

PSMA PET NEW MEDICARE ITEM

From 1 July 2022, two new items will be introduced for prostate-specific membrane antigen (PSMA) PET study, for patients with prostate cancer.

ITEM 61563 – INITIAL STAGING

The specialist or consultant physician is to record in the clinical notes and the request that the patient:

- Has intermediate to high-risk prostate adenocarcinoma (as defined below)
- ▶ Has previously been untreated; AND
- **▷** Is considered suitable for locoregional therapy with curative intent.

Benefits are payable once in a patient's lifetime.

- Patients with intermediate risk prostate adenocarcinoma can be defined as having at least one of the following risk factors in the absence of any high-risk features: PSA of 10-20 ng/ml, or Gleason score of 7 or International Society of Urological Pathology (ISUP) grade group 2 or 3, or Stage T2b.
- Patients with high-risk prostate adenocarcinoma can be defined as having at least one of the following risk factors: PSA >20 ng/ml, or Gleason score >7 or ISUP grade group 4 or 5, or Stage T2c or ≥ T3.

ITEM 61564 - RESTAGING

The specialist or consultant physician is to record in the clinical notes and the request that the patient:

- Has intermediate to high-risk prostate adenocarcinoma; AND
- ► Has undergone prior locoregional therapy and is considered suitable for further locoregional therapy.

Benefits are payable for a maximum of two services in a patient's lifetime.

This item can be claimed by patients with:

- ► A PSA increase of 2ng/ml above the nadir after radiation therapy; OR
- ▶ Failure of PSA levels to fall to undetectable levels; OR
- ▶ Rising PSA serum after a radical prostatectomy.



For more information on billing and PET/CT locations, visit our online Billing Guide **qldxray.com.au/billingguide**