



Attention - DO NOT enter patient data on this form if the header does not contain *preprinted* HALT PKD ID number, clinical center ID, and visit number.

Participant ID: \_\_\_\_\_ *haltid* Clinical Center: \_\_\_\_\_ *clinic* Date of Visit: \_\_\_\_/\_\_\_\_/\_\_\_\_  
*dvm / dvd / dvy*

*visit*

Missing Data Codes: A-Participant Refused B-Reading Not Possible C-Institutional Error

**UNMASKING DRUG FORM**

**Form #26**

*This form is to be entered in WDES by authorized personnel in order to unmask study treatment arm. The paper form is to be completed by designated personnel within 24 hours and stored in the participant's research chart.*

1. Reason for unmasking study treatment: *rsust*

**Pregnancy** Refer to the Manual of Procedures for guidelines requiring unmasking in the event of pregnancy.

**Intercurrent Illness** (Specify) *uillname* \_\_\_\_\_

**Other** (Explain) *uoreasn* \_\_\_\_\_

2. Date of last dose of study medication: \_\_\_\_/\_\_\_\_/\_\_\_\_  
*ldmm ldmd ldmy*

3. Method of Unmasking: *umeth*  **Contacted DCC** Date Contacted DCC \_\_\_\_/\_\_\_\_/\_\_\_\_  
*dccm dccd dccy*

**Other:** (Specify) \_\_\_\_\_ *uometh*

4. **Comments:** *cmmt* \_\_\_\_\_

**Optional Section: Not Data Entered**

**A. Treatment Arm:**

Study A, treat to standard BP ( $\leq 130/80$  mm Hg)

Study A, treat to aggressive BP ( $\leq 110/75$  mm Hg)

Study B, treat to standard BP ( $\leq 130/80$  mm Hg)

**B. Treatment Regimen:**

Participant Received:  ACE+ placebo  ACE+ARB

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HALT PKD staff member completing this form: \_\_\_\_\_ *cmidnum* Date: \_\_\_\_/\_\_\_\_/\_\_\_\_  
*Month cdm Day cdd Year cdy*

Reviewed by Study Investigator (signature required): \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_  
*pism Month pisd Day pisy Year*

Data Entry Status: Please check to indicate that the above information has been entered

Primary Entered by: \_\_\_\_\_ *deidnum* Date: \_\_\_\_/\_\_\_\_/\_\_\_\_  
*Month Day Year dem / ded / dey*