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

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

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CCS_10_2018 - Healthcare Cost and Utilization Project (HCUP); HCUP Clinical Classifications Software (CCS); October 2017; Rockville, MD; http://www.hcup-us.ahrq.gov/toolsofsoftware/ccs10/ccs10.jsp; ENG;

Contact: Agency for Healthcare Research and Quality; 5600 Fishers Lane; Mail Stop 07N94A; Rockville, MD; United States; 20857; 1-866-290-HCUP; (301) 594-1430; hcup@ahrq.gov; https://www.hcup-us.ahrq.gov/toolsofsoftware/ccs10/ccs10.jsp;

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CDT2019 (updated) - Code on Dental Procedures and Nomenclature 2019 (CDT 2019); 2019; Chicago, IL; American Dental Association;

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CHV2011_02 - Consumer Health Vocabulary; 2011; USA; ENG;

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CPM2003 - Medical Entities Dictionary (CPM); New York (NY); Columbia Presbyterian Medical Center; 2003; ENG;

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CPT2018 - Current Procedural Terminology (CPT); American Medical Association; 2018; Chicago, IL;

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CVX_2018_06_22_18_09_04 (updated) - National Center for Immunization and Respiratory Diseases; CVX Code Set; Atlanta, GA; June 22, 2018;

Contact: CDC, National Center for Immunization and Respiratory Diseases Immunization Information System Support Branch - Informatics; 1600 Clifton Road; Mailstop: E-62; Atlanta, GA; United States; 30333; iisinfo@cdc.gov; http://www2a.cdc.gov/vaccines/iis/iisstandards/vaccines.asp?rpt=cvx;

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DDB00 - Diseases Database 2000; London (England); Medical Object Oriented Software Enterprises Ltd.; 2000; May, 2000; http://www.diseasesdatabase.com/;

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Contact: Malcolm H. Duncan; Medical Object Oriented Software Enterprises Ltd; Unit 36c Maryat Square; Fulham; London; UK; SW6 6UA; 44 (0)20 7381 4220 Mobile: 07710 483088; mhduncan@compuserve.com; http://www.diseasesdatabase.com/

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DMDICD10_1995 - Internationale Klassifikation der Krankheiten 10 [German translation of ICD10]; Germany; Deutsches Institut fuer Medizinische Dokumentation und Information; 1998;

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Contact: Dr. Michael Schopen; Deutsches Institut fur Medizinische Dokumentation und Information (DIMDI); Postfach 420580, D-50899; Koln; Germany; 49-221-472-4252; 49-221-41-1429; schopen@dimdi.de;

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DMDUMD_1996 - Die Nomenklatur fuer Medizinprodukte UMDNS [German translation of UMDNS]; Germany; Deutsches Institut fuer Medizinische Dokumentation und Information; 1996;

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DRUGBANK5.0_2018_07_31 (updated) - DrugBank; 5.0; OMx Personal Health Analytics Inc.; July 31, 2018; Edmonton, Alberta, Canada; Wishart DS, Knox C, Guo AC, Shrivastava S, Hassanali M, Stothard P, Chang Z, Woolsey J. DrugBank: a comprehensive resource for in silico drug discovery and exploration. Nucleic Acids Res. 2006 Jan 1 34(Database issue): D668-72.;

Contact: OMx Personal Health Analytics Inc.; 301 - 10359 104 St.; Edmonton; Alberta; Canada; T5J 1B9; 312-440-7742; support@omx.io; http://www.drugbank.ca;
The American Medical Association's CPT™ codes in HCPCS have a Source Abbreviation of HCPT04. The American Dental Association's CDT codes in HCPCS have a Source Abbreviation of HCDT4.
HL7V3.0_2017_07 - Health Level Seven (HL7) Vocabulary; Ann Arbor, MI; July 2017; http://www.hl7.org/;

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

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HLREL_1998 - Dr. Henk Lamberts; University of Amsterdam; ICPC2E-ICD10 relationships from HLREL; 1998;

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Contact: Henk Lamberts; University of Amsterdam; H.Lamberts@AMC.UVA.NL;

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HPO2018_07_23 (updated) - The Human Phenotype Ontology (HPO); The Human Phenotype Ontology Consortium; July 23, 2018;

Contact: Dr. Peter N. Robinson; Professor of Computational Biology; The Jackson Laboratory for Genomic Medicine; 10 Discovery Drive; Farmington; CT; 06032; 860.837.2095 t, 860.990.3130 m; peter.robinson@jax.org; https://hpo.jax.org/app/contact;

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ICD10_1998 - International Statistical Classification of Diseases and Related Health Problems (ICD-10); 10th rev.; Geneva (Switzerland); World Health Organization; 1998;

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

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ICD10_2016 - International Statistical Classification of Diseases and Related Health Problems (ICD-10); Tenth Revision; Geneva, Switzerland; World Health Organization; 2016; ENG;

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

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ICD10PCS_2019 (updated) -...;Centers for Medicare and Medicaid Services;International Classification of Diseases, 10th Revision, Procedure Coding System;...;Baltimore, MD;2019;......

Contact: Pat Brooks;Senior Technical Advisor;Center for Medicare and Medicaid Services;7500 Security Blvd. C4-08-06;Baltimore;MD;United States;21244;...;patricia.brooks2@cms.hhs.gov;https://www.cms.gov/Medicare/Coding/ICD10/index.html

ICD10PCS_2018 -...;Centers for Medicare and Medicaid Services;International Classification of Diseases, 10th Revision, Procedure Coding System;...;Baltimore, MD;2018;......

Contact: Pat Brooks;Senior Technical Advisor;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/Medicare/Coding/ICD10/index.html

ICD9CM_2014 -...;National Center for Health Statistics (NCHS);ICD-9-CM: International Classification of Diseases, Ninth Revision, Clinical Modification;...;FY 2014 Medicare Addendum;United States;October 1, 2013;Baltimore, MD;......

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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.hhs.gov


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but have not yet been reviewed and approved by the producers of ICD-9-CM.

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

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**ICPC2EDUT_200203** - Hirs, W., H.W. Becker, C. van Boven, S.K. Oskam, I.M. Okkes, H. Lamberts.;;;;;International Classification of Primary Care 2E: 2nd ed. electronic. Dutch Translation;;;;2nd ed. ;Amsterdam;Department of General Practice, Academic Medical Center/University of Amsterdam, Dutch College of General Practitioners (NHG);March 2002;;;;;;;

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Contact: LOINC c/o Regenstrief Center for Biomedical Informatics;;The Regenstrief Institute, Inc;410 West 10th Street, Suite 2000;IU campus mail address:HS, 2000;Indianapolis;IN;United States;46202-3012;317-274-9300;1-317-274-9305;loinc@regenstrief.org;https://loinc.org/;

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Contact: Arthur Alberto Correa Treuherz; Terminologies and Classifications, Supervisor; TCS AFI BIREME PAHO WHO; Centro Latinoamericano y del Caribe de Informacion en Ciencias de la Salud; Rua Vergueiro, 1759 - 12. andar; Sao Paulo; SP; Brasil; 04101-000; 55 11 5576-9817;

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Contact: Chris Hui; U.S. National Library of Medicine; Bethesda; MD; United States; huic@mail.nlm.nih.gov;
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Contact: UMLS Support; National Library of Medicine; 8600 Rockville Pike; Bethesda; Maryland; United States; 20894; custserv@nlm.gov;

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Contact: Enterprise Vocabulary Services (EVS); National Cancer Institute; 240-276-5541, 1-888-478-4423 (toll free); ncicbiit@mail.nih.gov; https://cbiit.nci.nih.gov/support/support;
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Contact: Quick Medical Reference, First Databank; 1111 Bayhill Drive; San Bruno; CA; 94066;

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

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

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This release contains concepts created by the National Library of Medicine which express the meaning of a drug name in a normalized form. These concepts relate the names of orderable medications to a dose form and the components of those medications. For further discussion, see the article at:


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Contact: UMLS Support, National Library of Medicine; e-mail: NLM SNOMED CT Support

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SOP8 (updated) - ;;Public Health Data Standards Consortium (PHDSC), Standards Data Committee, Payer
Subcommittee;;Source of Payment Typology;;;Version 8;;PHDSC;;December, 2017;Baltimore, MD;;;;;;

Contact: Anna Orlova;PHDSC Executive Director;Public Health Data Standards Consortium (PHDSC);111 South Calvert Street;Suite 2700;Baltimore;MD;21202;410-385-5272;866-637-6526;aorlova@jhsph.edu;http://www.phdsc.org;

Read more information about this source

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

Contact: ;;;;;;;custserv@nlm.nih.gov;

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

Contact: UMLS Support;;;;;;custserv@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;kjh970203@kiom.re.kr;

Read more information about this source

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;(617) 732-6281;;;;greenes@harvard.edu;

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UMD2018 - ;;ECRI Institute;;The Universal Medical Device Nomenclature System;;;;Plymouth Meeting, PA;;2018;;;;;;

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 ext. 5891;1-610-834-1275;umdns@ecri.org;https://www.ecri.org/components/UMDNS/Pages/default.aspx;

Read more information about this source
USP 2018_08_27 (new) - USP Compendial Nomenclature; August, 2018; United States Pharmacopeia; August, 2018; 

Contact: Steven Emrick; Senior HIT Product Manager; United States Pharmacopeia; 12601 Twinbrook Parkway; Rockville; MD; US; 20852-1790; 1-800-227-8772; HealthcareQuality@usp.org; http://www.usp.org/health-quality-safety

Read more information about this source

USPMG 2017 - USP Medicare Model Guidelines; Version 7; United States Pharmacopeia; February 6, 2017; 

Contact: Donna Bohanon, RPh.; Scientific Liaison; United States Pharmacopeia; 12601 Twinbrook Parkway; Rockville; MD; US; 20852-1790; 1-800-227-8772; ModelGuidelines@usp.org; http://www.usp.org/usp-healthcare-professionals/usp-medicare-model-guidelines

Read more information about this source

UWDA173 - Jose Mejino, M.D.; University of Washington Digital Anatomist, (UWDA); Version 1.7.3; Seattle (WA); University of Washington; March, 2003;

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

Read more information about this source

VANDF 2018_08_01 (updated) - Veterans Health Administration National Drug File; August 01, 2018; 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; michael.lincoln@med.va.gov; http://www.pbm.va.gov/default.aspx

Read more information about this source

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; +46-18-656080;

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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 License Agreement;

2.5.3 not permit the End User to sub-license or transfer any of its rights under the sub-license (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-license with the Licensee, subject to the same restrictions as apply to sub-licensing the International Release under the Affiliate's license agreement with the Licensor);

2.5.4 terminate automatically upon termination of this License Agreement;

2.5.5 provide that the End User may apply directly to the Licensor upon receiving notice that the sub-license 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 license 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-license 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-license to the Licensor in accordance with clauses 7 and 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 License 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 License 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 license back to the Licensee from the Licensor or will procure from the Member a license 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 License 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 Licensor 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. If 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 License Agreement shall commence on the date on which it comes into effect in accordance with the notice at the beginning of this License Agreement, and shall continue until terminated in accordance with this clause 5.

5.2 Either party may terminate this License Agreement if the other party commits a material breach of any of its obligations under this License Agreement (which, in the case of the Licensee, shall include, without limitation, any failure to pay License Fees when due under clause 7) 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 Neither party may terminate this License Agreement except in accordance with this clause 5.

5.4 The Licensee may terminate this License Agreement by giving up to twelve (12) months' prior written notice to the Licensor.

5.5 Upon termination of this License Agreement in accordance with this clause 5, all licenses granted under this License Agreement shall automatically and immediately be revoked.

5.6 The Licensee shall, by no later than forty five (45) days after termination of this License 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.

5.7 The Licensee shall, as soon as reasonably practicable following either party giving a Termination Notice for any reason, and in any event by no later than ninety (90) days after such Termination Notice is given, give written notice of such termination to each End User that the Licensee reasonably believes to be a current user of a Licensee Product and
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5.8 The Licensee may not grant any new sub-license under clause 2.1.5 after either party has given notice under clauses 5.2 or 5.4.

5.9 The Licensor shall be entitled to publicize the termination of this License 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, 5.11, 5.12, 7, 8 and 10 to 14 inclusive shall survive termination of this License Agreement.

5.11 The Licensee shall, by no later than thirty (30) days after termination of this License Agreement for any reason, submit a statement of account in accordance with clause 7.3 in respect of all periods that have not previously been covered by a statement of account under that clause.

5.12 Any termination of this License Agreement, for any reason, is without prejudice to the accrued liabilities of each party as at the date of termination (including, without limitation, any liability of the Licensee to pay License Fees that has accrued as at the date of termination), or to the Licensee's obligation to pay License Fees arising from the statement of account submitted under clause 5.11.

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6.3 The Licensor may vary the terms of this License 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 License Agreement to continue subject to the variation, the Licensee may terminate this License Agreement in accordance with clause 5.4, and if the Licensee gives notice of such termination before the variation takes effect then the variation shall not apply as between the Licensor and the Licensee.

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7.2 All License Fees and other amounts payable to the Licensor under this License 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, at least once in each calendar year, submit a statement of account to the Licensor in such manner and form as the Licensor may prescribe from time to time, setting out the Licensee's activities in Non-Member
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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 License 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 from time to time.

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9.2 Conditions prescribed by a Member under clause 9.1 may:

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9.2.2 relate to the International Release, the Member's National Release or any part of either of them.

9.3 The Licensee shall notify the Licensor (and, if the Licensee's registered office or principal place of business is situated in a Member Territory, shall also notify the Member for that Member Territory) in writing before exercising its rights under this License Agreement in any Non-Member Territory in respect of which the Licensee has not previously given notice under this clause 9.3. The notice shall be in such form and manner as the Licensor may prescribe from time to time, and shall include such information about the Licensee's current and proposed activities in that Non-Member Territory as the Licensor may require (but the Licensor may require only the same kinds of information as it requires to be provided by new Affiliates proposing to use, license or deploy the International Release or Licensee Products in Non-Member Territories).
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10.2 As an Affiliate, the Licensee shall be entitled to participate in the Licensor's Vendor Liaison Forum, which is a forum in which the Licensee and other Affiliates may communicate with the Licensor and with each other. The Licensor may make Regulations from time to time governing the Licensee's participation in the Vendor Liaison Forum. New Regulations that the Licensor shall make from time to time governing participation in the Vendor Liaison Forum shall not remove the Licensee's right to participate in that forum.

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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 License Agreement, whether in contract, tort (including negligence), misrepresentation, breach of statutory duty or otherwise, shall not in any event exceed the License Fees paid by the Licensee in respect of that year.

12.4 Nothing in this License 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 License 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 License Agreement to any person to whom the Licensor transfers the Intellectual Property Rights in respect of which the licenses under this License Agreement are granted.

14. GENERAL PROVISIONS

14.1 This License Agreement contains the entire agreement between the parties relating to the subject matter of this License 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 License Agreement, it has not relied on any representation, warranty, collateral contract or other assurance made by or on behalf of the other party before the date of this License Agreement.

14.3 Except as provided in clause 6.3, this License Agreement may not be varied except in writing signed by both parties and expressed to vary this License Agreement.

14.4 Nothing in this License 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 License 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 License Agreement, or the legality, validity or enforceability in any other jurisdiction of that or any other term of this License Agreement.

14.6 The Licensee agrees that the Licensor may appoint third parties to process personal data provided by the Licensee to the Licensor under or in connection with this License Agreement (including without limitation payment details provided in connection with the payment of License Fees). In connection with any such appointment, personal data provided by the Licensee may be transferred to, and processed in, a country outside the European Economic Area (EEA). The laws governing the processing of personal data may be less stringent in such a country than in the member countries of the EEA.

15. GOVERNING LAW AND JURISDICTION

15.1 This License 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 License 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 License Agreement, the following defined terms have the following meanings:

<table>
<thead>
<tr>
<th>Term</th>
<th>Meaning</th>
</tr>
</thead>
<tbody>
<tr>
<td>Affiliate</td>
<td>an affiliate of the Licensor in accordance with the Licensor's Articles of Association;</td>
</tr>
<tr>
<td>Cross-Map</td>
<td>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);</td>
</tr>
<tr>
<td>Data Analysis System</td>
<td>a computer system that is used to analyze records or other data that is encoded using SNOMED CT, but not if that system is also a Data Creation System;</td>
</tr>
<tr>
<td>Data Creation System</td>
<td>a computer system that is used to create records or other data that is encoded using SNOMED CT;</td>
</tr>
<tr>
<td>Derivative</td>
<td>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;</td>
</tr>
<tr>
<td>End User</td>
<td>a third party user of a Licensee Product;</td>
</tr>
<tr>
<td>Extension</td>
<td>a work consisting of SNOMED CT Content alone that is supplementary to the SNOMED CT Core and that depends on the SNOMED CT Core;</td>
</tr>
<tr>
<td>Hospital</td>
<td>a health care body or organisation providing secondary and/or tertiary care;</td>
</tr>
<tr>
<td>Intellectual Property Rights</td>
<td>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;</td>
</tr>
<tr>
<td>International Release</td>
<td>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;</td>
</tr>
<tr>
<td>License Fees</td>
<td>the license fees set out in Appendix B (License Fees in Non-Member Territories);</td>
</tr>
<tr>
<td>Licensee Products</td>
<td>products distributed or licensed by the Licensee that(1) include or interoperate with the International Release (or any part of it) and/or any Extensions or Derivatives created by the Licensee under this License Agreement, or (2) read or write records or other data that is encoded using SNOMED CT;</td>
</tr>
<tr>
<td>Member</td>
<td>a member of the Licensor;</td>
</tr>
<tr>
<td>Member</td>
<td>a territory that is represented by a Member (as published by the Licensor from time to time);</td>
</tr>
<tr>
<td><strong>Territory</strong></td>
<td><strong>Namespace Identifier</strong></td>
</tr>
<tr>
<td>---------------</td>
<td>--------------------------</td>
</tr>
<tr>
<td></td>
<td>a code or that part of a code that identifies the organization responsible for creating and maintaining a Standards-Based Extension or a Standards-Based Derivative and is used as an element of SNOMED CT Identifiers;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>National Release</strong></th>
<th><strong>Non-Member Territory</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>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;</td>
<td>a territory that is not a Member Territory;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Practice</strong></th>
<th><strong>Qualifying Research Project</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>(a) a single department of a Hospital (subject to paragraph 2.2 of Appendix B); or (b) any health care body or organisation that provides principally primary care, including without limitation a pharmacy, an optician's facility, a physiotherapy centre, a general medical practice or a family medical practice;</td>
<td>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; (d) the results of the research are offered for publication in peer-reviewed public journals and are provided to the Licensor free of charge prior to publication;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Regulations</strong></th>
<th><strong>Relationship</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>regulations made by the Licensor;</td>
<td>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;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>SNOMED CT</strong></th>
<th><strong>SNOMED CT Content</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>the concept-based work of clinical nomenclature and classification with multiple hierarchies and semantic definitions known as SNOMED Clinical Terms (SNOMED CT);</td>
<td>terminological content, consisting of concepts, descriptions and Relationships, each of which is identified using a SNOMED CT Identifier;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>SNOMED CT Core</strong></th>
<th><strong>SNOMED CT Identifier</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>the SNOMED CT Content that is controlled, maintained and distributed by the Licensor from time to time;</td>
<td>a code, of a kind defined by the Licensor in Specifications, for identifying concepts, descriptions and Relationships;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Specification</strong></th>
<th><strong>Sponsored Territory</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>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;</td>
<td>a Non-Member Territory that has been recognized and designated by the Licensor as a sponsored territory (as published on the Licensor's web site);</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Standard</strong></th>
<th>****</th>
</tr>
</thead>
<tbody>
<tr>
<td>a Specification that is formally adopted by the Licensor;</td>
<td>****</td>
</tr>
</tbody>
</table>
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; and

Sub-Set

a sub-set of SNOMED CT Content that is grouped together for one or more purposes.

Appendix B

License Fees in Non-Member Territories

1. Introduction

1.1 This Appendix B sets out the license fees payable by the Licensee in respect of its activities in Non-Member Territories.

1.2 The license 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 license 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 License Agreement and shall not relieve the Licensee of any of its other obligations under this License Agreement.

1.4 Beginning in 2015, license fees payable by the Licensee in respect of its activities in Non-Member Territories for each financial year shall be adjusted by the same percentage as the General Assembly of the Licensor agrees to adjust the Aggregate Annual Fee (as defined in the Licensor's Articles of Association) relative to the Aggregate Annual Fee in the previous year.

1.5 The license fees in respect of Hospitals that are set out in this Appendix B apply only to Hospitals that are located on a single contiguous physical site. Any Hospital that is located on multiple physical sites shall be treated as falling within paragraph 4 of this Appendix B (and not within paragraphs 2 or 3).

1.6 For purposes of this Appendix B, if a Practice is located on multiple physical sites then each such site is treated as a separate Practice.

1.7 Notwithstanding anything else in this Appendix B, the deployment, distribution or licensing of any software that operates on a mobile device of any kind (including without limitation a mobile phone or tablet device), or any software or service that is accessed via the internet and enables users to extract or download any substantial portion of SNOMED CT, shall be treated as falling within paragraph 4 of this Appendix B (and not within paragraphs 2 or 3).

1.8 The Licensee's obligation to pay license fees in respect of any deployment of the International Release or any Licensee Product is not dependent on that deployment of the International Release or Licensee Product being used in a live or production environment.

1.9 In any case where the Licensee is exempt from the requirement to pay license fees by reason of a Licensee Product, a Data Analysis System or a Data Creation System being used exclusively in connection with a Qualifying Research Project, the Licensee shall report to the Licensor on the progress of that Qualifying Research Project in such manner as the Licensor may reasonably require. The Licensor may revoke the Licensee's exemption for Qualifying Research Projects provided in this Appendix B if the Licensee fails to comply with this paragraph 1.9.

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 part of it) 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) deploys, 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.

<table>
<thead>
<tr>
<th>Fee Band</th>
<th>Fee</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hospital in Band A Territory</td>
<td>US$ 1,772 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Hospital in Band B Territory</td>
<td>US$ 1,182 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Hospital in Band C Territory</td>
<td>US$ 591 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Practice in Band A, B or C Territory</td>
<td>US$ 591 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Hospital or Practice in Low Income Band</td>
<td>US $0 per annum baseline fee, adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Hospital or Practice in other territory</td>
<td>As per paragraph 5.2.</td>
</tr>
</tbody>
</table>

2.2 The total fees payable by the Licensee in respect of a number of Practices that are departments of a single Hospital shall not exceed the fee applicable to the Hospital itself. For purposes of this Appendix B, a Practice is treated as a department of a Hospital only if: (a) it is located on the premises of that Hospital; and (b) it is funded solely by that Hospital. In any case where either or both of the conditions in the preceding sentence are not met in respect of any Practice, fees shall be payable in respect of that Practice in addition to any fees that are payable in respect of any Hospital.

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 part of it) 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) deploys, 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 license of the International Release (or any part of it), a Licensee Product or 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:

<table>
<thead>
<tr>
<th>Fee Band</th>
<th>Fee</th>
</tr>
</thead>
<tbody>
<tr>
<td>Band A Territory</td>
<td>US$ 1,772 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>------------------</td>
<td>-------------------------------------------------------------</td>
</tr>
<tr>
<td>Band B Territory</td>
<td>US$ 1,182 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Band C Territory</td>
<td>US$ 591 per annum baseline fee adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Low Income Band</td>
<td>US$ 0 per annum baseline fee, adjusted as per paragraph 1.4</td>
</tr>
<tr>
<td>Other territory</td>
<td>As per paragraph 5.2.</td>
</tr>
</tbody>
</table>

4. **Other Activities**

4.1 The Licensee shall notify the Licensor in writing before deploying the International Release (or any part of it) or deploying, distributing or licensing any Licensee Product (in each case, other than exclusively in connection with Qualifying Research Projects) in, for use in, or to any person situated in, any Non-Member Territory in a manner that does not fall within paragraphs 2 to 3 of this Appendix B, explaining the Licensee's proposed activities.

4.2 Upon receiving notice from the Licensee under this paragraph 4, 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 license and reasonable fee in respect of the Licensee's proposed activities.

4.3 The Licensee shall be liable to pay such license fees as the Licensor may determine in accordance with this paragraph 4.

5. **Non-Member Territory Bandings**

5.1 The allocation of a Non-Member Territory into Band A, Band B, Band C, or Low Income Band 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.