

# ICD-9-CM Coordination and Maintenance Committee Meeting

December 6, 2002

## Diagnosis Agenda

Welcome and announcements Donna Pickett, MPH, RHIA Co-chair, ICD-9-CM C & M Committee	
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John Hart, M.D. American Academy of Neurology	
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## ICD-9-CM TIME LINE

## ICD-9-CM Coordination and Maintenance Committee Meeting

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- December 5-6, 2002 ICD-9-CM Coordination and Maintenance Committee meeting in the CMS auditorium. Diagnosis and procedure code revisions discussed are for potential implementation on October 1, 2003.
- January 10, 2003 Deadline for receipt of public comments on proposed code revisions discussed at the April and December 2002 ICD-9-CM Coordination and Maintenance Committee meetings. These proposals are being considered for implementation on October 1, 2003.
- February 3, 2003 Deadline for submission of proposals to CMS for procedures and NCHS for diagnoses for presentation at the April 3-4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting.
- March 2003 Tentative agenda for the Procedure part of the April 3-4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: <http://www.hcfa.gov/medicare/icd9cm.htm>
- Tentative agenda for the Diagnosis part of the April 3-4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on NCHS homepage as follows: <http://www.cdc.gov/nchs/icd9.htm>
- Federal Register Notice of April 3-4, 2003 ICD-9-CM Coordination and Maintenance Meeting and tentative agenda will be published.
- April 3-4, 2003 ICD-9-CM Coordination and Maintenance Committee Meeting in the CMS auditorium. Diagnosis code revisions discussed are for potential implementation on October 1, 2004. Procedure code revisions discussed will be for possible implementation October 1, 2003. Those procedure code proposals that cannot be resolved quickly will be considered for implementation on October 1, 2004.
- April 2003 Summary report of the Procedure part of the April 3, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: <http://www.hcfa.gov/medicare/icd9cm.htm>
- Summary report of the Diagnosis part of the April 4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting report will be posted on

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NCHS homepage as follows: <http://www.cdc.gov/nchs/icd9.htm>

- October 1, 2003 New and revised ICD-9-CM codes from the 2002 cycle go into effect.
- October 1, 2003 Deadline for submission of proposals to CMS for procedures and NCHS for diagnoses for presentation at the December 4-5, 2003 ICD-9-CM Coordination and Maintenance Committee meeting.
- November 2003 Tentative agenda for the Procedure part of the December 4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: <http://www.hcfa.gov/medicare/icd9cm.htm>
- Tentative agenda for the Diagnosis part of the December 5, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on NCHS homepage as follows: <http://www.cdc.gov/nchs/icd9.htm>
- Federal Register Notice of December 4-5, 2003 ICD-9-CM Coordination and Maintenance Meeting and tentative agenda to be published.
- Dec. 4-5, 2003 ICD-9-CM Coordination and Maintenance Committee Meeting. Code revisions discussed are for potential implementation on October 1, 2004. December 4 will be devoted to discussions of procedure codes. December 5 will be devoted to discussions of diagnosis codes.
- December 2003 Summary report of the Procedure part of the December 4, 2003 ICD-9-CM Coordination and Maintenance Committee meeting will be posted on CMS homepage as follows: <http://www.hcfa.gov/medicare/icd9cm.htm>
- Summary report of the Diagnosis part of the December 5, 2003 ICD-9-CM Coordination and Maintenance Committee meeting report will be posted on NCHS homepage as follows: <http://www.cdc.gov/nchs/icd9.htm>
- January 10, 2004 Deadline for receipt of public comments on proposed code revisions discussed at the April 3-4, 2003 and December 4-5, 2003 ICD-9-CM Coordination and Maintenance Committee meetings. These proposals are being considered for implementation on October 1, 2004.

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Septic shock and Sepsis

At the November 2, 2001 C & M meeting codes for systemic inflammatory response syndrome (SIRS) and severe sepsis were presented by Eli Lilly. Those new codes became effective October 1, 2002. To build on the new codes Lilly is requesting:

1. A unique code for septic shock.
2. Modifications to the index and tabular to distinguish the terms sepsis and septicemia.

The terms septic shock, severe sepsis, sepsis and septicemia are often used interchangeably by clinicians but are clinically distinct conditions that have very different clinical pictures and outcomes. Septic shock is a unique form of shock because of combined decreased systemic vascular resistance and the presence of myocardial performance. The clinical picture of severe sepsis, SIRS with organ dysfunction, must be present for a patient to advance to septic shock. Septic shock is the end point in the continuum from sepsis to severe sepsis to septic shock. Mortality from severe sepsis is between 20-50% of patients. When severe sepsis advances to septic shock the mortality rises to 50-60%. Because no unique code for septic shock exists, it is currently not possible to classify this component of a patient's illness.

The term septicemia is an imprecise term that simply means the presence of bacteria, or their toxins, in the bloodstream. This is equivalent to the term bacteremia (ICD-9-CM code 790.7). The term sepsis is specifically for an infection accompanied by the systemic inflammatory response (SIRS). Two or more of the SIRS criteria must be met for a septicemia to clinically be sepsis. Organ dysfunction must accompany the SIRS for the diagnosis of severe sepsis.

ICD-9-CM category 038 is titled Septicemia. Because of this title it is not possible to know if the patient's assigned codes from this category suffer from a simple septicemia or a true sepsis according to the definitions of the terms.

In addition to the new code for septic shock it is being proposed that the term sepsis be added as an inclusion under code 995.91, Systemic inflammatory response syndrome due to infectious process without organ dysfunction and that the term sepsis be indexed to 995.91. A note at both category 038 and at code 785.52 would be added to instruct users.

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**TABULAR MODIFICATIONS**

	038	Septicemia
Add		Note: A code for the corresponding systemic inflammatory response syndrome (SIRS), 995.91-995.92 must accompany a code from category 038. Sequencing is discretionary.
	785	Symptoms involving cardiovascular system
	785.5	Shock without mention of trauma
New code	785.52	Septic shock
Add		Note: code 785.52 must be used in conjunction with code 995.92, Systemic inflammatory response syndrome due to infectious process with organ dysfunction. Sequencing is discretionary.
	785.59	Other
		Shock:
Delete		septic

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995 Certain adverse effects not elsewhere classified

995.9 Systemic inflammatory response syndrome (SIRS)

Add 995.91 Systemic inflammatory response syndrome due to infectious process without organ dysfunction  
Sepsis

995.92 Systemic inflammatory response syndrome due to infectious process with organ dysfunction

Revise Use additional code to specify organ dysfunction, such as:  
Delete encephalopathy (348.31-348.32)  
~~heart failure (428.0-428.9)~~

995.94 Systemic inflammatory response syndrome due to noninfectious process with organ dysfunction

Revise Use additional code to specify organ dysfunction, such as:  
Delete encephalopathy (348.31-348.32)  
~~heart failure (428.0-428.9)~~

**INDEX MODIFICATION**

Revise Sepsis (generalized) (~~see also Septicemia) 038.9~~ 995.91

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Topic: Dementia

The American Academy of Neurology has requested new ICD-9-CM codes for two forms of dementia, Dementia with Lewy bodies and Frontotemporal dementia. Dementia classification has expanded with new knowledge of pathology and pathophysiology. Dementia and geriatric specialists are increasingly using new terms to describe their patients.

Dementia with Lewy bodies is a dementia with Parkinsonian motor features. Unexplained falls are a predominant early feature. Pathologically, it differs from Parkinson's disease in that the Lewy body intracellular inclusions are also found frontally, not just within the basal ganglia. This cerebral degeneration is also manifest by somewhat greater neuropsychiatric features than Alzheimer's with more prominent hallucinations. Patients with this form of dementia classically may worsen with phenothiazine-like medications which are used otherwise for psychoses.

Frontotemporal dementia differs from Alzheimer's disease by its frontal lobe personality changes, such as impulsivity, disinhibition, motor impersistence, loss of social awareness and lack of attention to personal hygiene. Language can be affected. Other cognition may not be as impaired as in Alzheimer's disease. One subset of Frontotemporal dementia is Pick's disease, that has its own code. Frontotemporal dementia would need to be excluded from Pick's disease.

### TABULAR MODIFICATIONS

	331	Other cerebral degenerations
	331.1	Pick's disease
Add		Excludes: Frontotemporal dementia (331.83)
	331.8	Other cerebral degenerations
New code	331.82	Dementia with Lewy bodies
Add		Lewy body dementia
Add		Lewy body disease
New code	331.83	Frontotemporal dementia
Add		Excludes: Pick's disease (331.1)

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Scrotal transposition

Scrotal transposition is a congenital anomaly where the scrotum is transposed above the penis. It looks like the penis and scrotum are reversed. The testes are also involved. It is treated surgically. There is no specific code for this anomaly. Such a code is requested by the American Urological Association.

### TABULAR MODIFICATION

752 Congenital anomalies of genital organs

752.8 Other specified anomalies of genital organs

Delete

~~Absence of:~~

~~prostate~~

~~spermatic cord~~

~~vas deferens~~

~~Anorchism~~

~~Aplasia (congenital) of:~~

~~prostate~~

~~round ligament~~

~~testicle~~

~~Atresia of:~~

~~ejaculatory duct~~

~~vas deferens~~

~~Fusion of testes~~

~~Hypoplasia of testis~~

~~Monorchism~~

~~Polyorchism~~

New code

752.81 Scrotal transposition

New code

752.89 Other specified anomalies of genital organs



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Topic: Peyronie's disease

Peyronie's disease is fibrosis of the cavernous sheaths leading to contracture of the investing fascia of the corpora resulting in a deviated and painful erection. The cause is unknown. The disease occurs in adult males. Resolution may occur spontaneously over many months. Minor disease that does not cause sexual dysfunction does not warrant treatment. Treatment results are unpredictable.

There is no unique code for Peyronie's disease in the ICD-9-CM. The American Urological Association is requesting a code.

TABULAR MODIFICATION

607 Disorders of penis

607.8 Other specified disorders of penis

New code                      607.85 Peyronie's disease

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Penile injury

Although penetrating and perforating injuries to the penis occur, severe injuries are more commonly caused by crushing blows and avulsion of the skin and genitalia. Avulsion injuries among factory and farm workers can happen when clothing is caught in machinery. Avulsion of the penis may also occur secondary to various devices (e.g., penile rings, vacuum cleaner attachments) or excessive trauma during sex. Avulsed skin should be conserved, cooled, and reapplied as quickly as possible. Debridement should be conservative. Skin grafting may be necessary. Even a completely transected penis may be reattached successfully.

The ICD-9-CM does not provide a code for injury of the penis. All injuries to the external genital organs are grouped under code 959.1, Injury of trunk. It has been requested that the external genital organs be separated out of 959.1 to allow the specific coding of penile injuries. Additional codes for other sites are also included with this modification.

### TABULAR MODIFICATIONS

959 Injury, other and unspecified

	959.1	Trunk
Delete		<del>Abdominal wall</del>
		<del>Back</del>
		<del>Breast</del>
		<del>Buttock</del>
		<del>Chest wall</del>
		<del>External genital organs</del>
		<del>Flank</del>
		<del>Groin</del>
		<del>Interscapular region</del>
		<del>Perineum</del>
New code	959.11	Breast
New code	959.12	Penis
New code	959.13	Scrotum
New code	959.14	Testes
New code	959.15	Vulva
New code	959.16	Buttock
New code	959.19	Injury of other sites of trunk

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Urgency of urination

The American Urological Association is requesting an ICD-9-CM code for the symptom of urgency. Urgency is the common symptom of feeling an intense need to urinate. It is now being inappropriately coded with the urge incontinence ICD-9-CM code. The symptom of urgency is as common as the diagnosis of frequency. The difference between urge incontinence and urgency is the intense feeling of having to urinate whereas urge incontinence is the intense feeling of having to urinate but not being able to make it to the restroom.

### TABULAR MODIFICATION

788 Symptoms involving urinary system

788.6 Other abnormality of urination

New code                      788.63 Urgency of urination

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Topic: Pre-operative insertion of ureteral stent for ureteral visualization

In many cases a urologist is called in to assist with an abdominal or pelvic procedure. The urologist may need to insert ureteral stents pre-operatively to assist the other surgeon in visualizing the ureters prior to open abdominal exploration or hysterectomy. There is no code that explains this situation. Such a code is being requested by the American Urological Association.

**TABULAR MODIFICATION**

593 Other disorders of kidney and ureter

593.8 Other specified disorders of kidney and ureter

New code                      593.83 Nonvisible ureter

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Impaired fasting glucose

On July 23, 1997 the Expert Committee on the Diagnosis and Classification of Diabetes mellitus released new recommendations which include a new stage of impaired glucose homeostasis called impaired fasting glucose (IFG). The Committee defined IFG as a random fasting glucose of  $\geq 110$ mg/dl but  $\leq 126$  mg/dl. This is distinct from an abnormal glucose tolerance (IGT) which is defined as a glucose obtained during an oral glucose tolerance test of  $\geq 140$ mg/dl but  $< 200$ mg/dl. These metabolic stages are intermediate between and separate from normal glucose homeostasis and diabetes.

Although there are usually no clinical signs or symptoms related to IFG, the clinical implications of early identification of patients at risk for diabetes and its associated cardiovascular disease are important. The first and second phase of insulin release are impaired in patients with IFG indicating beta cell dysfunction. Cardiovascular risk factors are significantly raised in all types of non-diabetic hyperglycemia. The greatest impact on mediating cardiovascular complications will occur in patients identified at the earliest stages of impaired glucose homeostasis.

The American Association of Clinical Endocrinologists has requested that a new code for IFG be created to allow for the identification of these patients.

### TABULAR MODIFICATIONS

	790	Nonspecific findings on examination of blood
Revise	790.2	Abnormal glucose <del>tolerance</del> test
Add		Excludes: dysmetabolic syndrome X (277.7)
Add		glycosuria (791.5)
New code	790.21	Impaired fasting glucose
New code	790.22	Abnormal glucose tolerance test

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Carnitine deficiency

Carnitine is an essential metabolic intermediary that is a critical factor for mitochondrial energy production and essential for normal biochemical function. The biochemical importance of L-carnitine is mainly due to its property of facilitating the transport of long-chain fatty acid inside the mitochondria for beta-oxidation. L-carnitine is also important for acting as an acyl group acceptor in order to maintain sufficient cellular levels of free coenzyme A (CoA) and for removing from the cell possible toxic acyl-groups that cannot be metabolized.

Carnitine deficiency occurs with certain genetic abnormalities, certain medical conditions, and from treatments for disease. Patients with carnitine deficiency can manifest with cardiomyopathy, Reye-like encephalopathy, hypoketotic hypoglycemia, hypotonia, muscle weakness, failure to thrive, and certain dialysis related problems such as intra-dialytic hypotension and erythropoietin-resistant anemia.

There are two forms of carnitine deficiency that are now recognized by the scientific community, primary and secondary. Primary carnitine deficiency is caused by a defect in the transporter responsible for the carnitine uptake from plasma to the tissues. Primary carnitine deficiency is a permanent condition that currently requires chronic therapy with L-carnitine. Secondary carnitine deficiency may be divided further into two other major categories, carnitine deficiency due to genetic defect of the metabolism and carnitine deficiency due to iatrogenic factors. Secondary carnitine deficiency is more common than primary carnitine deficiency and is mainly due to inborn errors of metabolism. Secondary carnitine deficiency may also result from medical conditions such as cirrhosis, Fanconi syndrome and HIV and treatments for diseases such as hemodialysis and e.g. valproic acid therapy.

Carnitine deficiency is an important disorder that deserves to be tracked and studied. No codes for disorders of carnitine metabolism exist in the ICD-9-CM. Sigma-Tau Pharmaceuticals, Inc. has requested that codes be created for these conditions. Additionally, since hemodialysis may be responsible for carnitine deficiency with an accompanying hypotension a code for hypotension of hemodialysis is also being requested.

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TABULAR MODIFICATIONS

277 Other and unspecified disorders of metabolism

Delete	277.8 Other specified disorders of metabolism <del>Hand-Schuller-Christian disease</del> <del>Histiocytosis (acute) (chronic)</del> <del>Histiocytosis X (chronic)</del> Excludes: <del>histiocytosis:</del> <del>acute differentiated progressive (202.5)</del> <del>X, acute (progressive) (202.5)</del>
New code	277.81 Primary carnitine deficiency
New code	277.82 Carnitine deficiency due to inborn errors of metabolism
New code	277.83 Iatrogenic carnitine deficiency Carnitine deficiency due to: Hemodialysis Valproic acid therapy
New code	277.84 Other secondary carnitine deficiency
New code	277.89 Other specified disorders of metabolism Hand-Schuller-Christian disease Histiocytosis (acute) (chronic) Histiocytosis X (chronic)  Excludes: histiocytosis: acute differentiated progressive (202.5) X, acute (progressive) (202.5)
	458 Hypotension
New code	458.3 Hypotension of hemodialysis Intra-dialytic hypotension

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Topic: Rhabdomyolysis

Rhabdomyolysis is an acute, sometimes fatal disease marked by destruction of skeletal muscle. Rarely, this may occur following strenuous exercise and in association with drugs that cause coma, such as alcohol, heroin, or cocaine. When associated with the crush syndrome, where muscle tissue disintegrates due to prolonged, continuous pressure, it is referred to as traumatic rhabdomyolysis. Due to severity of this condition, the Editorial Advisory Board of "Coding Clinic for ICD-9-CM" has requested that a unique code be created for rhabdomyolysis.

**TABULAR MODIFICATION**

728 Disorders of muscle, ligament, and fascia

728.8 Other disorders of muscle, ligament, and fascia

New code                      728.88 Rhabdomyolysis



## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Hypercoagulable states

The hypercoagulable states are a group of inherited and acquired disorders which cause an increased risk of thrombosis. The primary hypercoagulable states are for the most part inherited abnormalities of specific proteins, particularly the anticoagulant factors that normally break down blood clots. Some examples of primary hypercoagulable states include antithrombin III deficiency, protein C deficiency, protein S deficiency, activated protein C resistance (Factor V Leiden mutation), and prothrombin gene mutation. The secondary hypercoagulable states are a diverse group of mostly acquired disorders which predispose to thrombosis through complex and often multifactorial mechanisms. These mechanisms may broadly involve abnormalities of blood flow, of blood composition, and of vessel walls. Some disorders that can cause a secondary hypercoagulable state include malignancy, myeloproliferative disorders, antiphospholipid antibody syndrome, pregnancy, and trauma. Prognosis and treatment of a hypercoagulable state depends on the specific disorder involved. Anticoagulation therapy may be indicated short term, or for some cases, a lifetime.

### TABULAR MODIFICATION

	289 Other diseases of blood and blood-forming organs
	289.8 Other specified diseases of blood and blood-forming organs
Delete	<del>Hypergammaglobulinemia</del> <del>Myelofibrosis</del> <del>Pseudocholinesterase deficiency</del>
New code	289.81 Primary hypercoagulable state Activated protein C resistance Antithrombin III deficiency Lupus anticoagulant Protein C deficiency Protein S deficiency
	289.82 Secondary hypercoagulable state
New code	289.89 Other specified diseases of blood and blood-forming organs Hypergammaglobulinemia Myelofibrosis Pseudocholinesterase deficiency

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Topic: Hyperaldosteronism

The ICD-9-CM code 255.1, Hyperaldosteronism, includes several different states of aldosterone excess for which the diagnostic work up and treatment are distinct. Also important is the recent realization that the primary form of aldosteronism is responsible for the hypertension seen in 5-10% of patients with hypertension. These findings and changes in the terminology for states of excess aldosterone now merit the revision of the classification for such disorders.

The Endocrine Society requests that code 255.1, Hyperaldosteronism, be expanded at the 5<sup>th</sup> digit level to identify the different conditions currently included in 255.1

**TABULAR MODIFICATIONS**

	255 Disorders of adrenal glands
	255.1 Hyperaldosteronism
Delete	<del>Aldosteronism (primary) (secondary)</del> <del>Bartter's syndrome</del> <del>Conn's syndrome</del>
New code	255.10 Primary aldosteronism Aldosteronism NOS Hyperaldosteronism, unspecified
New code	255.11 Secondary aldosteronism
New code	255.12 Bartter's syndrome
New code	255.13 Glucocorticoid-remediable aldosteronism Familial aldosteronism type I
New code	255.14 Conn's syndrome

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Barrett's esophagus

Barrett's esophagus (BE) is a metaplastic disorder in which specialized columnar epithelium replaces healthy squamous epithelium. BE is an acquired condition, secondary to chronic gastroesophageal reflux damage to the esophageal mucosa. Its origin probably involves multipotential undifferentiated cells.

Currently, BE is indexed to code 530.2, Ulcer of esophagus. This code does not appropriately capture the inherent physiologic changes to the structure of the mucosal lining of the esophagus and lower esophageal junction. A unique code for BE is being requested by the American Gastroenterological Association (AGA).

### TABULAR MODIFICATION

530 Diseases of esophagus

530.8 Other specified disorders of esophagus

New code                      530.85 Barrett's esophagus

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Early satiety

The symptom of early satiety is common in many conditions, including serious conditions such as a brain tumor or hormonal problems. This symptom is not equivalent to anorexia, when a person chooses not to eat or who has no appetite. Early satiety occurs in persons who have hunger and are eating, but feel full quickly. A new code for this general symptom is being proposed.

### TABULAR MODIFICATION

780 General symptoms

780.9 Other general symptoms

New code                      780.94 Early satiety

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Topic: Encounter for emergency contraception

Codes exist for encounters for prescription for oral contraceptives and other forms of birth control but no code exists for the unique circumstances of an encounter for emergency contraception.

**TABULAR MODIFICATION**

V25 Encounter for contraceptive management

V25.0 General counseling and advice

New code                      V25.03 Encounter for emergency contraceptive counseling and prescription

**ICD-9-CM Coordination and Maintenance Committee Meeting**

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Topic: Thoracoscopic procedure converted to open procedure

A code exists for laparoscopic surgical procedure converted to open procedure, V64.4. A similar code for thoracoscopic procedure converted to open procedure has been requested.

**TABULAR MODIFICATION**

V64 Person encountering health services for specific procedures, not carried out

New code           V64.5 Thoracoscopic surgical procedure converted to open procedure

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Subaponeurotic hemorrhage (Subgaleal hemorrhage)

Caput succedaneum, cephalhematoma, chignon and subaponeurotic hemorrhage (Subgaleal hemorrhage-SGH) currently are all assigned to code 767.1, Injuries to scalp. The first three of these conditions are commonly encountered events after normal spontaneous vaginal delivery or uncomplicated forceps or vacuum extraction deliveries, and have no significant associated morbidity or mortality. SGH is a relatively rare event but one associated with high rates of morbidity and mortality. SGH has reported mortality rates of between 3% and 23%.

The frequency of this injury in the United States has increased as the percentage of vacuum extraction deliveries has increased. An increase in usage and injury is reflected in worldwide literature as well. Without a unique code for SGH there has been an inability to implement specific programs to decrease the incidence of this injury, as there is no accurate means to measure outcomes of any particular program. It is proposed that code 767.1 be expanded to create a unique code for SGH.

### TABULAR MODIFICATION

	767	Birth trauma
	767.1	Injuries to scalp
Delete		<del>Caput succedaneum</del>
		<del>Cephalhematoma</del>
		<del>Chignon (from vacuum extraction)</del>
		<del>Massive epicranial subaponeurotic hemorrhage</del>
New code	767.11	Epicranial subaponeurotic hemorrhage (massive)
Add		Subgaleal hemorrhage
New code	767.19	Other injuries to scalp
Add		Caput succedaneum
		Cephalhematoma
		Chignon (from vacuum extraction)

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Postpartum cardiomyopathy

Cardiomyopathy may occur in the postpartum period in women without pre-existing heart disease. It tends to affect multiparas, women over age 30, those carrying multiple fetuses and those whose pregnancy is complicated by pre-eclampsia. It has a mortality rate of 50% within 5 years and a high probability of recurrence in subsequent pregnancies, which are therefore contraindicated. Heart transplant is frequently required.

Postpartum cardiomyopathy is currently an inclusion under code 674.8, Other complications of the puerperium, not elsewhere classified. Due to the severity of the condition, it is being proposed that a unique code be created.

### TABULAR MODIFICATIONS

	674	Other and unspecified complications of the puerperium, not elsewhere classified
New code	674.5	Postpartum cardiomyopathy [2, 4]
Delete	674.8	Other Postpartum: <del>cardiomyopathy</del>



## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Late infant

When code 645, Late pregnancy, was expanded creating codes for post-term pregnancy, 645.1, those between 40 and 42 weeks, and prolonged pregnancy, 645.2, those over 42 weeks, a similar modification was not made to the corresponding newborn code, 766.2, Post-term infant, not "heavy for dates". Because of this, 766.2 does not correspond to either of the codes or definitions at 645. An expansion of code 766.2 is being proposed at the request of the Central Office on ICD-9-CM of the American Hospital Association to correct this discrepancy.

### TABULAR MODIFICATION

766 Disorders relating to long gestation and high birthweight

Revise	766.2	<del>Post-term infant, not "heavy for dates"</del> <u>Late infant, not "heavy-for-dates"</u> <del>Fetus or infant with gestation period of 294 days or more [42 or more completed weeks], not "heavy" or "large for dates"</del> <del>Postmaturity NOS</del>
New code Add	766.21	Post-term infant Infant with gestation period over 40 completed weeks to 42 completed weeks
New code Add Add	766.22	Prolonged gestation of infant Infant with gestation period over 42 completed weeks Postmaturity NOS

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Loss of consciousness in mild traumatic brain injury

In recent decades, public health and health care communities have become increasingly aware that the consequences of mild traumatic brain injury (MTBI) may not, in fact, be mild. Epidemiologic research has identified MTBI as a public health problem of large magnitude. Clinical research has provided evidence that these injuries can cause serious, lasting problems.

Responding to a mandate from Congress, the National Center for Injury Prevention and Control (NCIPC), one of the Centers for Disease Control and Prevention (CDC), Division of Injury and Disability Outcomes and Programs (DIDPO), and a group of experts have recommended methods to assess the incidence and prevalence of MTBI in the United States. For this purpose, a surveillance definition for MTBI was established. One of the many criteria for this definition is either observed or reported loss of consciousness (LOC) lasting 30 minutes or less.

The codes for concussion, which are the codes that are applicable for mild traumatic brain injury, indicate length of LOC. Currently, the shortest time frame for LOC for the concussion codes is less than one hour. This time frame does not conform to the surveillance definition established for MTBI of LOC lasting less than 30 minutes. It is being proposed that new codes be created to indicate LOC under 30 minutes and for between 31-59 minutes to conform to the surveillance definition while maintaining statistical integrity.

### TABULAR MODIFICATIONS

850 Concussion

850.1 With brief loss of consciousness

New code                      850.11 with loss of consciousness less than 30 minutes

New code                      850.12 with loss of consciousness from 31 to 59 minutes

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Cerebral infarct of unknown vessel

Often a patient with confirmed occlusion of his precerebral and cerebral arteries may have signs and symptoms of a cerebral infarction but it is not possible to determine the vessel responsible for the infarction. Because of this, use of the 5<sup>th</sup> digits for categories 433, Occlusion and stenosis of precerebral arteries, and 434, Occlusion of cerebral arteries, has been a problem. A 5<sup>th</sup> digit modifies the code to which it is assigned. 5<sup>th</sup> digit 1, with cerebral infarction, modifies codes that specify a vessel. If it can't be determined that the site of an occlusion is also the site responsible for the infarct then its is incorrect to use 5<sup>th</sup> digit 1. It is also a problem using 5<sup>th</sup> digit 0, without mention of infarction, since the patient clearly has had an infarct and it is documented as such. The use of code 436, Acute, but ill-defined, cerebrovascular disease, is always discouraged, and is illogical with a code that has a 5<sup>th</sup> digit stating without infarct.

A solution to this problem is being proposed. A new 5<sup>th</sup> digit for "with cerebral infarction of other or unspecified vessel" would be added to the 5<sup>th</sup> digit subclassifications for categories 433 and 434. The existing 5<sup>th</sup> digit 1 would be retitled to read "with cerebral infarction of this vessel. With this modification the use of a 5<sup>th</sup> digit indicating a cerebral infarction could still be used even when the site of the infarct is unknown.

### TABULAR MODIFICATIONS

#### 433 Occlusion and stenosis of precerebral arteries

The following fifth-digit subclassification is for use with category 433:

Revise	1 with cerebral infarction <u>of this vessel</u>
New	2 with cerebral infarction of other or unspecified vessel

#### 434 Occlusion of cerebral arteries

The following fifth-digit subclassification is for use with category 434:

Revise	1 with cerebral infarction <u>of this vessel</u>
New	2 with cerebral infarction of other or unspecified vessel

## ICD-9-CM Coordination and Maintenance Committee Meeting

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Topic: Hair tourniquet syndrome

Hair tourniquet syndrome, also referred to as hair strangulation, is the external constriction of a finger, toe, or the external genitalia by hair. The external constriction may also be due to thread, a ring, a rubber band, or any other object that can go around a body part. It is a relatively common finding in infants. It may be an emergency situation if the object is not removed and infection or amputation results. Hair is most commonly associated with toes and external genitalia while thread is more often found around fingers. There is an association with older frequently washed cloths and the wearing of mittens.

External constriction is classified as a superficial injury. However, there are no index entries to identify this type of injury and there is no external cause code to specify the object causing the constriction. Index entries for this syndrome were presented at the April 19, 2002 C & M meeting. External cause codes are now being proposed to allow for more specific classification of this syndrome.

### TABULAR MODIFICATION

E928 Other and unspecified environmental and accidental causes

New code                    E928.4 External constriction caused by hair

New code                    E928.5 External constriction caused by other object

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**Addenda  
TABULAR**

- 250 Diabetes mellitus
- 250.5 Diabetes with ophthalmic manifestations
- Use additional code to identify manifestations, as:  
diabetic  
retinal edema (362.10)
- Revise
- 278 Obesity and other hyperalimentation
- 278.0 Obesity
- 278.01 Morbid obesity  
Severe obesity
- Add
- 285 Other and unspecified anemia
- 285.2 Anemia in chronic illness
- 285.21 Anemia in end-stage renal disease  
Erythropoietin-resistant anemia (EPO resistant anemia)
- Add
- 491 Chronic bronchitis
- 491.2 Obstructive chronic bronchitis
- 491.21 With acute exacerbation  
~~Acute bronchitis with chronic obstructive pulmonary  
disease [COPD]~~
- Delete
- 593 Other disorders of kidney and ureter
- 593.9 Unspecified disorder of kidney and ureter  
Renal disease (chronic) NOS  
Renal insufficiency (acute) (chronic)
- Revise  
Add

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- 614 Inflammatory disease of ovary, fallopian tube, pelvic cellular tissue, and peritoneum
- 614.6 Pelvic peritoneal adhesions, female (postoperative) (postinfection)
- Add Excludes: pelvic peritoneal adhesions complicating pregnancy or labor (648.9)
- 625 Pain and other symptoms associated with female genital organs
- Add 625.4 Premenstrual tension syndrome  
Premenstrual dysphoric disorder
- 710 Diffuse diseases of connective tissue
- 710.1 Systemic sclerosis
- Revise Use additional code to identify...  
lung involvement (517.2)
- 785 Symptoms involving cardiovascular system
- 785.4 Gangrene
- Delete ~~Code first any associated underlying condition~~
- Add Code first the following conditions only:
- Add diabetes mellitus with peripheral circulatory disorders (250.7)
- Add Raynaud's syndrome (443.0)
- 965 Poisoning by analgesics, antipyretics, and antirheumatics
- 965.1 Salicylates
- Add Excludes: salicylates used for (as):  
antiplatelet (964.8)  
antithrombotic (964.8)

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## Addenda INDEX

- Aftercare...
  - fracture...
    - healing...
      - pathologic...
        - Add ankle V54.29
        - Add finger V54.29
        - Add foot V54.29
        - Add hand V54.29
        - Add toe(s) V54.29
        - Add wrist V54.29
      - traumatic
        - Add ankle V54.19
        - Add finger V54.19
        - Add foot V54.19
        - Add hand V54.29
        - Add toe(s) V54.19
        - Add wrist V54.19
- Add Allergy
  - milk protein 558.3
- Add Angina
  - accelerated 411.1
- Delete Bedsore 707.0
  - ~~with gangrene 707.0 [785.4]~~
- Revise Blue
  - toe syndrome –see Atherosclerosis 445.02
- Revise Burnett's syndrome (milk-alkali) 275.42
- Add Complication
  - mechanical
    - esophagostomy 996.59

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- Delete      ~~Decubital gangrene 707.0 [785.4]~~
- Delete      Decubitus (ulcer) 707.7  
~~with gangrene 707.7 [785.4]~~
- Revise      Diabetes...(poorly controlled) ...
- Revise      Disease  
                renal  
                chronic ~~–See Nephritis, chronic~~ 593.9
- Add          Findings, abnormal, without diagnosis...  
                stress test 794.39
- Revise      Fitz-Hugh and Curtis syndrome (~~gonococcal peritonitis~~) 098.86
- Add          due to:
- Add          Chlamydia trachomatis 099.56
- Add          Neisseria gonorrhoeae (gonococcal peritonitis ) 098.86
- Revise      Fracture  
                march...  
                fibula 733.93
- Revise      tibia 733.93
- Delete      Gangrene  
                abdominal wall  
                ~~arteriosclerosis 440.29 [785.4]~~
- Revise      Hypertension  
                with  
                heart involvement (conditions classifiable to ~~425.8...~~)...
- Revise      Hypertrophy  
                spinal ligament 724.8
- Revise      Keratoacanthoma (M8071/1) 238.2 -see Neoplasm, skin, uncertain behavior



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Laceration...  
Revise anus (sphincter) 879.6  
Add complicated 879.7  
uterus  
Revise obstetrical trauma NEC 665.5

Newborn  
Add affected by maternal abuse of drugs (gestational) (via placenta) (via breast milk)  
760.70

Obesity  
Add severe 278.01

Pregnancy...  
complicated by...  
Add adhesion, pelvic peritoneal 648.9  
Revise infection (conditions classifiable to 614.0-614.5, 614.7-614.9, 615)  
Revise inflammation (conditions classifiable to 614.0-614.5, 614.7-614.9, 615)  
Revise pelvic inflammatory disease (conditions classifiable to 614.0-614.5, 614.7-614.9, 615)  
Add pelvic peritoneal adhesion 648.9

Sore  
pressure  
Delete ~~with gangrene 707.0 [785.4]~~

Syndrome...  
Add Angelman 759.89  
blue  
Revise toe –see ~~Atherosclerosis~~ 445.02  
Revise Burnett's (milk-alkali) 275.42  
Revise Fitz-Hugh and Curtis (~~gonococcal peritonitis~~) 098.86  
Add due to:  
Add Chlamydia trachomatis 099.56  
Add Neisseria gonorrhoea (gonococcal peritonitis ) 098.86  
Add Mal de Debarquement 780.4  
Revise milk-alkali (milk drinkers') 275.42  
Add Smith-Magenis 758.3

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	Thrombosis...
Add	vein 453.8
Add	deep 453.8
	Ulcer...
Delete	<del>with gangrene 707.9 [785.4]</del>
	decubitus...
Delete	<del>with gangrene 707.0 [785.4]</del>
	lower extremity...
Delete	<del>with gangrene...707.10 [785.4]</del>
	decubitus...
Delete	<del>with gangrene 707.0 [785.4]</del>
	skin...
Delete	<del>with gangrene 707.9 [785.4]</del>
	decubitus...
Delete	<del>with gangrene 707.0 [785.4]</del>
	lower extremity...
Delete	<del>with gangrene 707.10 [785.4]</del>
Delete	arteriosclerotic 440.24
	Vasculopathy
Add	cardiac allograft 996.83

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**ADDENDA**

**Index to External Causes**

Cold  
self-inflicted...  
Revise suicidal E958.3 (~~herded~~)

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**Table of Drugs and Chemicals**

Aspirin	<del>965.1</del>	<del>E850.3</del>	<del>E935.3</del>	<del>E950.0</del>	<del>E962.0</del>	<del>E980.0</del>
for (as)						
analgesia	965.1	E850.3	E935.3	E950.0	E962.0	E980.0
antiplatelet	964.8	E858.2	E934.8	E950.4	E962.0	E980.4
antipyretic	965.1	E850.3	E935.3	E950.0	E962.0	E980.0
antirheumatics	965.1	E850.3	E935.3	E950.0	E962.0	E980.0
antithrombotic	964.8	E858.2	E934.8	E950.4	E962.0	E980.4
thrombocytopenia (essential)	964.8	E858.2	E934.8	E950.4	E962.0	E980.4
Add Botox	975.3	E858.6	E945.3	E950.4	E962.0	E980.4