



# Cinvanti® (aprepitant) (Intravenous)

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10/2023, 04/2024, 04/2025

## I. Length of Authorization

Coverage is provided for 6 months and may be renewed.

## **II.** Dosing Limits

#### Max Units (per dose and over time) [HCPCS Unit]:

520 billable units per 28 days

## III. Initial Approval Criteria <sup>1</sup>

Coverage is provided in the following conditions:

Patient is at least 18 years of age; AND

#### Universal Criteria 1

- Patient is not taking pimozide concurrently; AND
- Patient must have failed or experienced intolerable side effects to a fosaprepitant product (e.g., Emend, Fosaprepitant, Focinvez, etc.) prior to consideration of Cinvanti; AND

#### Prevention of Chemotherapy induced Nausea and vomiting (CINV) † ‡ 1-5

- Patient is receiving highly and/or moderately emetogenic anticancer chemotherapy (HEC\*/MEC\*\*); AND
  - Used in combination with a 5-HT3 antagonist (e.g., ondansetron, granisetron, palonosetron, etc.);
  - Used in combination with a corticosteroid such as dexamethasone; AND
  - Used with or without olanzapine (applies to HEC only); OR
- Patient experienced emesis during a previous cycle of anticancer chemotherapy with a 3-drug regimen (olanzapine or NK-1 receptor antagonist-containing regimen);
  - Used in combination with olanzapine, 5HT3 antagonist and dexamethasone as a component of a 4-drug regimen if not previously given

#### \*Highly emetogenic chemotherapy (HEC):

	Highly Emetogenic C	Chemotherapy (HEC) <sup>3</sup>	
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 considered HEC in certain patients <sup>3</sup>			
Dactinomycin	Daunorubicin	Idarubicin	Irinotecan
Methotrexate ≥250mg/m <sup>2</sup>	Oxaliplatin	Trabectedin	
The following regimens can be considered HEC <sup>3</sup>			
FOLFOX	FOLFIRI	FOLFIRINOX; FOLFOXIRI	AC (any anthracycline + cyclophosphamide)

## \*\*Moderately emetogenic chemotherapy (MEC):

Moderately Emetogenic Chemotherapy (MEC) <sup>3</sup>				
Aldesleukin >12-15 million IU/m²	Amifostine >300 mg/m <sup>2</sup>	Bendamustine	Busulfan	
Clofarabine	Cytarabine >200 mg/m²	Dinutuximab	Dual-drug liposomal encapsulation of cytarabine and daunorubicin	
Irinotecan (liposomal)	Lurbinectedin	Melphalan <140 mg/m <sup>2</sup>	Mirvetuximab soravtansine-gynx	
Naxitamab-gqgk	Romidepsin	Temozolomide		

† FDA Approved Indication(s); ‡ Compendia Recommended Indication(s); ♠ Orphan Drug

## IV. Renewal Criteria <sup>1</sup>

Coverage can be renewed based upon the following criteria:

- Patient continues to meet the universal and other indication-specific relevant criteria such as concomitant therapy requirements (not including prerequisite therapy), performance status, etc. identified in section III; AND
- Beneficial response as evidenced by reduction in nausea and/or vomiting; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe hypersensitivity reactions (including anaphylaxis), etc.

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## V. Dosage/Administration <sup>1</sup>

Indication	Dose
Prevention of chemotherapy- induced nausea and vomiting (CINV)	Administer as either a 30 minute infusion or 2 minute injection  HEC (Single Dose Regimen)  130 mg intravenously (IV) on Day 1 approximately 30 minutes prior to chemotherapy  MEC (3-Day Regimen with oral aprepitant)  100 mg IV on Day 1 approximately 30 minutes prior to chemotherapy followed by oral aprepitant (80mg) on Days 2 and 3.
	MEC (Single-dose Regimen)  – 130 mg IV on Day 1 approximately 30 minutes prior to chemotherapy

## VI. Billing Code/Availability Information

#### **HCPCS Code:**

J0185 – Injection, aprepitant, 1 mg; 1 billable unit = 1 mg

#### NDC:

Cinvanti 130 mg/18 mL injectable emulsion single-dose vial: 47426-0201-xx

#### VII. References

- 1. Cinvanti [package insert]. San Diego, CA; Heron Therapeutics, Inc.; March 2024. Accessed February 2025.
- 2. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) for Aprepitant. National Comprehensive Cancer Network, 2025. 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. February 2025.
- 3. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Antiemesis. Version 2.2024. National Comprehensive Cancer Network, 2025. 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 February 2025.
- 4. 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.
- 5. 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
TEO 00ED	Adverse effect of unspecified drugs, medicaments and biological substances, subsequent
T50.905D encounter	
T50.905S	Adverse effect of unspecified drugs, medicaments and biological substances, sequela
Z51.11	Encounter for antineoplastic chemotherapy
Z51.12	Encounter for antineoplastic immunotherapy

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

The preceding information is intended for non-Medicare coverage determinations. 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 Determinations (NCDs) and/or Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. Local Coverage Articles (LCAs) may also exist for claims payment purposes or to clarify benefit eligibility under Part B for drugs which may be self-administered. The following link may be used to search for NCD, LCD, or LCA documents <a href="https://www.cms.gov/medicare-coverage-database/search.aspx">https://www.cms.gov/medicare-coverage-database/search.aspx</a>. Additional indications, including any preceding information, may be applied 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)		







Medicare Part B Administrative Contractor (MAC) Jurisdictions			
Jurisdiction	Applicable State/US Territory	Contractor	
N (9)	FL, PR, VI	First Coast Service Options, Inc.	
J (10)	TN, GA, AL	Palmetto GBA	
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA	
, ,	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	

