

F240

Hemodialysis Fistula Maturation Study (HFM/MANVAS Studies) Baseline Drop Out/Not Eligible to be Followed Form (Form # 240)

This form should be completed for any patient who will not be part of the follow up cohort.

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1. Identification Number PID 2. Alphacode AC 3. Visit Number VIST 4. Date Dropped Out/Not Eligible: mm/dd/yyyy VISIT_DT

5. Primary Reason for Baseline DropoutPRIM_RSN

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

- 1 = A different type of access, not an AVF, was created because the artery was not suitable
- 2 = A different type of access, not an AVF, was created because the vein was not suitable
- 3 = A different type of access, not an AVF, was created for some other reason (Describe in Q7 text)
- 4 = A different type of access, not an AVF, was created. The reason is unknown.
- 5 = An AVF was created but a two stage procedure is planned
- 6 = Patient received a transplant
- 7 = Patient decided to go on peritoneal dialysis instead
- 8 = No access was created. There is a known reason, other than transplant or peritoneal dialysis (Describe in Q7 text).
- 9 = No access was created. The reason is not known.
- 10 = Pre-operative US was not performed on the arm where the study fistula was created
- 11 = Pre-operative US was done on the arm where the study fistula was created but was inadequate or **unreadable** and could not be repeated before surgery
- 12 = Missing data for FMD and venous plethysmography vascular function testing
- 13 = Missing data for venous plethysmography and arterial pulse wave velocity vascular function testing
- 14 = Missing data for FMD and arterial pulse wave velocity vascular function testing
- 15 = Missing data for all three vascular function studies
- 16 = Surgery was performed by a surgeon not participating in the study
- 17 = Patient is not yet on chronic dialysis and was over 80 years of age at access creation surgery
- 21 = More than 30 days passed after surgery was done and no Form 230 Surgery Notification Form in the database to confirm that a single stage AVF was created
- 31 = At the time of surgery, learned that the patient will not be available for the 2-week ultrasound studies
- 32 = At the time of surgery, learned that the patient will not be available for the 6-week ultrasound studies
- 41 = Local IRB consent form guidelines. Too much time passed between the time the consent form was signed and the time of surgery, and a new consent could not be obtained
- 51 = Patient changed mind/decided not to participate (on day of surgery or before)
- 61 = Patient was lost to follow up before surgery was done
- 71 = Other unusual logistic reason (Describe in Q7 text)
- 72 = Other unusual medical reason (Describe in Q7 text)

81 = Study team preference
99 = Study enrollment period ended

6. Secondary Reason for Baseline Dropout SEC_RSN
(Use codes from Q5)

7. If the primary reason for Baseline Dropout in Q5 is:
3 = A different type of access, not an AVF, was created for some other reason
8 = no access created for some other known reason
71 = other unusual logistic reason
72 = other unusual medical reason

Please describe what happened: DESCRIBE

200. Date this form completed (mm/dd/yyyy)..... COMP_DT

201. Username of person completing/reviewing completeness of this form.....COMP_USER

<p>Clinical Center Use Only</p> <p>Date Form Entered (mm/dd/yyyy) ____/____/____ ENTER_DT</p> <p>Username of person entering this form____ ENTER_USER</p>
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