PET FACILITY ID #:

REGISTRY CASE #:

PATIENT NAME:

Your patient had a PET scan on mm/dd/yyyy. [Date will automatically be filled.]

You previously indicated that the PET scan was done for **restaging of** cancer type [Will automatically be filled in from data supplied on Pre-PET form.] to assess for

* new osseous metastatic disease as a site of recurrence or
* progression of known osseous metastatic disease.  
  [Reason will automatically be filled in from data supplied on Pre-PET form.]
* After reviewing the PET report, please complete the following questions and return the form to the PET Facility.
* This form must be entered into the database within 30 days of the PET scan.

1. COMPARED TO YOUR PRE-PET ASSESSMENT, WHAT IS YOUR IMPRESSION OF THE EXTENT OF THE PATIENT'S CANCER?

* More extensive
* No change
* Less extensive

1. YOUR POST-PET WORKING CLINICAL STAGING IS: (SELECT *ONLY* ONE)

* No evidence of disease / In remission
* Low probability of local recurrence or metastases
* Local recurrence
* Metastatic (distant) with a single suspected site
* Metastatic (distant) with a multiple suspected sites

1. DID THE PET SCAN ENABLE YOUR PATIENT TO AVOID ANY
2. noninvasive diagnostic tests?

* Yes
* No

1. any invasive procedures?

* Yes
* No

1. IN LIGHT OF THE PET FINDINGS, WHICH ONE OF THE FOLLOWING ARE YOU PLANNING OR HAVE YOU ALREADY DONE AS THE NEXT STEP IN YOUR CURRENT MANAGEMENT STRATEGY? (check only one)

* Observation (with close follow-up)
* Additional Imaging
  + **If additional imaging is selected, please indicate which specific type of imaging you would order next.** *(check one)*
  + ❑ Plain radiographs
  + ❑ Body CT (neck, chest, and/or abdomen/pelvis)
  + ❑ Extremity CT
  + ❑ Body MRI (spine, neck, chest, and/or abdomen/pelvis)
  + ❑ Extremity MRI
  + ❑ FDG-PET
  + ❑ Other, specify:
* Tissue Biopsy (surgical, percutaneous, or endoscopic).

[Note: If concurrent biopsy and a surgical procedure are planned, then mark “treatment” below. ]

* Supportive care only (e.g., pain management, hospice care)
* Treatment for the Cancer   
    
  If treatment was selected, answer the questions below:
  1. Treatment Goal: (check one)
* Curative
* Palliative
  1. Treatment will be directed to: (check all that apply)
* Primary tumor and/or locoregional disease
* Non-osseous distant metastatic disease
* Osseous distant metastatic disease
  1. Type(s): (check all that apply)
* Surgery
* Radiation
* Chemotherapy (including biologic modifiers)
* Hormonal therapy
* Bisphosphonate therapy
* Immunotherapy (e.g., sipuleucel T (Provenge®) for prostate cancer)
* Radiopharmaceutical therapy (strontium-89, samarium-153, etc.)
* Other

Specify other treatment type:

1. I HAVE READ THE REFERRING PHYSICIAN INFORMATION STATEMENT AND:

* I DO give my consent for the inclusion of data collected for this patient in NOPR research.
* I DO NOT give my consent for the inclusion of data collected for this patient in NOPR research.

1. NAME OF PERSON SUBMITTING THIS FORM

First Name: Last Name: Date:

1. **PHYSICIAN ATTESTATION OF DATA ACCURACY**

By signing below I verify that, to the best of my knowledge, the information on this form is accurate.

Physician Signature: Date:

Printed Name of Physician:

**Thank you for your assistance.**

**PRA Disclosure Statement**

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0938-0968. The time required to complete this information collection is estimated to average five (5) minutes per response, including the time to review instructions, search existing data resources, gather the data needed, and complete and review the information collection. If you have comments concerning the accuracy of the time estimate(s) or suggestions for improving this form, please write to: CMS, 7500 Security Boulevard, Attn: PRA Reports Clearance Officer, Mail Stop C4-26-05, Baltimore, Maryland 21244-1850.