Clinical Policy: Total Parenteral Nutrition and Intradialytic Parenteral Nutrition
Reference Number: CP.PHAR.205
Effective Date: 05/16
Last Review Date: 05/17

See Important Reminder at the end of this policy for important regulatory and legal information.

Description
Parenteral nutrition (PN) is the intravenous administration of an artificially prepared solution of nutrients that bypasses the gastrointestinal tract and that meets the nutritional requirements of a patient. PN is necessary when enteral nutrition is incapable of meeting the needs of the patient’s gastrointestinal tract. This policy describes the medical necessity requirements for two types of PN, (A) total parenteral nutrition (TPN), in which all of the necessary macronutrients and micronutrients are supplied to the patient, and (B) intradialytic parenteral nutrition (IDPN), in which nutrition is supplied to end-stage renal disease (ESRD) patients undergoing dialysis as an alternative to regularly scheduled TPN.

*Please see CP.MP.34 Hyperemesis Gravidarum Treatment regarding use of TPN in pregnancy.

Policy/Criteria
I. It is the policy of health plans affiliated with Centene Corporation® that the following are medically necessary for members when meeting the associated indications:

A. Total Parenteral Nutrition, when all the following criteria are met:
   1. Documentation of failure of enteral (i.e. oral or tube feeding) nutrition, as shown by any of the following:
      a. Weight loss > 10% of ideal body weight in 3 months, or > 20% of usual body weight;
      b. Total protein < 6 g/dL in the past 4 weeks;
      c. Serum albumin < 3.4 g/dL in the past 4 weeks;
   2. Evidence of structural or functional bowel disease that makes oral or tube feedings inappropriate, or a condition in which the gastrointestinal tract is non-functioning for a period of time, including, but not necessarily limited to, any of the following:
      a. Crohn’s disease;
      b. Short bowel syndrome;
      c. Single or multiple fistulae (entercolic, entervesical, or enterocutaneous);
      d. CNS disorder resulting in swallowing difficulties and high risk of aspiration;
      e. Obstructing stricture;
      f. Motility disorder;
      g. Newborn anomalies of the gastrointestinal tract which prevent or contraindicate oral feedings such as tracheoesophageal fistula, gastroschisis, omphalocele, or massive intestinal atresia;
      h. Infants and young children who fail to thrive due to cardiac or respiratory disease, short bowel syndrome, malabsorption or chronic idiopathic diarrhea;
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i. Prolonged paralytic ileus following a major surgical procedure or multiple injuries.

Initial approval duration for TPN is for 3 months. Continued approval duration is 6 months, given that the member has no evidence of unacceptable complications from treatment, and documentation supports positive response to therapy.

B. Intradialytic Parenteral Nutrition, when all the following criteria are met:
   1. Meets TPN criteria in section A;
   2. Patient has ESRD;
   3. Patient is undergoing hemodialysis;
   4. IDPN is offered as an alternative to regularly scheduled TPN.

Initial approval duration for IDPN is for 3 months. Continued approval duration is 6 months, given that the member has no evidence of unacceptable complications from treatment and documentation supports positive response to therapy.

II. It is the policy of health plans affiliated with Centene Corporation® that the following indications are experimental/investigational:

A. Total Parenteral Nutrition:
   1. Children who were previously well nourished or mildly malnourished, who are undergoing oncologic treatment associated with a low nutrition risk (e.g. less advanced disease, less intense cancer treatments, advanced disease in remission during maintenance treatment);
   2. Patients with advanced cancer whose malignancy is documented as unresponsive to chemotherapy or radiation therapy;
   3. Patients for whom liver transplantation is not feasible and whose prognosis will not change in spite of TPN therapy;

B. Intradialytic Parenteral Nutrition, when any of the following criteria are met:
   1. IDPN treatments offered in addition to regularly scheduled infusions of TPN;
   2. IDPN treatments in patients who are suffering from acute kidney injury and who do not have ESRD.

Background

Total Parenteral Nutrition

TPN is the delivery of macronutrients (i.e. proteins, fats, and carbohydrates) and micronutrients (i.e. vitamins, minerals, and trace elements) intravenously. TPN is indicated in situations for which the gastrointestinal tract is incapable of digesting nutrients through enteral (oral or feeding tube) nutrition. Short-term TPN is delivered peripherally through a subclavian, internal jugular, or a femoral central venous catheter while long-term TPN requires a tunneled central venous catheter, such as a Hickman, Groshong catheter, or an implanted infusion port.¹

Some of the advantages of TPN include the ease of administration, easier correction of fluid and electrolyte disturbances, and the ability to manage nutrition in the setting of mucositis. However, some disadvantages of TPN include, catheter-associated infections, fluid overload,
hyperglycemia, catheter-associated thrombosis, hepatic thrombosis, hepatic dysfunction, blood electrolyte abnormalities, and enterocyte atrophy.²

**American Gastroenterological Association**
Long-term PN is indicated for patients with prolonged gastrointestinal tract failure that prevents the absorption of adequate nutrients to sustain life.⁷

**Intradialytic Parenteral Nutrition**
Malnutrition presents an ongoing concern with patients receiving chronic hemodialysis or peritoneal dialysis. Malnutrition can occur in between 20-70% of patients, and there is a positive association with length of time on dialysis and increasing decline in nutritional parameters. IDPN is delivered during dialysis for patients who continue to lose weight or have very low serum albumin (< 3.4 g/dL) despite oral supplements and for those with severe gastroparesis who may be unable to tolerate oral supplements. However, IDPN only provides 70% of the nutrients to the patient because of the loss into the dialysate.³

Several societies published position guidelines that favor the use of IDPN in specific situations. **American Society for Parenteral and Enteral Nutrition**
IDPN should be reserved for patients that are incapable of meeting their nutritional needs orally and who are not candidates for enteral nutrition or TPN because of gastrointestinal intolerance, venous access problems, or other reasons.⁴

**European Society for Clinical Nutrition and Metabolism**
IDPN is indicated in undernourished patients undergoing hemodialysis with poor compliance to oral nutritional supplements and not requiring TPN.⁵

**National Kidney Foundation/Dialysis Outcomes Quality Initiative**
These guidelines indicates that IDPN is appropriate if an intervention is combined with oral nutritional supplements to help meet the dietary requirements of patients.⁶

**Coding Implications**
This clinical policy references Current Procedural Terminology (CPT®). CPT® is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2016, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

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HCPCS Codes | Description
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B4164 – B5200 | Parenteral nutrition solutions and supplies
B9004 | Parenteral nutrition infusion pump, portable
B9006 | Parenteral nutrition infusion pump, stationary
B9998 | NOC for enteral supplies
B9999 | NOC for parenteral supplies

ICD-10-CM Diagnosis Codes that Support Coverage Criteria

| ICD-10-CM Code | Description |
--- | ---
N18.6 | End stage renal disease
Z99.2 | Dependence on renal dialysis

| Reviews, Revisions, and Approvals | Date | Approval Date |
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Policy developed. | 04/16 | 05/16 |
Added ICD-10 codes | 08/16 | 08/16 |
References reviewed and updated. Added 3 month time period for weight loss >10% of ideal body weight. Added that protein and albumin labs should be from last 4 weeks. | 05/17 | 05/17 |

References

Important Reminder
This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

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Note: For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

Note: For Medicare members, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs, LCDs, and Medicare Coverage Articles should be reviewed prior to applying the criteria set forth in this clinical policy. Refer to the CMS website at for additional information.