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 *initial staging of cancer type* [Cancer type 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

2. DID THE PET SCAN ENABLE YOUR PATIENT TO AVOID ANY

a. noninvasive diagnostic tests?

b. any invasive procedures?

- □ Yes
- 🗆 No

_ □ Yes

□ No

- 3. YOUR POST-PET WORKING CLINICAL SUMMARY STAGING IS? (You must check only one)
 - □ No evidence of disease / In remission
 - Localized only
 - □ Regional by direct extension
 - □ Metastatic (distant) with a single suspected site
 - □ Metastatic (distant) with multiple suspected sites
 - □ Unknown or uncertain

- 4. 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)
 - o D Plain radiographs
 - o Body CT (neck, chest, and/or abdomen/pelvis)
 - o 🛛 Extremity CT
 - o Body MRI (spine, neck, chest, and/or abdomen/pelvis)
 - o 🛛 Extremity MRI
 - o 🛛 FDG-PET
 - o 🖵 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:

- a. **Treatment Goal:** (check one)
 - □ Curative
 - □ Palliative
- b. **Treatment will be directed to:** (check all that apply)
 - Primary tumor and/or locoregional disease
 - □ Non-osseous distant metastatic disease
 - Osseous distant metastatic disease
- C. **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:_____

5. 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.

6. NAME OF PERSON SUBMITTING THIS FORM

First Name:	Last Name:	Date:	/	/
			/	/

7. 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

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