee
Hospital in Band A Territory	US\$ 1,500 per annum baseline fee adjusted as per paragraph 1.4
Hospital in Band B Territory	US\$ 1,000 per annum baseline fee adjusted as per paragraph 1.4
Hospital in Band C Territory	US\$ 500 per annum baseline fee adjusted as per paragraph 1.4
Practice in Band A, B or C Territory	US\$ 500 per annum baseline fee adjusted as per paragraph 1.4
Hospital or Practice in Low Income Band	US\$ 0 per annum baseline fee adjusted as per paragraph 1.4
Hospital or Practice in other territory	As per paragraph 6.2.

2.2 The total fees payable by the Licensee in respect of a number of Practices in a single hospital shall not exceed the fee applicable to the hospital itself.

3. Data Analysis Systems

- 3.1 The Licensee shall pay the fees set out in paragraph 3.4 if the Licensee:
 - (a) deploys the International Release or any Licensee Product that contains the International Release (or any part of it) in a Data Analysis System in a Non-Member Territory, unless that Data Analysis System is used exclusively in connection with a Qualifying Research Project; or
 - (b) distributes or licenses a Licensee Product that is or includes a Data Analysis System in a Non-Member Territory, unless that Licensee Product is used exclusively in connection with a Qualifying Research Project.
- 3.2 The fees set out in paragraph 3.4 apply in respect of each deployment, distribution or licence of a Data Analysis System, and vary according to the Non-Member Territory in which the deployment, distribution or licensing takes place.
- 3.3 If any Data Analysis System to which the fees in paragraph 3.4 apply consists of more than one database, the fees applicable to that Data Analysis System shall be multiplied by the number of databases in that Data Analysis System.
- 3.4 The fees under this paragraph 3 are as follows:

Non-Member Territory (fee bands)	Associated License Fee
Band A Territory	US\$ 1,500 per annum baseline fee adjusted as per paragraph 1.4
Band B Territory	US\$ 1,000 per annum baseline fee adjusted as per paragraph 1.4
Band C Territory	US\$ 500 per annum baseline fee adjusted as per paragraph 1.4
Hospital or Practice in Low Income Band	US\$ 0 per annum baseline fee adjusted as per paragraph 1.4
Other territory	As per paragraph 5.2.

4. Other Activities

- 4.1 The Licensee shall notify the Licensor in writing before deploying the International Release or distributing or licensing Licensee Products (in each case, other than exclusively in connection with Qualifying Research Projects) in any Non-Member Territory in a manner that does not fall within paragraphs 2 to 4 of this Appendix B, explaining the Licensee's proposed activities.
- 4.2 Upon receiving notice from the Licensee under this paragraph 5, the Licensor may request, and the Licensee shall provide, such additional information in relation to the Licensee's proposed activities as the Licensor considers reasonably necessary to determine an appropriate licence and reasonable fee in respect of the Licensee's proposed activities.
- 4.3 The Licensee shall be liable to pay such licence fees as the Licensor may determine in accordance with this paragraph 5.

5. Non-Member Territory Bandings

- 5.1 The allocation of a Non-Member Territory into Band A, Band B or Band C shall be as determined by the Licensor (based on the Non-Member Territory's relative Gross National Income (GNI) or other measure adopted by the Licensor) and published by the Licensor on its web site.
- 5.2 The Licensee shall notify the Licensor in writing before carrying out any activity of a kind described in paragraphs 2 or 3 of this Appendix B in a Non-Member Territory that has not been allocated by the Licensor under paragraph 5.1. Upon receiving notice from the Licensee under this paragraph 5.2, the Licensor shall allocate the Non-Member Territory as described in paragraph 5.1.

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