Field Name	Field Description
Prior Authorization	Dendritic Cell Tumor Peptide Immunotherapy
Group Description	Dendritic Cen Tumor Teptide immunotnerapy
Drugs	Provenge (sipuleucel-T)
Covered Uses	Medically accepted indications are defined using the following
	sources: the Food and Drug Administration (FDA), Micromedex,
	American Hospital Formulary Service (AHFS), United States
	Pharmacopeia Drug Information for the Healthcare Professional
	(USP DI), the Drug Package Insert (PPI), or disease state specific
	standard of care guidelines.
Exclusion Criteria	Small cell/neuroendocrine prostate cancer
Required Medical	See "Other Criteria"
Information	
Age Restrictions	See "Other Criteria"
Prescriber	Prescriber must be an oncologist or urologist
Restrictions	
Coverage Duration	<u>3 doses per lifetime</u>
Other Criteria	**Drug is being requested through the member's medical benefit**
	Initial Authorization: • Metastatic castrate resistant (hormone-refractory) prostate cancer (mCRPC) (consistent with medical chart history) • Evidenced by soft tissue and/or bony metastases • Patient does NOT have • M0CRPC (defined as CRPC whose only evidence of disseminated disease is an elevated serum PSA) is not authorized • Visceral metastases (e.g. liver, lung, adrenal, peritoneal, brain) • Patient is not currently being treated with systemic immunosuppressants (e.g. chemotherapy, corticosteroids) or, if the patient is being treated with immunosuppressants, the prescriber has provided a valid medical reason for combination therapy • Eastern Cooperative Oncology Group (ECOG) score 0-1 • Serum testosterone <50 ng/dL (e.g. castration levels of testosterone) • Predicted survival of at least six months
Revision/Review Date 5/2022	 <u>Reauthorization:</u> <u>Treatment exceeding 3 doses per lifetime will not be</u> <u>authorized</u>

Medical Director/clinical reviewer must override criteria when, in his/her
professional judgement, the requested item is medically necessary.