

DAC Study Form 302 - Fistula Study Dropout Form

This Form is completed when it is determined that a patient who was enrolled in the Baseline of the DAC Fistula Full Scale Study will not be randomized.

- 1. Patient Identification Number..... _ _ _ _ _
- 2. Name Code..... _ _ _ _ _
- 3. Date of Pre-Randomization Dropout _ _ / _ _ / _ _ _ _ _
- 4. Primary Reason for Pre-Randomization Dropout _ _

Note: These choices are in rank order. Please enter the first reason that applies.

- 1 = Graft was placed instead
- 2 = Fistula was not placed for some reason other than transplant or peritoneal dialysis
- 4 = We could not determine within 1 day what type of access was placed
- 5 = We could not get the blood results from the lab on time
- 6 = Patient ineligible due to taking aspirin before randomization
- 7 = Patient ineligible due to out of range lab values which have been entered into the study database
- 8 = Patient ineligible due to starting anti-platelet or anti-thrombotic therapy
- 9 = Fistula clotted prior to randomization
- 10 = Logistics reasons
- 20 = More than 45 days passed since Baseline data collection, and new data could not be collected
- 21 = More than 90 days passed since the consent form was signed, and a new consent could not be obtained
- 22 = Patient received a transplant
- 23 = Patient decided to go on peritoneal dialysis instead
- 24 = Patient ineligible due to developing an exclusion criteria
- 29 = Patient was lost to follow-up
- 30 = Patient Preference
- 31 = Study Team Preference
- 32 = Study enrollment period ended

- 5. Secondary Reason for Pre-Randomization Dropout _ _
(Use codes from q.4)
- 6. If q.4 or q.5 is "Logistics reasons", or if q.4 = "Developed an exclusion criteria", describe what happened.

201. Date this form completed _ _ / _ _ / _ _ _ _ _

202. User ID of person completing this form..... _ _ _ _ _

<i>Clinical Center Use Only</i>
Date Form Entered _ _ / _ _ / _ _ _ _ _
Person Entering this Form _____