

## **Baseline Evaluation (Adult)**

Patient ID \_\_\_\_\_ - \_\_\_ ID \_\_\_ - \_\_\_\_ Date of Evaluation: DOEDATE

## SECTION I: ADVERSE EFFECTS

1. Does the patient currently have any of the following:

 Has the patient experienced any adverse events (reportable at the level of detail of an adverse event), since the last protocol visit? AE

□ Yes (Complete an Adverse Events form, if SAE complete the MedWatch form too) □ No

## SECTION II: CONCOMITANT MEDICATIONS

1. Has there been any change (start or stop) in prescription medications since the last protocol visit? CONMED

□ Yes □ No If Yes, update the Concomitant Medication Log

3. Is the patient currently taking vitamins or minerals? If Yes, (check all that apply)				Yes INo IUnknown MEDVIT				
	□ Multi-vitamin VITMULT	□ Vitamin D VITD	□ Vitamin E VITE	□ Folate VITFOL	□ Iron VITFE	□ Calcium VITCA	□ Other VITOTH	
SECTION III: STUDY MEDICATION (do not complete for Control patients) 1. Was counseling on adherence provided during visit? MATI						Confirm acceptable method of contraception, when applicable		
2. Was study drug (entecavir) dispensed?							J	

SECTION IV: PHYSICAL ASSESSMENT

1. Weight: WGT 1 Ibs. 2 kg WLBKG INot done

2. Blood pressure: **BPS** / **BPD** mmHg Not done

## SECTION V: BIOSPECIMENS

 1. Were samples obtained at this visit?
 □ Yes
 □ No
 BIOSPEC

 If Yes, (check all that apply):
 □ NIDDK repository
 □ Central lab
 □ Genetics
 □ Immunology study

 NIDDKR
 CLAB
 GEN
 IMM

Data collector initials: DCID

Date data collection completed (mm/dd/yy): DCM / DCD / DCY

If No, complete the Off Protocol form