Discontinuation of Treatment or Study



Patient ID _____ - ___ ID ___ - ____ Date Form Completed: DFCDATE

Instruction: Complete this form when the patient prematurely discontinues study medication, study participation or both.

- 1. Time period: 1 Treatment 2 Follow-up TIMEP
- 2. Is this a discontinuation in study medication and/or study participation? (check all that apply)

□ Study medication (complete Section I) SMED

□ Study participation (complete Section II) **SPART**

SECTION I: STUDY MEDICATION

1. Reason(s) for discontinuing study medication(s) (check all that apply):	
Withdrawal of informed consent RMWCONS	Hypersensitivity reaction RMHYPS
□ Neutropenia RMNEUT	Pulmonary function impairment RMPF
Hepatic decompensation RMHDC	□ Anemia RMANEM
Autoimmune hepatitis RMAUTO	□ Renal function impairment RMRF
Pregnancy RMPREG	Ophthalmologic disorder RMOPH
□ Psoriatic lesion RMPSOR	Mitochondrial toxicity RMMIT
Hypoglycemia, hyperglycemia or diabetes mellitus RMDIAB	Grade IV toxicity RMTOX4
Thyroid disorder/dysfunction RMTHYD	□ Virological non-response RMVNRSP
Decrease in bone mineral density RMBONE	Partial virological response RMPVRSP
Depression or other psychiatric or mood disorder RMPSY	□ Virological breakthrough RMVBRK
□ Adverse event other than those listed RMAE , specify	RMAES
□ Investigator discretion RMINV, explainRMINVS	
2. Date of last dose of tenofovir (mm/dd/yy): LDTM / LDTD / LDTY	
3. Date of last dose of peginterferon (mm/dd/yy): LDPM / LDPD / LDPY	
SECTION II: STUDY PARTICIPATION	

- 1. Reason(s) for discontinuing study participation (check all that apply):
 - □ Patient lost to follow-up **RSLFUP**
 - □ Withdrawal of informed consent **RSWCONS**
 - □ Patient on alternate therapy for HBV **RSHBVTX**

□ Investigator discretion RSINV, explain ______RSINVS_____

Other RSOTH, specify _____RSOTHS_____

- 2. Date of withdrawal (or date considered to be withdrawn) (mm/dd/yy): WDM / WDD/ WDY
- 3. Date of last contact (mm/dd/yy): LCM / LCD / LCY