

Dose Change

Patient ID	-	ID	_	
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Instructions:

Complete one line for each <u>prescribed</u> dose change (reduction or increase) in one or both study medications. Record the newly prescribed dose for one or both study medications.

If only one study medication has been changed, record the current dose of the other study medication.

The Date of Change is the date that the first changed dose is taken.

If a Dose Reduction, record the reason for the dose reduction, otherwise leave that column blank.

Date of Change (mm/dd/yy)	Tenofovir DF Prescribed Dose	Peginterferon Prescribed Dose	Reason for Dose Reduction (check all that apply)	System ID
DCHM/D/Y	1 □ 300 mg daily 2 □ 300 mg q 48 hrs 3 □ 300 mg q 72 hrs 4 □ 300 mg q 96 hrs 5 □ Discontinued TENDOSE	1 □ 180 μg 2 □ 135 μg 3 □ 90 μg 4 □ 45 μg 5 □ Discontinued PEGDOSE	 □ CrCl ≥ 30 - 50 ml/min DRCR50 □ CrCl < 30 ml/min DRCR30 □ Reduction in platelet count DRPLAT □ Reduction in neutrophil count DRNEUT □ Lab result (other than platelet or neutrophil) DRLAB □ Adverse reaction (complete AE form) DRAE □ Emotional symptoms DRSYMP □ Pregnant DRPREG □ Patient preference DRPATNT □ Other, DROTH specify: DROTHS 	SYSID
/	1 □ 300 mg daily 2 □ 300 mg q 48 hrs 3 □ 300 mg q 72 hrs 4 □ 300 mg q 96 hrs 5 □ Discontinued	1 □ 180 μg 2 □ 135 μg 3 □ 90 μg 4 □ 45 μg 5 □ Discontinued	 □ CrCl ≥ 30 - 50 ml/min □ CrCl < 30 ml/min □ Reduction in platelet count □ Reduction in neutrophil count □ Lab result (other than platelet or neutrophil) □ Adverse reaction (complete AE form) □ Emotional symptoms □ Pregnant □ Patient preference □ Other, specify: 	
	1 □ 300 mg daily 2 □ 300 mg q 48 hrs 3 □ 300 mg q 72 hrs 4 □ 300 mg q 96 hrs 5 □ Discontinued	1 □ 180 µg 2 □ 135 µg 3 □ 90 µg 4 □ 45 µg 5 □ Discontinued	 □ CrCl ≥ 30 - 50 ml/min □ CrCl < 30 ml/min □ Reduction in platelet count □ Reduction in neutrophil count □ Lab result (other than platelet or neutrophil) □ Adverse reaction (complete AE form) □ Emotional symptoms □ Pregnant □ Patient preference □ Other, specify: 	