

Progr Opti		Therapy	ICD-10 Code	ICD-10 Descriptions	HCPCS Code	HCPCS Descriptions	CPT Code	CPT Description
	GTS-5	Luxturna	H35.50 H35.52 H35.54	Unspecified hereditary retinal dystrophy Other indexing guidance for H35.50: Leber's congenital amaurosis Best's disease Pigmentary retinal dystrophy Retinitis pigmentosa Dystrophies primarily involving the retinal pigment epitheliu	J3398	Injection, Luxturna (voretigene nepar- vovec-rzyl), 1 billion vector genomes	67036 67299	Vitrectomy, mechanical, pars plana approach Unlisted procedure, posterior segment
			G12.0	Infantile spinal muscular atrophy, type 1			96365	IV infusion, for therapy, prophylaxis, or diagnosis; initial, up to one
		Zolgensma	G12.1	Other inherited spinal muscular atrophy	J3399	Injection, Zolgensma (onasemnogene abeparvovec-xioi), per treatment up to 5×1015 vector genomes	99218- 99220	Initial observation care, per day, for the evaluation and management of a patient, which requires these 3 key components: a detailed or comprehensive history, a detailed or comprehensive examination, and medical decision-making that is
			G12.25	Progressive spinal muscular atrophy			99234- 99236	straightforward or of low complexity, moderate complexity, or high complexity Observation care admission and discharge services for the evaluation and management of a patient including admission and discharge on the same date, which requires these 3 key components: detailed or comprehensive history, a detailed or comprehensive examination, and medical decision-making that is straightforward or of low complexity, moderate complexity, or high complexity
			G12.8 G12.9	Other spinal muscular atrophies and related syndromes Spinal muscular atrophy, unspecified				
			G12.0	Infantile spinal muscular atrophy, type 1				
		Spinraza	G12.1	Other inherited spinal muscular atrophy	J2326 J3490	INJECTION, NUSINERSEN, 0.1 MG Unclassified drugs	96450	Chemotherapy administration, into central nervous system (CNS) (e.g. intrathecal), requiring spinal puncture Chemotherapy administration, intravenous infusion technique; up to 1 hour, single
			G12.25 G12.8 G12.9	Progressive spinal muscular atrophy Other spinal muscular atrophies and related syndromes Spinal muscular atrophy, unspecified				
			D56	Thalassemia				
		Zynteglo	D56.1 D56.2	Beta Thalassemia Delta-beta Thalassemia	J3590	Unclassified biologics	96413	or initial substance/drug
			D56.3 D56.4 D56.5 D56.8	Thalassemia minor Hereditary persistence of fetal hemoglobin (HPFH) Hemoglobin E-beta thalassemia Other thalassemia	J3393	Injection, betibeglogene autotemcel, per treatment (Eff. 7/1/2024)	96415	Chemotherapy administration, intravenous infusion technique; each additional hour
			D56.9	Thalassemia, unspecified				Chemotherapy administration, intravenous infusion technique; up to 1 hour, single
		Skysona	E71.52 E71.520	X-linked adrenoleukodystrophy Childhood cerebral X-linked adrenoleukodystrophy	J3490 J3590	Unclassified drugs	96413	or initial substance/drug
			E71.521 E71.522 E71.528	Adolescent X-linked adrenoleukodystrophy Adrenomyeloneuropathy Other X-linked adrenoleukodystrophy		Unclassified biologics	96415	Chemotherapy administration, intravenous infusion technique; each additional hour
-			E71.529	X-linked adrenoleukodystrophy, unspecified type				
		Hemgenix	D67	Hereditary factor IX deficiency	- J1411	Injection, etranacogene dezapar- vovec-drlb, per therapeutic dose	96365	Infusion first hour 96366 Infusion each additional hour
			D68	Other coagulation defects				
					C9399	Unclassified drugs or biologicals	96365	Intravenous infusion, for therapy, prophylaxis, or diagnosis (specify substance or drug); initial, up to 1 hour
10					E0780	Ambulatory infusion pump, mechanical, reusable, for infusion less than 8 hours	96366	Intravenous infusion, for therapy, prophylaxis, or diagnosis (specify substance or drug); each additional hour
GTS-15			D66 Hereditary factor VIII deficiency		J3490	Unclassified drugs	99221	Initial hospital inpatient or observation care, per day, for the evaluation and management of a patient, which requires a medically appropriate history and/ or examination and straightforward or lowlevel medical decision-making. When using total time on the date of the encounter for code selection, 40 minutes must be met or exceeded.
		Roctavian		J3590	Unclassified biological	99222	Initial hospital inpatient or observation care, per day, for the evaluation and management of a patient, which requires a medically appropriate history and/ or examination and moderate level of medical decision-making. When using total time on the date of the encounter for code selection, 55 minutes must be met or exceeded. Initial hospital inpatient or observation care, per day, for the evaluation and manage-	
						99223 99234	ment of a patient, which requires a medically appropriate history and/or examination and high level of medical decision making. When using total time on the date of the encounter for code selection, 75 minutes must be met or exceeded. Hospital inpatient or observation care, for the evaluation and management of a patient including admission and discharge on the same date, which requires a medically appropriate history and/or examination and straightforward or low level of medical decisionmaking. When using total time on the date of the encounter for	
		Lyfgenia	D57	Sickle-cell disorders	C9399	Unclassified drugs or biologicals		code selection, 45 minutes must be met or exceeded.
			D57.0-D57.09	Hb-SS disease with crisis	J3490	Unclassified drugs	96413	Chemotherapy administration, intravenous infusion technique; up to 1 hour, single or initial substance/drug
			D57.1 D57.2-D57.219 D57.3	Sickle-cell disease without crisis Sickle-cell/Hb-C disease Sickle-cell trait	J3590 - J3394	Unclassified biologics Injection, lovotibeglogene autotemcel, per treatment (Eff. 7/1/2024)	96415	Chemotherapy administration, intravenous infusion technique; each additional hour
			D57.4-D57.459 D57.8-D57.819	Sickle-cell thalassemia Other sickle-cell disorders				
		Casgevy	D57 D57.0-D57.09	Sickle-cell disorders Hb-SS disease with crisis	C9399 J3490	Unclassified drugs or biologicals Unclassified drugs		Hematopoietic progenitor cell (HPC); autologous transplantation (Casgevy)
			D57.0-D57.09	Sickle-cell disease without crisis	30 100	Shokusamuu uruga		
			D57.2-D57.219	Sickle-cell/Hb-C disease	J3590		38241	
			D57.3 D57.4-D57.459	Sickle-cell trait Sickle-cell thalassemia		Unclassified biologics		
			D57.8-D57.819	Other sickle-cell disorders				
		Elevidys	G71.01	Duchenne or Becker muscular dystrophy	C9399 J3490	Unclassified drugs or biologicals Unclassified drugs	TBD	
		Lenmeldy	E75.25	Matachramatic laukadustranhy	J3590 J3590	Unclassified biologics		
		25lioldy	C90.0	Metachromatic leukodystrophy Multiple Myeloma and malignant plasma cell neoplasms	30090	Unclassified biologics Carvykti (Ciltacabtagene autoleucel),		
		Carvykti*	C90.00	Multiple Myeloma not having achieved remission	00055	up to 100 million autologous b-cell maturation antigen (BCMA) directed car-positive t cells, including leukaphere- sis and dose preparation procedures, per	0540T	Infusion of modified cells
			C90.01	Multiple myeloma in remission	Q2056			
		Abecma*	C90.02	Multiple myeloma in relapse	Q2055	therapeutic dose. Abecma (idecabtagene vicleucel), up to 460 million autologous B-cell maturation antigen (BCMA) directed CAR-positive T cells, including leukapheresis and dose preparation procedures, per therapeutic dose.	0540T	Infusion of modified cells
			C90.00	Multiple Myeloma and malignant plasma cell neoplasms Multiple Myeloma not having achieved remission Multiple myeloma in remission				
			C90.01	Multiple myeloma in relapse	-			
					J3590			
		Rethymic*	C90.02	Multiple myeloma in relapse	J3590 C9399	dose.		

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Resource for Spinraza CPT code: https://www.cms.gov/medicare-coverage-database/view/article.aspx?articleid=58579