

HEALTH PLAN OF SAN JOAQUIN		
Subject: (CSHCN) Children with Special Health Care Needs – Identification of and Coordination of Care with VMRC/Early Start Program		
Department: Medical Management - Utilization		Policy #: UM51
Applies to: Medi-Cal		Scope: UM
Effective Date: 2/1/96	Revised Date: 10/08	Approved by: <i>Signature on file</i> Medical Director

DEFINITION:

Medical Home: (Defined by the Academy of Pediatrics AAP The Medical Home. PEDIATRICS Vol.110 No. 1 July 2002, pp. 184-186) - *A medical home is a provider that is accessible, family-centered, continuous, comprehensive, coordinated, compassionate, and culturally effective. The medical home implies a partnership of joint responsibility and trust between the physician and the family to address all aspects of the child and family needs.*

POLICY

- A. Health Plan of San Joaquin (HPSJ) will make every effort to add those children under age 21 who meet the definition of Children with Special Health Care Needs (CSHCN).
 - 1. Provide for coordination and continuity of care
 - 2. Ensure direct access to specialists
 - 3. Assess and improve the quality of care provided
- B. HPSJ will maintain Memorandum of Understanding (MOU) with agencies specializing in providing specialty care for this population. Local agencies are:
 - 1. California Children Services (CCS)
 - 2. Valley Mountain Regional Center (VMRC)
 - 3. San Joaquin County Mental Health (SJCMMH)
 - 4. Public Health (PH)

PROCEDURE

- A. PCP Identification and Referral Procedures for CSHCN
 - 1. The PCP will schedule an Initial Health Assessment (IHA) on all new members assigned, within specific time frames.

2. The IHA will be within 60 days of enrollment for children under the age of 18 months.
 3. The IHA will be within 120 days of enrollment for children/members over the age of 18 months.
 4. Developmental assessments are preformed as part of the Initial Health Assessment (IHA), or during routine health assessment exams., Children/members age birth to 21 identified as having significant developmental delay, or are at risk for development disability, (based on the qualifying risk factors – Attachment A or with a diagnosis listed on – Attachment B), will be scheduled for a comprehensive evaluation by the PCP within 60 days to determine a course of treatment and possible referrals.
 5. The PCP, is responsible for providing and/or arranging all medically necessary diagnostic, specialty or therapeutic services to evaluate, correct and/or ameliorate a suspected or confirmed condition. Therefore, upon identification of a CSHCN, he/she will assume the role as the member's medical home and will be responsible for monitoring, care coordination, and case management requirements in relation to the member's ongoing special health care and related needs. Documentation of all tests, referrals to specialists, consultation notes, follow up services and recommendations, etc., pertaining to the member are to be filed as part of the member's medical record.
 6. If the PCP feels unable to perform this function, HPSJ will assist in locating a PCP that will accept the responsibility and function as the member's PCP and medical home. The member and all medical records will then be transferred to the care of that physician.
 7. Under special circumstances, a specialist may serve as the PCP.
 8. The PCP may refer the member to SJCMH, CCS or VMRC for evaluation of newly diagnosed or ongoing specialty care and needs. All medical records pertinent to the member's condition will be sent to agency that the member is being referred.
 9. In situations where the child is eligible for both CCS and VMRC services, the primary referral will be to CCS. A referral may be made to CCS and VMRC simultaneously if both medical and early intervention services appear to be necessary.
- B. HPSJ Identification of CSHCN
1. The Utilization Management staff will identify CSHCN through the following:
 2. Referral / Authorization requests submitted by the PCP or Specialists with any of the qualifying diagnosis codes from Attachment B.

3. DRE, the HPSJ electronic authorization system, will flag any requests submitted that have been identified as a CSHCN diagnosis. (Attachment B)
 4. Medical record review with documentation supporting any of the clinical indications from Attachment A or B.
 5. Quarterly reports run using the qualifying diagnosis and aid codes.
- C. Case Management
1. Coordination of services to the member is essential, and the HPSJ Case Manager will be involved to see that;
 2. A primary Case Manager or Caregiver is assigned to coordinate services.
 3. A list of resources and material is available in-network as well as out-of-network that will assist in accessing resources.
 4. Timely sharing or team conferences with in-plan or out-of-plan agencies/providers of care are established so that all caregivers responsible for the member's care are up to date on the current status of the case. These conferences must be consistent with laws to protect the confidentiality of medical information.
 5. Family or guardians of the member are involved in the decision-making processes concerning the member's health in collaboration with all appropriate caregivers.

REFERENCE

- A. Title 22 Section 51184 and 51340
- B. DHS Contract Exhibit A, Attachment 11, Section 7
- C. HPSJ Policy and Procedure QA 22
- D. HPSJ Policy and Procedure UM 53
- E. MOU with Mental Health, CCS, VMRC and Public Health

Created by/Date	Revised by/Date	Revised by/Date	Revised by/Date	Revised by/Date	Revised by/Date
2/1996	M. Jordan 8/2004	M. Jordan RN 2/06	Donna 6/26/08 Reviewed by Joy Scott 10/08		

Attachment A

Qualifying risk factors for service for the Early Start Program include:

A. Medical

- Very low birth weight (1500 grams).
- Prematurity (<32 weeks).
- Metabolic Problems (i.e., hypoglycemia, hypocalcemia)
- CNS infection/abnormality
- Seizure activity during first weeks of life.
- Serious biomedical insult (i.e., CNS bleed)
- Multiple congenital abnormalities requiring specific services.
- Positive neonatal tox screen/drug withdrawal.
- Prolonged hypoxemia
- Hyperbilirubinemia
- Prenatal exposure to teratogens
- Significant failure to thrive.

B. Clinical/Behavioral

- Infant born to DD parent
- Persistent tonal problems

C. Developmental Delay

- Cognitive
- Physical
- Communication
- Social/emotional
- Adaptive/self-help skills

		Attachment B
		QUALIFYING DIAGNOSIS as indicated by DHS
DIAGNOSIS	DESCRIP	
20801	ACT LEUK UNS CL W RMSON	
2080	ACT LEUK UNS CL W/O RMSN	
20800	ACT LEUK UNS CL W/O RMSN	
20401	ACT LYM LEUK W RMSION	
20400	ACT LYM LEUK W/O RMSION	
20501	ACT MYL LEUK W RMSION	
20500	ACT MYL LEUK W/O RMSION	
2040	ACUTE LYMPHOID LEUKEMIA	
2050	ACUTE MYELOID LEUKEMIA	
2983	ACUTE PARANOID REACTION	
2969	AFFECT PSYCHOSES NEC/NOS	
296	AFFECTIVE PSYCHOSES	
29699	AFFECTIVE PSYCHOSES NEC	
29690	AFFECTIVE PSYCHOSIS NOS	
31501	ALEXIA	
95217	ANTERIOR CORD SYN/T7-T12	
95202	ANTERIOR CORD SYND/C1-C4	
95207	ANTERIOR CORD SYND/C5-C7	
95212	ANTERIOR CORD SYND/T1-T6	
284	APLASTIC ANEMIA	
2849	APLASTIC ANEMIA NOS	
2848	APLASTIC ANEMIAS NEC	
3151	ARITHMETICAL DISORDER	
V08	ASYMP HIV INFECTN STATUS	
29682	ATYPICAL DEPRESSIVE DIS	
29681	ATYPICAL MANIC DISORDER	
74913	BILAT CLEFT LIP-COMplete	
74914	BILAT CLEFT LIP-INCOMPL	
74903	BILAT CLEFT PALATE-COMPL	
74904	BILAT CLEFT PALATE-INC	
74923	BILAT CLFT PALAT/LIP-COM	
74924	BILAT CLFT PALAT/LIP-INC	
29656	BIPOL AFF DEPR-FULL REM	
29655	BIPOL AFF DEPR-PART REM	
29646	BIPOL AFF MANIC-FULL REM	
29645	BIPOL AFF MANIC-PART REM	
29653	BIPOL AFF, DEPR-SEVERE	
29643	BIPOL AFF, MANIC-SEVERE	
29640	BIPOL AFF, MANIC-UNSPEC	
29663	BIPOL AFF, MIXED-SEVERE	

29660	BIPOL AFF, MIXED-UNSPEC
29666	BIPOL AFF, MIX-FULL REM
29665	BIPOL AFF, MIX-PART REM
29654	BIPOL DEPR-SEV W PSYCH
29644	BIPOL MANIC-SEV W PSYCH
29664	BIPOL MIXED-SEV W PSYCH
29650	BIPOLAR AFF, DEPR-UNSPEC
29641	BIPOLAR AFF, MANIC-MILD
29661	BIPOLAR AFF, MIXED-MILD
29651	BIPOLAR AFFEC, DEPR-MILD
29652	BIPOLAR AFFEC, DEPR-MOD
29642	BIPOLAR AFFEC, MANIC-MOD
29662	BIPOLAR AFFEC, MIXED-MOD
2965	BIPOLAR AFFECT, DEPRESS
29670	BIPOLAR AFFECTIVE DISORDER, UN
2967	BIPOLAR AFFECTIVE NOS
2964	BIPOLAR AFFECTIVE, MANIC
2966	BIPOLAR AFFECTIVE, MIXED
33381	BLEPHAROSPASM
5792	BLIND LOOP SYNDROME
33393	BNIGN SHUDDERING ATTACKS
V424	BONE TRANSPLANT STATUS
95204	C1-C4 SPIN CORD INJ NEC
95200	C1-C4 SPIN CORD INJ NOS
95209	C5-C7 SPIN CORD INJ NEC
95205	C5-C7 SPIN CORD INJ NOS
745	CARDIAC SEPTAL CLOS ANOM
9524	CAUDA EQUINA INJURY
34460	CAUDA EQUINA SYND NOS
3446	CAUDA EQUINA SYNDROME
5790	CELIAC DISEASE
95218	CENTRAL CORD SYN/T7-T12
95203	CENTRAL CORD SYND/C1-C4
95208	CENTRAL CORD SYND/C5-C7
95213	CENTRAL CORD SYND/T1-T6
3303	CERB DEG CHLD IN OTH DIS
3308	CEREB DEGEN IN CHILD NEC
3309	CEREB DEGEN IN CHILD NOS
3302	CEREB DEGEN IN LIPIDOSIS
330	CEREBRAL DEGEN IN CHILD
3301	CEREBRAL LIPIDOSES
3438	CEREBRAL PALSY NEC
3439	CEREBRAL PALSY NOS
9520	CERVICAL SPINAL CORD INJ
29980	CHILD PSYCHOS NEC-ACTIVE
29981	CHILD PSYCHOS NEC-RESID
29990	CHILD PSYCHOS NOS-ACTIVE

29991	CHILD PSYCHOS NOS-RESID
3335	CHOREA NEC
20811	CHR LEUK UNS CL W RMSON
20810	CHR LEUK UNS CL W/O RMSN
20411	CHR LYM LEUK W RMSION
20410	CHR LYM LEUK W/O RMSION
2041	CHR LYMPHOID LEUKEMIA
20511	CHR MYL LEUK W RMSION
20510	CHR MYL LEUK W/O RMSION
2081	CHRONIC LEUKEMIA NOS
2051	CHRONIC MYELOID LEUKEMIA
7491	CLEFT LIP
74910	CLEFT LIP NOS
7490	CLEFT PALATE
749	CLEFT PALATE & CLEFT LIP
74925	CLEFT PALATE & LIP NEC
74920	CLEFT PALATE & LIP NOS
74900	CLEFT PALATE NOS
7492	CLEFT PALATE W CLEFT LIP
7450	COMMON TRUNCUS
7453	COMMON VENTRICLE
74510	COMPL TRANSPOS GREAT VES
95201	COMPLETE LES CORD/C1-C4
95206	COMPLETE LES CORD/C5-C7
95211	COMPLETE LES CORD/T1-T6
95216	COMPLETE LES CORD/T7-T12
7464	CONG AORTA VALV INSUFFIC
7463	CONG AORTA VALV STENOSIS
7468	CONG HEART ANOMALY NEC
74689	CONG HEART ANOMALY NEC
7469	CONG HEART ANOMALY NOS
3590	CONG HERED MUSC DYSTRPHY
7466	CONG MITRAL INSUFFICIENC
74601	CONG PULMON VALV ATRESIA
74602	CONG PULMON VALVE STENOS
74681	CONG SUBAORTIC STENOSIS
7461	CONG TRICUSP ATRES/STEN
2840	CONGEN APLASTIC ANEMIA
7465	CONGEN MITRAL STENOSIS
3430	CONGENITAL DIPLEGIA
74686	CONGENITAL HEART BLOCK
3431	CONGENITAL HEMIPLEGIA
3433	CONGENITAL MONOPLÉGIA
3432	CONGENITAL QUADRIPLEGIA
3154	COORDINATION DISORDER
31540	COORDINATION DISORDER
7457	COR BILOCULARE

74682	COR TRIATRIATUM
V425	CORNEA TRANSPLANT STATUS
74685	CORONARY ARTERY ANOMALY
74512	CORRECT TRANSPOS GRT VES
3330	DEGEN BASAL GANGLIA NEC
2963	DEPR PSYCH, RECUR EPISOD
2962	DEPR PSYCH, SINGL EPISOD
29626	DEPR PSYCHOS-FULL REMISS
29625	DEPR PSYCHOS-PART REMISS
29624	DEPR PSYCHOS-SEV W PSYCH
29621	DEPRESS PSYCHOSIS-MILD
29623	DEPRESS PSYCHOSIS-SEVERE
29620	DEPRESS PSYCHOSIS-UNSPEC
29622	DEPRESSIVE PSYCHOSIS-MOD
3159	DEVELOPMENT DELAY NOS
3158	DEVELOPMENT DELAYS NEC
31531	DEVELOPMENT LANGUAGE DIS
31502	DEVELOPMENTAL DYSLEXIA
2502	DIAB W HYPEROSMOLAR COMA
2506	DIAB W NEUROLOGIC MANIF
2505	DIAB W OPHTHALMIC MANIF
2504	DIAB W RENAL MANIFEST
250	DIABETES MELLITUS
2500	DIABETES MELLITUS UNCOMP
2507	DIABETES W CIRCULAT DIS
2509	DIABETES W COMPLIC NOS
2501	DIABETES W KETOACIDOSIS
2508	DIABETES W MANIFEST NEC
2503	DIABETES WITH COMA NEC
3442	DIPLEGIA OF UPPER LIMBS
29910	DISINTEGR PSYCH-ACTIVE
29911	DISINTEGR PSYCH-RESIDUAL
2991	DISINTEGRATIVE PSYCHOSIS
25071	DMI CIRC NT ST UNCNRDL
25073	DMI CIRC UNCNRDL
25023	DMI HPROMLR UNCONTROLD
25021	DMI HPRSM NT ST UNCNRDL
25011	DMI KETO NT ST UNCNRDL
25013	DMI KETOACD UNCONTROLD
25061	DMI NEURO NT ST UNCNRDL
25063	DMI NEURO UNCNRDL
25031	DMI O CM NT ST UNCNRDL
25051	DMI OPHTH NT ST UNCNRDL
25053	DMI OPHTH UNCNRDL
25033	DMI OTH COMA UNCONTROLD
25081	DMI OTH NT ST UNCNRDL
25083	DMI OTH UNCNRDL

25043	DMI RENAL UNCINTRLD
25041	DMI RENL NT ST UNCINTRLD
25091	DMI UNSPF NT ST UNCINTRLD
25093	DMI UNSPF UNCINTRLD
25001	DMI WO CMP NT ST UNCINTRL
25003	DMI WO CMP UNCINTRLD
25070	DMII CIRC NT ST UNCINTRLD
25072	DMII CIRC UNCINTRLD
25022	DMII HPROMLR UNCONTROLD
25020	DMII HPRSM NT ST UNCINTRL
25010	DMII KETO NT ST UNCINTRLD
25012	DMII KETOACD UNCONTROLD
25060	DMII NEURO NT ST UNCINTRL
25062	DMII NEURO UNCINTRLD
25030	DMII O CM NT ST UNCINTRLD
25050	DMII OPHTH NT ST UNCINTRL
25052	DMII OPHTH UNCINTRLD
25032	DMII OTH COMA UNCONTROLD
25080	DMII OTH NT ST UNCINTRLD
25082	DMII OTH UNCINTRLD
25042	DMII RENAL UNCINTRLD
25040	DMII RENL NT ST UNCINTRLD
25090	DMII UNSPF NT ST UNCINTRL
25092	DMII UNSPF UNCINTRLD
25000	DMII WO CMP NT ST UNCINTR
25002	DMII WO CMP UNCINTRLD
9521	DORSAL SPINAL CORD INJUR
74511	DOUBLE OUTLET RT VENTRIC
2998	EARLY CHLD PSYCHOSES NEC
2999	EARLY CHLD PSYCHOSIS NOS
7462	EBSTEIN'S ANOMALY
74569	ENDOCARD CUSHION DEF NEC
74560	ENDOCARD CUSHION DEF NOS
7456	ENDOCARD CUSHION DEFECTS
2823	ENZYME DEFIC ANEMIA NEC
34571	EPIL PAR CONT W INTR EPI
34570	EPIL PAR CONT W/O INT EP
34580	EPILEP NEC W/O INTR EPIL
34590	EPILEP NOS W/O INTR EPIL
3457	EPILEPS PARTIAL CONTINUA
345	EPILEPSY
3458	EPILEPSY NEC
34581	EPILEPSY NEC W INTR EPIL
3459	EPILEPSY NOS
34591	EPILEPSY NOS W INTR EPIL
2981	EXCITATIV TYPE PSYCHOSIS
333	EXTRAPYRAMIDAL DIS NEC

33399	EXTRAPYRAMIDAL DIS NEC
33390	EXTRAPYRAMIDAL DIS NOS
3339	EXTRPYRMIDAL DIS NEC/NOS
3593	FAMIL PERIODIC PARALYSIS
3420	FLACCID HEMIPLEGIA
34202	FLCCD HMIPLG NONDMNT SDE
34201	FLCCD HMIPLGA DOMNT SIDE
34200	FLCCD HMIPLGA UNSPF SIDE
33389	FRAGM TORSION DYSTON NEC
3338	FRAGMNT TORSION DYSTONIA
34511	GEN CNV EPIL W INTR EPIL
34510	GEN CNV EPIL W/O INTR EP
3451	GEN CONVULSIVE EPILEPSY
34501	GEN NONCONV EP W INTR EP
3450	GEN NONCONVULS EPILEPSY
34500	GEN NONCV EP W/O INTR EP
2822	GLUTATHIONE DIS ANEMIA
3453	GRAND MAL STATUS
28261	HB-S DISEASE W/O CRISIS
28262	HB-S DISEASE WITH CRISIS
V421	HEART TRANSPLANT STATUS
V422	HEART VALVE TRANSPLANT
342	HEMIPLEGIA
3429	HEMIPLEGIA NOS
2827	HEMOGLOBINOPATHIES NEC
2828	HERED HEMOLYTIC ANEM NEC
2829	HERED HEMOLYTIC ANEM NOS
3591	HERED PROG MUSC DYSTRPHY
2821	HEREDIT ELLIPTOCYTOSIS
282	HEREDIT HEMOLYTIC ANEMIA
2820	HEREDITARY SPHEROCYTOSIS
0429	HTLV-III/LAV AIDS W/WO OTHER C
0421	HTLV-III/LAV CAUSING OTHER SPE
0422	HTLV-III/LAV W/SPEC MALIGNANT
0420	HTLV-III/LAV WITH SPECIFIED CO
042	HUMAN IMMUNO VIRUS DIS
3334	HUNTINGTON'S CHOREA
7467	HYPOPLAS LEFT HEART SYND
3336	IDIOPAT TORSION DYSTONIA
34561	INF SPASM W INTRACT EPIL
34560	INF SPASM W/O INTR EPIL
2990	INFANTILE AUTISM
29900	INFANTILE AUTISM-ACTIVE
29901	INFANTILE AUTISM-RESID
343	INFANTILE CEREBRAL PALSY
3434	INFANTILE HEMIPLEGIA
3456	INFANTILE SPASMS

3596	INFL MYOPATHY IN OTH DIS
74683	INFUNDIB PULMON STENOSIS
5798	INTEST MALABSORPTION NEC
5799	INTEST MALABSORPTION NOS
5793	INTEST POSTOP NONABSORB
579	INTESTINAL MALABSORPTION
V4284	INTESTINES
V420	KIDNEY TRANSPLANT STATUS
20891	LEUKEMIA NOS W REMISSION
20890	LEUKEMIA NOS W/O REMSION
2088	LEUKEMIA-UNSPEC CELL NEC
2089	LEUKEMIA-UNSPEC CELL NOS
208	LEUKEMIA-UNSPECIF CELL
3300	LEUKODYSTROPHY
V427	LIVER TRANSPLANT STATUS
34481	LOCKED-IN STATE
9522	LUMBAR SPINAL CORD INJUR
V426	LUNG TRANSPLANT STATUS
204	LYMPHOID LEUKEMIA
2048	LYMPHOID LEUKEMIA NEC
2049	LYMPHOID LEUKEMIA NOS
1917	MAL NEO BRAIN STEM
1945	MAL NEO CAROTID BODY
1915	MAL NEO CEREB VENTRICLE
1916	MAL NEO CEREBELLUM NOS
1948	MAL NEO ENDOCRINE NEC
1949	MAL NEO ENDOCRINE NOS
1914	MAL NEO OCCIPITAL LOBE
194	MAL NEO OTHER ENDOCRINE
1946	MAL NEO PARAGANGLIA NEC
1913	MAL NEO PARIETAL LOBE
1912	MAL NEO TEMPORAL LOBE
1918	MALIG NEO BRAIN NEC
1919	MALIG NEO BRAIN NOS
1911	MALIG NEO FRONTAL LOBE
1941	MALIG NEO PARATHYROID
1943	MALIG NEO PITUITARY
1944	MALIGN NEO PINEAL GLAND
1940	MALIGN NEOPL ADRENAL
1910	MALIGN NEOPL CEREBRUM
191	MALIGNANT NEOPLASM BRAIN
74687	MALPOSITION OF HEART
2960	MANIC DIS, SINGL EPISODE
29606	MANIC DIS-FULL REMISSION
29601	MANIC DISORDER-MILD
29602	MANIC DISORDER-MOD
29603	MANIC DISORDER-SEVERE

29600	MANIC DISORDER-UNSPEC
29605	MANIC DIS-PARTIAL REMISS
29604	MANIC DIS-SEVERE W PSYCH
2961	MANIC, RECURRENT EPISODE
29689	MANIC-DEPRESSIVE NEC
2968	MANIC-DEPRESSIVE NEC/NOS
29680	MANIC-DEPRESSIVE NOS
319	MENTAL RETARDATION NOS
3155	MIXED DEVELOPMENT DIS
34432	MNPLG LWR LMB NONDMNT SD
34442	MNPLG UPR LMB NONDMNT SD
3180	MOD MENTAL RETARDATION
3445	MONOPLÉGIA NOS
3443	MONOPLÉGIA OF LOWER LIMB
3444	MONOPLÉGIA OF UPPER LIMB
34431	MONPLGA LWR LMB DMNT SDE
34430	MONPLGA LWR LMB UNSP SDE
34441	MONPLGA UPR LMB DMNT SDE
34440	MONPLGA UPR LMB UNSP SDE
359	MUSCULAR DYSTROPHIES
205	MYELOID LEUKEMIA
2058	MYELOID LEUKEMIA NEC
2059	MYELOID LEUKEMIA NOS
2053	MYELOID SARCOMA
20531	MYL SRCOMA W RMSION
20530	MYL SRCOMA W/O RMSION
3332	MYOCLONUS
3595	MYOPATHY IN ENDOCRIN DIS
3598	MYOPATHY NEC
3599	MYOPATHY NOS
3592	MYOTONIC DISORDERS
34461	NEUROGENIC BLADDER
33392	NEUROLEPTIC MALGNT SYND
74684	OBSTRUCT HEART ANOM NEC
V4282	ORGAN OR TISSUE REPLACED BY TR
V4283	ORGAN OR TISSUE REPLACED BY TR
V4281	ORGAN OR TISSUEREPLACED BY TRA
V42	ORGAN TRANSPLANT STATUS
33384	ORGANIC WRITERS' CRAMP
33382	OROFACIAL DYSKINESIA
74561	OSTIUM PRIMUM DEFECT
34282	OT SP HMIPLG NONDMNT SDE
34281	OT SP HMIPLGA DOMNT SIDE
34280	OT SP HMIPLGA UNSPF SIDE
3152	OTH LEARNING DIFFICULTY
20881	OTH LEUK UNS CL W RMSON
20880	OTH LEUK UNS CL W/O RMSN

20481	OTH LYM LEUK W RMSION
20480	OTH LYM LEUK W/O RMSION
20581	OTH MYL LEUK W RMSION
20580	OTH MYL LEUK W/O RMSION
298	OTH NONORGANIC PSYCHOSES
344	OTH PARALYTIC SYNDROMES
34489	OTH SPCF PARALYTIC SYND
29880	OTHER AND UNSPECIFIED REACTIVE
746	OTHER CONGEN HEART ANOM
318	OTHER MENTAL RETARDATION
V4289	OTHER ORGAN OR TISSUE REPLACED
34409	OTHER QUADRIPLEGIA
31800	OTHER SPECIFIED MENTAL RETARDA
5794	PANCREATIC STEATORRHEA
3449	PARALYSIS NOS
3448	PARALYTIC SYNDROMES NEC
3441	PARAPLEGIA NOS
34551	PART EPIL W INTR EPIL
34550	PART EPIL W/O INTR EPIL
3455	PARTIAL EPILEPSY NEC
3452	PETIT MAL STATUS
3182	PROFOUND MENTAL RETARDAT
31820	PROFOUND MENTAL RETARDATION
2984	PSYCHOGEN PARANOID PSYCH
3454	PSYCHOMOTOR EPILEPSY
299	PSYCHOSES OF CHILDHOOD
2989	PSYCHOSIS NOS
34541	PSYMOTR EPIL W INTR EPIL
34540	PSYMOTR EPIL W/O INT EPI
74609	PULMONARY VALVE ANOM NEC
74600	PULMONARY VALVE ANOM NOS
7460	PULMONARY VALVE ANOMALY
3440	QUADRIPLEGIA NOS
34400	QUADRIPLEGIA, UNSPECIFD
34401	QUADRPLG C1-C4, COMPLETE
34402	QUADRPLG C1-C4, INCOMPLT
34403	QUADRPLG C5-C7, COMPLETE
34404	QUADRPLG C5-C7, INCOMPLT
2980	REACT DEPRESS PSYCHOSIS
2988	REACT PSYCHOSIS NEC/NOS
2982	REACTIVE CONFUSION
31509	READING DISORDER NEC
31500	READING DISORDER NOS
29634	REC DEPR PSYCH-PSYCHOTIC
31532	RECEPTIVE LANGUAGE DISORDER (M
29636	RECUR DEPR PSYC-FULL REM
29633	RECUR DEPR PSYCH-SEVERE

29635	RECUR DEPR PSYC-PART REM
29611	RECUR MANIC DIS-MILD
29612	RECUR MANIC DIS-MOD
29613	RECUR MANIC DIS-SEVERE
29610	RECUR MANIC DIS-UNSPEC
29616	RECUR MANIC-FULL REMISS
29615	RECUR MANIC-PART REMISS
29614	RECUR MANIC-SEV W PSYCHO
29631	RECURRE DEPR PSYCHOS-MILD
29632	RECURRE DEPR PSYCHOS-MOD
29630	RECURRE DEPR PSYCHOS-UNSP
9523	SACRAL SPINAL CORD INJUR
20821	SBAC LEUK UNS CL W RMSON
20820	SBAC LEUK UNS CL W/O RMS
20421	SBAC LYM LEUK W RMSION
20420	SBAC LYM LEUK W/O RMSION
20521	SBAC MYL LEUK W RMSION
20520	SBAC MYL LEUK W/O RMSION
7455	SECUNDUM ATRIAL SEPT DEF
7458	SEPTAL CLOSURE ANOM NEC
7459	SEPTAL CLOSURE ANOM NOS
3181	SEVERE MENTAL RETARDAT
31810	SEVERE MENTAL RETARDATION
2826	SICKLE-CELL ANEMIA
28269	SICKLE-CELL ANEMIA NEC
28260	SICKLE-CELL ANEMIA NOS
2825	SICKLE-CELL TRAIT
28263	SICKLE-CELL/HB-C DISEASE
V423	SKIN TRANSPLANT STATUS
33383	SPASMODIC TORTICOLLIS
3421	SPASTIC HEMIPLEGIA
315	SPECIFIC DEVELOP DELAYS
3150	SPECIFIC READING DIS
31539	SPEECH/LANGUAGE DIS NEC
3153	SPEECH/LANGUAGE DISORDER
74101	SPIN BIF W HYDRCEPH-CERV
74102	SPIN BIF W HYDRCEPH-DORS
74103	SPIN BIF W HYDRCEPH-LUMB
74100	SPIN BIF W HYDROCEPH NOS
9528	SPIN CORD INJ-MULT SITE
7410	SPINA BIF W HYDROCEPHAL
741	SPINA BIFIDA
7419	SPINA BIFIDA
74190	SPINA BIFIDA
74191	SPINA BIFIDA-CERV
74192	SPINA BIFIDA-DORSAL
74193	SPINA BIFIDA-LUMBAR

952	SPINAL CORD INJ W/O FX
9529	SPINAL CORD INJURY NOS
34212	SPSTC HMIPLG NONDMNT SDE
34211	SPSTC HMIPLGA DOMNT SIDE
34210	SPSTC HMIPLGA UNSPF SIDE
33391	STIFF-MAN SYNDROME
2042	SUBAC LYMPHOID LEUKEMIA
2052	SUBACUT MYELOID LEUKEMIA
2082	SUBACUTE LEUKEMIA NOS
3337	SYMPTOM TORSION DYSTONIA
95214	T1-T6 SPIN CORD INJ NEC
95210	T1-T6 SPIN CORD INJ NOS
95219	T7-T12 SPIN CORD INJ NEC
95215	T7-T12 SPIN CORD INJ NOS
7452	TETRALOGY OF FALLOT
2824	THALASSEMIAS
3333	TICS OF ORGANIC ORIGIN
3594	TOXIC MYOPATHY
V428	TRANSPLANT STATUS NEC
V429	TRANSPLANT STATUS NOS
74519	TRANSPOS GREAT VESS NEC
7451	TRANSPOS OF GREAT VESSEL
3331	TREMOR NEC
5791	TROPICAL SPRUE
74921	UNIL CLEFT PALAT/LIP-COM
74922	UNIL CLEFT PALAT/LIP-INC
74911	UNILAT CLEFT LIP-COMPL
74912	UNILAT CLEFT LIP-IMCOMPL
74901	UNILAT CLEFT PALATE-COMP
74902	UNILAT CLEFT PALATE-INC
20491	UNS LYM LEUK W RMSION
20490	UNS LYM LEUK W/O RMSION
20591	UNS MYL LEUK W RMSION
20590	UNS MYL LEUK W/O RMSION
34291	UNSP HEMIPLGA DOMNT SIDE
34290	UNSP HEMIPLGA UNSPF SIDE
34292	UNSP HMIPLGA NONDMNT SDE
31900	UNSPECIFIED MENTAL RETARDATION
29890	UNSPECIFIED PSYCHOSIS
7454	VENTRICULAR SEPT DEFECT