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**Participant ID**

**Variable #** 1 **Usage Notes:** none  
**Sas Name:** ID **Categories:** Study: Administration  
**Sas Label:** Participant ID  
**Type:** Continuous

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**Visit type for HRT adherence period**

**Variable #** 2 **Usage Notes:** none  
**Sas Name:** ADHVTyp **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Visit type for HRT adherence period  
**Type:** Categorical  
**Values**  
3 Annual Visit

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**Visit year for HRT adherence period**

**Variable #** 3 **Usage Notes:** none  
**Sas Name:** ADHVY **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Visit year for HRT adherence period  
**Type:** Continuous

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**Days from rand to start of HRT adherence period**

Days from randomization to start of HRT adherence period.

**Variable #** 4 **Usage Notes:** none  
**Sas Name:** STARTDY **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Days from rand to start of HRT adherence period  
**Type:** Continuous

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**Days from randomization to end of adherence period**

Days from randomization to end of HRT adherence period.

**Variable #** 5 **Usage Notes:** none  
**Sas Name:** ENDDY **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Days from rand to end of HRT adherence period  
**Type:** Continuous

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**HRT medication adherence rate for the period**

**Variable #** 6 **Usage Notes:** none  
**Sas Name:** ADHRATE **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** HRT medication adherence rate for the period  
**Type:** Continuous

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**Was adherence collection performed during period**

Were HRT pill bottles collected to allow adherence determination for period? If not, no adherence rate can be calculated.

**Variable #** 7

**Usage Notes:** none

**Sas Name:** COLLECT

**Categories:** Study Interventions: HRT Intervention/Management

**Sas Label:** Was adherence collection performed during period

**Type:** Categorical

**Values**

0	No
1	Yes

**Participant inactive in intervention during period**

Was the participant inactive in the HRT intervention (i.e. not taking study pills) for all or part of the period.

**Variable #** 8

**Usage Notes:** none

**Sas Name:** STOPHRT

**Categories:** Study Interventions: HRT Intervention/Management

**Sas Label:** Participant inactive in HRT intervenin during period

**Type:** Categorical

**Values**

0	No
1	Yes

**Participant resumed HRT intervention during period**

Did the participant resume HRT intervention (start taking study pills) during this period after having stopped?

**Variable #** 9

**Usage Notes:** none

**Sas Name:** RESUMEHRT

**Categories:** Study Interventions: HRT Intervention/Management

**Sas Label:** Participant resumed HRT intervention during period

**Type:** Categorical

**Values**

0	No
1	Yes

**Participant lost-to-follow-up during period**

Did the participant have a status of lost-to-follow-up during all or part of this HRT adherence period?

**Variable #** 10

**Usage Notes:** none

**Sas Name:** LOST

**Categories:** Study Interventions: HRT Intervention/Management

**Sas Label:** Participant lost-to-follow-up during period

**Type:** Categorical

**Values**

0	No
1	Yes



**Participant deceased during period**

Did the participant have a status of deceased during all or part of this HRT adherence period?

**Variable #** 11 **Usage Notes:** none  
**Sas Name:** DEAD **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Participant deceased during period  
**Type:** Categorical

**Values**

0	No
1	Yes

**Were open label HRT meds dispensed during period**

Were any open label HRT medications dispensed to the participant during this period?

**Variable #** 12 **Usage Notes:** Open label medications are not factored into the adherence rate calculation.  
**Sas Name:** OPENLABEL **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Were open label HRT meds dispensed during period  
**Type:** Categorical

**Values**

0	No
1	Yes

**Participant switched from E-alone to E+P in period**

Participant was switched from the unopposed estrogen study group to the estrogen+progesterone study group during this period (January 1995), due to PEPI trial results indicating long-term adherence to estrogen was not feasible in women with a uterus.

**Variable #** 13 **Usage Notes:** none  
**Sas Name:** ERT2PERT **Categories:** Study Interventions: HRT Intervention/Management  
**Sas Label:** Participant was switched from E-alone to E+P in period  
**Type:** Categorical

**Values**

0	No
1	Yes