TRICARE POLICY MANUAL 6010.54-M, AUGUST 1, 2002

ADMINISTRATIVE

Chapter 1
Section 3.1

RARE DISEASES

ISSUE DATE: May 18, 1994

AUTHORITY: 32 CFR 199.2(b) and 32 CFR 199.4(g)(15)

I. DESCRIPTION

TRICARE defines a rare disease as any disease or condition that affects less than 200,000 persons in the United States.

II. POLICY

- A. Coverage for treatment of rare diseases may be considered on a case-by-case basis. Case-by-case review is not required for drugs, devices, medical treatments, and procedures that have already been established as safe and effective for treatment of rare diseases.
- B. In reviewing the case, any or all of the following sources may be used to determine if the proposed benefit is considered safe and effective.
 - 1. Trials published in refereed medical literature.
 - 2. Formal technology assessments.
 - 3. National medical policy organization positions.
 - 4. National professional associations.
 - 5. National expert opinion organizations.
- C. If case review indicates that the proposed benefit for a rare disease is safe and effective for that disease, benefits may be allowed. If benefits are denied, an appropriate appealing party may request an appeal.
- D. Off-label use of rituximab may be considered for cost-sharing for the treatment of recurrent nodular CD20 positive lymphocyte predominant Hodgkin's disease. The effective date is January 1, 2003.
- E. Off-label use of rituximab may be considered for cost-sharing in reducing proteinuria for the treatment of Immunoglobulin A (IgA) nephropathy (proliferative glomerulonephritis). The effective date is May 1, 2007.

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- F. Effective May 13, 2009, Intraperitoneal Hyperthermic Chemotherapy (IPHC) (CPT¹ procedure codes 77600, 77605, and 96445) in conjunction with cytoreductive surgery or peritonectomy for treatment of pseudomyxoma peritonei resulting from appendiceal carcinoma may be covered on a case-by-case basis for adult patients when all of the following criteria are met:
 - 1. There is no evidence of distant metastasis.
 - 2. There is evidence of low histological aggressiveness of the disease.
 - 3. The patient has not undergone preoperative systemic chemotherapy.
 - 4. The patient's condition does not preclude major surgery.
- 5. The chemotherapeutic agents used are Mitomycin C, Cisplatin (also known as Cisplatinum), or Fluorouracil.
- G. External Infusion Pumps (EIPs) for insulin may be considered for cost-sharing when the diagnosis is Cystic Fibrosis-Related Diabetes (CFRD) with fasting hyperglycemia. See Chapter 8, Section 2.3 for policy regarding EIPs. Effective January 21, 2009.
- H. Post-operative proton beam radiosurgery/radiotherapy (CPT¹ procedures codes 77520, 77522, 77523, and 77525) may be considered for cost-sharing when the diagnosis is sacral chordoma. See Chapter 5, Section 3.1 for policy regarding proton beam radiosurgery/radiotherapy.
- l. Extracorporeal photopheresis (CPT¹ procedure code 36522) may be considered for cost-sharing when the diagnosis is Bronchiolitis Obliterans Syndrome (BOS) that is refractory to immunosuppressive drug treatment. See Chapter 4, Section 9.2 for policy regarding photopheresis.

III. EXCLUSIONS

- A. Intracranial angioplasty with stenting (CPT¹ procedure code 61635) of the venous sinuses for treatment of pseudotumor cerebri (also known as idiopathic intracranial hypertension and benign intracranial hypertension) is unproven.
 - B. Off-label use of rituximab for pediatric Immunoglobulin A (IgA) is unproven.
- C. Proton Beam Therapy (PBT)/radiosurgery/radiotherapy for the treatment of thymoma is unproven.

- END -

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