

Registration Header *(Information in this box below may be pre-printed by some groups.)*

Coordinating Group Protocol No. _____	Coordinating Group Code _____
Protocol Title _____	
Patient Study ID _____	Participating Group Code _____
Patient Medical Record Number _____	Other Patient ID _____
Institution / Affiliate _____	Physician of Record _____

Protocol Administration

IRB/REB Approval Date	<input type="text"/>	Person Completing Form, Last Name _____					
Date Informed Consent Signed	<input type="text"/>	Person Completing Form, First Name _____					
Projected Start Date of Treatment	<input type="text"/>	Person Completing Form, Phone (____) _____					
Date of Registration	<input type="text"/>	Person Completing Form, Fax (____) _____					
	MM	DD	YYYY				

Patient Demographics / Pre-Treatment Characteristics

Patient Name <i>(initials acceptable)</i>	<input style="width:100%;" type="text"/>					
Patient Birth Date	<input type="text"/>	<input type="text"/>	<input type="text"/>	Patient Gender	<input type="checkbox"/> Male	<input type="checkbox"/> Female
	MM	DD	YYYY			
Patient Race <i>(check all that apply)</i> <i>(U.S. and Canada only)</i>	<input type="checkbox"/> White	<input type="checkbox"/> Black or African American	<input type="checkbox"/> Asian	<input type="checkbox"/> Native Hawaiian or other Pacific Islander	<input type="checkbox"/> American Indian or Alaska Native	<input type="checkbox"/> Unknown
Patient Ethnicity <i>(U.S. and Canada only)</i>	<input type="checkbox"/> Hispanic or Latino	<input type="checkbox"/> Not Hispanic or Latino	<input type="checkbox"/> Unknown			
Patient Social Security Number <i>(USA only)</i>	<input type="text"/>	<input type="text"/>	<input type="text"/>			
Patient ZIP Code <i>(USA)</i>	<input style="width:100%;" type="text"/>	Country of Residence <i>(if not USA)</i>	<input style="width:100%;" type="text"/>			
Patient Height <i>(cm)</i>	<input type="text"/>	Patient Weight <i>(kg)</i>	<input type="text"/>	Body Surface Area <i>(m²)</i>	<input type="text"/>	<input type="text"/>
Performance Status <i>(check one)</i>	<input type="checkbox"/> 0 = Fully active, able to carry on all pre-disease performance without restriction (Karnofsky 90 - 100) <input type="checkbox"/> 1 = Restricted in physically strenuous activity but ambulatory (K 70 - 80) <input type="checkbox"/> 2 = Ambulatory and capable of all selfcare but unable to carry out any work activities (K 50 - 60) <input type="checkbox"/> 3 = Capable of only limited selfcare, confined to bed or chair more than 50% of waking hours (K 30 - 40) <input type="checkbox"/> 4 = Completely disabled (K 10 - 20)					
Method of Payment <i>(check one)</i> <i>(USA only)</i>	<input type="checkbox"/> Private Insurance <input type="checkbox"/> Medicare and Private Insurance <input type="checkbox"/> Self pay (no insurance) <input type="checkbox"/> Medicare <input type="checkbox"/> Military or Veterans Sponsored NOS <input type="checkbox"/> No means of payment (no insurance) <input type="checkbox"/> Medicaid <input type="checkbox"/> Military Sponsored (including CHAMPUS & TRICARE) <input type="checkbox"/> Other <input type="checkbox"/> Medicaid and Medicare <input type="checkbox"/> Veterans Sponsored <input type="checkbox"/> Unknown					

Certification of Eligibility *(This section may be linked to a separate eligibility checklist at a group's discretion; it is provided to indicate that the investigator has reviewed all eligibility criteria.)*

In the opinion of the investigator, is the patient eligible?	<input type="checkbox"/> Yes <input type="checkbox"/> No <i>(if No, the patient should not be registered)</i>
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Protocol Design

Stratification Factors <i>(protocol specific)</i>	_____
_____	_____
Assigned Treatment Arm <i>(protocol specific)</i>	_____
_____	_____

Initial Patient Consent for Specimen Use (Optional – for specimen banking studies only)

Patient's Initial Consent given for specimen use for research on the patient's cancer? Yes No

Patient's Initial Consent given for specimen use for research unrelated to the patient's cancer? Yes No

Patient's Initial Consent given for further contact regarding specimen? Yes No

Date of Consent for Specimen Use

MM
DD
YYYY