

September 15-16, 2010 Diagnosis Agenda, Part 1 of 2

Welcome and announcements
Donna Pickett, MPH, RHIA
Co-Chair, ICD-9-CM Coordination and Maintenance Committee

Diagnosis Topics:

Infection Following Transfusion	9
Mikhail Menis, PharmD, MS	
Center for Biologics Evaluation & Research (CBER)	
Food and Drug Administration (FDA)	
Anaphylactic Reaction, and Other Serum Reaction	10
Mikhail Menis, PharmD, MS	
Center for Biologics Evaluation & Research (CBER)	
Food and Drug Administration (FDA)	
Mesh erosion/Mesh exposure	16
Sage Claydon, MD	
American College of Obstetricians and Gynecologists (ACOG)	
Malnutrition	18
Gordon L. Jensen, MD, Ph,D	
Penn State University	
Co-Chair American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.) Task Force	
Lymphangioleiomyomatosis	23
Francis X. McCormack, MD	
University of Cincinnati	
Representing the American Thoracic Society	
Dementia unspecified with and without behavioral disturbance	24
Elective c-sections prior to 39 weeks	25
Jon Hathaway, MD	
American College of Obstetricians and Gynecologists (ACOG)	
Personal history of gestational diabetes	26
Encounter for fetal viability ultrasound/personal history of ectopic pregnancy	27

Adult Pulmonary Langerhans Cell Histiocytosis (PLCH)	28
Francis X. McCormack, MD	
University of Cincinnati	
Representing the American Thoracic Society	
Acquired absence of joint	30
Glaucoma severity staging	32
Corticobasal degeneration	36
Pulmonary Arteriovenous Malformation and Pulmonary Atresia	37
Complications of stem cell transplant	39
Pseudobulbar Affect	40
Reportable malignant skin cancers	42
ICD-9-CM Proposed Addenda	45
ICD-10-CM	
Opioids expansion for ICD-10-CM	53
ICD-10-CM Weeks of gestation of pregnancy	63
ICD-10-CM Weeks of gestation for newborn	66
Benign neoplasm of genitourinary organs	68
Urethral False Passage	69
Nodular prostate	70
Inflammatory disease of the prostate	71
Cyst of the prostate	72
Acquired and congenital torsion of the penis	73
Cyst of the epididymis	74
Hidden penis	75
Personal history of malignant neoplasm of ureter	76
Visual agnosia and related conditions (ICD-10-CM only)	77
Displacement/dislocation of internal hip prosthesis titles (ICD-10-CM Only)	78
Gastroparesis	79
ICD-10-CM Proposed Addenda	80

ICD-9-CM TIMELINE

A timeline of important dates in the ICD-9-CM process is described below:

September	15 –	16,
2010		

ICD-9-CM Coordination and Maintenance Committee meeting.

Those who wish to attend the ICD-9-CM Coordination and Maintenance Committee meeting **must have registered for the meeting online by September 10, 2010.** You must bring an official form of picture identification (such as a drivers license) in order to be admitted to the building.

October 2010

Summary report of the Procedure part of the September 15 – 16, 2010 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

Summary report of the Diagnosis part of the September 15–16, 2010 ICD-9-CM Coordination and Maintenance Committee meeting report will be posted on NCHS homepage as follows: http://www.cdc.gov/nchs/icd.htm

October 1, 2010

New and revised ICD-9-CM codes go into effect along with DRG changes. Final addendum posted on web pages as follows:

Diagnosis addendum - http://www.cdc.gov/nchs/icd.htm
Procedure addendum at http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

October 8, 2010

Deadline for receipt of public comments on proposed code revisions discussed at the September 15-16, 2010 ICD-9-CM Coordination and Maintenance Committee meetings for implementation on April 1, 2011.

November 2010

Any new ICD-9-CM codes required to capture new technology that will be implemented on the following April 1 will be announced. Information on any new codes to be implemented April 1, 2011 will be posted on the following websites: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes http://www.cdc.gov/nchs/icd.htm

November 12, 2010 Deadline for receipt of public comments on revisions to the General Equivalence Maps (GEMs) discussed at the September

15, 2010 ICD-9-CM Coordination and Maintenance Committee.

November 19, 2010 Deadline for receipt of public comments on proposed code

revisions discussed at the September 15-16, 2010 ICD-9-CM Coordination and Maintenance Committee meetings for

implementation on October 1, 2011.

January 7, 2011 Deadline for requestors: Those members of the public requesting

that topics be discussed at the March 9 – March 10, 2011 ICD-9-CM Coordination and Maintenance Committee meeting must have their requests to CMS for procedures and NCHS for

diagnoses by this date.

February 2011 Draft agenda for the Procedure part of the March 9, 2011 ICD-9-

CM Coordination and Maintenance Committee meeting posted on

CMS homepage as follows:

http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

Draft agenda for the Diagnosis part of the March 10, 2011 ICD-9-CM Coordination and Maintenance Committee meeting posted on

NCHS homepage as follows:

http://www.cdc.gov/nchs/icd.htm

Federal Register notice of March 9 – March 10, 2011 ICD-9-CM Coordination and Maintenance Committee Meeting will be

published.

February 11, 2011 On-line registration opens for the March 9 – 10, 2011

ICD-9-CM Coordination and Maintenance Committee meeting at:

http://www.cms.hhs.gov/apps/events

March 2011 Because of increased security requirements, those wishing to

attend the March 9 – March 10, 2011 ICD-9-CM Coordination and Maintenance Committee meeting must register for the meeting

online at:

http://www.cms.hhs.gov/apps/events

Attendees must register online by March 4, 2011 failure to do so

may result in lack of access to the meeting.

March 9 – 10 ICD-9-CM Coordination and Maintenance Committee 2011 meeting. April 1, 2011 Any new ICD-9-CM codes required to capture new technology will be implemented. Information on any new codes implemented on April 1, 2011 previously posted in early November 2010 will be on the following websites: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes http://www.cdc.gov/nchs/icd.htm http://www.cms.hhs.gov/MLNGenInfo April 1, 2011 Deadline for receipt of public comments on proposed code revisions discussed at the March 9-10, 2011 ICD-9-CM Coordination and Maintenance Committee meetings for implementation on October 1, 2011. Notice of Proposed Rulemaking to be published in the Federal April 2011 Register as mandated by Public Law 99-509. This notice will include the final ICD-9-CM diagnosis and procedure codes for the upcoming fiscal year. It will also include proposed revisions to the DRG system on which the public may comment. The proposed rule can be accessed at: http://www.cms.hhs.gov/AcuteInpatientPPS/IPPS/list.asp April 2011 Summary report of the Procedure part of the March 9, 2011 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes Summary report of the Diagnosis part of the March 10, 2011 ICD-9-CM Coordination and Maintenance Committee meeting report will be posted on NCHS homepage as follows: http://www.cdc.gov/nchs/icd.htm June 2011 Final addendum posted on web pages as follows: Diagnosis addendum at - http://www.cdc.gov/nchs/icd.htm Procedure addendum at http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes July 15, 2011 Those members of the public requesting that topics be discussed at the September 14 - 15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting must have their requests to CMS for procedures and NCHS for diagnoses.

August 1, 2011

Hospital Inpatient Prospective Payment System final rule to be published in the Federal Register as mandated by Public Law 99-509. This rule will also include all the final codes to be

implemented on October 1, 2011.

This rule can be accessed at:

http://www.cms.hhs.gov/AcuteInpatientPPS/IPPS/list.asp

August 2011

Tentative agenda for the Procedure part of the September 14 – 15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage at - http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

Tentative agenda for the Diagnosis part of the September 14 – 15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on NCHS homepage at - http://www.cdc.gov/nchs/icd.htm

Federal Register notice for the September 14 –15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting will be published. This will include the tentative agenda.

August 12, 2011

On-line registration opens for the September 14-15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting at: http://www.cms.hhs.gov/apps/events

September 9, 2011

Because of increased security requirements, those wishing to attend the September 14 - 15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting must register for the meeting online at:

http://www.cms.hhs.gov/apps/events

Attendees must register online by September 9, 2011; failure to do so may result in lack of access to the meeting.

September 14 –15, 2011

ICD-9-CM Coordination and Maintenance Committee meeting.

Those who wish to attend the ICD-9-CM Coordination and Maintenance Committee meeting must have registered for the meeting online by September 9, 2011. You must bring an official form of picture identification (such as a drivers license) in order to be admitted to the building.

\triangle	h ~ "	20	11
Octo	ber	ZU	тт

Summary report of the Procedure part of the September 14 – 15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

Summary report of the Diagnosis part of the September 14–15, 2011 ICD-9-CM Coordination and Maintenance Committee meeting report will be posted on NCHS homepage as follows: http://www.cdc.gov/nchs/icd.htm

October 1, 2011

New and revised ICD-9-CM codes go into effect along with DRG changes. Final addendum posted on web pages as follows:

Diagnosis addendum - http://www.cdc.gov/nchs/icd.htm
Procedure addendum at http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes

October 7, 2011

Deadline for receipt of public comments on proposed code revisions discussed at the September 14-15, 2011 ICD-9-CM Coordination and Maintenance Committee meetings for implementation on April 1, 2012.

November 2011

Any new ICD-9-CM codes required to capture new technology that will be implemented on the following April 1 will be announced. Information on any new codes to be implemented April 1, 2012 will be posted on the following websites: http://www.cms.hhs.gov/ICD9ProviderDiagnosticCodes http://www.cdc.gov/nchs/icd.htm

November 18, 2011

Deadline for receipt of public comments on proposed code revisions discussed at the September 14-15, 2011 ICD-9-CM Coordination and Maintenance Committee meetings for implementation on October 1, 2012.

Contact Information

Mailing address:

National Center for Health Statistics ICD-9-CM Coordination and Maintenance Committee 3311 Toledo Road, Room 2402 Hyattsville, Maryland 20782

Fax: (301) 458-4022

Donna Pickett (301) 458-4434

E-mail: dfp4@cdc.gov

David Berglund (301) 458-4095

E-mail zhc2@cdc.gov

Lizabeth Fisher (301) 458-4091

E-mail <u>llw4@cdc.gov</u>

Traci Ramirez (301) 458-4454

E-mail tfr4@cdc.gov

NCHS Classifications of Diseases web page:

http://www.cdc.gov/nchs/icd9.htm

Please consult this web page for updated information

Infection Following Transfusion

A previous proposal was presented Sept. 2009 to create a code for transfusion-transmitted infections at subcategory 999.3. However, concerns were raised at that time related to the existing use additional code note at 999.3, in that it would imply that this code should precede the code for HIV disease.

As previously described, there are a number of infectious organisms (including bacteria, viruses, and parasites, among others) that may be transmitted through transfusion of blood or blood products (which include whole blood, RBCs, plasma, and platelets, among others). To address previous concerns, it is proposed to add a note to make it explicit that HIV disease is to be coded first, if present.

Currently there are no specific ICD-9-CM diagnosis codes for infections following transfusion. A request was received from the Food and Drug Administration (FDA) Center for Biologics Evaluation and Research (CBER) to create a unique code for infection following transfusion, as proposed here.

References

U.S. Biovigilance Network (http://www.cdc.gov/nhsn/PDFs/hemovigModuleProtocol_current.pdf)
Canadian transfusion safety surveillance system
(http://www.phac-aspc.gc.ca/hcai-iamss/tti-it/index-eng.php)

TABULAR MODIFICATIONS

042 Human immunodeficiency virus [HIV] disease

Use additional code, if applicable, to identify infection following transfusion, infusion, or injection of blood and blood products (999.32)

999 Complications of medical care, not elsewhere classified

999.3 Other infection

Add	Code first, if applicable, human immunodeficiency virus (HIV) disease (042)		
New code	999.32	Infection following transfusion, infusion, or injection of blood and blood products	
Revise	999.39	Infection following other infusion, injection, transfusion, or vaccination	

Anaphylactic Reaction, and Other Serum Reaction

An anaphylactic reaction, or anaphylaxis, is a type of allergic hypersensitivity reaction, which may be caused by a number of different allergens. It may cause itching, hives, and wheezing, among other symptoms, and in severe cases, can cause a drop in blood pressure, which is anaphylactic shock.

In ICD-9-CM, codes for anaphylactic reactions have been titled with the term anaphylactic shock, with inclusion terms and indexing making it clear that all such reactions are included, even without shock. However, this titling is somewhat confusing for clinicians. Therefore, it has been proposed that these codes be retitled, to make it more clear what is included. Parts of these changes were proposed independently by the American Academy of Pediatrics, and by the Food and Drug Administration (FDA) Center for Biologics Evaluation and Research (CBER).

It was proposed by FDA CBER to add specific codes for anaphylactic reactions due to administration of blood and blood products, and for anaphylactic reactions due to vaccination.

Anaphylactic reactions are a type I hypersensitivity reaction, involving IgE antibodies, and occuring immediately with exposure. Other serum reactions can also occur, including type II and type III hypersensitivity reactions, which both involve IgG antibodies.

There are certain types of transfusion reactions and drug reactions that exemplify type II hypersensitivity reactions. Examples of type III hypersensitivity reactions include serum sickness and Arthus reaction. Serum sickness is a hypersensitivity reaction to a protein in serum, generally occuring one to three weeks after exposure. It may involve fever, itching, and rash, potentially with other manifestations, some of which include joint pain and swollen lymph nodes.

It was also proposed by FDA CBER to add specific codes for other serum reactions due to administration of blood and blood products, and for other serum reactions due to vaccination.

TABULAR MODIFICATIONS

713 Arthropathy associated with other disorders classified elsewhere

713.6 Arthropathy associated with hypersensitivity reaction

Code first underlying disease, as: serum sickness (999.5<u>1-999.59</u>)

Revise

785 Symptoms involving cardiovascular system

785.5 Shock without mention of trauma

785.59 Other

Excludes: shock (due to):

Revise due to serum (999.4<u>1-999.49</u>)

958 Certain early complications of trauma

958.4 Traumatic shock

Excludes: shock:

Revise due to serum (999.4<u>1-999.49</u>)

995 Certain adverse effects not elsewhere classified

Revise 995.0 Other anaphylactic shock_reaction

Add Anaphylactic shock NOS or due to adverse effect of correct medicinal

substance properly administered

Revise Excludes: anaphylactic reaction to serum (999.4<u>1-999.49</u>)

Revise anaphylactic shock <u>or reaction</u> due to adverse food reaction (995.60-

995.69)

995.1 Angioneurotic edema

Excludes: urticaria:

Revise due to serum (999.5<u>1-999.59</u>)

Revise 995.6 Anaphylactic shock reaction due to adverse food reaction
Revise Anaphylactic reaction shock due to adverse food reaction
Revise Anaphylactic shock or reaction due to nonpoisonous foods

Revise 995.60 <u>Anaphylactic reaction</u> <u>Dd</u>ue to unspecified food

Revise 995.61 <u>Anaphylactic reaction</u> <u>Ddue to peanuts</u>

Revise 995.62 <u>Anaphylactic reaction</u> <u>Pd</u>ue to crustaceans

Revise 995.63 <u>Anaphylactic reaction Pdue to fruits and vegetables</u>

Revise 995.64 <u>Anaphylactic reaction</u> <u>Ddue</u> to tree nuts and seeds

Revise 995.65 Anaphylactic reaction Ddue to fish

Revise 995.66 <u>Anaphylactic reaction</u> <u>Pd</u>ue to food additives

Revise 995.67 <u>Anaphylactic reaction</u> <u>Pdue to milk products</u>

Revise 995.68 <u>Anaphylactic reaction</u> <u>Ddue to eggs</u>

Revise 995.69 <u>Anaphylactic reaction</u> <u>Pdue to other specified food</u>

995.7 Other adverse food reactions, not elsewhere classified

Revise Excludes: anaphylactic <u>reaction or</u> shock due to adverse food reaction (995.6)

998 Other complications of procedures, NEC

998.0 Postoperative shock

Excludes: shock:

Revise anaphylactic due to serum (999.4<u>1-999.49</u>)

999 Complications of medical care, not elsewhere classified

Revise 999.4 Anaphylactic shock reaction due to serum

Add Allergic shock

Revise Anaphylactic reaction shock due to serum

Add Anaphylaxis

Add Excludes: reaction:

Add allergic NOS (995.0)
Add anaphylactic:
Add NOS (995.0)

Add due to drugs and chemicals (995.0)

New Code 999.41 Anaphylactic reaction due to administration of blood and blood

products

New Code 999.42 Anaphylactic reaction due to vaccination

New Code 999.49 Anaphylactic reaction due to other serum

999.5 Other serum reaction

New Code 999.51 Other serum reaction due to administration of blood and blood

products

New Code 999.52 Other serum reaction due to vaccination

New Code 999.59 Other serum reaction

INDEX MODIFICATIONS

Allergy, allergic (reaction) 995.3 anaphylactic reaction or shock 995.0 Revise due to food - see Anaphylactic reaction or shock, due to, food Revise serum (prophylactic) (therapeutic) 999.5 anaphylactic reaction or shock 999.4 shock (anaphylactic) (due to adverse effect of correct medicinal substance properly administered) 995.0 Revise food - see Anaphylactic reaction or shock, due to, food from serum or immunization 999.5 anaphylactic 999.4 Anaphylactic reaction or shock or reaction (correct substance properly Revise administered) 995.0 due to Add chemical - see Table of Drugs and Chemicals Add correct medicinal substance properly administered 995.0 Add drug or medicinal substance Add correct substance properly administered 995.0 Add overdose or wrong substance given or taken 977.9 Add specified drug - see Table of Drugs and Chemicals Add following sting(s) 989.5 food 995.60 additives 995.66 crustaceans 995.62 eggs 995.68 fish 995.65 fruits 995.63 milk products 995.67 nuts (tree) 995.64 peanuts 995.61 seeds 995.64 specified NEC 995.69 tree nuts 995.64 vegetables 995.63 immunization 999.4 overdose or wrong substance given or taken 977.9 specified drug - see Table of Drugs and Chemicals serum 999.4 serum 999.4

Anaphylactoid <u>reaction or</u> shock <u>or reaction</u> - see Anaphylactic <u>reaction or</u> shock Anaphylaxis - see Anaphylactic <u>reaction or</u> shock

Arthritis, arthritic (acute) (chronic) (subacute) 716.9 due to or associated with

serum sickness 999.5 [713.6] Arthritis, cont. serum (nontherapeutic) (therapeutic) 999.5 [713.6] Arthus phenomenon 995.21 due to serum 999.5 Complications vaccination 999.9 anaphylaxis NEC 999.4 protein sickness 999.5 reaction (allergic) 999.5 serum 999.5 serum intoxication, sickness, rash, or other serum reaction NEC 999.5 shock (allergic) (anaphylactic) 999.4 Effect, adverse NEC foodstuffs anaphylactic reaction or shock due to food NEC - see Anaphylactic reaction or shock, due to food 995.60 Food anaphylactic shock - see Anaphylactic reaction or shock, due to food History (personal) of anaphylactic reaction or shock V15.09 anaphylactic - see Shock, anaphylactic Anaphylactic reaction food - see also Allergy, food anaphylactic shock - see Anaphylactic reaction or shock, due to, food serum (prophylactic) (therapeutic) 999.5 immediate 999.4 Shock 785.50 allergic - see Shock, anaphylactic Anaphylactic reaction and shock anaphylactic 995.0 - see Anaphylactic reaction and shock chemical - see Table of Drugs and Chemicals correct medicinal substance properly administered 995.0 drug or medicinal substance correct substance properly administered 995.0 overdose or wrong substance given or taken 977.9 specified drug - see Table of Drugs and Chemicals following sting(s) 989.5 food - see Anaphylactic shock, due to, food immunization 999.4

Revise

Revise

Revise

Revise

Delete

Delete Delete

Delete

Delete

Delete

Delete Delete

Delete

Delete serum 999.4
Revise anaphylactoid - see Shock, anaphylactic Anaphylactic reaction and shock

Mesh erosion/Mesh exposure

This proposal was presented at the September 2009 ICD-9-CM Coordination and Maintenance Committee meeting. There were comments made at the meeting and received in writing following the meeting concerning the definitions and use of these terms. Some indicated that the terms erosion and exposure may be used interchangeably. Others were concerned that the terms may have different meanings if these codes were going to be used for mesh complications in other surgical specialties (such as for hernia surgery). The original requestor, the American College of Obstetricians and Gynecologists (ACOG), has asked to have this proposal presented again to both help answer questions about the terms and to propose an option to locate these codes Chapter 10, Diseases of the Genitourinary System.

Previous proposal background statement:

An effective surgical treatment for vaginal vault prolapse is abdominal sacral colpopexy. In this procedure, a graft is used to suspend the upper vagina to the anterior longitudinal ligament of the sacrum. Synthetic graft material used to suspend the apex of the vagina to the anterior longitudinal ligament of the sacrum has been associated with mesh erosion and subsequent pelvic infection (due to the erosion into surrounding organs or tissue). Treatment for the erosion usually requires surgical removal of the mesh. Exposure of the mesh, into the vagina, can also occur. Though this is a less severe condition which can be treated sometimes nonsurgically it does have potential for infection to develop.

Previous coding advice has been to assign code 996.76, Other complications due to genitourinary device, implant, and graft for the erosion and/or code 996.65, Infection and inflammatory reaction due to other genitourinary device, implant and graft if an infection occurred from the mesh.

Mesh and mesh patches are also used to repair ventral (incisional) hernias caused by thinning or stretching of scar tissue that forms after surgery. Mesh erosion is a known complication that results in bowel perforations and/or, chronic intestinal fistulae.

The following tabular modifications were proposed at the September 2009 ICD-9-CM C&M Committee meeting :

TABULAR MODIFICATIONS

Option 1:

996 Complications peculiar to certain specified procedures

Add Excludes: mesh erosion (998.84) Add mesh exposure (998.85)

998 Other complications of procedures, NEC

998.8 Other specified complications of procedures, not elsewhere classified

New code 998.84 Mesh erosion

New code 998.85 Mesh exposure

Option 2:

TABULAR MODIFICATIONS

629 Other disorders of female genital organs

New subcategory 629.3 Mechanical complication of implanted vaginal mesh

New code 629.31 Erosion of implanted vaginal mesh to surrounding organ or

tissue

New Code 629.32 Exposure of implanted vaginal mesh into vagina

996 Complications peculiar to certain specified procedures

996.7 Other complications of internal (biological) (synthetic) prosthetic device,

implant, and graft

Use additional code to identify complication, such as:

Add implanted vaginal mesh erosion or exposure (629.31-629.32)

Malnutrition

The American Dietetic Association (ADA) and the American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.) have requested several new codes and instructional notes be added to the ICD-9-CM to update the current classification of malnutrition. The existing ICD-9-CM codes for malnutrition are outdated and do not reflect the current standard of care or understanding of malnutrition-disease interaction. Thus, the existing ICD-9-CM malnutrition codes are inconsistently applied by clinicians and facilities across health care settings. Inconsistency in the recognition and documentation of malnutrition in adults is of concern and can significantly impact patient health, safety, quality of life, and health care costs.

Patients with acute disease or injuries, or those with chronic disease, are at high risk for malnutrition. Inconsistencies in the documentation of malnutrition limit our ability to quantify and benchmark malnutrition's overall incidence, prevalence and impact on our population's health and our nation's health care resources utilization.

TABULAR MODIFICATIONS

Delete	260 Kwashiorkor Nutritional edema with dyspigmentation of skin and hair		
Add	Excludes: severe malnutrition related to acute injury, illness and other of (262.11-262.14, 262.19)	disorders	
Delete Delete Delete	261 Nutritional marasmus Nutritional atrophy Severe calorie deficiency Severe malnutrition NOS		
Add	Excludes: severe malnutrition related to acute injury, illness and other of (262.11-262.14, 262.19)	disorders	
Revise Delete	262 Other <u>specified</u> severe protein-calorie malnutrition Nutritional edema without mention of dyspigmentation of skin and hair		
New subcatego	y 262.1 Severe malnutrition in injury, illness or other disorders		
New code	262.11 Severe malnutrition in acute injury		

Code first the related acute injury, such as:

Traumatic brain injury (850-854)

Burns (940-949)

New code 262.12 Severe malnutrition in acute illness

Code first the related acute illness, such as:

Acute pancreatitis (577.0) Pneumonia (480-486)

Transplant complications (996.80-996.89)

New code 262.13 Severe malnutrition in chronic illness

Code first the related chronic illness, such as:

Chronic pancreatitis (577.1) Crohn's (555.0-555.9)

Frailty (797)

Malignant neoplasm (140-195) Ulcerative colitis (556.0-556.9)

New code 262.14 Severe malnutrition in environmental and social circumstances

Code first the related abuse and neglect, such as: Adult abuse and neglect (995.80-995.89)

Child abuse and neglect (995.50-995.59)

New code 262.19 Severe malnutrition related to other disorders

Code first the related disorder, such as:

Anorexia nervosa (307.1) Bulimia nervosa (307.51)

Frailty (797)

263 Other and unspecified protein-calorie malnutrition

Add Excludes: severe malnutrition related to acute injury, illness and other disorders

(262.11-262.14, 262.19)

782 Symptoms involving skin and other integumentary tissue

782.3 Edema

Delete Excludes: nutritional edema (260, 262)

INDEX MODIFICATIONS

Anasarca 782.3

Revise nutritional 262 – see Malnutrition

Revise Athrepsia 261 – see Malnutrition Atrophy, infantile 261 - see Disease, motor neuron Revise nutritional 261 – see Malnutrition Revise Cachexia 799.4 Revise due to malnutrition - see Malnutrition Deficiency, deficient Revise calorie, severe 261 – see Malnutrition edema 262 – see Malnutrition Revise Revise multiple, syndrome 260 – see Malnutrition Revise protein 260 – see Malnutrition syndrome, multiple 260 - see Malnutrition Revise Deprivation protein (familial) (kwashiorkor) 260 - see Malnutrition Revise Development arrested 783.40 due to malnutrition (protein-calorie) 263.2 – see Malnutrition Revise Revise Lancereaux's (diabetes mellitus with marked emaciation) 250.8 [261] [263.8] Revise due to secondary diabetes 249.8 [261] [263.8] Revise secondary Revise Lancereaux's (diabetes mellitus with marked emaciation) 249.8 [261] [263.8] Disease... wasting NEC 799.4 Revise due to malnutrition 261 - see Malnutrition Dwarf, dwarfism 259.4 Revise nutritional 263.2 – see Malnutrition Dystrophy... Revise due to malnutrition 263.9 – see Malnutrition Revise nutritional 263.9 – see Malnutrition Edema... Revise famine 262 - see Malnutrition inanition 262 - see Malnutrition Revise Revise nutritional (newborn) 262 – see Malnutrition with dyspigmentation, skin and hair 260 - see Malnutrition Revise starvation 262 - see Malnutrition Revise Emaciation (due to malnutrition) 261 – see Malnutrition Revise Famine 994.2 Revise edema 262 – see Malnutrition Hydrops... Revise nutritional 262 – see Malnutrition Revise Hypoproteinosis 260 – see Malnutrition Inanition 263.9 – see Malnutrition Revise

Kwashiorkor (marasmus type) 260

due to

with edema 262 - see Malnutrition

malnutrition 263.9 – see Malnutrition

Revise

Revise

Revise

Revise Lancereaux's diabetes (diabetes mellitus with marked emaciation) 250.8 [261] [263.8]

Revise due to secondary diabetes 249.8 [261] [263.8]

Malnutrition (calorie) 263.9

degree

 Revise
 first 263.1 263.9

 Revise
 second 263.0 263.9

 Revise
 third 262 263.9

Revise mild (protein) NEC 263.1

Add due to specified underlying condition – see Malnutrition, related to, by

cause

moderate (protein) 263.0

Add due to specified underlying condition – see Malnutrition, related to, by

cause

Revise severe 261 263.9

Revise protein-calorie <u>NEC</u> 262

Add due to specified underlying condition – see Malnutrition, related to,

by cause

Add due to – see Malnutrition, related to

Revise malignant 260 263.9

mild (protein) 263.1

Add due to specified underlying condition – see Malnutrition, related to, by

cause

moderate (protein) 263.0

Add due to specified underlying condition – see Malnutrition, related to, by

cause

Revise protein 260 <u>263.9</u>

mild 263.1

Add due to specified underlying condition – see Malnutrition, related to, by

cause moderate 263.0

Add due to specified underlying condition – see Malnutrition, related to, by

cause

protein-calorie 263.9

Revise severe NEC 262

specified type NEC 263.8

Add due to specified underlying condition – see Malnutrition, related to, by

cause

Add related to (due to)

Add acute illness NEC 262.12 Add acute injury 262.11 Add chronic illness 262.13

Add environmental circumstances 262.14

Add injury, acute 262.11

Add social circumstances 262.14

Add specified NEC 262.19

Revise severe 261 263.9

Add due to specified underlying condition – see Malnutrition, related to, by

cause

Revise Pedatrophia 261 – see Malnutrition

Revise Pluricarential syndrome of infancy 260 – see Malnutrition

Revise Plurideficiency syndrome of infancy 260 – see Malnutrition

Revise Plurideficiency syndrome of infancy 260 – see Malnutrition

Polycarential syndrome of infancy 260 – see Malnutrition

Revise Prekwashiorkor 260 – see Malnutrition

Protein

Revise deficiency 260 – see Malnutrition
Revise malnutrition 260 – see Malnutrition

Retardation

growth (physical) in childhood 783.43

Revise due to malnutrition 263.2 – see Malnutrition

physical 783.43

Revise due to malnutrition 263.2 – see Malnutrition

Starvation ...

Revise edema 262 – see Malnutrition

Syndrome...

multiple

Revise deficiency 260 – see Malnutrition

Revise pluricarential of infancy 260 – see Malnutrition
Revise plurideficiency of infancy 260 – see Malnutrition
Revise polycarential of infancy 260 – see Malnutrition

Wasting

disease 799.4

Revise due to malnutrition 261 – see Malnutrition
Revise extreme (due to malnutrition) 261 – see Malnutrition

Lymphangioleiomyomatosis

Lymphangioleiomyomatosis, also known as lymphangiomyomatosis or LAM,. is a rare, frequently fatal lung disease that affects women almost exclusively. It may occur sporadically (not inherited), and also in patients with tuberous sclerosis complex (inherited). LAM is characterized by the infiltration of the lung with neoplastic smooth muscle cells of unknown origin, and cystic destruction of lung tissue.

There is no specific ICD-9-CM code for LAM. Many clinicians use code 171.9, Malignant neoplasm of connective and other soft tissue, site unspecified. However, this is a poor fit, since the 'malignant' description is controversial or inaccurate. The ICD-9-CM has an index entry "Lymphangiomyomatosis," with instruction to "see Neoplasm, connective tissue, uncertain behavior." This would lead to code 238.1, Neoplasm of uncertain behavior of other and unspecified sites and tissues, Connective and other soft tissue. A specific code would facilitate epidemiological, clinical, comparative effectiveness and cost effectiveness research.

The clinical behavior of LAM is more consistent with an interstitial lung disease, rather than a neoplasm. Thus, it has been requested that this be moved to category 516. The American Thoracic Society (ATS) and the American College of Chest Physicians (ACCP) have requested creation of a specific code for LAM, as well as for a number of other interstitial lung diseases.

References:

McCormack FX. Lymphangioleiomyomatosis: a clinical update. Chest. 2008

Feb;133(2):507-16. Review. PubMed PMID: 18252917

This reference can be found at link: http://chestjournal.chestpubs.org/content/133/2/507.full.pdf+html

or PUBMED at: http://www.ncbi.nlm.nih.gov/pubmed/18252917.

TABULAR MODIFICATIONS

516 Other alveolar and parietoalveolar pneumonopathy

New code 516.4 Lymphangioleiomyomatosis Lymphangiomyomatosis

23

Dementia unspecified with and without behavioral disturbance

NCHS has received a request from the West Palm, Florida Veterans Affairs Medical Center, requesting that fifth digits be added at 294.8, Other persistent mental disorders due to conditions classified elsewhere, to identify the presence or absence of behavioral disturbance when the etiology of the dementia is not known. The request is intended to parallel the fifth digits at subcategory 294.1, Dementia in conditions classified elsewhere. It was noted that providers will document dementia with behavioral disturbance but are not always able to identify the etiology of the dementia.

The American Academy of Neurology and the American Psychiatric Association support new codes to identify unspecified dementia with or without behavioral disturbance.

TABULAR MODIFICATIONS

ORGANIC PSYCHOTIC CONDITIONS (290-294)

294 Persistent mental disorders due to conditions classified elsewhere

New subcategory 294.2 Dementia, unspecified

Dementia NOS

Excludes: mild memory disturbances, not amounting to dementia (310.8)

294.20 Dementia unspecified without behavioral disturbance

294.21 Dementia unspecified with behavioral disturbance

Aggressive behavior Combative behavior Violent behavior Wandering off

294.8 Other persistent mental disorders due to conditions classified elsewhere

Amnestic disorder NOS

Dementia NOS

Epileptic psychosis NOS

Mixed paranoid and affective organic psychotic states

Elective c-sections prior to 39 weeks

The American College of Obstetricians and Gynecologists (ACOG) has indicated that one of the new markers of quality involves looking at elective deliveries done prior to 39 weeks gestation. In reviewing these cases planned repeat cesarean section deliveries done prior to 39 weeks fall into this category. However, many times these deliveries occur earlier because the woman presents at 37-38 weeks gestation in labor and the physician determines that is best to deliver at that time rather than try to take measures to wait until the 39th week. Using code 654.2x, Previous cesarean delivery does not completely describe this scenario. They are requesting a new code that will specify the onset of labor for this time period with delivery. Existing codes for onset of labor only apply to labor prior to 37 weeks, or false labor and neither of these apply in this situation. The following modifications are proposed:

TABULAR MODIFICATIONS

644 Early or threatened labor

New code 644.3 Late preterm onset of labor, with delivery

[0-1]

Late preterm onset of labor, after 37 completed weeks of gestation but before 39 weeks gestation, with delivery

Onset (spontaneous) of labor after 37 completed weeks of gestation but before 39 weeks gestation, with delivery

Personal history of gestational diabetes

The American College of Obstetrics and Gynecology (ACOG) has requested a new code for patients with a personal history of gestational diabetes. Currently this is indexed to V12.2, Personal history endocrine, metabolic, and immunity disorders.

TABULAR MODIFICATIONS

V12 Personal history of certain other diseases

V12.2 Personal history endocrine, metabolic, and immunity disorders

New code V12.21 Personal history of gestational diabetes

New code V12.29 Personal history of other endocrine, metabolic, and immunity

disorders

Encounter for fetal viability ultrasound/personal history of ectopic pregnancy

The American College of Obstetrics and Gynecology (ACOG) has received many requests asking for the most appropriate diagnosis coding for encounters to check fetal viability, using ultrasound, especially when pregnancy has been confirmed in the early weeks of pregnancy. Patients previously confirmed as pregnant in very early weeks may return within a few weeks and if the fetal heartbeat cannot be heard an ultrasound may be necessary to confirm that the pregnancy is viable.

In addition, ACOG would find it useful to have a code to allow tracking of patients who have had a previous ectopic pregnancy.

TABULAR MODIFICATIONS

- V23 Supervision of high-risk pregnancy
 - V23.4 Pregnancy with other poor obstetric history
 - V23.42 Pregnancy with history of ectopic pregnancy
 - V23.43 Pregnancy with inconclusive fetal viability
 Encounter to determine fetal viability of pregnancy

Adult Pulmonary Langerhans Cell Histiocytosis (PLCH)

Adult Pulmonary Langerhans Cell Histiocytosis (PLCH) is a rare interstitial lung disorder of unknown etiology that occurs almost exclusively in smokers, with an average peak incidence of 20-40 years. In adults, pulmonary involvement with Langerhans' cell histiocytosis usually occurs as a single-system disease and is characterized by focal Langerhans' cell granulomas infiltrating and destroying distal bronchioles. The typical high resolution CT pattern combines small nodules with or without cavitation and cysts predominantly involving upper lungs with relative sparing of the lung bases. Adult PLCH is distinct from the multisystem histiocytic disorders that occur in children and adolescents (named Hand-Schüller-Christian disease, Letterer-Siwe disease, and histiocytosis X), in which pulmonary disease is rarely the focus.

Adult PLCH is not currently indexed in ICD-9-CM. Histiocysosis is included at code 277.89, Other specified disorders of metabolism. Another potential code is 202.5, Letterer-Siwe disease, which includes acute differentiated progressive histiocytosis, and acute histiocytosis X (progressive); this is a pediatric form of histiocytosis. While some may use either of these codes, due to the interstitial lung involvement, ATS-ACCP would currently code adult PLCH to 516.8, Other specified alveolar and parietoalveolar pneumonopathies.

The clinical behavior of adult PLCH is consistent with an interstitial lung disease, rather than a neoplasm or a metabolic disorder. Thus, it has been requested that this be classified to category 516.

The American Thoracic Society (ATS) and the American College of Chest Physicians (ACCP) have requested creation of a specific code for adult PLCH, as well as for a number of other interstitial lung diseases. A specific code for adult PLCH will distinguish the histiocytic disease which occurs in adult smokers from the neoplastic histiocytic diseases that occur in children, and will facilitate epidemiological, clinical, comparative effectiveness and cost effectiveness research.

Reference:

Tazi A. Adult pulmonary Langerhans' cell histiocytosis. Eur Respir J. 2006 Jun;27(6):1272-85. Review. PubMed PMID: 16772390. This article can be accessed here:

http://erj.ersjournals.com/cgi/reprint/27/6/1272

TABULAR MODIFICATIONS

202 Other malignant neoplasms of lymphoid and histiocytic tissue

202.5 Letterer-Siwe disease

Add Excludes: Adult Pulmonary Langerhans Cell Histiocytosis (516.5)

277 Other and unspecified disorders of metabolism

277.8 Other specified disorders of metabolism

277.89 Other specified disorders of metabolism

Excludes: histiocytosis:

Add Adult pulmonary Langerhans cell (516.5)

516 Other alveolar and parietoalveolar pneumonopathy

New code 516.5 Adult Pulmonary Langerhans Cell Histiocytosis

Adult PLCH

Acquired absence of joint

This request has been presented previously at the March 2008, March 2009, and March 2010 ICD-9-CM Coordination and Maintenance Committee meetings. The request originated from a query reviewed by the Editorial Advisory Board for Coding Clinic for ICD-9-CM regarding coding advice for patient encounters for joint replacement surgery, following previous explantation of a joint prosthesis. Last March it was proposed to create codes to show the status of acquired absence of a joint which would include joint prosthesis explantation status. This proposal was favorably commented upon by participants. It was further reviewed by the American Academy of Orthopaedic Surgeons (AAOS) who asked that in order to make these codes useful for tracking patients who have had their joint prosthesis explanted (with or without an antibioticimpregnated cement spacer) to add the inclusion notes under the codes at proposed new subcategory V88.2, Acquired absence of joint. The clinical rationale given for these changes is that that there are many other congenital and acquired causes of destruction of the hip or knee joint (including trauma, neoplasm, infection, etc). It is important to clearly specify the cause of the acquired absence of the joint, which is explantation of joint prosthesis. The revised tabular modifications reflecting comments received after the March 2010 ICD-9-CM C&M Committee meeting and the AAOS recommendation is below:

ICD-9-CM TABULAR MODIFICATIONS

V54 Other orthopedic aftercare

V54.8 Other orthopedic aftercare

New code

V54.82 Aftercare following explantation of joint prosthesis
Aftercare following explantation of joint prosthesis, staged
procedure
Encounter for joint prosthesis insertion following prior

explantation of joint prostness insertion following prior

V88 Acquired absence of other organs and tissue

New subcategory V88.2 Acquired absence of joint

Acquired absence of joint following prior explantation of joint

prosthesis

Joint prosthesis explantation status

New code V88.21 Acquired absence of hip joint

Acquired absence of hip joint following explantation of joint prosthesis, with or without presence of antibiotic-

impregnated cement spacer

New code V88.22 Acquired absence of knee joint

Acquired absence of knee joint following explantation of joint prosthesis, with or without presence of antibiotic-

impregnated cement spacer

New code V88.29 Acquired absence of other joint

Acquired absence of other joint following explantation of joint prosthesis, with or without presence of antibiotic-

impregnated cement spacer

Glaucoma severity staging

Revise

Glaucoma is characterized by optic nerve damage that results in visual field loss as the disease progresses to more advanced stages. The most common types of glaucoma are Primary Open Angle Glaucoma (POAG), Primary Angle closure glaucoma, pigmentary glaucoma, corticosteroid-induced glaucoma, pseudoexfoliation glaucoma, and glaucoma associated with ocular inflammation, ocular vascular disorders, and ocular trauma. Patients present at vastly different stages of disease, and typically treatment at early stages of disease results in better outcomes and uses fewer resources than patients who present with more severe stages of glaucoma.

The American Academy of Ophthalmology has requested that new codes be created as it would be important to be able to capture the stage of disease when coding for the most commonly encountered types of glaucoma.

Several other revisions are also requested by the American Academy of Ophthalmology that are consistent with knowledge and terminology currently in use. Modern glaucoma evaluation now entails the assessment of clinical factors in determining risk for the development of open angle glaucoma. Open angle glaucoma suspect patients are commonly stratified into low risk and high risk. Similarly, the angle closure code revisions are also requested to be consistent with current worldwide terminology.

TABULAR MODIFICATIONS

Open angle with borderline findings, low risk

365.0 Borderline glaucoma [glaucoma suspect]

365.01

Add	Open angle, low risk
Delete	Open anngle with:
Delete	borderline intraocular pressure
Delete	cupping of optic discs
	365.02 Anatomical narrow angle
Add	Primary angle closure suspect
New code	365.05 Open angle with borderline findings, high risk
	Open angle, High Risk
New code	365.06 Primary angle closure without glaucoma damage
	365.1 Open-angle glaucoma
	365.10 Open-angle glaucoma, unspecified
	Wide-angle glaucoma NOS
Add	Use additional code to identify glaucoma stage (365.71-365.74)
	365.11 Primary open angle glaucoma
	Chronic simple glaucoma
Add	Use additional code to identify glaucoma stage (365.71-365.74)

		365.12	Low-tension glaucoma
Add Add		LISE addit	Normal tension glaucoma tional code to identify glaucoma stage (365.71-365.74)
Auu		Ose addit	tional code to identify gladcoma stage (505.71-505.74)
		365.13	Pigmentary glaucoma
Add		Use addit	tional code to identify glaucoma stage (365.71-365.74)
	365.2	Primary a	angle-closure glaucoma
		365.20	Primary angle-closure glaucoma, unspecified
Add		Use addit	tional code to identify glaucoma stage (365.71-365.74)
		365.22	Acute angle clecure glaucema
Add		303.22	Acute angle-closure glaucoma Acute angle-closure glaucoma crisis
Add			Acute angle-closure glaucoma attack
A -I -I		365.23	Chronic angle-closure glaucoma
Add Add		libb addi	Primary angle closure glaucoma tional code to identify glaucoma stage (365.71-365.74)
Add		OSC dddii	tional code to identify gladcoma stage (303.71 303.74)
	365.3	3 Cortico:	steroid-induced glaucoma
		365.31	Glaucomatous stage
Add		Use addit	tional code to identify glaucoma stage (365.71-365.74)
	365.5	Glaucom	a associated with disorders of the lens
		365.52	Pseudoexfoliative glaucoma
Add		Use addit	tional code to identify glaucoma stage (365.71-365.74)
	365.6	Glaucom	a associated with other ocular disorders
		365.62	Glaucoma associated with ocular inflammations
Add			tional code to identify glaucoma stage (365.71-365.74)
			, see
		365.63	Glaucoma associated with vascular disorders
Add	Us	e additiona	al code to identify glaucoma stage (365.71-365.74)
		365.65	Glaucoma associated with ocular trauma
Add		Use addit	tional code to identify glaucoma stage (365.71-365.74)
New subcategory	365.7		coma stage
		-365.63, 3	ated type of glaucoma (365.11-365.13, 365.26, 365.31, 365.52, 65.65
	555.52	505.05, 5	

New code 365.71 Mild stage glaucoma

Early stage glaucoma

New code 365.72 Moderate stage glaucoma

New code 365.73 Severe stage glaucoma

Advanced stage glaucoma

End-stage glaucoma

New code 365.74 Indeterminate stage glaucoma

Glaucoma stage NOS

PERSONS WITH POTENTIAL HEALTH HAZARDS RELATED TO PERSONAL AND FAMILY HISTORY (V10-V19)

V19 Family history of other conditions

New subcategory V19.1 Other eye disorders New code V19.11 Glaucoma

New code V19.19 Other specified eye disorder

INDEX MODIFICATIONS

Glaucoma

Add

angle closure 365.20 Add attack 365.22 crisis 365.22 Add Add primary angle closure glaucoma 365.23 Add angle recession 365.65 Add exfoliation 365.52 Add inflammatory 365.62 Add neovascular 365.63 Add normal tension 365.12 Revise open angle 365.10 with Add abnormal optic disc appearance or asymmetry 365.01 thin central corneal thickness (pachymetry) 365.01Add Add phacoanaphylactic 365.59 Add phacomorphic, acute 365.22 Add phacomorphic, borderline 365.06 Add pigment dispersion 365.13 secondary NEC 365.60 Add due to Add surgery 365.60 Add steroids 365.31 steroid induced 365.31 Add Add uveitic 365.62 Plateau iris syndrome 364.82 Add without glaucoma 365.06

with glaucoma 365.23

Corticobasal degeneration

Corticobasal degeneration (CBD) is a neurodegenerative disease that is manifest by both movement disorder and cognitive impairment. The cognitive symptoms resemble those of the frontotemporal dementias, especially loss of executive function, visuospatial and number processing, and language impairment. The movement disorder most often presents asymmetrically and may include akinetic-rigid syndrome, myoclonus or dystonia. Patients may also have alien limb syndrome, apraxia and cortical sensory loss. Because CBD is clinically and pathologically different than the neurodegenerative diseases with currently assigned ICD codes, and also is not a degenerative disease primarily of the basal ganglia, the American Academy of Neurology (AAN) proposes the following new code:

TABULAR MODIFICATIONS

331 Other cerebral degenerations

New code 331.6 Corticobasal degeneration

Pulmonary Arteriovenous Malformation and Pulmonary Atresia

A pulmonary arteriovenous malformation (AVM) is an abnormal communication between pulmonary arteries and pulmonary veins. While these are most often congenital, they can also be acquired. These may also be called pulmonary arteriovenous aneurysms, or pulmonary arteriovenous fistulae.

Most often, a pulmonary AVM is small, and will usually not present until adulthood, even when congenital. However, in some cases these can become serious problems, and they may present with cyanosis, heart failure, dyspnea, and even respiratory failure. Pulmonary arteriovenous malformation is not indexed in ICD-9-CM. However, pulmonary arteriovenous aneurysm is included at 747.3, Anomalies of pulmonary artery. Pulmonary circulation anomalies are also indexed to 747.3, thus including those that do not affect the main pulmonary artery.

Narowing of the pulmonary artery is called coarctation or stenosis. Complete failure of the pulmonary valve to form, with the origin of the pulmonary artery not connecting to the heart, may be called pulmonary artery atresia or agenesis; this is generally seen only along with a patent ductus arteriosis, that transfers blood from the left heart circulation to the pulmonary circulation.

The question of coding for a pulmonary AVM was raised through the Coding Clinic Editorial Advisory Board, and it is proposed to create a specific code for pulmonary AVM, and a code for Pulmonary artery coarctation and atresia, and a code for Other anomalies of pulmonary artery and pulmonary circulation.

TABULAR MODIFICATIONS

042 Human immunodeficiency virus [HIV] disease

Use additional code, if applicable, to identify infection following transfusion, infusion, or injection of blood and blood products (999.32)

417 Other diseases of pulmonary circulation

417.0 Arteriovenous fistula of pulmonary vessels

Revise Excludes: congenital arteriovenous fistula (747.32)

417.1 Aneurysm of pulmonary artery

Revise Excludes: congenital aneurysm (747.39)

Add congenital arteriovenous aneurysm (747.32)

747 Other congenital anomalies of circulatory system

747.3 Anomalies of pulmonary artery and pulmonary circulation

Delete Agenesis of pulmonary artery
Anomaly of pulmonary artery

Atresia of pulmonary artery
Coarctation of pulmonary artery
Hypoplasia of pulmonary artery
Stenosis of pulmonary artery
Pulmonary arteriovenous aneurysm

New code 747.31 Pulmonary artery coarctation and atresia

Agenesis of pulmonary artery Atresia of pulmonary artery Coarctation of pulmonary artery Hypoplasia of pulmonary artery Stenosis of pulmonary artery

New code 747.32 Pulmonary arteriovenous malformation

Pulmonary arteriovenous aneurysm

Excludes: acquired pulmonary arteriovenous fistula (417.0)

New code 747.39 Other anomalies of pulmonary artery and pulmonary

circulation

Anomaly of pulmonary artery

747.6 Other anomalies of peripheral vascular system

Excludes: anomalies of:

Revise pulmonary artery (747.39)

747.8 Other specified anomalies of circulatory system

747.89 Other

Excludes: congenital aneurysm:

Revise pulmonary (747.3<u>9</u>)

Add arteriovenous (747.32)

Complications of stem cell transplant

The ICD-9-CM currently does not have a unique code for complications of stem cell transplants. During recent discussions of the AHA Editorial Advisory Board meeting for *Coding Clinic for ICD-9-CM* on the code assignment for complications of stem cell transplants, it was suggested that either a new code be created or that entry be added to the Alphabetical Index at existing code 996.85, Complications of transplanted organ, bone marrow.

The greatest concentration of blood stem cells is in the bone marrow. However, it is possible to move blood stem cells out of the bone marrow into the bloodstream or "peripheral blood" where they can be collected and used instead of bone marrow for the transplant. Umbilical cord blood also contains blood stem cells that can be used for transplant.

With the availability of the stem cell growth factors granulocyte-macrophage colony-stimulating factor (GM-CSF) and granulocyte colony-stimulating factor (G-CSF), most hematopoietic stem cell transplantation procedures are now performed using stem cells collected from the peripheral blood, rather than from the bone marrow. Collecting peripheral blood stem cells provides a bigger graft, does not require that the donor be subjected to general anesthesia to collect the graft, results in a shorter time to engraftment, and may provide for a lower long-term relapse rate.

A stem cell transplant poses many risks of complications, some potentially fatal. Complications that can arise with a stem cell transplant include: graft-versus-host disease, stem cell (graft) failure, organ damage, cataracts, and secondary cancers.

The two options are:

TABULAR MODIFICATIONS

Option 1:

996.8 Complications of transplanted organ

New code 996.88 Stem cell

bone marrow peripheral blood umbilical cord

Option 2:

996.8 Complications of transplanted organ

996.85 Bone marrow

Add Stem cell transplant (bone marrow) (peripheral blood) (umbilical cord)

Pseudobulbar Affect

Avanir Pharmaceuticals has requested that a unique code be created for pseudobulbar affect (PBA). Currently, PBA is indexed to code 310.8 "Other specified nonpsychotic mental disorders following organic brain damage, which includes several unrelated conditions, including mild memory disturbance, amnesia, and postencephalitic syndrome. A unique code would assist in recognition and diagnosis of PBA and would improve the epidemiologic tracking of this distinct neurologic condition.

PBA is a neurologic condition caused by underlying structural damage in the brain which triggers involuntary, frequent and disruptive outbursts of crying or laughing. PBA episodes typically occur out of proportion or incongruent to the patient's underlying emotional state. The pathophysiology of PBA is widely believed to involve injury to the neurologic pathways that regulate affect.

PBA occurs secondary to neurologic disease or injury and has been reported to occur in 49% of patients with amyotrophic lateral sclerosis (ALS), in 10% of patients with multiple sclerosis (MS), in 11% of patients 1 year after suffering a stroke, and in 11% of patients after a traumatic brain injury. PBA has also been reported to occur secondary to certain other neurological conditions. The etiology of PBA is not completely understood but the symptoms are similar across patient populations.

Despite the high prevalence of PBA among persons with underlying neurologic conditions, it is under recognized and undertreated in neurological, psychiatric, and general medical settings. Due to the symptomatology, PBA is often mistaken for psychiatric disorders such as depression, bipolar disorder, schizophrenia and generalized anxiety disorder.

In summary, PBA is a prevalent neurologic condition associated with significant functional impairment and worsened quality of life for patients and their caregivers. Avanir believes that a PBA-specific ICD-9-CM code would help to increase recognition, improve diagnosis and ultimately advance the management of PBA.

The creation of a unique code for PBA has support from the following organizations: Multiple Sclerosis Association of America, Brain Injury Association of America, and the Physician Foundation at California Pacific Medical Center.

TABULAR MODIFICATIONS

310 Specific nonpsychotic mental disorders due to brain damage

310.8 Other specified nonpsychotic mental disorders following organic brain damage

Delete Mild memory disturbance Delete Postencephalitic syndrome

Delete Other focal (partial) organic psychosyndromes

Excludes: memory loss of unknown cause (780.93)

New code 310.81 Pseudobulbar affect

Involuntary emotional expression disorder

Code first underlying cause, such as:

amyotrophic lateral sclerosis (335.20)

late effect of cerebrovascular accident (438.89) late effect of traumatic brain injury (907.0)

mutiple sclerosis (340)

New code 310.89 Other

> Mild memory disturbance Postencephalitic syndrome

Other focal (partial) organic psychosyndromes memory loss of unknown cause (780.93) Excludes:

Reportable malignant skin cancers

The New York State Cancer Registry has requested that the ICD-9-CM be expanded in some way to allow for the classification of reportable skin cancers that are currently included under category 173, Other malignant neoplasm of skin.

The vast majority of skin cancers are either basal or squamous cell, neither of which are reportable conditions to central cancer registries. These histologies are included in category 173. Precedent for including histologic type has already been set with malignant melanoma, indexing of Kaposi's sarcoma, and most recently new codes for Merkel cell carcinoma.

Currently, case finding for reportable cancers is done by a combination of automated and manual methods. In hospitals, the medical record index which generally utilizes ICD-9-CM codes is used to filter records for review by tumor registrars. Registrars must review all records with a code from category 173 to identify reportable cases.

Laboratories and other non-hospital facilities are starting to submit reports automatically to central registries. The ICD9-CM codes are commonly used to filter reports for automatic creation of HL-7 records. Because of the difficulty in distinguishing reportable skin cancers from non-reportable skin cancers the facilities are transmitting all skin cancers to central registries. This places an additional burden on central registries and also results in the transmission of confidential patient information on patients whose information should not be reported.

Due to the volume of data that has been collected over the years on codes under category 173 it seems best to leave basal and squamous cell classified under category 173. Creating a new category in ICD-9-CM at this late date in its use seems problematic. It is being proposed that codes under category 173 be expanded to the 5th digit level to allow for the differentiation of reportable and non-reportable (basal and squamous cell) skin cancers.

Once a final design for this expansion is completed and a valid, comprehensive list of the reportable skin cancers is compiled, the best method for indexing these new codes in the Index to Diseases and, specifically, the Neoplasm table will be determined.

TABULAR MODIFICATIONS

173 Other malignant neoplasm of skin

Includes: malignant neoplasm of:

sebaceous glands

sudoriferous, sudoriparous glands

sweat glands

Excludes: Kaposi's sarcoma (176.0-176.9)

malignant melanoma of skin (172.0-172.9) skin of genital organs (184.0-184.9, 187.1-187.9)

173.0 Skin of lip

New code	173.00 Basal cell and squamous cell carcinoma of skin of lip Malignant neoplasm of skin of lip NOS
New code	173.09 Other malignant neoplasm of skin of lip
173.	1 Eyelid, including canthus
New code	173.10 Basal cell and squamous cell carcinoma of skin of eyelid, including canthus Malignant neoplasm of skin of eyelid (canthus) NOS
New code	173.19 Other malignant neoplasm of skin of eyelid, including canthus
173.	2 Skin of ear and external auditory canal
New code	173.20 Basal cell and squamous cell carcinoma of skin of ear and external auditory canal Malignant neoplasm of skin of ear NOS Malignant neoplasm of skin of auditory canal NOS
New code	173.29 Other malignant neoplasm of skin of ear and external auditory canal
173.	3 Skin of other and unspecified parts of face
New code	173.30 Basal cell and squamous cell carcinoma of skin of other and unspecified parts of face
New code	Malignant neoplasm of skin of face NOS 173.39 Other malignant neoplasm of skin of other and unspecified parts of face
173.	4 Scalp and skin of neck
New code	173.40 Basal cell and squamous cell carcinoma of scalp and skin of neck Malignant neoplasm of scalp NOS
New code	Malignant neoplasm of skin of neck NOS 173.49 Other malignant neoplasm of scalp and skin of neck
173.	5 Skin of trunk, except scrotum
New code	173.50 Basal cell and squamous cell carcinoma of skin of trunk, except scrotum
New code	Malignant neoplasm of skin of trunk NOS 173.59 Other malignant neoplasm of skin of trunk, except scrotum

173.6 Skin of upper limb, including shoulder

New code 173.60 Basal cell and squamous cell carcinoma of skin of upper

limb, including shoulder

Malignant neoplasm of skin of upper limb NOS

New code 173.69 Other malignant neoplasm of skin of upper limb,

including shoulder

173.7 Skin of lower limb, including hip

New code 173.70 Basal cell and squamous cell carcinoma of skin of lower

limb, including hip

Malignant neoplasm of skin of lower limb NOS

New code 173.79 Other malignant neoplasm of skin of lower limb,

including hip

173.8 Other specified sites of skin

New code 173.80 Basal cell and squamous cell carcinoma of other

specified sites of skin

New code 173.89 Other malignant neoplasm of other specified sites of

skin

173.9 Skin, site unspecified

New code 173.90 Basal cell and squamous cell carcinoma of skin

Malignant neoplasm of skin NOS

Skin cancer NOS

New code 173.99 Other malignant neoplasm of skin

ICD-9-CM TABULAR LIST OF DISEASES

PROPOSED ADDENDA (Effective October 1, 2011)

249 Secondary diabetes mellitus

249.8 Secondary diabetes mellitus with other specified manifestations

Use additional code to identify manifestation, as:

any associated ulceration (707.10 707.9) (707.10-707.19,

707.8, 707.9)

250 Diabetes mellitus

250.8 Diabetes mellitus with other specified manifestations

Use additional code to identify manifestation, as:

Revise any associated ulceration (707.10-707.9) (707.10-707.19,

707.8, 707.9)

Revise Chapter

Revise

Title 5. MENTAL AND BEHAVIORAL DISORDERS (290-319)

430 Subarachnoid hemorrhage

Add Excludes: berry aneurysm, nonruptured (437.3)

440 Atherosclerosis

440.2 Of native arteries of the extremities

440.23 Atherosclerosis of the extremities with ulceration

Revise Use additional code for any associated ulceration (707.10-707.9)

(707.10-707.19, 707.8, 707.9)

440.24 Atherosclerosis of the extremities with gangrene

Revise Use additional code for any associated ulceration (707.10-707.9)

(707.10-707.19, 707.8, 707.9)

	459	Other disorders of circulatory system
		459.8 Other specified disorders of circulatory system
		459.81 Venous (peripheral) insufficiency, unspecified
Revise		Use additional code for any associated ulceration (707.10-707.9) <u>(707.10-707.19, 707.8, 707.9)</u>
	518	Other diseases of lung
		518.3 Pulmonary eosinophilia
Add		Excludes: pulmonary infiltrate NOS (793.1)
	536	Disorders of function of stomach
		536.3 Gastroparesis
Revise		Code first, if applicable, underlying disease, such as: diabetes mellitus (249.6, 250.6)
	569	Other disorders of intestine
		569.4 Other specified disorders of rectum and anus
		569.49 Other
Revise		Use additional code for <u>any</u> associated fecal incontinence (787.60-787.63)
	618	Genital prolapse
		618.0 Prolapse of vaginal walls without mention of uterine prolapse
		618.04 Rectocele
Revise		Use additional code for <u>any</u> associated fecal incontinence (787.60-787.63)

	718	Other derangement of joint
Revise		718.6 Unspecified intrapelvic protrusion of acetabulum [0,5]
	793	Nonspecific (abnormal) findings on radiological and other examination of body structure
Add		793.1 Lung field Pulmonary infiltrate NOS
	999	Complications of medical care, not elsewhere classified
		999.6 ABO incompatibility reaction due to transfusion of blood or blood products
Revise		Excludes: minor blood group antigens reactions (Duffy) (E) (K(ell)) (Kidd) (Lewis) (M) (N) (P) (S) (999.89) (999.75-999.79)
Revise		EMENTARY CLASSIFICATION OF FACTORS INFLUENCING HEALTH S AND CONTACT WITH HEALTH SERVICES (V01-V90) <u>(V01-V91)</u>

ICD-9-CM INDEX TO DISEASES AND INJURIES

PROPOSED ADDENDA (Effective October 1, 2011)

Absence (organ or part) (complete or partial)

artery ...747.60

Revise <u>brain 747.81</u>

Automatism 348.89

Add with temporal sclerosis 348.81

Revise AIPHI (acute idiopathic pulmonary hemorrhage in infants (over 28 days old) 786.31

Add Anaplasmosis, human 082.49

Aneurysm... 442.9

berry (congenital) (ruptured) (see also Hemorrhage, subarachnoid) 430

Add nonruptured 437.3

brain 437.3

berry (congenital) (ruptured) (see also Hemorrhage, subarachnoid) 430

Add nonruptured 437.3

Complications

vaccination 999.9

reaction (allergic) 999.5

Delete Herxheimer's 995.0

Dense

Revise breast(s) - omit code 793.82

Disease, diseased - see also Syndrome

Fournier's disease (idiopathic gangrene) 608.83

Add female 616.89

iron

Revise metabolism (see also Hemochromatosis) 275.03 275.09

Add microvillus atrophy 751.5 Add polyethylene 996.45

Revise Sweeley-Klionsky 272.4 <u>272.7</u>

Disorder Add alcohol-induced mood 291.89 mood- (see also Disorder, bipolar) 296.90 Add alcohol-induced 291.89 Embolism 444.9 vein 453.9 upper extremity (acute) 453.83 Revise Delete acute 453.83 deep 453.82 Delete Delete superficial 453.81 Add deep 453.82 Add superficial 453.81 Encephalopathy (acute) 348.30 due to drugs - (see also Table of Drugs and Chemicals) 348.39 349.82 Revise metabolic (see also Delirium) 348.31 Add drug induced 349.82 Fournier's disease (idiopathic gangrene) 608.83 female 616.89 Add Fracture ... 829.0 femur, femoral (closed) 821.00 neck (closed) 820.8 Add stress 733.96 shaft (lower third) (middle third) (upper third) 821.01 Add stress 733.97 march 733.95 Add femoral neck 733.96 Add pelvis 733.98 Add shaft of femur 733.97 pelvis, pelvic (bone(s)) (with visceral injury) (closed) 808.8 Add stress 733.98 stress Add femoral neck 733.96 Add pelvis 733.98 shaft of femur 733.97 Add

Hemochromatosis (acquired) (liver) (myocardium) (secondary) 275.03

Revise	Herxheimer's reaction	995.0 995.91
--------	-----------------------	-------------------------

	Hypertension, hypertensive	Malignant	Benign	Unspecified
	pulmonary (artery) (secondary)	-	-	416.8
Revise	with cor pulmonale (chronic)	_		416.8
Delete	acute	-		415.0
Add	cor pulmonale (chronic)	-	-	416.8
Add	acute	-	-	415.0
Add	right heart ventricular strain/fai	lure -	-	416.8
Add	acute			415.0

Infiltrate, infiltration

Revise lung (see also Infiltrate, pulmonary) 518.3 793.1

Revise pulmonary 518.3 793.1

Injury 959.9

internal 869.0 lung 861.20

Add aspiration 507.0

Intrauterine contraceptive device

reinsertion V25.13

Delete and reinsertion V25.13

removal V25.12

Add and reinsertion V25.13

Leukoencephalopathy (see also Encephalitis) 323.9

Add arteriosclerotic 437.0

Add Lipodermatosclerosis 729.39

Add Melanocytosis, neurocutaneous 757.33

Add MVID (microvillus inclusion disease) 751.5

Osteomalacia 268.2

Add oncogenic 275.8

Pain

Delete temporomaxillary joint 524.62

Pancytopenia

with

Add myelodysplastic syndrome – see Syndrome, myelodysplastic

Revise Prognathism (mandibular) (maxillary) 524.00 524.10

Reaction

Revise Herxheimer's 995.0 995.91

Septum, septate (congenital) - see also Anomaly, specified type NEC

Revise uterus (see also Double, uterus) (complete) (partial) 752.35

Siderosis (lung) (occupational) 503

Add CNS 437.8

Syndrome

myelodysplastic 238.75

Add lesions, low grade 238.72

superior

Add semi-circular canal dehiscence 386.8

Tear

Add annular fibrosis 722.51

Thrombosis, thrombotic...453.9

Revise iliac (acute) (vein) 453.89 453.41

Revise chronic 453.79 453.51

vein 453.9

Revise upper extremity (acute) 453.83

Delete acute 453.83
Delete deep 453.82
Delete superficial 453.81

Add deep 453.82 Add superficial 453.81

ICD-9-CM ALPHABETIC INDEX TO EXTERNAL CAUSES OF INJURY PROPOSED ADDENDA (Effective October 1, 2011)

Blast (air) in

war operations E993.9

Revise from nuclear explosion - see War operation<u>s</u>, injury due to,

nuclear weapons

War operations (during hostilities) (injury) (by) (in) E995.9

blast (air) (effects) E993.9

Revise from nuclear explosion see War operations, injury due to, nuclear

weapons

TABLE OF DRUGS AND CHEMICALS PROPOSED ADDENDA (Effective October 1, 2011)

Revise	Cleaner, cleansing agent <u>, NEC</u>						
	type not specified	989.89	E861.9	-	E950.9	E962.1	E980.9
Add	specified type NEC	989.89	E861.3	-	E950.9	E962.1	E980.9

Opioids expansion for ICD-10-CM

Opioid drugs have benefit when used properly and are a necessary component of pain management for certain patients. Opioid drugs have serious risks when used improperly, as well as certain risks when used properly. In the past, the Food and Drug Administration (FDA), drug manufacturers, and others have taken a number of steps to prevent misuse, abuse and accidental overdose of these drugs, including providing additional warnings in product labeling, implementing risk management plans, conducting inter-agency collaborations and issuing direct communications to both prescribers and patients. Despite these efforts, the rates of misuse and abuse, and of accidental overdose of opioids, have risen over the past decade. The FDA believes that establishing risk evaluation and mitigation strategies (REMS) for opioids will reduce these risks while ensuring that patients with legitimate need for these drugs will continue to have appropriate access.

A discussion regarding the lack of specificity available in data collected on the use of opioids occurred during an exchange between the FDA and an Industry Working Group held on December 4, 2009. (FDA Public Meeting on Risk Evaluation and Mitigation Strategies (REMS) for Certain Opioids) The discussion focused on developing metrics to assess the effectiveness of a class wide REMS program to mitigate the risks of abuse, misuse, and overdose of extended-release opioids. The issue was addressed again during the most recent FDA long-acting opioid REMS Advisory Committee meeting which was held July 22 & 23 2010, as well as the FDA REMS public meeting on July 27 & 28 2010.

Covidien Pharmaceuticals, a member of the industry working group and a manufacturer of certain opioid agents, submitted a request to NCHS for an expansion to the ICD-10-CM for various opioids categories. The proposal includes creating 7th characters for category F11, Opioid related disorders, to identify the type of opioid, an expansion of the subcategories under category T40, Poisoning by, adverse effect of and underdosing of narcotics and psychodysleptics [hallucinogens], with specific codes for the various opioid molecules, and creating a unique code under category T39, Poisoning by, adverse effect of and underdosing of nonopioid analgesics, antipyretics and antirheumatics, for acetaminophen with instructional notes to allow for the coding of NSAID/opioid combinations. These changes will allow for:

Differentiating long-acting vs. short-acting opioids:

o FDA mandate class-wide long-acting opioid REMS which may result in what is commonly known as the "balloon" effect – putting regulatory pressure on one side of the pain medication spectrum (long-acting opioids) will result in the expanded use on the other side of the spectrum (immediate-release or short-acting opioids).

Granularity of opioid molecules:

 Able to compare molecule to molecule through various public databases (i.e. FDA AERS, DAWN, AAPCC, etc.) to track trends of misuse, abuse, and deaths related to pharmaceutical opioid products.

Differentiating class of drugs by formulations (i.e. NSAID):

o FDA mandated class-wide medication guide to be dispensed with all NSAID products, which include oral, topical, and other forms of this drug class. The class-wide NSAID medication guide address risks mainly associated with oral formulation (i.e. ulcers and bleeding). To reduce the risk of inadvertently misuse

of non-oral NSAID products this level of granularity may support class-wide medication guide may not mitigate the risks the FDA intended to address.

The following modifications are being proposed and would become effective with the implementation of the ICD-10-CM.

TABULAR MODIFICATIONS

F11 Opioid related disorders

Add The appropriate 7th character is to be added to each code from subcategory F11

- 0 buprenorphine
- 1 codeine
- 2 fentanyl
- 3 hydrocodone
- 4 hydromorphone
- 5 meperidine
- 6 methadone
- 7 morphine
- 8 oxycodone
- 9 oxymorphone
- a tramadol
- b other opioid
- T39 Poisoning by, adverse effect of and underdosing of nonopioid analgesics, antipyretics and antirheumatics
 - T39.1 Poisoning by, adverse effect of and underdosing of 4-Aminophenol derivatives

	derivativ	es	es				
Add	Code	e first appropriate code from category T40, Poisoning by, adverse effect of and underdosing of narcotics and psychodysleptics [hallucinogens], if 4-Aminophenol derivative is a combination 4-Aminophenol/opioid product					
New subcategory	T39.11	Poisoning bacetaminop	by, adverse effect of and underdosing of othen				
New code			Poisoning by acetaminophen, accidental unintentional)				
New code		T39.112 P	Poisoning by acetaminophen, intentional self-harm				
New code		T39.113 P	Poisoning by acetaminophen, assault				
New code		T39.114 P	Poisoning by acetaminophen, undetermined				
New code		T39.115 A	Adverse effect of acetaminophen				
New code		T39.116 U	Inderdosing of acetaminophen				
New subcategory	T39.19	Poisoning b	by, adverse effect of and underdosing of other 4-				

Aminophenol derivatives

New code		Т39	9.191	Poisoning by other 4-Aminophenol derivatives, accidental (unintentional) Poisoning by other 4-Aminophenol derivatives
New code		Т39	9.192	NOS Poisoning by other 4-Aminophenol derivatives,
New code		Т39	9.193	intentional self-harm Poisoning by other 4-Aminophenol derivatives, assault
New code		T39	9.194	Poisoning by other 4-Aminophenol derivatives, undetermined
New code New code				Adverse effect of other 4-Aminophenol derivatives Underdosing of other 4-Aminophenol derivatives
Add	T39.3	anti-inflamm Code firs eff [ha	atory of a	rse effect of and underdosing of other nonsteroidal drugs [NSAID] opriate code from category T40, Poisoning by, adverse and underdosing of narcotics and psychodysleptics ogens], if NSAID is a combination NSAID/opioid
T40	Poison	ing by, adverse	e effec	t of and underdosing of narcotics and
Add		underdosing	ode fro of non	gens] m category T39, Poisoning by, adverse effect of and opioid analgesics, antipyretics and antirheumatics, if mbination NSAID/opioid product
Revise	T40.0	Poisoning by, natural origir		rse effect of and underdosing of opium <u>and other</u> n alkaloids
			_	dverse effect of and underdosing of short acting rigin opium alkaloids
New subcategory New code		T40.01 Poi	isoning	by, adverse effect of and underdosing of opium Poisoning by opium, accidental (unintentional)
New code		T4(N N12	Poisoning by opium NOS Poisoning by opium, intentional self-harm
New code				Poisoning by opium, assault
New code				Poisoning by opium, undetermined
New code				Adverse effect of opium
New code		T40	0.016	Underdosing of opium
New subcategory			_	by, adverse effect of and underdosing of short
			_	prphine
		Excludes2:	poiso	ning by, adverse effect of and underdosing of long
New code		T40	0.021	acting morphine (T40.18-) Poisoning by short acting morphine, accidental (unintentional)
New code		T40	0.022	Poisoning by short acting morphine NOS Poisoning by short acting morphine, intentional self-

				harm
New code			T40.023	Poisoning by short acting morphine, assault
New code				Poisoning by short acting morphine, undetermined
New code				Adverse effect of short acting morphine
New code				Underdosing of short acting morphine
New subcategory		T40.03		g by, adverse effect of and underdosing of codeine
New code			-	Poisoning by codeine, accidental (unintentional)
				Poisoning by codeine NOS
New code			T40.032	Poisoning by codeine, intentional self-harm
New code			T40.033	Poisoning by codeine, assault
New code			T40.034	Poisoning by codeine, undetermined
New code			T40.035	Adverse effect of codeine
New code			T40.036	Underdosing of codeine
New subcategory		T40.09	Poisoning	g by, adverse effect of and underdosing of other
			natural o	rigin opium alkaloids
			Poiso	ning by, adverse effect of and underdosing of
				thebaine
New code			T40.091	Poisoning by other natural origin opium alkaloids,
				accidental (unintentional)
				Poisoning by other natural origin opium alkaloids
				NOS
New code			T40.092	Poisoning by other natural origin opium alkaloids,
				intentional self-harm
New code			T40.093	Poisoning by other natural origin opium alkaloids,
				assault
New code			T40.094	Poisoning by other natural origin opium alkaloids,
				undetermined
New code				Adverse effect of other natural origin opium alkaloids
New code				Underdosing of other natural origin opium alkaloids
Revise	T40.1		-	rse effect of and underdosing of heroin <u>and short</u>
	Eval. d		-	tic opioid derivatives
	EXCIUO	esz: poisc		dverse effect of and underdosing of long acting semi-
Nowsubsatagons		T40.11	•	opioid derivatives (T40.2-)
New subcategory New code		140.11		g by and adverse effect of heroin
New code			140.111	Poisoning by heroin, accidental (unintentional)
New code			T40.112	Poisoning by heroin NOS
New code				Poisoning by heroin, intentional self-harm
New code				Poisoning by heroin, assault Poisoning by heroin, undetermined
New code				Adverse effect of heroin
New code				Underdosing of heroin
New subcategory		T40.12		g by, adverse effect of and underdosing of short
ivew subcategory		140.12	-	dromorphone
New code				Poisoning by short acting hydromorphone, accidental
IVEVV COUC			170.121	(unintentional)
				Poisoning by short acting hydromorphone NOS
New code			T40 122	Poisoning by short acting hydromorphone,
				. olooling by short dethis hydrolliorpholic,

			intentional self-harm
New code			Poisoning by short acting hydromorphone, assault
New code			Poisoning by short acting hydromorphone,
			undetermined
New code			Adverse effect of short acting hydromorphone
New code			Underdosing of short acting hydromorphone
New subcategory	T40.13	_	by, adverse effect of and underdosing of short
		acting oxy	
New code		T40.131	Poisoning by short acting oxycodone, accidental
			(unintentional)
Newcoods		T40 122	Poisoning by short acting oxycodone NOS
New code			Poisoning by short acting oxycodone, intentional self-harm
New code			Poisoning by short acting oxycodone, assault
New code			Poisoning by short acting oxycodone, assault
New code			Adverse effect of short acting oxycodone
New code			Underdosing of short acting oxycodone
New subcategory	T40.14		by, adverse effect of and underdosing of short
New Subcategory	140.14	•	drocodone
New code			Poisoning by short acting hydrocodone, accidental
New code		140.141	(unintentional)
			Poisoning by short acting hydrocodone NOS
New code		T40.142	
			self-harm
New code		T40.143	Poisoning by short acting hydrocodone, assault
New code		T40.144	Poisoning by short acting hydrocodone,
			undetermined
New code		T40.145	Adverse effect of short acting hydrocodone
New code		T40.146	Underdosing of short acting hydrocodone
New subcategory	T40.15	Poisoning	by, adverse effect of and underdosing of short
		acting oxy	ymorphone
New code		T40.151	
			(unintentional)
			Poisoning by short acting oxymorphone NOS
New code		T40.152	Poisoning by short acting oxymorphone, intentional
		T40 450	self-harm
New code			Poisoning by short acting oxymorphone, assault
New code		140.154	Poisoning by short acting oxymorphone,
Nowcodo		T40 155	undetermined
New code			Adverse effect of short acting oxymorphone
New code	T40 10		Underdosing of short acting oxymorphone
New subcategory	T40.19	_	s by, adverse effect of and underdosing of other short
New code		_	mi-synthetic opioid derivatives
ivew code		140.131	Poisoning by other short acting semi-synthetic opioid derivatives, accidental (unintentional)
			Poisoning by other short acting semi-synthetic
			opioid derivatives NOS
			opiola derivatives 1803

New code			T40.192	Poisoning by other short acting semi-synthetic opioid derivatives, intentional self-harm
New code			T40.193	Poisoning by other short acting semi-synthetic opioid derivatives, assault
New code			T40.194	Poisoning by other short acting semi-synthetic opioid derivatives, undetermined
New code			T40.195	Adverse effect of other short acting semi-synthetic opioid derivatives
New code			T40.196	Underdosing of other short acting semi-synthetic opioid derivatives
Revise	T40.2		-	rse effect of and underdosing of long acting semi-
	Eval. d	-	opioid de	
	EXCIUO	esz: poisc		dverse effect of and underdosing of short acting thetic opioid derivatives (T40.1-)
New subcategory		T40.21	•	g by, adverse effect of and underdosing of long acting
ivew subcutegory		140.21	hydromo	
			•	oning by, adverse effect of and underdosing of CR
				hydromorphone
New code			T40.211	Poisoning by long acting hydromorphone, accidental
				(unintentional)
				Poisoning by long acting hydromorphone NOS
New code			T40.212	Poisoning by long acting hydromorphone, intentional
				self-harm
New code				Poisoning by long acting hydromorphone, assault
New code			T40.214	Poisoning by long acting hydromorphone, undetermined
New code			T40.215	Adverse effect of long acting hydromorphone
New code			T40.216	Underdosing of long acting hydromorphone
New subcategory		T40.22	Poisoning oxycodor	g by, adverse effect of and underdosing of long acting
				oning by, adverse effect of and underdosing of CR
				oxycodone
New code			T40.221	Poisoning by long acting oxycodone, accidental
				(unintentional)
				Poisoning by long acting oxycodone NOS
New code			T40.222	Poisoning by long acting oxycodone, intentional
				self-harm
New code				Poisoning by long acting oxycodone, assault
New code				Poisoning by long acting oxycodone, undetermined
New code				Adverse effect of long acting oxycodone
New code				Underdosing of long acting oxycodone
New subcategory		T40.23		g by, adverse effect of and underdosing of long acting
			morphine	
			Poisc	oning by, adverse effect of and underdosing of CR
Newcoods			T40 224	morphine
New code			140.231	Poisoning by long acting morphine, accidental
				(unintentional)

			Poisoning by long acting morphine NOS
New code		T40.232	Poisoning by long acting morphine, intentional self- harm
New code		T40.233	Poisoning by long acting morphine, assault
New code			Poisoning by long acting morphine, undetermined
New code			Adverse effect of long acting morphine
New code			Underdosing of long acting morphine
New subcategory	T40.24		g by, adverse effect of and underdosing of long acting
3 ,		buprenoi	
New code		T40.241	
			Poisoning by long acting buprenorphine NOS
New code		T40.242	Poisoning by long acting buprenorphine, intentional self-harm
New code		T40.243	Poisoning by long acting buprenorphine, assault
New code		T40.244	Poisoning by long acting buprenorphine, undetermined
New code		T40.245	Adverse effect of long acting buprenorphine
New code			Underdosing of long acting buprenorphine
New subcategory	T40.25		g by, adverse effect of and underdosing of long
- '			ymorphone
		Poisc	oning by, adverse effect of and underdosing of CR
			oxymorphone
New code		T40.251	Poisoning by long acting oxymorphone, accidental (unintentional)
			Poisoning by long acting oxymorphone NOS
New code		T40.252	Poisoning by long acting oxymorphone, intentional self-harm
New code		T40.253	Poisoning by long acting oxymorphone, assault
New code		T40.254	Poisoning by long acting oxymorphone, undetermined
New code		T40.255	Adverse effect of long acting oxymorphone
New code		T40.256	Underdosing of long acting oxymorphone
New subcategory	T40.29	Poisoning	g by, adverse effect of and underdosing of other long
		acting se	mi-synthetic opioid derivatives
New code		T40.291	Poisoning by other long acting semi-synthetic opioid derivatives, accidental (unintentional)
New code		T40.292	Poisoning by other long acting semi-synthetic opioid derivatives, intentional self-harm
New code		T40.293	Poisoning by other long acting semi-synthetic opioid derivatives, assault
New code		T40.294	Poisoning other long acting semi-synthetic opioid derivatives, undetermined
New code		T40.295	Adverse effect of other long acting semi-synthetic opioid derivatives
New code		T40.296	Underdosing of other long acting semi-synthetic opioid derivatives

Revise	T40.3 Poisoning by, adverse effect of and underdosing of methadone and other long acting synthetic opioids						
	Excludes 2: poisoning by, adverse effect of and underdosing of short acting						
	synthetic opioids (T40.4-)						
New subcategory	T4	0.31	•	Poisoning by, adverse effect of and underdosing of			
		0.0_	methado				
New code				Poisoning by methadone, accidental (unintentional)			
				Poisoning by methadone NOS			
New code			T40.312	Poisoning by methadone, intentional self-harm			
New code				Poisoning by methadone, assault			
New code				Poisoning by methadone, undetermined			
New code				Adverse effect of methadone			
New code				Underdosing of methadone			
New subcategory	T4	0.32		g by, adverse effect of and underdosing of long acting			
σ ,			fentanyl				
				oning by, adverse effect of and underdosing of			
				transdermal fentanyl (fentanyl patch)			
New code			T40.321	Poisoning by long acting fentanyl, accidental			
				(unintentional)			
New code			T40.322	Poisoning by long acting fentanyl, intentional self-			
				harm			
New code			T40.323	Poisoning by long acting fentanyl, assault			
New code			T40.324	Poisoning by long acting fentanyl, undetermined			
New code			T40.325	Adverse effect of long acting fentanyl			
New code			T40.326	Underdosing of long acting fentanyl			
New subcategory	T4	0.33	Poisoning	g by, adverse effect of and underdosing of long			
			acting tra	amadol			
			Poisc	oning by, adverse effect of and underdosing of CR			
				tramadol			
New code			T40.331	Poisoning by long acting tramadol, accidental			
				(unintentional)			
				Poisoning by long acting tramadol NOS			
New code			T40.332	Poisoning by long acting tramadol, intentional self-			
				harm			
New code				Poisoning by long acting tramadol, assault			
New code				Poisoning by long acting tramadol, undetermined			
New code				Adverse effect of long acting tramadol			
New code	Τ.4	0.20		Underdosing of long acting tramadol			
New subcategory	14	0.39		g by, adverse effect of and underdosing of other long			
			υ,	nthetic opioids			
			POISC	oning by, adverse effect of and underdosing of levo			
			alphacetylmethadol (LAAM)				
			PUISC	oning by, adverse effect of and underdosing of levorphanol			
New code			T40.391	Poisoning by other long acting synthetic opioids,			
NEW COUC			140.331	accidental (unintentional)			
				Poisoning by other long acting synthetic opioids			
				i organing by other long acting synthetic opiolos			

				NOS
New code			T40.392	Poisoning by other long acting synthetic opioids,
				intentional self-harm
New code			T40.393	Poisoning by other long acting synthetic opioids, assault
New code			T40.394	Poisoning by other long acting synthetic opioids, undetermined
New code			T40.395	Adverse effect of other long acting synthetic opioids
New code				underdosing of other long acting synthetic opioids
Revise	T40.4	Poisoning	g by, adver	rse effect of and underdosing of short acting synthetic
		<u>opioids</u>		
	Exclud	es2: poiso	ning by, a	dverse effect of and underdosing of long acting
			•	opioids (T40.3-)
New subcategory		T40.41		g by, adverse effect of and underdosing of short
			acting fer	<i>,</i>
New code			T40.411	Poisoning by short acting fentanyl, accidental (unintentional)
New code			T40.412	Poisoning by short acting fentanyl, intentional self-harm
New code			T40.413	
New code				Poisoning by short acting fentanyl, undetermined
New code				Adverse effect of short acting fentanyl
New code				Underdosing of short acting fentanyl
New subcategory		T40.42		g by, adverse effect of and underdosing of short
			acting me	
New code			_	Poisoning by short acting meperidine, accidental (unintentional)
New code			T40.422	Poisoning by short acting meperidine, intentional self-harm
New code			T40.423	Poisoning by short acting meperidine, assault
New code				Poisoning by short acting meperidine, undetermined
New code				Adverse effect of short acting meperidine
New code			T40.426	Underdosing of short acting meperidine
New subcategory		T40.43	Poisoning	g by, adverse effect of and underdosing of short
			acting tra	imadol
New code			T40.431	Poisoning by short acting tramadol, accidental (unintentional)
				Poisoning by short acting tramadol NOS
New code			T40.432	Poisoning by short acting tramadol, intentional self-harm
New code			T40.433	Poisoning by short acting tramadol, assault
New code			T40.434	Poisoning by short acting tramadol, undetermined
New code				Adverse effect of short acting tramadol
New code				Underdosing of short acting tramadol
New subcategory		T40.49		g by, adverse effect of and underdosing of other short
			-	nthetic opioids
				ning by, adverse effect of and underdosing of short

				acting butorphanol	
			Poisoning by, adverse effect of and underdosing of short		
			acting dextropropoxyphene		
			Poiso	oning by, adverse effect of and underdosing of short	
				acting pentazocine	
New code			T40.491	Poisoning by other short acting synthetic opioids,	
				accidental (unintentional)	
				Poisoning by other short acting synthetic opioids NOS	
New code			T40.492	Poisoning by other short acting synthetic opioids, intentional self-harm	
New code			T40.493	Poisoning by other short acting synthetic opioids, assault	
New code			T40.494	Poisoning by other short acting synthetic opioids,	
				undetermined	
New code			T40.495	Adverse effect of other short acting synthetic opioids	
New code			T40.496	Underdosing of other short acting synthetic opioids	
	T40.5	Poisonin	g by, adve	rse effect of and underdosing of cocaine	
New subcategory		T40.51	Poisoning	g by, adverse effect of and underdosing of powder	
			cocaine		
New code			T40.511	Poisoning by powder cocaine, accidental	
				(unintentional)	
				Poisoning by powder cocaine NOS	
New code				Poisoning by powder cocaine, intentional self-harm	
New code			T40.513	· · · · · · · · · · · · · · · · · · ·	
New code			T40.514		
New code				Adverse effect of powder cocaine	
New code				Underdosing of powder cocaine	
New subcategory		T40.52		g by, adverse effect of and underdosing of crack	
New code			cocaine	Deigning by erack cossine assidental (unintentional)	
new code			140.521	Poisoning by crack cocaine, accidental (unintentional) Poisoning by crack cocaine NOS	
New code			T40 522	Poisoning by crack cocaine NOS Poisoning by crack cocaine, intentional self-harm	
New code				Poisoning by crack cocaine, intentional sen-narm	
New code				Poisoning by crack cocaine, assault Poisoning by crack cocaine, undetermined	
New code				Adverse effect of crack cocaine	
New code				Underdosing of crack cocaine	
11311 0000				22.2.2.200 3. 3. 3. 3. 3. 3. 3. 3. 3. 3. 3. 3. 3.	

ICD-10-CM Weeks of gestation of pregnancy

The American Medical Association's Physician Consortium for Performance Improvement® (PCPI) convened a Maternity Care Work Group (MCWG) that included members from health plans, hospitals, patients, consumers and health professionals, to develop physician-level performance measures. During the meeting the work group reached consensus on the potential usefulness of having additional ICD-10-CM codes for specific gestational age. This information is considered important for quality improvement and measurement purposes. The benefits of documenting specific gestational age within coding includes the ability to risk stratify and/or adjust for measurement as well as the ability to identify preterm and late pre-term births.

The work group submitted a proposal to the American College of Obstetricians and Gynecologists' (ACOG) Committee on Coding and Nomenclature who then submitted the proposal to the ICD-9-CM Coordination and Maintenance Committee (C&M) for consideration for inclusion in ICD-10-CM (once it is implemented) since it is not possible to add this amount of detail to ICD-9-CM.

The codes in ICD-10-CM Chapter 15, Pregnancy, childbirth and the puerperium, do include the trimester of the pregnancy. This designation was included in the ICD-10-CM at the request of ACOG during the initial development of ICD-10-CM in 1994. Though trimester is still considered important, the specific gestational age is now considered a more precise indicator of risk.

The initial proposal of the work group included indicating days within the week of gestation as well as the method used (such as ultrasound or estimated date of conception) in determining gestation. These methods cannot be included within an ICD status code. However, the National Center for Health Statistics (NCHS) recognizes the importance of this information in ICD-10-CM and has designed the proposal to fit within the structure of the classification. This proposal has been accepted by ACOG and the MCWG. This proposal does not alter the existing Chapter 15 codes. Instead, it provides a secondary status code to be used in conjunction with a Chapter 15 code to identify the specific gestational age. Should this information not be available in the medical record and unspecified code within each trimester subcategory is provided to indicate that.

ICD-10-CM TABULAR MODIFICATIONS

CHAPTER 15

Pregnancy, childbirth and the puerperium (O00-O99)

Note: Trimesters are counted from the first day of the last menstrual period. They are defined as

follows:

1st trimester- less than 14 weeks 0 days

2nd trimester- 14 weeks 0 days to less than 28 weeks 0 days

3rd trimester- 28 weeks 0 days until delivery

Add Use an additional code from category Z35, Weeks of gestation, to identify the specific week the

pregnancy

New category Z35 Weeks of gestation

Note: Codes from category Z35 are for use on the maternal record only in conjunction with a code from Chapter 15, Pregnancy, childbirth and the puerperium, to indicate the weeks of gestation of the pregnancy. They are never for use on a newborn record.

New subcategory	Z35.1	First trim	ester, weeks 0-14
New code		Z35.10	First trimester, unspecified week
New code		Z35.11	First trimester, week 0 to week 7
New code		Z35.12	First trimester, week 8 to week 14
New subcategory	Z35.2	Second to	rimester, weeks 15-21 and unspecified
New code		Z35.20	Second trimester, unspecified week
New code		Z35.21	Second trimester, week 15
New code		Z35.22	Second trimester, week 16
New code		Z35.23	Second trimester, week 17
New code		Z35.24	Second trimester, week 18
New code		Z35.25	Second trimester, week 19
New code		Z35.26	Second trimester, week 20
New code		Z35.27	Second trimester, week 21
New subcategory	Z35.3	Second to	rimester, weeks 22-28
New code		Z35.31	Second trimester, week 22
New code		Z35.32	Second trimester, week 23
New code		Z35.33	Second trimester, week 24
New code		Z35.34	Second trimester, week 25
New code		Z35.35	Second trimester, week 26
New code		Z35.36	Second trimester, week 27
New code		Z35.37	Second trimester, week 28
New subcategory	Z35.4	Third trin	nester, weeks 29-35 and unspecified
New code	233.4	Z35.40	Third trimester, unspecified week
New code		Z35.40	Third trimester, wispectified week Third trimester, week 29
New code		Z35.41	Third trimester, week 29 Third trimester, week 30
New code		Z35.42 Z35.43	Third trimester, week 30 Third trimester, week 31
New code		Z35.43	Third trimester, week 31 Third trimester, week 32
New code		Z35.44 Z35.45	Third trimester, week 32 Third trimester, week 33
New code		Z35.45	Third trimester, week 33 Third trimester, week 34
New code		Z35.40	Third trimester, week 34 Third trimester, week 35
New code		233.47	Tillia tillilester, week 55
New subcategory	Z35.5	Third trin	nester, weeks 36-40
New code		Z35.51	Third trimester, week 36
New code		Z35.52	Third trimester, week 37
New code		Z35.53	Third trimester, week 38
New code		Z35.54	Third trimester, week 39
New code		Z35.55	Third trimester, week 40

New subcategory	Z35.6	Over 40 c	completed weeks of gestation
New code		Z35.61	41 weeks gestation
New code		Z35.62	42 weeks gestation
New code		Z35.63	43 weeks gestation

ICD-10-CM Weeks of gestation for newborn

The National Association of Children's Hospitals and Related Institutions (NACHRI) and the American Academy of Pediatrics (AAP) request that the weeks of gestation codes in ICD-10-CM be revised and expanded to even more accurately reflect prematurity of the newborn then the current codes in ICD-9-CM (765.2x).

The current ICD-9-CM code set has been very helpful to both NACHRI and the Academy in analyses of premature newborn patients using the birthweight diagnosis codes, the gestational age diagnosis codes, and various combinations of these diagnosis codes. These are the most fundamental descriptors of premature newborns. It is especially striking to see how much mortality varies within a birthweight range when further stratified by gestational age. Each week of gestational age makes a big difference in terms of outcomes: perinatal complications, in-hospital mortality, length of stay, and longer term morbidity and mortality. The differences are especially pronounced for the more premature newborns. We believe an expansion of these codes will be in the best interest of these children.

AAP and NACHRI believe that detailed analysis would be much improved if the prematurity codes in ICD-10-CM are expanded. For example, it would be more difficult to differentiate mortality for a 24 week newborn (which is higher) versus a 25 or 26 week newborn since these weeks of ages are all included at code P07.22, Extreme immaturity of newborn, 24-26 completed weeks.

Each week of gestational age makes a big difference in terms of outcomes: perinatal complications, in-hospital mortality, length of stay, and longer term morbidity and mortality. The differences are especially pronounced for the more premature newborns. NACHRI and AAP believe an expansion of P07.2 and P07.3 will be in the best interest of these children. With the additional available space, the expansion of this code set will allow for even better morbidity and mortality analysis. To this end NACHRI and AAP recommend the following expansion of the codes to become effective for the 2013 version of ICD-10-CM.

TABULAR MODIFICATIONS

P07.2 Extreme immaturity of newborn

Less than 28 completed weeks (less than 196 completed days) of gestation

P07.20 Extreme immaturity of newborn, unspecified weeks

Add Gestation less than 28 completed weeks

Revise P07.21 Extreme immaturity of newborn, gestational age less than 23 24

completed weeks

Revise P07.22 Extreme immaturity of newborn, gestational age 23 24-26 completed

weeks

Revise	P07.23 Extreme immaturity of newborn, gestational age 24 27 completed weeks
New code	P07.24 Extreme immaturity of newborn, gestational age 25 completed weeks
New code	P07.25 Extreme immaturity of newborn, gestational age 26 completed weeks
New code	P07.26 Extreme immaturity of newborn, gestational age 27 completed weeks
Revise P07.3	Other preterm [premature] newborn [other]
	28 completed weeks or more but less than 37 completed weeks (196 completed days but less than 259 completed days) of gestation.
	Prematurity NOS
Revise	P07.30 Other preterm Preterm newborn, unspecified weeks
Revise weeks	P07.31 Other preterm Preterm newborn, gestational age 28-31 completed
Revise	P07.32 Other preterm Preterm newborn, gestational age 29 32-36 completed weeks
New code	P07.33 Preterm newborn, gestational age 30 completed weeks
New code	P07.34 Preterm newborn, gestational age 31 completed weeks
New code	P07.35 Preterm newborn, gestational age 32 completed weeks
New code	P07.36 Preterm newborn, gestational age 33 completed weeks
New code	P07.37 Preterm newborn, gestational age 34 completed weeks
New code	P07.38 Preterm newborn, gestational age 35 completed weeks
New code	P07.39 Preterm newborn, gestational age 36 completed weeks

Benign neoplasm of genitourinary organs

The American Urological Association (AUA) has requested that benign neoplasms of the genitourinary organs be given more specific codes as it would be helpful to identify specific areas where neoplasm are most prevalent. For example, angiomyolipomas, a benign neoplasm, are a common occurrence in male urology patients in their 50's. When an angiomyolipoma grows to over 4 cm, the treatment options can either be non-surgical or surgical. While benign, they can cause other symptoms or spontaneous bleeding. Currently, the only unique code for benign neoplasm involving the genitourinary organs is that for the spermatic cord, D17.6, Benign lipomatous neoplasm of spermatic cord.

TABULAR MODIFICATIONS

D17.7 Benign lipomatous neoplasm of other sites

New code	D17./1 Benign lipomatous neoplasm of kidney
New code	D17.72 Benign lipomatous neoplasm of other genitourinary organ
New code	D17.79 Benign lipomatous neoplasm of other sites

Benign lipomatous neoplasm of peritoneum Benign lipomatous neoplasm of retroperitoneum

Urethral False Passage

In ICD-9-CM, there was a specific diagnosis code for urethral false passage (599.4). This code was included in the original WHO ICD-9 but was not included in the WHO ICD-10. A false urethral passage is a generally a traumatic occurrence caused by instrumentation or catheterization and causes a by-pass of the normal urethra. The intervention of a false urethral passage is either by catheter drainage or by allowing the false passage to heal on its own. The American Urological Association (AUA) feels that this is an occurrence that happens frequently enough in urology patients that there should be a separate diagnosis code in ICD-10 and not included in the N36.0 as an inclusion term.

TABULAR MODIFICATIONS

Other diseases of the urinary system (N30-N39)

N36 Other disorders of the urethra

N36.0 Urethral fistula

Delete False urethral passage

New code N36.3 Urethral false passage

Nodular prostate

A nodular prostate appears on initial examination where the urologist can feel a hardened area in the prostate. This occurs frequently in male patients and can represent several diagnoses when more tests are completed. These additional diagnoses can be prostatic stones, granulomatous prostatitis or prostate cancer. Unique codes have existed in ICD-9-CM since October 1, 2003 when distinctions were added: code 600.10, Nodular prostate without urinary obstruction and code 600.11, Nodular prostate with urinary obstruction. The American Urological Association (AUA) requests that nodular prostate continue to have its own ICD-10-CM diagnosis codes but change the descriptor to follow the current nomenclature to include "with" and "without lower urinary tract symptoms" as suggested above to maintain the ability to track the different diagnosis of nodular prostate which is a significantly different diagnosis from hypertrophy and hyperplasia.

TABULAR MODIFICATIONS

Diseases of male genital organs (N40-N51)

N40 Enlarged prostate (EP)

New code N40.2 Nodular prostate with lower urinary tract symptoms

(LUTS)

New code N40.3 Nodular prostate without lower urinary tract symptoms

(LUTS)

Inflammatory disease of the prostate

Prostatitis is an inflammation of the prostate gland in males. It is a common problem for men at any age but specifically for men in their 20's and 30's. The American Urological Association (AUA) now questions the codes created for prostatitis without and with hematuria. The urologist does not look for hematuria with prostatitis. The AUA believes that the suggested changes to the inflammatory disease of the prostate section is better served by eliminating the "with and without hematuria". Please note that this deletion is being proposed now before the freeze and implementation of ICD-10-CM. As has been the established practice no codes will be deleted from ICD-10-CM after October 1, 2013.

TABULAR MODIFICATIONS

Delete Delete	N41.0	Acute prostatitis N41.00 Acute prostatitis without hematuria N41.01 Acute prostatitis with hematuria
	N41.1	Chronic prostatitis
Delete		N41.10 Chronic prostatitis without hematuria
Delete		N41.11 Chronic prostatitis with hematuria

Cyst of the prostate

A cyst in the prostate is a fluid filled cavity. For the most part, the cyst is asymptomatic but may lead to perineal discomfort. In ICD-9-CM, cyst of the prostate was given a diagnosis code of 600.3 (October 1, 2000). In ICD-10 CM, cyst of the prostate is included under code N42.89, "Other specified disorders of the prostate." The American Urological Association (AUA) believes that a separate designation of cyst of prostate should be included in ICD-10-CM in order to capture this urologic diagnosis which occurs frequently.

TABULAR MODIFICATIONS

N42 Other and unspecified disorders of prostate

N42.8 Other specified disorders of prostate

New code N42.83 Cyst of prostate

N42.89Other specified disorders of prostate

Delete Cyst of prostate

Acquired and congenital torsion of the penis

The American Urological Association (AUA) believes that there should be ICD-10-CM codes to incorporate both the congenital penile torsion and the acquired penile torsion. The AUA believes that these diagnoses should be given their own codes given the fact that the congenital torsion occurs in pediatric patients and requires that the torsion of the penis be separated out from the N48.89 other specified disorders of the penis and Q55.69 Other congenital malformation of penis NOS. The congenital torsion of the penis occurs in pediatric patients in the first trimester in-utero and is a common occurrence in pediatric patients. The urologist can decide to correct the problem based on the severity. In the acquired torsion, the twist may occur as a result of the repair of a hypospadias condition of the penis.

TABULAR MODIFICATIONS

N48 Other disorders of penis

N48.8 Other specified disorders of penis

New code N48.82 Acquired torsion of penis, acquired

Acquired torsion of penis NOS

Excludes 1: Congenital torsion of penis (Q55.63)

Q55 Other congenital malformations of male genital organs

Q55.6 Other congenital malformations of penis

New code Q55.63 Congenital torsion of penis

Excludes 1: Acquired torsion of penis (N48.82)

Cyst of the epididymis

The American Urological Association (AUA) has proposed the addition of an ICD-10-CM code to track the occurrence of cysts in the male epididymis. As with the cyst of the prostate, the cyst of the epididymis occurs frequently in male urology patients. It is a fluid filled growth on the epididymis that may remain asymptomatic or may cause perineal pain.

TABULAR MODIFICATIONS

N50 Other and unspecified disorders of male genital organs New code N50.3 Cyst of epididymis

Hidden penis

A unique was added to ICD-9-CM effective October 1, 1996, for hidden penis (752.65). There is no code in ICD-10-CM to report this diagnosis except for the other congenital malformation of penis NOS code. This is an occurrence that frequently appears in babies and young children as well as adults and the American Urological Association (AUA) recommends that it is appropriate to have a separate diagnosis code for this anomaly in ICD-10-CM.

TABULAR MODIFICATIONS

Q55 Other congenital malformations of male genital organs

Q55.6 Other congenital malformations of penis
New code
Q55.64 Hidden penis
Buried penis
Concealed penis

Personal history of malignant neoplasm of ureter

Cancer of the ureter is a frequent occurrence in the urologic patient. A patient with a personal history of ureteral cancer needs to monitored closely as there can be a recurrence on the contralateral ureter or may even appear as a secondary cancer in the bladder. The American Urological Association (AUA) believes that there is a significant enough occurrence of cancer of the ureter to warrant a code for personal history and therefore there should be a separate code for neoplasm of ureter.

TABULAR MODIFICATIONS

Z85 Personal history of malignant neoplasm

Z85.5 Personal history of malignant neoplasm of urinary tract

New code Z85.54 Personal history of malignant neoplasm of ureter

Visual agnosia and related conditions (ICD-10-CM only)

Currently the conditions visual agnosia and prosopagnosia are included in ICD-10-CM code H53.16, Psychophysical visual disturbances. These were included here as a carryover of addenda changes done recently to ICD-9-CM. However, these are symbolic dysfunctions and more appropriately categorized as such. Simultanagnosia (asimultagnosia) is a symbolic dysfunction in which the patient can process individual elements of a visual presentation, but not the whole presentation at once.

The American Academy of Neurology (AAN) proposes deleting the terms prosopagnosia and visual object agnosia from code H53.16, and adding a unique code for visual agnosia. These proposed changes are shown below.

ICD-10-CM TABULAR MODIFICATIONS

H53 Visual disturbances

H53.1 Subjective visual disturbances

H53.16 Psychophysical visual disturbances

Delete <u>Prosopagnosia</u>

Delete Visual object agnosia

R48 Dyslexia and other symbolic dysfunctions, not elsewhere classified

R48.1 Agnosia

Revise Excludes1: visual object agnosia (H53.16) (R48.3)

New code R48.3 Visual agnosia

Prosopagnosia

Simultanagnosia (asimultagnosia)

Displacement/dislocation of internal hip prosthesis titles (ICD-10-CM Only)

The terms "dislocation" and "instability" are specific and clinically relevant for the knee. Though complete dislocation of prosthetic knees is extremely rare instability of prosthetic knees is a common indication for revision surgery. For this reason the American Academy of Orthopaedic Surgeons (AAOS) is requesting the following revisions to ICD-10-CM code titles:

TABULAR MODIFICATIONS

T84. Complications of internal orthopedic prosthetic devices, implants and grafts

T84.0 Mechanical complication of internal joint prosthesis

T84.02 Dislocation of internal joint prosthesis
Instability of internal joint prosthesis
Subluxation of internal joint prosthesis

T84.022 Dislocation Instability of internal right knee prosthesis

T84.023 Dislocation Instability of internal left knee prosthesis

Gastroparesis

A request has been received from Clarity Coding to add a unique code to ICD-10-CM for gastroparesis. The condition is currently indexed to K31.89, Other diseases of stomach and duodenum.

Gastroparesis is a chronic disorder of the stomach characterized by abnormal motility and delayed gastric emptying. Because the stomach cannot properly macerate food or propel the bolus into the small intestine, the patient's digestive and nutritional health suffer.

Symptoms of gastroparesis include early satiety, bloating, epigastric and upper abdominal pain, chronic nausea, and frequent vomiting. In some patients, nausea and vomiting are intractable. Gastroparesis can lead to repeated episodes of dehydration and electrolyte imbalance as well as severe nutritional compromise and extreme weight loss.

The most common cause of gastroparesis is diabetes. Other causes include infections, endocrine disorders, connective tissue disorders such as scleroderma, autoimmune conditions, neuromuscular diseases, cancer, some forms of chemotherapy and radiation therapy, and surgery of the upper intestinal tract. It's believed that, in many of these scenarios, gastroparesis may be associated with some level of vagal nerve damage, because this nerve regulates gastrointestinal peristalsis.

Although gastroparesis is frequently due to diabetes or another known underlying condition, in about one-third to one-half of cases no underlying diagnosis is identified. Some research indicates that idiopathic gastroparesis is at least as common as diabetic gastroparesis.

A unique code for gastroparesis was added to ICD-9-CM effective October 1, 1994. NCHS agrees with the importance of this code being included in ICD-10-CM.

K31 Other diseases of stomach and duodenum
Includes: functional disorders of stomach
Excludes2: diabetic gastroparesis (E08.43, E09.43, E10.43, E11.43, E13.43)
diverticulum of duodenum (K57.00-K57.11)

K31.4 Gastroparesis
Gastroparalysis

Code first underlying disease, if known, such as: anorexia nervosa (F50.0-) diabetes mellitus (E08.43, E09.43) scleroderma (M34.-)

ICD-10-CM TABULAR PROPOSED ADDENDA ITEMS

G31 Other degenerative diseases of nervous system, not elsewhere classified

G31.0 Frontotemporal dementia

G31.01 Pick's disease

Delete Circumscribed brain atrophy
Add Primary progressive aphasia

ICD-10-CM INDEX PROPOSED ADDENDA ITEMS

Atrophy, atrophic (of)

- brain (cortex) (progressive) G31.9

Revise -- <u>frontotemporal</u> circumscribed G31.01 [F02.80]

Revise Gerstmann's syndrome (developmental) F81.2 R48.8

Add - developmental F81.2

Isoimmunization NEC - see also Incompatibility

- affecting management of pregnancy (ABO) (with hydrops fetalis) O36.11-

Add -- anti-c sensitization O36.09-Add -- anti-C sensitization O36.09-Add -- anti-e sensitization O36.09-

Revise -- anti-E sensitization O36.19- <u>O36.09</u>-