Comments:	- Affix label here-						
	Clinical Center/ID:						
	First NameM.I						
	Last Name						
1. Contact Date: (M/D/Y)	(Complete Question 5 before interview.)						
• • • • •	5. Dosage/Adherence						
 Staff Person: Contact Type: 	5.1. Taking Standard WHI Dosage:						
<u> </u>							
☐ ₂ Mail ☐ ₈ Other	Yes Unable to do						
4. Visit Type:							
☐2 Semi-Annual # L	5.2. Taking Altered Dosage:						
	No Adherence rate						
o	Yes Unable to do						
Non-Routine							
 Refer to the Hysterectomy Status in WHILMA: If the Hysterectomy Status is "Yes", mark YES in 6 and go to 6.1. If the Hysterectomy Status is "No", ask, "Have you had a hysterectomy?" If the participant reports a hysterectomy, mark YES in 6 and go to 6.1. (Contact the CCC before dispensing any study pills.) If participant says she has not had a hysterectomy, mark NO in 6 and go to 6.2. 							
6. Has the participant had a hysterectomy?							
your last contact?"							
\square_0 No \longrightarrow Go to Question :	7.						
	7 and refer to Clinic Practitioner.						
6.2. Review Form 53 - HRT Calenda last contact?"	Review Form 53 - HRT Calendar if available. "Have you had any vaginal bleeding since your						
· ·	'						
"These next questions are about your vaginal bleeding."							
<u> </u>	6.3. "How heavy was it?" (Use the heaviest time since the previous contact.) Spotting - Approx. 1 pad's worth/day Moderate - Approx. 4-7 pads' worth/day						
<u> </u>	3						
Light - Approx. 2-3 pads' v	7						
	6.4. "When did the bleeding start?" (Use the earliest time since the previous contact.)						
	(M/D/Y)						
	"Did the bleeding start and stop again?" ☐ No ☐ Yes						
U I	0 1						
The state of the s	(100071)						

7.	"Since your last contact, have you had any breast tenderness?"	11.	"Since your last contact, has you had any of the following			
	□ ₀ No	11.1	"Endometrial hyperplasia"	□ ₀ No	1 Yes	
	7.1 "Was your breast tenderness mild, moderate, or severe?"	11.2	"High triglycerides in your blood (triglycerides are not the same as cholesterol)"	□ ₀ No	☐ ₁ Yes	
	☐2 Moderate ☐3 Severe		11.3 If yes: "Were your triglycerides over 1,000 (mg/dl)?"	□ ₀ No	T ₁ Yes	
	"Since your last contact, have you had any operations on or noticed any <u>other</u> changes in your breasts (new lumps, nipple discharge, or skin changes)?"	11.4	"Blood clot to your leg or lung"	□ ₀ No	☐ ₁ Yes	
		11.5	"Melanoma of skin"	□ ₀ No	☐ ₁ Yes	
	□ ₀ No	11.6	"Heart attack or stroke"	□ ₀ No	1 Yes	
	Yes Refer to Clinic Practitioner.	11.7	"Meningioma, or tumors in the brain"	□ ₀ No	1 Yes	
oth	u may have already answered these questions on er forms, but I'd like to recheck these items to make	11.8	"Breast cancer"	□ ₀ No	☐ ₁ Yes	
sure it is safe for you to stay on your study pills."	11.9	"Gall bladder disease"	□ ₀ No	☐ ₁ Yes		
9.	"What was the date of your last mammogram?"	11.10	"Problems with your pancreas"	□ ₀ No	1 Yes	
10.	Month Year "Are you now taking, or has your doctor prescribed,	11.11	l"Transient ischemic attack (TIA or "mini- stroke")"	No	Yes	
10.1	any:" "Corticosteroids (such as Prednisone, Decadron, Medrol in pill form)?" Yes		2"Sudden, serious changes in your eyes or vision" fer any "Yes" responses in 11.	No 1 - 11.12 to 0	Yes	
10.2	"Blood thinning medications of the such as Coumadin or Warfarin)?" No the such as Coumadin or Warfarin)?"	12.	•			
	"Other than your WHI study pills, are you now taking, or has your doctor prescribed, any hormones such as:"					
10.3	"Estrogen?" D ₀ No D ₁ Yes					
10.4	"Progesterone?"					
10.5	"Testosterone?"					
10.6	"Tamoxifen, Raloxifene					
Ref	er any "Yes" responses in 10.1 - 10.6 to CP.					

 13. Resulting action from participant reports of symptoms or concerns in items 6-12. (This item must be completed. Mark all that apply.) 	14.4. "People miss taking their study pills for many reasons. If there were days you did not take the pills, what were the reasons you didn't?" (Mark all that apply.) 1 Took all pills every day
Participant advised to return to clinic for evaluation. Date and time of next appointment:	Experienced symptomsForgot pill(s)
Consulting gynecologist notified. A Participant referred to primary physician:	 ☐ 4 Forgot bottle ☐ 5 Needed/Took a break ☐ 6 Afraid of health problems
Physician: Medications changed or stopped (complete Form	Family/Friend recommendation MD recommendation
54 – Change of Medications) Other (Specify):	Didn't have any pills 88 Other
14. "I'd like to talk with you about your HRT study pills."	14.5. Strategies to improve adherence (Refer to forms instructions for specific examples.)
14.1. "Since your last contact, how often did you take the study pills? Would you say" (Mark the response most often true.) (Read responses to participant.)	Ask participant to describe reason(s) given.Provide reassurance, using validation, review of facts
□ "Not at all"	Recommend palliative measures, using specific examples.
☐ 1 "Less than once per week" ☐ 2 "1 - 2 days per week" ☐ 3 "3 - 4 days per week"	Recommend steps to improve adherence, such as ways to deal with problem at home, self-motivation, mobilizing social support
□ 3	Put issues into perspective- emphasize safety of study, importance of WHI in answering health problems
14.2. "It is common for people to miss taking pills. About how many days have you missed taking your pills in the last month?" (Use best estimate.)	Use local CC guidelines to determine if referral to CP or other specialist is needed.
days in the last month	15.1 Should participant be put on Intensive Adherence Program (IAP)? (See instructions for entry criteria.)
14.3. "What helped you remember to take your pills?"	Yes \longrightarrow 15.2 Date to be recontacted (M/D/Y)
	16.1 Should participant be recontacted in one month by phone for clinical follow-up?
	No
	Yes -> 16.2 Date to be recontacted (M/D/Y)

WHI	Form 10 - HRT Management and Safety Interview	Ver. 7.2
	17. Comments:	