



PET/CT Imaging Request

Patient Identification

Or ID Sticker

Surname:

First name:

Date of Birth: / /

Address:

Phone No: (h)

(m)

(w)

PET scan results required by: / /

Patient Information

• Patient status at present OP **or** IP

If IP, where?

• Diabetic No IDDM NIDDM

• Is patient claustrophobic? Yes **or** No

PET Low dose non diagnostic CT included

FDG DOTATATE PSMA FET

NaF Other

Clinical details

Clinical Indication

Staging Restaging Other

Site of disease Histopath

Additional Diagnostic Imaging Required

CT Brain Neck Chest Abdomen/Pelvis Extremities

MRI

Other (please specify modality and region e.g. Ultrasound pelvis)

Recent Comparison Imaging (Please attach results)

CT Date/where:

MRI Date/where:

Please ensure patient brings films with them for their appointment.

Recent biopsy/scope (Please attach results)

Biopsy Date/where:

Scope Date/where:

Referring Consultant/Specialist

(Must be Specialist Referred for MC Rebate)

Doctor's name:

Provider No:

Phone contact:

Signature: / /
Date

Address for Films & Report:

Fax (if required):

Rare and Uncommon Cancer Types

- **61614 (Nov 1, 2024)** Whole body FDG PET study for the evaluation of suspected residual, metastatic or recurrent cancer for a patient who is undergoing or suitable for active therapy.
- **61612** Whole body FDG PET study for the initial staging of eligible cancer types, for a patient who is considered suitable for active therapy, if: the eligible cancer type is a rare or uncommon cancer and a typically FDG-avid cancer; and there is at least a 10% likelihood that the PET study result will inform a significant change in management for the patient. Applicable once per cancer diagnosis.

PSMA Prostate Ca

- **61563** Whole body PSMA PET study performed for the initial staging of intermediate to high-risk prostate adenocarcinoma, for a previously untreated patient who is considered suitable for locoregional therapy with curative intent.
- **61564** Whole body PSMA PET study performed for the restaging of recurrent prostate adenocarcinoma, for a patient who has undergone prior loco regional therapy and is considered suitable for further locoregional therapy to determine appropriate therapeutic pathways and timing of treatment initiation. 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.

Alzheimer's Disease

- **61560** Performed for diagnosis if clinical evaluation of the patient by a specialist, or in consultation with a specialist, is equivocal, and the patient has not had this service performed for diagnosis or management in the previous 12months. Applicable not more than 3 times per lifetime.

Breast

- **61524** Whole body 18F-FDG PET study performed for the staging of locally advanced (Stage III) breast cancer in a patient considered potentially suitable for active therapy.
- **61525** Whole body 18F-FDG PET study performed for the evaluation of suspected metastatic or suspected locally or regionally recurrent breast carcinoma in a patient considered suitable for active therapy.

Lymphoma

- **61620** Initial staging of newly diagnosed or previously untreated Hodgkin's or non-Hodgkin's lymphoma.
- **61622** Assess response to first line therapy either during treatment or within three months of completing definitive first line treatment for Hodgkin's or non-Hodgkin's lymphoma.
- **61628** Restaging following confirmation of recurrence of Hodgkin's or non-Hodgkin's lymphoma.
- **61632** Assess response to second-line chemotherapy when stem cell transplantation is being considered, for Hodgkin's or non-Hodgkin's lymphoma.

Brain

- **61538** Evaluation of suspected residual or recurrent malignant brain tumour based on anatomical imaging findings, after definitive therapy (or during ongoing chemotherapy) in patients who are considered suitable for further active therapy.
- **61559** Study of the brain, performed for the evaluation of refractory epilepsy which is being evaluated for surgery.

Head and neck

- **61598** Staging of biopsy-proven newly diagnosed or recurrent head and neck cancer.
- **61604** Evaluation of patients with suspected residual head and neck cancer after definitive treatment, and who are suitable for active therapy.

Metastatic SCC unknown primary

- **61610** Evaluation of metastatic squamous cell carcinoma of unknown primary site involving cervical nodes.

Lung

- **61523** Evaluation of a solitary pulmonary nodule where the lesion is considered unsuitable for transthoracic fine needle aspiration biopsy, or for which an attempt at pathological characterisation has failed.
- **61529** Staging of proven non-small cell lung cancer, where curative surgery or radiotherapy is planned.

Upper GIT

- **61577** Staging of proven oesophageal or GEJ carcinoma, in patients considered suitable for active therapy.

Lower GIT

- **61541** Evaluation (following initial therapy) of suspected residual, metastatic or recurrent colorectal carcinoma in patients considered suitable for active therapy.

Gynaecological

- **61571** Primary staging of patients with histologically proven carcinoma of the uterine cervix, at FIGO stage IB2 or greater, prior to planned radical radiation therapy or combined modality therapy with curative intent.
- **61575** Staging of patients with confirmed local recurrence of carcinoma of the uterine cervix considered suitable for salvage pelvic chemoradiotherapy or pelvic exenteration with curative intent.
- **61565** Evaluation (following initial therapy) of suspected residual, metastatic or recurrent ovarian carcinoma in patients considered suitable for active therapy.

Melanoma

- **61553** Evaluation (following initial therapy) of suspected metastatic or recurrent malignant melanoma in patients considered suitable for active therapy.

Sarcoma

- **61640** Initial staging of patients with biopsy-proven bone or soft tissue sarcoma (excluding gastrointestinal stromal tumour) considered by conventional staging to be potentially curable.
- **61646** Evaluation of patients with suspected residual or recurrent sarcoma (excluding gastrointestinal stromal tumour) after the initial course of definitive therapy to determine suitability for subsequent therapy with curative intent.

DOTATATE Whole body Ga-DOTA-peptide PET study

- **61647** Staging of a suspected gastro-entero-pancreatic (GEP) neuroendocrine tumour. Assessment of resectability of metastatic GEP neuroendocrine tumour.

MEDICARE NON-REBATEABLE INDICATIONS

- Brain FET-PET for primary or metastatic disease