# **Fosaprepitant:**

# Emend®; Fosaprepitant Ψ; Focinvez Ψ (Intravenous)

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# I. Length of Authorization

Coverage is provided for 6 months and may be renewed.

# **II.** Dosing Limits

- A. Quantity Limit (max daily dose) [NDC Unit]:
  - 150 mg single-dose vial: 3 vials per 7 days
- B. Max Units (per dose and over time) [HCPCS Unit]:
  - 450 billable units (450 mg) per 7 days

# III. Initial Approval Criteria 1-3

Coverage is provided in the following conditions:

• Patient is at least 6 months of age; **AND** 

# Universal Criteria 1-7

• Patient is not taking pimozide concurrently; **AND** 

## Prevention of Chemotherapy-Induced Nausea and Vomiting (CINV) †

- Patient is receiving highly and/or moderately emetogenic chemotherapy (see HEC/MEC list below);
- Must be used in combination with a 5-HT<sub>3</sub> antagonist such as ondansetron, granisetron, palonosetron, etc.; **AND**
- Must be used in combination with a corticosteroid such as dexamethasone (Note: Only applicable to adult patients)

# **Highly Emetogenic Chemotherapy (HEC)**



Carboplatin	Carmustine	Cisplatin	Cyclophosphamide			
Dacarbazine	Doxorubicin	Epirubicin	Fam-trastuzumab deruxtecan-nxki			
Ifosfamide	Mechlorethamine	Melphalan ≥140 mg/m²	Sacituzumab govitecan- hziy			
Streptozocin						
	The following can be consider	dered HEC in certain patient	s			
Dactinomycin	Daunorubicin	Idarubicin	Irinotecan			
Methotrexate ≥250mg/m²	Oxaliplatin	Trabectedin				
	Moderately Emetoger	nic Chemotherapy (MEC)				
Aldesleukin >12-15 million IU/m²	Amifostine >300mg/m <sup>2</sup>	Bendamustine	Busulfan			
Clofarabine	Cytarabine >200mg/m <sup>2</sup>	Dinutuximab	Dual-drug liposomal encapsulation of cytarabine and daunorubicin			
Irinotecan Liposomal	Lurbinectedin	Melphalan <140 mg/m²	Naxitamab-gqgk			
Romidepsin	Temozolomide					
The following regimens can be considered HEC						
FOLFOX	FOLFIRI	FOLFIRINOX; FOLFOXIRI	AC (any anthracycline +			

 $\ \, \dagger \ \, \text{FDA-Approved Indication(s);} \ \, \textbf{$\updownarrow$ Compendia Recommended Indication(s);} \ \, \boldsymbol{\Phi} \ \, \text{Orphan Drug}$ 

# IV. Renewal Criteria 1-5

Coverage can be renewed based upon the following criteria:

- Patient continues to meet universal and other indication-specific relevant criteria such as concomitant therapy requirements (not including prerequisite therapy), performance status, etc. identified in section III; **AND**
- Disease response; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe hypersensitivity reactions, severe infusion site reactions, etc.

# V. Dosage/Administration 1-3

Indication	Dose
marcation	5030



Prevention of Chemotherapy-Induced Nausea Pediatric dosing: and Vomiting (CINV)

## Adult dosing:

Administer 150 mg intravenously (IV) over 20 to 30 minutes on Day 1

Age	Single-Day Chemotherapy Regimen	Single or Multi-Day Chemotherapy Regimens (oral formulations may be given as an alternative on Days 2-3)	
12 to 17 years	150 mg IV on Day 1	115 mg IV on Day 1, then 80 mg IV/PO on Days 2-3	
2 to < 12 years	4 mg/kg (maximum dose 150 mg) IV on Day 1	3 mg/kg (maximum dose 115 mg) on Day 1, then 2 mg/kg (maximum dose 80 mg) IV/PO on Days 2-3	
6 months to $<2$ years (patient $\ge 6 \text{ kg}$ )	5 mg/kg (maximum dose 150 mg) IV on Day 1		

<sup>\*</sup>Infusion should be completed 30 minutes prior to chemotherapy.

#### VI. **Billing Code/Availability Information**

## HCPCS Code:

- J1453 Injection, fosaprepitant, 1 mg; 1 billable unit = 1 mg (Emend Only)
- J1456 Injection, fosaprepitant (teva), not therapeutically equivalent to J1453, 1 mg; 1 billable unit = 1 mg  $\Psi$
- J3490 Unclassified drugs (Focinvez Only) **Ψ**

## NDC:

- Emend\* 150 mg powder for injection, single-dose vial: 00006-3061-xx
- Fosaprepitant 150 mg powder for injection, single-dose vial: 00591-4385-xx Ψ
- Focinvez 150 mg/50 mL (3 mg/mL) ready-to-use injection solution in a single-dose vial: 82243-1001-xx **Ψ** 
  - \*Available as a multi-sourced generic.
- \$\mathbf{P}\$ Designated products approved by the FDA as a 505(b)(2) NDA of the innovator product. These products are not rated as therapeutically equivalent to their reference listed drug in the Food and Drug Administration's (FDA) Orange Book and are therefore considered single source products based on the statutory definition of "single source drug" in section 1847A(c)(6) of the Act. For a complete list of all approved 505(b)(2) NDA products please reference the latest edition of the Orange Book:

Approved Drug Products with Therapeutic Equivalence Evaluations / Orange Book / FDA

#### VII. References

- 1. Emend [package insert]. Whitehouse Station, NJ; Merck & Co., Inc.; May 2022. Accessed March 2023.
- 2. Fosaprepitant [package insert]. North Wales, PA; Teva Pharmaceuticals USA, Inc.; September 2019. Accessed March 2023.



- 3. Focinvez [package insert]. North Brunswick, NJ; Spes Pharm., Inc.; August 2023. Accessed August 2023.
- 4. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) for Fosaprepitant. National Comprehensive Cancer Network, 2023. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. March 2023.
- 5. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Antiemesis. Version 1.2023. National Comprehensive Cancer Network, 2023. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed March 2023.
- 6. Roila F, Molassiotis A, Herrstedt J, et al. MASCC and ESMO Consensus Guidelines for the Prevention of Chemotherapy and Radiotherapy-Induced Nausea and Vomiting: ESMO Clinical Practice Guidelines. Ann Oncol (2016) 27 (suppl 5): v119-v133.
- 7. Hesketh PJ, Kris MG, Basch E, et al. Antiemetics: American Society of Clinical Oncology Guideline Update. J Clin Oncol. 2020 Aug 20;38(24):2782-2797. doi: 10.1200/JCO.20.01296.

# Appendix 1 – Covered Diagnosis Codes

ICD-10	ICD-10 Description	
R11.0	Nausea	
R11.10	Vomiting, unspecified	
R11.11	Vomiting without nausea	
R11.12	Projectile vomiting	
R11.2	Nausea with vomiting, unspecified	
T45.1X5A	Adverse effect of antineoplastic and immunosuppressive drugs, initial encounter	
T45.1X5D	Adverse effect of antineoplastic and immunosuppressive drugs, subsequent encounter	
T45.1X5S	Adverse effect of antineoplastic and immunosuppressive drugs, sequela	
T45.95XA	Adverse effect of unspecified primarily systemic and hematological agent, initial encounter	
T45.95XD	Adverse effect of unspecified primarily systemic and hematological agent, subsequent encounter	
T45.95XS	Adverse effect of unspecified primarily systemic and hematological agent, sequela	
T50.905A	Adverse effect of unspecified drugs, medicaments and biological substances, initial encounter	
T50.905D	Adverse effect of unspecified drugs, medicaments and biological substances, subsequent encounter	
T50.905S	Adverse effect of unspecified drugs, medicaments and biological substances, sequela	



ICD-10	ICD-10 Description
Z51.11	Encounter for antineoplastic chemotherapy
Z51.12	Encounter for antineoplastic immunotherapy

# Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD), Local Coverage Determinations (LCDs) and Local Coverage Articles (LCAs) may exist and compliance with these policies is required where applicable. They can be found at: <a href="https://www.cms.gov/medicare-coverage-database/search.aspx">https://www.cms.gov/medicare-coverage-database/search.aspx</a>. Additional indications may be covered at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCD/LCA): N/A

	Medicare Part B Administrative Contractor (MAC) Jurisdictions				
Jurisdiction	Applicable State/US Territory	Contractor			
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC			
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC			
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)			
6	MN, WI, IL	National Government Services, Inc. (NGS)			
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.			
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)			
N (9)	FL, PR, VI	First Coast Service Options, Inc.			
J (10)	TN, GA, AL	Palmetto GBA, LLC			
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA, LLC			
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.			
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)			
15	KY, OH	CGS Administrators, LLC			