## MEDICAID POLICY INFORMATION SHEET

#### Policy Analyst: Phillip Kurdunowicz

#### Phone Number: 517-284-1199

Initial 🗌	Public Comment 🖂	Final 🖂	

### Brief description of policy:

The Michigan Department of Health and Human Services (MDHHS) issued bulletin MSA 19-35 to enact new work requirements for Medicaid beneficiaries who have Healthy Michigan Plan (HMP) health care coverage. This new bulletin rescinds the work requirements but maintains some of the other components of the original bulletin. The new bulletin also contains updated information on the exemption of medically frail beneficiaries from cost sharing obligations.

## Reason for policy (problem being addressed):

MDHHS is issuing this policy in response to the court order from the United States District Court for the District of Columbia in Young et al v. Azar et al. The court vacated the approval of the work requirements as part of the Special Terms and Conditions of the Section 1115 Demonstration Waiver Amendment. MDHHS is stopping the implementation of work requirements in response to the court order.

#### Budget implication:

budget neutral
will cost MDHHS \$29.2 million gross, \$14.6 million General Fund, and is budgeted in current appropriation
will save MDHHS \$

#### Is this policy change mandated per federal requirements?

No.

## Does policy have operational implications on other parts of MDHHS?

Yes, the Economic Stability Administration (ESA) will also be impacted by this change. ESA field staff and Central Office staff have assumed new responsibilities related to administering work requirements, and ESA will need to provide updated information to staff and make other organizational adjustments in response to this change.

#### Does policy have operational implications on other departments?

The Department of Treasury currently assists with MDHHS with pursuing tax off-sets for beneficiaries who have not paid required MI Health Account co-pays and fees. MDHHS does not anticipate that this new bulletin will impact this process because the bulletin does not change current policy and operational processes for cost-sharing for HMP beneficiaries.

MDHHS also partnered with the Department of Labor and Economic Opportunity (LEO) and local Michigan Works! agencies to support beneficiaries with the reporting process and offer job search and other employment supports to beneficiaries. LEO and the local Michigan Works!

## MEDICAID POLICY INFORMATION SHEET

agencies will need to provide updated information to staff and make other organizational adjustments in response to this change.

# Summary of input:

controversial Several stakeholders have expressed support for work requirements and are not supportive of stopping the implemention process.

acceptable to most/all groups

limited public interest/comment

# Supporting Documentation:

State Plan Amendment Required: Yes 🛛 No		Public Notice Required:	Yes	🛛 No		
If Yes, please p	rovide status:					
Approved	Pending	🗌 De	nied	lf yes,		
Date:	Approval	Date:		Submission Date:		

DRAFT FOR PUBLIC COMMENT		
Michigan Department of		
Health and Human Services	Project Number: 2020-HMP	Date: April 28, 2020
Comments Due: Jun	e 2, 2020	
Proposed Effective Date: Mai	rch 4, 2020	
Direct Comments To: Phi	l Kurdunowicz	
Address:		
E-Mail Address: Kur	<u>dunowiczp@michigan.gov</u>	
<b>Phone:</b> 517	2-284-1199	Fax:

Policy Subject: Rescinding Work Requirements for the Healthy Michigan Plan

Affected Programs: Healthy Michigan Plan

**Distribution:** All Providers, Bridges Eligibility Manual (BEM) and Bridges Administrative Manual (BAM) Holders

**Summary:** The Michigan Department of Health and Human Services (MDHHS) issued bulletin MSA 19-35 to enact new work requirements for Medicaid beneficiaries who have Healthy Michigan Plan (HMP) health care coverage. This new bulletin rescinds the work requirements but maintains some of the other components of the original bulletin. The new bulletin also contains updated information on the exemption of medically frail beneficiaries from cost sharing obligations.

**Purpose:** MDHHS is issuing this policy in response to the court order from the United States District Court for the District of Columbia in Young et al v. Azar et al. The court vacated the approval of the work requirements as part of the Special Terms and Conditions of the Section 1115 Demonstration Waiver Amendment. MDHHS is stopping the implementation of work requirements in response to the court order.

**Cost Implications:** \$29.2 million gross, \$14.6 million General Fund

Potential Hearings & Appeal Issues: Yes.

State Plan Amendment Required: Yes 🗌 No 🖂 If yes, date submitted:		o 🖂	Public Notice Required: Yes 🗌 Submitted date:	No 🖂	
Triba	Tribal Notification: Yes 🗌 No 🖂 - Date:				
THIS	SECTION COMPLETED BY RECEIVER				
	Approved		No Comments		
			See Comments Below		
	Disapproved		See Comments in Text		
Signa	ature:		Phone Number		

Signature Printed:		
Bureau/Administration (please print)	Date	
Comment001		Revised 6/16





Bulletin Number:	MSA 20-10
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- **Distribution:** All Providers, Bridges Eligibility Manual (BEM) and Bridges Administrative Manual (BAM) Holders
  - **Issued:** April 28, 2020
  - Subject: Rescinding Work Requirements for the Healthy Michigan Plan
  - Effective: March 4, 2020

# Programs Affected: Healthy Michigan Plan

The purpose of this policy is to rescind the work requirements described in Bulletin MSA 19-35, issued December 2, 2019. Bulletin MSA 19-35 describes work requirements that were implemented for Medicaid beneficiaries who have Healthy Michigan Plan (HMP) health care coverage. Pursuant to the court order from the United States District Court for the District of Columbia in Young et al. v. Azar et al., the Michigan Department of Health and Human Services (MDHHS) is stopping the implementation of work requirements.

HMP changes described in Section I (subsections A-C) and Section IV of Bulletin MSA 19-35 are rescinded. HMP changes described in Section II (subsections A-B) and Section III of Bulletin MSA 19-35 remain in effect. HMP changes described in the following attachments to Bulletin MSA 19-35 also remain in effect:

- Attachment I: Policy and Operational Process Document: Healthy Behavior Requirements
- Attachment II: Policy and Operational Process Document: MI Health Account Co-Pays and Fees for HMP Beneficiaries

MDHHS is also amending Attachment III: "Policy and Operational Process Document: Identification of Medically Frail Beneficiaries" of Bulletin MSA 19-35 in accordance with the court order. Because MDHHS is stopping the implementation of work requirements, MDHHS will no longer use the medically frail identification process to exempt beneficiaries from the work requirements. However, MDHHS will still use this process to exempt beneficiaries who are medically frail from cost-sharing for the HMP program. MDHHS will also still allow beneficiaries to self-report that they are medically frail through several methods, which are described in the "Policy and Operational Process Document: Identification of Medically Frail Beneficiaries."

MDHHS is implementing this bulletin effective March 4, 2020, the date of the court order from the United States District Court for the District of Columbia.

## **Public Comment**

The public comment portion of the policy promulgation process is being conducted concurrently with the implementation of the changes noted in this bulletin. Any interested party wishing to comment on the changes may do so by submitting comments to <u>KurdunowiczP@michigan.gov</u>. Please include "Rescinding Work Requirements for the Healthy Michigan Plan" in the subject line.

Comments received will be considered for revisions to the changes implemented by this bulletin.

## **Manual Maintenance**

Providers should retain this bulletin until applicable information has been incorporated into the Bridges Administration Manual (<u>http://www.mfia.state.mi.us/olmweb/ex/html/</u>) and the Bridges Eligibility Manual (<u>http://www.mfia.state.mi.us/olmweb/ex/html/</u>).

# Questions

Any questions regarding this bulletin should be directed to Provider Inquiry, Department of Health and Human Services, P.O. Box 30731, Lansing, Michigan 48909-8231, or e-mailed to <u>ProviderSupport@michigan.gov</u>. When you submit an e-mail, be sure to include your name, affiliation, NPI number, and phone number so you may be contacted if necessary. Providers may phone toll-free 1-800-292-2550.

Approved

Kate Massey, Director Medical Services Administration

# Policy and Operational Process Document: Identification of Medically Frail Beneficiaries

# BACKGROUND

Healthy Michigan Plan (HMP) beneficiaries who are considered medically frail in accordance with 42 CFR 440.315(f) are exempt from cost sharing obligations for the HMP program. Beneficiaries who are medically frail may have any of the following:

- A physical, mental, or emotional health condition that limits a daily activity (like bathing, dressing, daily chores, etc.)
- A physical, intellectual, or developmental disability that impairs the ability to perform one or more activities of daily living
- A physical, mental, or emotional health condition that requires frequent monitoring
- A disability determination based on Social Security criteria (SSDI)
- A chronic substance use disorder
- A serious and complex medical conditions or special medical needs
- Is in a nursing home, hospice, or is receiving home help services
- Is homeless
- Is a survivor of domestic violence

Beneficiaries identified as medically frail will remain exempt for 12 months from the reported date or until their next health care coverage redetermination date, whichever comes first. Based upon the results of the redetermination, the exemptions may either be extended or will end. Medically frail beneficiaries will be identified by the following methods:

# Self-Identification

Individuals may report and self-attest to their medically frail status through any of the medical assistance program applications: Application for Health Coverage & Help Paying Costs (DCH-1426), the all programs Assistance Application (MDHHS-1171), and the online MI Bridges Application. A telephone option for application is also available.

Application processes include questions that allow individuals to report and attest that they have physical, mental, or emotional health conditions that limit their daily activities, such as bathing, dressing, or daily chores; or limit their ability to work, attend school, or take care of daily needs. Additionally, questions will allow the individual to report they reside in a medical or nursing facility.

If beneficiaries become medically frail during a period of eligibility, they may update their application information. In addition to the paper and online applications, beneficiaries may also call the Beneficiary Help Line (telephone option) to report and attest to the medically frail exemption at any time. Beneficiaries should also notify their specialist within 10 days of any changes in their circumstances which may affect their eligibility for assistance.

# **Retrospective Claims Analysis**

When available, MDHHS will review health care claims data available within the Community Health Automated Medicaid Processing System (CHAMPS) from the preceding 12 months for the presence of select diagnosis codes to identify beneficiaries considered medically frail. The list of codes is included as Appendix A. MDHHS may pursue updates to this list on an annual basis. The claims data to be reviewed include the following:

- a. ICD-10 diagnosis codes (over 350 codes selected) that identify:
  - Individuals with disabling mental disorders;
  - Individuals with serious and complex medical conditions; and
  - Individuals with a physical, intellectual or developmental disability that significantly impairs their ability to perform one or more activities of daily living.
- b. Whether a beneficiary is in a nursing home or hospice or is receiving services through the Adult Home Help program or Children's Special Health Care Services program.
- c. Whether a beneficiary is homeless.
- d. Whether a beneficiary is a survivor of domestic violence.

#### CODE DESCRIPTION A170 **TUBERCULOUS MENINGITIS** A171 MENINGEAL TUBERCULOMA A1781 TUBERCULOMA OF BRAIN AND SPINAL CORD A1782 TUBERCULOUS MENINGOENCEPHALITIS A1783 **TUBERCULOUS NEURITIS** A1789 OTHER TUBERCULOSIS OF NERVOUS SYSTEM A179 TUBERCULOSIS OF NERVOUS SYSTEM UNSPECIFIED A1801 TUBERCULOSIS OF SPINE A1802 TUBERCULOUS ARTHRITIS OF OTHER JOINTS A1803 TUBERCULOSIS OF OTHER BONES A1809 OTHER MUSCULOSKELETAL TUBERCULOSIS A1810 TUBERCULOSIS OF GENITOURINARY SYSTEM UNSPECIFIED A1811 TUBERCULOSIS OF KIDNEY AND URETER A1812 TUBERCULOSIS OF BLADDER A1813 TUBERCULOSIS OF OTHER URINARY ORGANS A1814 TUBERCULOSIS OF PROSTATE A1815 TUBERCULOSIS OF OTHER MALE GENITAL ORGANS A1816 TUBERCULOSIS OF CERVIX A1817 TUBERCULOUS FEMALE PELVIC INFLAMMATORY DISEASE A1818 TUBERCULOSIS OF OTHER FEMALE GENITAL ORGANS A182 TUBERCULOUS PERIPHERAL LYMPHADENOPATHY A1831 **TUBERCULOUS PERITONITIS** A1832 TUBERCULOUS ENTERITIS A1839 **RETROPERITONEAL TUBERCULOSIS** A184 TUBERCULOSIS OF SKIN AND SUBCUTANEOUS TISSUE A1850 TUBERCULOSIS OF EYE UNSPECIFIED

# Appendix A: Medically Frail Diagnosis Codes

CODE	DESCRIPTION
A1851	TUBERCULOUS EPISCLERITIS
A1852	TUBERCULOUS KERATITIS
A1853	TUBERCULOUS CHORIORETINITIS
A1854	TUBERCULOUS IRIDOCYCLITIS
A1859	OTHER TUBERCULOSIS OF EYE
A186	TUBERCULOSIS OF INNER MIDDLE EAR
A187	TUBERCULOSIS OF ADRENAL GLANDS
A1881	TUBERCULOSIS OF THYROID GLAND
A1882	TUBERCULOSIS OF OTHER ENDOCRINE GLANDS
A1883	TUBERCULOSIS OF DIGESTIVE TRACT ORGANS NEC
A1884	TUBERCULOSIS OF HEART
A1885	TUBERCULOSIS OF SPLEEN
A1889	TUBERCULOSIS OF OTHER SITES
B20	HUMAN IMMUNODEFICIENCY VIRUS HIV DISEASE
B900	SEQUELAE OF CENTRAL NERVOUS SYSTEM TUBERCULOSIS
B901	SEQUELAE OF GENITOURINARY TUBERCULOSIS
B902	SEQUELAE OF TUBERCULOSIS OF BONES AND JOINTS
B908	SEQUELAE OF TUBERCULOSIS OF OTHER ORGANS
D5700	HB-SS DISEASE WITH CRISIS, UNSPECIFIED
D5701	HB-SS DISEASE WITH ACUTE CHEST SYNDROME
D5702	HB-SS DISEASE WITH SPLENIC SEQUESTRATION
D571	SICKLE-CELL DISEASE WITHOUT CRISIS
D5720	SICKLE-CELL/HB-C DISEASE WITHOUT CRISIS
D57211	SICKLE-CELL/HB-C DISEASE WITH ACUTE CHEST SYNDROME
D57212	SICKLE-CELL/HB-C DISEASE WITH SPLENIC SEQUESTRATION
D57219	SICKLE-CELL/HB-C DISEASE WITH CRISIS, UNSPECIFIED
D5740	SICKLE-CELL THALASSEMIA WITHOUT CRISIS
D57411	SICKLE-CELL THALASSEMIA WITH ACUTE CHEST SYNDROME
D57412	SICKLE-CELL THALASSEMIA WITH SPLENIC SEQUESTRATION
D57419	SICKLE-CELL THALASSEMIA WITH CRISIS, UNSPECIFIED
D5780	OTHER SICKLE-CELL DISORDERS WITHOUT CRISIS
D57811	OTHER SICKLE-CELL DISORDERS WITH ACUTE CHEST SYNDROME
D57812	OTHER SICKLE-CELL DISORDERS WITH SPLENIC SEQUESTRATION
D57819	OTHER SICKLE-CELL DISORDERS WITH CRISIS, UNSPECIFIED
D808	OTHER IMMUNODEF W/PREDOMINANTLY ANTIBODY DEFECTS
D809	IMMUNODEF W/PREDOMINANTLY ANTIBODY DEFECTS UNS
D810	SEVERE COMBINED IMMUNODEF W/RETICULAR DYSGENESIS
D811	SEVERE COMBINED IMMUNODEF LOW T & B-CELL NUMBERS
D812	SEVERE COMBINED IMMUNODEF W/NORMAL B-CELL NUMBRS
D8130	ADENOSINE DEAMINASE DEFICIENCY, UNSPECIFIED
D8131	SEVERE COMBINED IMMUNODEF DUE TO ADENOSINE DEAMINASE DEFIC
D8132	ADENOSINE DEAMINASE 2 DEFICIENCY
D8139	OTHER ADENOSINE DEAMINASE DEFICIENCY
D814	NEZELOFS SYNDROME

CODE	DESCRIPTION
D815	PURINE NUCLEOSIDE PHOSPHORYLASE DEFICIENCY
D816	MAJ HISTOCOMPATIBILITY COMPLX CLASS I DEFICIENCY
D817	MAJ HISTOCOMPATIBILTY COMPLX CLASS II DEFICIENCY
D81810	BIOTINIDASE DEFICIENCY
D81818	OTHER BIOTIN-DEPENDENT CARBOXYLASE DEFICIENCY
D81819	BIOTIN-DEPENDENT CARBOXYLASE DEFICIENCY UNS
D8189	OTHER COMBINED IMMUNODEFICIENCIES
D819	COMBINED IMMUNODEFICIENCY UNSPECIFIED
D820	WISKOTT-ALDRICH SYNDROME
D821	DI GEORGES SYNDROME
D823	IMMUNODEFIC FLW HEREDITARY DEFECT RESPONS TO EBV
D828	IMMUNODEFIC ASSOCIATED W/OTH SPEC MAJOR DEFECT
D829	IMMUNODEFICIENCY ASSOCIATED W/MAJOR DEFECTS UNS
D830	CVI W/PREDOMINANT ABN OF B-CELL NUMBERS & FUNCT
D831	CVI W/PREDOMINANT IMMUNOREGULATORY T-CELL D/O
D832	CVI WITH AUTOANTIBODIES TO B- OR T-CELLS
E701	OTHER HYPERPHENYLALANINEMIAS
E7502	TAY-SACHS DISEASE
E7521	FABRY-ANDERSON DISEASE
E7522	GAUCHER DISEASE
E7523	KRABBE DISEASE
E75240	NIEMANN-PICK DISEASE TYPE A
E75241	NIEMANN-PICK DISEASE TYPE B
E75242	NIEMANN-PICK DISEASE TYPE C
E75243	NIEMANN-PICK DISEASE TYPE D
E75248	OTHER NIEMANN-PICK DISEASE
E75249	NIEMANN-PICK DISEASE UNSPECIFIED
E7525	METACHROMATIC LEUKODYSTROPHY
E7529	OTHER SPHINGOLIPIDOSIS
E840	CYSTIC FIBROSIS WITH PULMONARY MANIFESTATIONS
E8419	CYSTIC FIBROSIS W/OTH INTESTINAL MANIFESTATIONS
E848	CYSTIC FIBROSIS WITH OTHER MANIFESTATIONS
E849	CYSTIC FIBROSIS UNSPECIFIED
E8840	MITOCHONDRIAL METABOLISM DISORDER UNSPECIFIED
F0150	VASCULAR DEMENTIA WITHOUT BEHAVIORAL DISTURBANCE
F0151	VASCULAR DEMENTIA WITH BEHAVIORAL DISTURBANCE
F0280	DEMENTIA OTH DZ CLASS ELSW W/O BEHAVRL DISTURB
F0281	DEMENTIA OTH DISEAS CLASS W/BEHAVIORAL DISTURB
F0390	UNSPEC DEMENTIA WITHOUT BEHAVIORAL DISTURBANCE
F0391	UNSPECIFIED DEMENTIA WITH BEHAVIORAL DISTURBANCE
F04	AMNESTIC DISORDER DUE KNOWN PHYSIOLOGICAL COND
F060	PSYCHOTIC DISORDER W HALLUCIN DUE TO KNOWN PHYSIOL CONDITION
F061	CATATONIC DISORDER DUE TO KNOWN PHYSIOLOGICAL CONDITION
F062	PSYCHOTIC DISORDER W DELUSIONS DUE TO KNOWN PHYSIOL COND

CODE	DESCRIPTION
F0631	MOOD DISORDER DUE TO KNOWN PHYSIOL COND W DEPRESSV FEATURES
F0632	MOOD DISORD D/T PHYSIOL COND W MAJOR DEPRESSIVE-LIKE EPSD
F0633	MOOD DISORDER DUE TO KNOWN PHYSIOL COND W MANIC FEATURES
F0634	MOOD DISORDER DUE TO KNOWN PHYSIOL COND W MIXED FEATURES
F064	ANXIETY DISORDER DUE TO KNOWN PHYSIOLOGICAL CONDITION
F200	PARANOID SCHIZOPHRENIA
F201	DISORGANIZED SCHIZOPHRENIA
F202	CATATONIC SCHIZOPHRENIA
F203	UNDIFFERENTIATED SCHIZOPHRENIA
F205	RESIDUAL SCHIZOPHRENIA
F2081	SCHIZOPHRENIFORM DISORDER
F2089	OTHER SCHIZOPHRENIA
F209	SCHIZOPHRENIA UNSPECIFIED
F21	SCHIZOTYPAL DISORDER
F22	DELUSIONAL DISORDERS
F23	BRIEF PSYCHOTIC DISORDER
F24	SHARED PSYCHOTIC DISORDER
F250	SCHIZOAFFECTIVE DISORDER BIPOLAR TYPE
F251	SCHIZOAFFECTIVE DISORDER DEPRESSIVE TYPE
F258	OTHER SCHIZOAFFECTIVE DISORDERS
F259	SCHIZOAFFECTIVE DISORDER UNSPECIFIED
F28	OTH PSYCHOT D/O NOT DUE SUBSTANCE/PHYSIOLOG COND
F29	UNS PSYCHOSIS NOT DUE SUBSTANCE/PHYSIOLOG COND
F3012	MANIC EPISODE WITHOUT PSYCHOTIC SYMPTOMS, MODERATE
F3013	MANIC EPISODE, SEVERE, WITHOUT PSYCHOTIC SYMPTOMS
F302	MANIC EPISODE, SEVERE WITH PSYCHOTIC SYMPTOMS
F3112	BIPOLAR DISORD, CRNT EPISODE MANIC W/O PSYCH FEATURES, MOD
F3113	BIPOLAR DISORD, CRNT EPSD MANIC W/O PSYCH FEATURES, SEVERE
F312	BIPOLAR DISORD, CRNT EPISODE MANIC SEVERE W PSYCH FEATURES
F3132	BIPOLAR DISORDER, CURRENT EPISODE DEPRESSED, MODERATE
F314	BIPOLAR DISORD, CRNT EPSD DEPRESS, SEV, W/O PSYCH FEATURES
F315	BIPOLAR DISORD, CRNT EPSD DEPRESS, SEVERE, W PSYCH FEATURES
F3162	BIPOLAR DISORDER, CURRENT EPISODE MIXED, MODERATE
F3163	BIPOLAR DISORD, CRNT EPSD MIXED, SEVERE, W/O PSYCH FEATURES
F3164	BIPOLAR DISORD, CRNT EPISODE MIXED, SEVERE, W PSYCH FEATURES
F321	MAJOR DEPRESSIVE DISORDER, SINGLE EPISODE, MODERATE
F322	MAJOR DEPRESSV DISORD, SINGLE EPSD, SEV W/O PSYCH FEATURES
F323	MAJOR DEPRESSV DISORD, SINGLE EPSD, SEV W/O PSTCHTEATORES
F331	MAJOR DEPRESSIVE DISORDER, RECURRENT, MODERATE
F332	MAJOR DEPRESSVE DISORDER, RECORRENT, MODERATE MAJOR DEPRESSV DISORDER, RECURRENT SEVERE W/O PSYCH FEATURES
F333	MAJOR DEPRESSV DISORDER, RECORRENT SEVERE W/O PSYCH FEATORES MAJOR DEPRESSV DISORDER, RECURRENT, SEVERE W PSYCH SYMPTOMS
F333 F4001	AGORAPHOBIA WITH PANIC DISORDER
F4001 F410	
	PANIC DISORDER [EPISODIC PAROXYSMAL ANXIETY]
F4311	POST-TRAUMATIC STRESS DISORDER, ACUTE

CODE	DESCRIPTION
F4312	POST-TRAUMATIC STRESS DISORDER, CHRONIC
F440	DISSOCIATIVE AMNESIA
F441	DISSOCIATIVE FUGUE
F444	CONVERSION DISORDER WITH MOTOR SYMPTOM OR DEFICIT
F445	CONVERSION DISORDER WITH SEIZURES OR CONVULSIONS
F446	CONVERSION DISORDER WITH SENSORY SYMPTOM OR DEFICIT
F447	CONVERSION DISORDER WITH MIXED SYMPTOM PRESENTATION
F4481	DISSOCIATIVE IDENTITY DISORDER
F4522	BODY DYSMORPHIC DISORDER
F481	DEPERSONALIZATION-DEREALIZATION SYNDROME
F600	PARANOID PERSONALITY DISORDER
F601	SCHIZOID PERSONALITY DISORDER
F71	MODERATE INTELLECTUAL DISABILITIES
F72	SEVERE INTELLECTUAL DISABILITIES
F73	PROFOUND INTELLECTUAL DISABILITIES
F801	EXPRESSIVE LANGUAGE DISORDER
F840	AUTISTIC DISORDER
F845	ASPERGERS SYNDROME
F848	OTHER PERVASIVE DEVELOPMENTAL DISORDERS
F849	PERVASIVE DEVELOPMENTAL DISORDER UNSPECIFIED
F952	TOURETTES DISORDER
G041	TROPICAL SPASTIC PARAPLEGIA
G114	HEREDITARY SPASTIC PARAPLEGIA
G1221	AMYOTROPHIC LATERAL SCLEROSIS
G130	PARANEOPLASTIC NEUROMYOPATHY AND NEUROPATHY
G131	OTH SYSTEM ATROPHY PRIM AFFECT CNS NEOPLASTIC DZ
G231	PROGRESSIVE SUPRANUCLEAR OPHTHALMOPLEGIA
G300	ALZHEIMERS DISEASE WITH EARLY ONSET
G301	ALZHEIMERS DISEASE WITH LATE ONSET
G308	OTHER ALZHEIMERS DISEASE
G309	ALZHEIMERS DISEASE UNSPECIFIED
G3109	OTHER FRONTOTEMPORAL DEMENTIA
G3183	DEMENTIA WITH LEWY BODIES
G35	MULTIPLE SCLEROSIS
G450	VERTEBRO-BASILAR ARTERY SYNDROME
G451	CAROTID ARTERY SYNDROME HEMISPHERIC
G452	MULTIPLE & BILATERAL PRECEREBRAL ARTERY SYND
G453	AMAUROSIS FUGAX
G454	TRANSIENT GLOBAL AMNESIA
G458	OTH TRANSIENT CERBRL ISCHEMIC ATTACKS & REL SYND
G459	TRANSIENT CEREBRAL ISCHEMIC ATTACK UNSPECIFIED
G460	MIDDLE CEREBRAL ARTERY SYNDROME
G461	ANTERIOR CEREBRAL ARTERY SYNDROME
G462	POSTERIOR CEREBRAL ARTERY SYNDROME

CODE	DESCRIPTION
G7100	MUSCULAR DYSTROPHY, UNSPECIFIED
G7101	DUCHENNE OR BECKER MUSCULAR DYSTROPHY
G7102	FACIOSCAPULOHUMERAL MUSCULAR DYSTROPHY
G7109	OTHER SPECIFIED MUSCULAR DYSTROPHIES
G731	LAMBERT-EATON SYNDROME IN NEOPLASTIC DISEASE
G800	SPASTIC QUADRIPLEGIC CEREBRAL PALSY
G801	SPASTIC DIPLEGIC CEREBRAL PALSY
G802	SPASTIC HEMIPLEGIC CEREBRAL PALSY
G803	ATHETOID CEREBRAL PALSY
G804	ATAXIC CEREBRAL PALSY
G808	OTHER CEREBRAL PALSY
G809	CEREBRAL PALSY UNSPECIFIED
G8100	FLACCID HEMIPLEGIA AFFECTING UNSPECIFIED SIDE
G8101	FLACCID HEMIPLEGIA AFFECTING RIGHT DOMINANT SIDE
G8102	FLACCID HEMIPLEGIA AFFECTING LEFT DOMINANT SIDE
G8103	FLACCID HEMIPLEGIA AFFECTING RT NONDOMINANT SIDE
G8104	FLACCID HEMIPLEGIA AFFECTING LT NONDOMINANT SIDE
G8110	SPASTIC HEMIPLEGIA AFFECTING UNSPECIFIED SIDE
G8111	SPASTIC HEMIPLEGIA AFFECTING RIGHT DOMINANT SIDE
G8112	SPASTIC HEMIPLEGIA AFFECTING LEFT DOMINANT SIDE
G8113	SPASTIC HEMIPLEGIA AFFECTING RT NONDOMINANT SIDE
G8114	SPASTIC HEMIPLEGIA AFFECTING LT NONDOMINANT SIDE
G8190	HEMIPLEGIA UNS AFFECTING UNSPECIFIED SIDE
G8191	HEMIPLEGIA UNS AFFECTING RIGHT DOMINANT SIDE
G8192	HEMIPLEGIA UNS AFFECTING LEFT DOMINANT SIDE
G8193	HEMIPLEGIA UNS AFFECTING RIGHT NONDOMINANT SIDE
G8194	HEMIPLEGIA UNS AFFECTING LEFT NONDOMINANT SIDE
G8220	PARAPLEGIA UNSPECIFIED
G8221	PARAPLEGIA COMPLETE
G8222	PARAPLEGIA INCOMPLETE
G8250	QUADRIPLEGIA UNSPECIFIED
G8251	QUADRIPLEGIA C1-C4 COMPLETE
G8252	QUADRIPLEGIA C1-C4 INCOMPLETE
G8253	QUADRIPLEGIA C5-C7 COMPLETE
G8254	QUADRIPLEGIA C5-C7 INCOMPLETE
G830	DIPLEGIA OF UPPER LIMBS
G8310	MONOPLEGIA LOWER LIMB AFFECTING UNSPECIFIED SIDE
G8311	MONOPLEGIA LOWER LIMB RIGHT DOMINANT SIDE
G8312	MONOPLEGIA LOWER LIMB LEFT DOMINANT SIDE
G8313	MONOPLEGIA LOWER LIMB RIGHT NONDOMINANT SIDE
G8314	MONOPLEGIA LOWER LIMB LEFT NONDOMINANT SIDE
G8320	MONOPLEGIA UPPER LIMB AFFECTING UNSPECIFIED SIDE
G8321	MONOPLEGIA UPPER LIMB RIGHT DOMINANT SIDE
G8322	MONOPLEGIA UPPER LIMB LEFT DOMINANT SIDE

CODE	DESCRIPTION
G8323	MONOPLEGIA UPPER LIMB RIGHT NONDOMINANT SIDE
G8324	MONOPLEGIA UPPER LIMB LEFT NONDOMINANT SIDE
G8330	MONOPLEGIA UNS AFFECTING UNSPECIFIED SIDE
G8331	MONOPLEGIA UNS AFFECTING RIGHT DOMINANT SIDE
G8332	MONOPLEGIA UNS AFFECTING LEFT DOMINANT SIDE
G8333	MONOPLEGIA UNS AFFECTING RIGHT NONDOMINANT SIDE
G8334	MONOPLEGIA UNS AFFECTING LEFT NONDOMINANT SIDE
H4930	TOTAL EXTERNAL OPHTHALMOPLEGIA UNSPECIFIED EYE
H4931	TOTAL EXTERNAL OPHTHALMOPLEGIA RIGHT EYE
H4932	TOTAL EXTERNAL OPHTHALMOPLEGIA LEFT EYE
H4933	TOTAL EXTERNAL OPHTHALMOPLEGIA BILATERAL
H4940	PROGRESSIVE EXTERNAL OPHTHALMOPLEGIA UNS EYE
H4941	PROGRESSIVE EXTERNAL OPHTHALMOPLEGIA RIGHT EYE
H4942	PROGRESSIVE EXTERNAL OPHTHALMOPLEGIA LEFT EYE
H4943	PROGRESSIVE EXTERNAL OPHTHALMOPLEGIA BILATERAL
H5120	INTERNUCLEAR OPHTHALMOPLEGIA UNSPECIFIED EYE
H5121	INTERNUCLEAR OPHTHALMOPLEGIA RIGHT EYE
H5122	INTERNUCLEAR OPHTHALMOPLEGIA LEFT EYE
H5123	INTERNUCLEAR OPHTHALMOPLEGIA BILATERAL
H52511	INTERNAL OPHTHALMOPLEGIA COMPLETE TOTAL RT EYE
H52512	INTERNAL OPHTHALMOPLEGIA COMPLETE TOTAL LT EYE
H52513	INTERNAL OPHTHALMOPLEGIA COMPLETE TOTAL BILAT
H52519	INTERNAL OPHTHALMOPLEGIA COMPLETE TOTAL UNS EYE
I120	HYPERTENSIVE CKD W/STAGE 5 CKD OR ESRD
11311	HTN HEART & CKD W/O HF W/STAGE 5 CKD OR ESRD
l132	HTN HEART & CKD W/HF W/STAGE 5 CKD OR ESRD
169351	HEMIPLEGIA FLW CEREBRAL INFARCT AFF RT DOM SIDE
169352	HEMIPLEGIA FLW CEREBRAL INFARCT AFF LT DOM SIDE
169353	HEMIPLEGIA FLW CEREBRAL INFARCT AFF RT NON-DOM
169354	HEMIPLEGIA FLW CEREBRAL INFARCT AFF LT NON-DOM
169359	HEMIPLEGIA FLW CEREBRAL INFARCT AFFCT UNS SIDE
M623	IMMOBILITY SYNDROME PARAPLEGIC
N184	CHRONIC KIDNEY DISEASE STAGE 4 SEVERE
N185	CHRONIC KIDNEY DISEASE STAGE 5
N186	END STAGE RENAL DISEASE
Q050	CERVICAL SPINA BIFIDA WITH HYDROCEPHALUS
Q051	THORACIC SPINA BIFIDA WITH HYDROCEPHALUS
Q052	LUMBAR SPINA BIFIDA WITH HYDROCEPHALUS
Q053	SACRAL SPINA BIFIDA WITH HYDROCEPHALUS
Q054	UNSPECIFIED SPINA BIFIDA WITH HYDROCEPHALUS
Q055	CERVICAL SPINA BIFIDA WITHOUT HYDROCEPHALUS
Q056	THORACIC SPINA BIFIDA WITHOUT HYDROCEPHALUS
Q057	LUMBAR SPINA BIFIDA WITHOUT HYDROCEPHALUS
Q058	SACRAL SPINA BIFIDA WITHOUT HYDROCEPHALUS

Q059     SPINA BIFIDA UNSPECIFIED       Q900     TRISOMY 21, NONMOSAICISM (MEIOTIC NONDISJUNCTION)       Q901     TRISOMY 21, TRANSLOCATION       Q902     DOWN SYNDROME, UNSPECIFIED       Q911     TRISOMY 21, TRANSLOCATION       Q902     DOWN SYNDROME, UNSPECIFIED       Q911     TRISOMY 18, MOSAICISM (MEIOTIC NONDISJUNCTION)       Q912     TRISOMY 18, TRANSLOCATION       Q913     TRISOMY 18, UNSPECIFIED       Q914     TRISOMY 13, MOSAICISM (MEIOTIC NONDISJUNCTION)       Q915     TRISOMY 13, MOSAICISM (MEIOTIC NONDISJUNCTION)       Q916     TRISOMY 13, NOSAICISM (MITOTIC NONDISJUNCTION)       Q917     TRISOMY 13, UNSPECIFIED       WHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTIC       Q920     NONDISJUNCTION)       Q921     WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)       Q921     WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)       Q922     PARTIAL TRISOMY       Q926     MARKER CHROMOSOMES IN NABNAL INDIVIDUAL       Q926     MARKER CHROMOSOMES IN ABNORMAL INDIVIDUAL       Q927     TRIPIOIDY AND POLYPIOIDY       Q928     OTHER SPECIFIED TRISOMY 67 AUTOSOMES, UNSPECIFIED <th>CODE</th> <th>DESCRIPTION</th>	CODE	DESCRIPTION
Q901TRISOMY 21, MOSAICISM (MITOTIC NONDISJUNCTION)Q909DOWN SYNDROME, UNSPECIFIEDQ910TRISOMY 21, TRANSLOCATIONQ911TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q912TRISOMY 18, NONSAICISM (MEIOTIC NONDISJUNCTION)Q913TRISOMY 18, TRANSLOCATIONQ914TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q915TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q916TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q917TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q918TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ9262MARKER CHROMOSOMES IN ABNORMAL INDIVIDUALQ9263OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ928OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOM OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MITOTICQ931NONDISJUNCTION)Q932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ9331VELO-CARDIO-FACIAL SYNDROMEQ934OTHER MICRODELETIONSQ935DELETIONS FROM THE AUTOSOMESQ936OTHER MICRODELETIONSQ937DELETIONS FROM THE AUTOSOMESQ938OTHER MICRODELETIONS <td>Q059</td> <td>SPINA BIFIDA UNSPECIFIED</td>	Q059	SPINA BIFIDA UNSPECIFIED
0902     TRISOMY 21, TRANSLOCATION       0909     DOWN SYNDROME, UNSPECIFIED       0910     TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)       0911     TRISOMY 18, MOSAICISM (MITOTIC NONDISJUNCTION)       0912     TRISOMY 18, MOSAICISM (MITOTIC NONDISJUNCTION)       0913     TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)       0914     TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)       0915     TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)       0916     TRISOMY 13, TRANSLOCATION       0917     TRISOMY 13, UNSPECIFIED       0918     WHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTIC       0920     DOPLICATIONSOME TRISOMY, NOSAICISM (MITOTIC NONDISJUNCTION)       0921     WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)       0922     PARTIAL TRISOMY       09261     MARKER CHROMOSOMES IN NORMAL INDIVIDUAL       09262     MARKER CHROMOSOMES IN NORMAL INDIVIDUAL       09262     MARKER CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTIC       0927     TRIBOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIED       0928     OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMES       0929     TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIED <t< td=""><td>Q900</td><td>TRISOMY 21, NONMOSAICISM (MEIOTIC NONDISJUNCTION)</td></t<>	Q900	TRISOMY 21, NONMOSAICISM (MEIOTIC NONDISJUNCTION)
Q909DOWN SYNDROME, UNSPECIFIEDQ910TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q911TRISOMY 18, MOSAICISM (MITOTIC NONDISJUNCTION)Q912TRISOMY 18, TRANSLOCATIONQ913TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q914TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q915TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q916TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ927TRIFLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MITOTICQ930NONDISJUNCTION)Q932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ9331VELO-CARDIO-FACIAL SYNDROMEQ9341VELO-CARDIO-FACIAL SYNDROMEQ935DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ936OTHER MICRODELETIONSQ937DELETIONS ROM THE AUTOSOMESQ938OTHER MICRODELETIONSQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMES	Q901	TRISOMY 21, MOSAICISM (MITOTIC NONDISJUNCTION)
Q909DOWN SYNDROME, UNSPECIFIEDQ910TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q911TRISOMY 18, MOSAICISM (MITOTIC NONDISJUNCTION)Q912TRISOMY 18, TRANSLOCATIONQ913TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q914TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q915TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q916TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ927TRIFLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MITOTICQ930NONDISJUNCTION)Q932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ9331VELO-CARDIO-FACIAL SYNDROMEQ9341VELO-CARDIO-FACIAL SYNDROMEQ935DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ936OTHER MICRODELETIONSQ937DELETIONS ROM THE AUTOSOMESQ938OTHER MICRODELETIONSQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMESQ939DELETIONS ROM THE AUTOSOMES	Q902	TRISOMY 21, TRANSLOCATION
Q910TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q911TRISOMY 18, NOSAICISM (MITOTIC NONDISJUNCTION)Q912TRISOMY 18, TRANSLOCATIONQ913TRISOMY 18, UNSPECIFIEDQ914TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q915TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q916TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ9262MARKER CHROMOSOMES IN ABNORMAL INDIVIDUALQ927TRIPLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ930NONDISJUNCTION)WHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTICQ930NONDISJUNCTION)WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTICQ931NONDISJUNCTION)Q932CHROMOSOME MONOSOMY, MOSAICISM (MITOTICQ933Q9389OTHER PELACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ934OTHER DELETIONS FROM THE AUTOSOMESQ9389OTHER DELETIONS FROM THE AUTOSOMESQ9389OTHER DELETIONS FROM THE AUTOSOMESQ9389OTHER DELETIONS FROM THE AUTOSOMESQ9329FRAGILEX CHROMOSOME REARRANGEMENT IN ABNORMAL INDIVIDUALQ952BALANCED SEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIV		
Q911   TRISOMY 18, MOSAICISM (MITOTIC NONDISJUNCTION)     Q912   TRISOMY 18, TRANSLOCATION     Q913   TRISOMY 18, UNSPECIFIED     Q914   TRISOMY 13, MOSAICISM (MEIOTIC NONDISJUNCTION)     Q915   TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)     Q916   TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)     Q917   TRISOMY 13, UNSPECIFIED     WHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTIC     Q920   NONDISJUNCTION)     Q921   WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)     Q922   PARTIAL TRISOMY     Q9261   MARKER CHROMOSOMES IN NORMAL INDIVIDUAL     Q9262   MARKER CHROMOSOMES IN NORMAL INDIVIDUAL     Q9263   DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTS     Q9264   MARKER CHROMOSOMES IN ABNORMAL INDIVIDUAL     Q927   TRIPLOIDY AND POLYPLOIDY     Q928   OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMES     Q929   TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIED     Q930   NONDISJUNCTION)     WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTIC     Q931   NONDISJUNCTION)     Q932   CHROMOSOME MONOSOMY, MOSAICISM (MITOTIC     Q933   OTHER PELECIF	Q910	TRISOMY 18, NONMOSAICISM (MEIOTIC NONDISJUNCTION)
Q912   TRISOMY 18, TRANSLOCATION     Q913   TRISOMY 18, UNSPECIFIED     Q914   TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)     Q915   TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)     Q916   TRISOMY 13, TRANSLOCATION     Q917   TRISOMY 13, UNSPECIFIED     WHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTIC     Q920   NONDISJUNCTION)     Q921   WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)     Q922   PARTIAL TRISOMY     Q925   DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTS     Q9261   MARKER CHROMOSOMES IN NORMAL INDIVIDUAL     Q927   TRIPLOIDY AND POLYPLOIDY     Q928   OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMES     Q929   TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIED     WHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTIC     Q930   NONDISJUNCTION)     WHOLE CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOME     Q932   CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOME     Q933   OTHER MICRODELETIONS     Q934   VELO-CARDIO-FACIAL SYNDROME     Q935   DELETION SROM THE AUTOSOMES     Q936   OTHER MICRODELETIONS		
Q913   TRISOMY 18, UNSPECIFIED     Q914   TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)     Q915   TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)     Q916   TRISOMY 13, TRANSLOCATION     Q917   TRISOMY 13, UNSPECIFIED     WHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTIC     Q920   NONDISJUNCTION)     Q921   WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)     Q922   PARTIAL TRISOMY     Q925   DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTS     Q9261   MARKER CHROMOSOMES IN NORMAL INDIVIDUAL     Q9262   MARKER CHROMOSOMES IN ABNORMAL INDIVIDUAL     Q927   TRIPLOIDY AND POLYPLOIDY     Q928   OTHER SPECIFIED TRISOMY OF AUTOSOMES, UNSPECIFIED     WHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTIC     Q930   NONDISJUNCTION)     WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTIC     Q931   WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTIC     Q931   WOLE CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOME     Q932   CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOME     Q9331   VELO-CARDIO-FACIAL SYNDROME     Q9332   CHERTONS FROM THE AUTOSOMES     Q9389   OTHER DELETI	-	
Q914TRISOMY 13, NONMOSAICISM (MEIOTIC NONDISJUNCTION)Q915TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q916TRISOMY 13, UNSPECIFIEDQ917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ927TRIPLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMIS AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MITOTICQ930NONDISJUNCTION)Q932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ933DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ934Q934Q935DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ938OTHER MICROBELETIONSQ938OTHER MICROBELETIONSQ939DELETION FROM AUTOSOMES, UNSPECIFIEDQ938OTHER DELETIONS FROM THE AUTOSOMESQ939DELETION FROM AUTOSOMES, UNSPECIFIEDQ932BALANCED AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ935BALANCED AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ936GTHER MICRODELETIONSQ937DELETION FROM AUTOSOMES, UNSPECIFIEDQ938OTHER DELETIONS FROM THE AUTOSOMESQ939DELETION FROM AUTOSOMES ROMAL RARANGEMENT IN ABNORMAL INDIVIDUALQ938 </td <td>-</td> <td></td>	-	
Q915TRISOMY 13, MOSAICISM (MITOTIC NONDISJUNCTION)Q916TRISOMY 13, TRANSLOCATIONQ917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ927TRIPLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMY OF AUTOSOMES, UNSPECIFIEDQ929TRISOMY AND PARTIAL TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTICQ930NONDISJUNCTION)Q931WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTICQ932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ933DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ9341VELO-CARDIO-FACIAL SYNDROMEQ9352BALANCED EX/AUTOSOMAS, UNSPECIFIEDQ938OTHER MICRODELETIONSQ939DELETION FROM AUTOSOMES, UNSPECIFIEDQ932BALANCED AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ932EALANCED DEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ932EALANCED SEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ932GHER DILETION FROM THE AUTOSOMESQ339DELETION FROM AUTOSOMES, UNSPECIFIEDQ932BALANCED SEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ932FRAGILE X CHROMOSOMER53	-	
Q916TRISOMY 13, TRANSLOCATIONQ917TRISOMY 13, UNSPECIFIEDWHOLE CHROMOSOME TRISOMY, NONMOSAICISM (MEIOTICQ920NONDISJUNCTION)Q921WHOLE CHROMOSOME TRISOMY, MOSAICISM (MITOTIC NONDISJUNCTION)Q922PARTIAL TRISOMYQ925DUPLICATIONS WITH OTHER COMPLEX REARRANGEMENTSQ9261MARKER CHROMOSOMES IN NORMAL INDIVIDUALQ9262MARKER CHROMOSOMES IN ABNORMAL INDIVIDUALQ927TRIPLOIDY AND POLYPLOIDYQ928OTHER SPECIFIED TRISOMIES AND PARTIAL TRISOMIES OF AUTOSOMESQ929TRISOMY AND PARTIAL TRISOMY OF AUTOSOMES, UNSPECIFIEDWHOLE CHROMOSOME MONOSOMY, NONMOSAICISM (MEIOTICQ930NONDISJUNCTION)WHOLE CHROMOSOME MONOSOMY, MOSAICISM (MITOTICQ931NONDISJUNCTION)Q932CHROMOSOME REPLACED WITH RING, DICENTRIC OR ISOCHROMOSOMEQ933DELETIONS WITH OTHER COMPLEX REARRANGEMENTSQ9381VELO-CARDIO-FACIAL SYNDROMEQ9382OTHER MICRODELETIONSQ9383OTHER MICRODELETIONSQ939DELETION FROM AUTOSOMES, UNSPECIFIEDQ952BALANCED SEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ953BALANCED AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ952RAGILE X CHROMOSOMER4181AGE-RELATED COGNITIVE DECLINER532FUNCTIONAL QUADRIPLEGIAR54AGE-RELATED COGNITIVE DECLINER532FUNCTIONAL QUADRIPLEGIAR54ADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XAADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER </td <td></td> <td></td>		
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Q953BALANCED SEX/AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUALQ992FRAGILE X CHROMOSOMER4181AGE-RELATED COGNITIVE DECLINER532FUNCTIONAL QUADRIPLEGIAR54AGE-RELATED PHYSICAL DEBILITYT7411XAADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	Q939	DELETION FROM AUTOSOMES, UNSPECIFIED
Q992FRAGILE X CHROMOSOMER4181AGE-RELATED COGNITIVE DECLINER532FUNCTIONAL QUADRIPLEGIAR54AGE-RELATED PHYSICAL DEBILITYT7411XAADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	Q952	BALANCED AUTOSOMAL REARRANGEMENT IN ABNORMAL INDIVIDUAL
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R532FUNCTIONAL QUADRIPLEGIAR54AGE-RELATED PHYSICAL DEBILITYT7411XAADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	Q992	FRAGILE X CHROMOSOME
R54AGE-RELATED PHYSICAL DEBILITYT7411XAADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER		AGE-RELATED COGNITIVE DECLINE
R54AGE-RELATED PHYSICAL DEBILITYT7411XAADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	R532	FUNCTIONAL QUADRIPLEGIA
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T7411XDADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTERT7411XSADULT PHYSICAL ABUSE, CONFIRMED, SEQUELAT7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	T7411XA	ADULT PHYSICAL ABUSE, CONFIRMED, INITIAL ENCOUNTER
T7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	T7411XD	ADULT PHYSICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER
T7421XAADULT SEXUAL ABUSE, CONFIRMED, INITIAL ENCOUNTERT7421XDADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER	T7411XS	ADULT PHYSICAL ABUSE, CONFIRMED, SEQUELA
T7421XD ADULT SEXUAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER		
		ADULT SEXUAL ABUSE, CONFIRMED, SEQUELA

CODE	DESCRIPTION
T7431XA	ADULT PSYCHOLOGICAL ABUSE, CONFIRMED, INITIAL ENCOUNTER
T7431XD	ADULT PSYCHOLOGICAL ABUSE, CONFIRMED, SUBSEQUENT ENCOUNTER
T7431XS	ADULT PSYCHOLOGICAL ABUSE, CONFIRMED, SEQUELA
T7451XA	ADULT FORCED SEXUAL EXPLOITATION, CONFIRMED, INIT
T7451XD	ADULT FORCED SEXUAL EXPLOITATION, CONFIRMED, SUBS
T7451XS	ADULT FORCED SEXUAL EXPLOITATION, CONFIRMED, SEQUELA
T7611XA	ADULT PHYSICAL ABUSE, SUSPECTED, INITIAL ENCOUNTER
T7611XD	ADULT PHYSICAL ABUSE, SUSPECTED, SUBSEQUENT ENCOUNTER
T7611XS	ADULT PHYSICAL ABUSE, SUSPECTED, SEQUELA
T7621XA	ADULT SEXUAL ABUSE, SUSPECTED, INITIAL ENCOUNTER
T7621XD	ADULT SEXUAL ABUSE, SUSPECTED, SUBSEQUENT ENCOUNTER
T7621XS	ADULT SEXUAL ABUSE, SUSPECTED, SEQUELA
T7631XA	ADULT PSYCHOLOGICAL ABUSE, SUSPECTED, INITIAL ENCOUNTER
T7631XD	ADULT PSYCHOLOGICAL ABUSE, SUSPECTED, SUBSEQUENT ENCOUNTER
T7631XS	ADULT PSYCHOLOGICAL ABUSE, SUSPECTED, SEQUELA
T7651XA	ADULT FORCED SEXUAL EXPLOITATION, SUSPECTED, INIT
T7651XD	ADULT FORCED SEXUAL EXPLOITATION, SUSPECTED, SUBS
T7651XS	ADULT FORCED SEXUAL EXPLOITATION, SUSPECTED, SEQUELA
Z510	ENCOUNTER FOR ANTINEOPLASTIC RADIATION THERAPY
Z5111	ENCOUNTER FOR ANTINEOPLASTIC CHEMOTHERAPY
Z5112	ENCOUNTER FOR ANTINEOPLASTIC IMMUNOTHERAPY
Z590	HOMELESSNESS
Z6911	ENCNTR FOR MNTL HLTH SERV FOR VICTIM OF SPOUS OR PRTNR ABUSE
Z6981	ENCOUNTER FOR MENTAL HEALTH SERVICES FOR VICTIM OF OTH ABUSE
Z7682	AWAITING ORGAN TRANSPLANT STATUS