INTERMACS® Screening Log

Inclusion: Patient must meet all inclusion criteria:

If patient meets all inclusion criteria then check 'ALL' inclusion reasons below*

Patient receives a durable mechanical circulatory support device (MCSD)

which is FDA approved

Implanted on or after March 1, 2006 (The device does not need to be the

first implant for the patient)

Patient signed informed consent for the registry

Exclusion: Any exclusion will disqualify the patient for entry into INTERMACS®

If patient meets 'ANY' exclusion criteria then check any of the appropriate exclusion reasons below (check all that apply):*

Patient receives a durable mechanical circulatory support device (MCSD)

which is not FDA approved

Patient is incarcerated (prisoner)

Patient did not sign the informed consent.

Select reasons why patient was not consented: *

Too sick pre-implant and died early post implant

Missed opportunity to consent

Patient refused

Patient is unable to communicate in English

If Patient is included, please complete INTERMACS® required screening information below:

Device type: *

Device brand: *

Device brand (RVAD):*

Specify:*

Specify RVAD:*

Implant date:* mm/dd/yyyy

If Patient is excluded, please complete INTERMACS® required screening information below:

Implant date:*

mm/dd/yyyy

Device type:*

Device brand: *

Device brand (RVAD):*

Specify: *

Specify RVAD:*

Age range (years):*

Race: select as many as

apply: *

American Indian or Alaska Native

Asian

African-American or Black

Hawaiian or other Pacific Islander

White

Unknown/Undisclosed Other/none of the above

Ethnicity: Hispanic or Latino:	YES NO	v3.0
Gender:*	Male Female	
Did death occur within 2 days post implant?*	YES NO	
Is this VAD an investigational device?*	YES NO	
Is patient involved in a VAD related study?:*	YES NO UNK	
What is the name of the study?:*		