I C	ICCTG
C T	PROTOCOL #

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Clinical Center Stop Point

Patient ID:		
Patient Initials:		
Clinical Center:		
Contact Week:		
Date: /	_/	
RC ID:	year	

(Research Coordinator completed when patient stops taking the study medication AND at the week 24 visit.)

1.	Which stop	point does the clinic staff feel the patient has reached? (Check only one)		
	$\square_{\scriptscriptstyle 1}$	Completed the study and will NOT continue in post-treatment follow-up period.		
	\square_2	Completed the study and WILL CONTINUE in post-treatment follow-up period (receiving masked study drug). (Do not complete question #2 or #3 on this form.)		
	\square_3	Use of unacceptable concomitant medication and recorded on Medication Diary Record (DIARYREC) as MED #		
		Date: / / year Please specify:		
	$\square_{\scriptscriptstyle 4}$	Positive pregnancy test (Indicate date of test): //		
	\square_{5}	Two consecutive abnormal LFT tests (2.5 x ULN) OR two consecutive blood coagulation tests outside the institution's limits of normal as defined in the protocol. (Indicate date of tests below) (Must be recorded on Lab Results form)		
		///		
	\square_{6}	Adverse event as determined by P.I. and recorded on AE/SAE form as AE #		
		Date of Onset: // month day year Please specify:		
	\square_{7}	Transfer to another clinical site (complete Patient Transfer form)		
	\square_{8}	Patient dissatisfied with treatment Please specify:		
	\square_{9}	Patient no longer interested in participating (for reasons other than treatment) Please specify reason:		
	\square_{10}	Other Please specify reason:		
2.	Date patier	at received final dose of white study medication: Date: //		
3.	Date patier	at received final dose of green study medication: Date: / /		