Quick Reference Guide to High-Cost Medications and Therapies

July 2024

Working to Help Contain Costs Through Awareness

Awareness is essential to smart claims management. With new high-cost drugs and therapies entering the market regularly, it's important to know which are influencing claim costs most significantly when talking with self-funded clients, their plan administrators, and the stop loss carrier protecting their plans. Being on the same page helps to optimize results.

In this document, we share what HM Insurance Group (HM) views as having a noteworthy influence on catastrophic claim costs, as well as what we anticipate will make an impact in the future.

When costs are presented, keep in mind that medications and therapies are always patient-specific, particularly in terms of dosing levels that relate to size and weight. This means actual costs can vary. In addition, many drugs are FDA-approved for more than one disease state, requiring different regimens, strengths, and lengths of therapy based on diagnosis — all of which affect costs.

What follows is a cross-sectional view of some of the more frequently seen, highest cost medications identified, including gene therapies, in 2023 and 2024 as well as those in the pipeline.

Cost assumptions: Costs listed within this document are for the medications or therapies only. They are general and point-in-time. They reflect a dose for an "average" patient and do not take into consideration dosing variances based on a patient's size or weight. They also do not reflect any treatment expansions that may have been approved following this publication or the costs related to that expanded use. Costs shown are to be viewed as a basic guide.

Approved Gene Therapies >\$2 Million Per Treatment*

(All listed gene therapies are once-in-a-lifetime treatments.)

Drug/Therapy	Disease	Cost	
Beqvez™	Hemophilia B	\$3,500,000	
Casgevy™	Sickle Cell Disease, Beta Thalassemia	\$2,200,000	
Elevidys®	Duchenne Muscular Dystrophy	\$3,200,000	
Hemgenix®	Hemophilia B	\$3,500,000	
Lenmeldy™	Metachromatic Leukodystrophy	\$4,250,000	
Lyfgenia™	Sickle Cell Disease	\$3,100,000	
Rethymic™	Congenital Athymia	\$2,800,000	
Roctavian®	Hemophilia A	\$2,900,000	
Skysona®	Cerebral Adrenoleukodystrophy	\$3,000,000	
Zolgensma®	Spinal Muscular Atrophy (SMA)	\$2,125,000	
Zynteglo®	Beta Thalassemia	\$2,800,000	





Medications Exceeding \$1 Million Per Plan Year in Adults*

(Based on average adult. Not every medication may be used on an ongoing basis every time.)

Drug/Therapy	Disease	Cost	
Aldurazyme®	Hurler Disease (MPS Type I)	\$1,000,000	
Amondys 45®	Duchenne Muscular Dystrophy	\$2,000,000+	
Carbaglu®	N-acetylglutamate Synthase Enzyme Deficiency	\$3,500,000+	
Danyelza®	Neuroblastoma	\$2,000,000 – \$3,000,000	
Elaprase®	Hunter Syndrome (MPS Type II)	\$1,375,000	
Elzonris®	Blastic Plasmacytoid Dendritic Cell Neoplasm	\$2,800,000+	
Exondys 51®	Duchenne Muscular Dystrophy	\$2,500,000	
Folotyn®	Peripheral T-Cell Lymphoma	\$1,350,000	
Gamifant®	Primary Hemophagocytic Lymphohistiocytosis	\$650,000+/week	
Hemlibra®	Hemophilia A	\$1,000,000+	
Kanuma®	Lysosomal Acid Lipase Deficiency	\$1,275,000+	
Kimmtrak®	Uveal Melanoma	\$1,200,000	
Korlym®	Cushing Syndrome	\$1,100,000	
Lamzede®	Alpha-Mannosidosis	\$2,000,000+	
Luxturna®	Inherited Retinal Dystrophy	\$1,020,000	
Myalept®	Leptin Deficiency	\$2,050,000	
Procysbi®	Nephropathic Cystinosis	\$1,675,000	
Ravicti®	Urea Cycle Disorders	\$1,650,000	
Ryplazim®	Plasminogen Deficiency Type 1	\$2,400,000	
Scemblix®	Chronic Myelogenous/ Myeloid Leukemia	\$1,500,000+	
Strensiq®	Hypophosphatasia	\$2,000,000	
Veopoz™	CHAPLE Disease	\$1,800,000	
Vyondys 53®	Duchenne Muscular Dystrophy	\$2,300,000	
Xenpozyme®	Niemann-Pick Disease	\$2,750,000+	



\$500,000 to \$1 Million Medications Per Plan Year in Adults*

(Based on average adult. Not every medication may be used on an ongoing basis every time.)

Drug/Therapy	Disease	Cost	
Abecma®	Multiple Myeloma (cancer)	\$500,000+ (once)	
Adcetris®	Lymphomas (multiple)	\$38,000 – \$50,000/dose	
Advate®	Hemophilia A	\$75,000+/month	
Amtagvi®	Melanoma	\$620,000 (once)	
Amvuttra®	Polyneuropathy of Amyloidosis	\$143,000/dose	
BeneFix®	Hemophilia B	\$70,000/month	
Breyanzi®	Large B-Cell Lymphoma (cancer)	\$500,000+ (once)	
Brineura®	Batten Disease	\$75,000/month	
Carvykti®	Multiple Myeloma (cancer)	\$500,000+ (once)	
Cerezyme®	Gaucher Disease Type I	\$50,000/month	
Crysvita®	X-Linked Hypophosphatemia	\$45,000/month	
Evrysdi®	Spinal Muscular Atrophy	\$36,000/month	
Fabrazyme®	Fabry Disease	\$55,000/month	
Givlaari®	Porphyria	\$50,000/month	
Kymriah®	Leukemia/Lymphoma (multiple)	\$500,000+ (once)	
Nexviazyme®	Pompe Disease	\$800,000 - \$1,000,000	
Padcev®	Urothelial Cancer	\$72,000/ treatment cycle	
Ruconest®	Hereditary Angioedema	\$70,000 – \$140,000/month	
Rylaze®	Acute Lymphoblastic Leukemia/ Lymphoma	\$30,000 – \$50,000/dose	
Soliris®	Multiple	\$70,000/month	
Spinraza®	Spinal Muscular Atrophy	\$135,000 – \$150,000/dose	
Takhzyro®	Hereditary Angioedema	\$60,000/month	
Talvey®	Multiple Myeloma	\$55,000+/month	
Targretin®	Cutaneous T-Cell Lymphoma	\$72,000/month	
Tecartus®	Acute Lymphoblastic Leukemia/ Mantle Cell Lymphoma	\$500,000+ (once)	
Tecvayli®	Multiple Myeloma	\$50,000+/month	
Tegsedi®	Polyneuropathy	\$50,000/month	
Ultomiris®	Multiple	\$85,000/dose	
Vyjuvek™	Epidermolysis Bullosa	\$631,000/year	
Yervoy®	Multiple	\$40,000/dose	

Assessing Outcomes of Recent Developments

A number of groundbreaking therapies have recently hit the market, offering high hopes and high costs. While these do deliver new options for the management of life-threatening illnesses, they also come with extremely high price tags and additional potential risks for patients — both of which can significantly impact outcomes for self-funded employers. HM Insurance Group works to be a resource to our clients who want to gain more control of the costs associated with high-cost pharmaceutical therapies. See what we've learned.

Gene Therapies: What to Consider

Gene therapies come with very high costs. Things that can be done to assist in managing them include:

- Anticipating and identifying the potential risk prior to drug/ therapy availability.
- · Working with HM to determine if possible risk may be reduced.
- Notifying HM upon receipt of precertification/prior authorization requests.
- Implementing cost containment strategies specific to what that manufacturer offers.
- Knowing the availability of value-based contracts or outcomes-based contracts.
- Understanding that not all gene therapies are "curative" or "one and done" and working to ensure the client's plan document is reflective of such concepts.

Pipeline Therapies to Watch in 2024/Early 2025

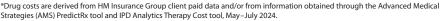
Drug/Therapy	Disease	Estimated Cost	When to Watch
PTC-AADC (eladocagene exuparvovec)	AADC Deficiency	\$3,000,000+ (once)	11/13/2024
RGX-121	Mucopolysaccharidosis II	\$3,000,000 (once)	First Half 2025
RP-L102	Fanconi Anemia	\$2,000,000 – \$3,000,000 (once)	First Half 2025
EB-101 (PZ-cel)	Dystrophic Epidermolysis Bullosa	\$1,000,000 – \$2,000,000 (once)	First Half 2025
DTX-401	Von Gierke Disease	\$1,500,000 – \$2,500,000 (once)	First Half 2025

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For more information, please contact **HMPharmacyServices@hmig.com**.



^{**}AM Best Company, September 2023

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