Post-Scan F-18 Fluoride PET Scan

	PET FACILITY ID #:				
REGISTRY CASE #:					
	PATIENT NAME:				
Yo	ur patient had a PET scan on mm/dd/yyyy. [Date	will automatically be filled.]			
	ou previously indicated that the PET scan was filled in from data supplied on Pre-PET fo	s done for <i>restaging of</i> cancer type [Will automatically orm.] to assess for			
	 new osseous metastatic disease as a sit progression of known osseous metastati [Reason will automatically be filled in 				
•		following questions and return the form to the PET Facility.			
1.	COMPARED TO YOUR PRE-PET ASSESS EXTENT OF THE PATIENT'S CANCER? More extensive No change Less extensive	SMENT, WHAT IS YOUR IMPRESSION OF THE			
2.	YOUR POST-PET WORKING CLINICAL S ☐ No evidence of disease / In remission ☐ Low probability of local recurrence or ☐ Local recurrence ☐ Metastatic (distant) with a single susp ☐ Metastatic (distant) with a multiple su	r metastases Dected site			
3.	DID THE PET SCAN ENABLE YOUR PATIENT TO AVOID ANY				
	a. noninvasive diagnostic tests?	a. any invasive procedures?			
	□ Yes	□ Yes			
	□ No	□ No			

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4.	4. IN LIGHT OF THE PET FINDINGS, WHICH ONE OF THE FOLLOWING ARE YOU PLANNIN OR HAVE YOU ALREADY DONE AS THE NEXT STEP IN YOUR CURRENT MANAGEMEN' STRATEGY? (check only one)					
	☐ Observation (with close follow-up)					
	□ Addi	itional Imaging				
	o If additional imaging is selected, please indicate which specific type of imaging you would order next. (check one)					
	。 O	☐ Plain radiographs				
	0	☐ Body CT (neck, chest, and/or abdomen/pelvis)				
	0	☐ Extremity CT				
o Body MRI (spine, neck, chest, and/or abdomen/pelvis)		☐ Body MRI (spine, neck, chest, and/or abdomen/pelvis)				
	0					
	0					
	0	☐ Other, specify:				
	□ Trea	[Note: If concurrent biopsy and a surgical procedure are planned, then mark "treatment" below.] portive care only (e.g., pain management, hospice care) tment for the Cancer atment was selected, answer the questions below:				
		reatment Goal: (check one)				
	_	Curative				
☐ Palliative						
b. Treatment will be directed to: (check all that apply)						
\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		- -				
		Non-osseous distant metastatic disease				
		Osseous distant metastatic disease				
	L	OSSEOUS distant metastatic disease				
	c. T	ype(s): (check all that apply)				
] Surgery				
] Radiation				
		Chemotherapy (including biologic modifiers)				
] Hormonal therapy				

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	☐ Bisphosphonate therap	У					
	$\ \square$ Immunotherapy (e.g., sipuleucel T (Provenge $^{ ext{@}}$) for prostate cancer)						
	☐ Radiopharmaceutical th	nerapy (strontium-89	9, 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 NOP						
	research.						
6.	NAME OF PERSON SUBMITTING THIS FORM						
	First Name: La	ast Name:		Date:	/	_/	
7.	PHYSICIAN ATTESTATION OF DA	ATA 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.						
	you ioi your accidantor						

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