Collection of this information is authorized under 21 CFR 312.57. This information is collected to ensure compliance with Food and Drug Administration (FDA) requirements for NCI as an IPÖ sponsor and that investigational agents are under the control and accounted for by competent authority. The information may be disclosed to researchers for investigational purposes, sponsors of clinical trials and their company collaborators, the applicable Institutional Review Board, NCI, FDA and Department of Health and Human Services. Submission of this information is voluntary, however, in order for you to conduct a study in accordance with relevant, current protocols, you must complete all fields.

Form Approved: OMB No. 0925-0613 Expires: 01-31-2025

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Investigational Agent Accountability Record Oral agents ONLY		National Cancer Institute CONTI Division of Cancer Treatment and Diagnosis CONTI Cancer Therapy Evaluation Program SATEL		PAGE NO. CONTROL RECORD SATELLITE RECORD		
Name of Institution:	Investigator	Name:			CTEP Investigator IDK	
Protocol Title:	NCI Protocol No:		Local Protocol No:	Dispensing Area:		
Agent Name:	Dose Form	and StrengthK		Bottle size (e.g., # tablets/bo	ottle):	

Line No. Date	Patient's Initials	Patient's ID No.	Dose	Quantity Dispensed or Received	Balance Forward	d Manufacturer and Lot No.	Recorder's Initials	Expiration Date (if available)	Date Patient Returned	Quantity Patient Returned	Recorder's Initials	
					Balance							
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