Intermacs

Screening Log

Implant Date
MM/DD/YYYY

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

Device type

- ○ LVAD
- ○ RVAD
- ○ Both (LVAD + RVAD in the same OR visit)
- ○ Total Artificial Heart

Please remember to fill out the RHF adverse event form

Device brand

- ○ Berlin Heart EXCOR (paracorporeal)
- ○ Medtronic HVAD
- ○ HeartMate II LVAS
- ○ HeartMate III
- ○ HeartMate IP
- ○ HeartMate VE
- ○ HeartMate XVE
- ○ Micromed DeBakey VAD - Child
- ○ Novacor PC
- ○ Novacor PCq
- ○ Thoratec IVAD
- ○ Thoratec PVAD
- ○ Other, Specify

Specify brand

Device brand (RVAD)

Specify brand (RVAD)

Age Range

- ○ 19 to 39
- ○ 40 to 59
- ○ 60 to 79
- ○ 80+

Race	☐ 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 ○ Unknown
Gender	○ Male ○ Female ○ Unspecified
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 ○ Unknown
What is the name of the study?	
Is this an industry sponsored post approval study?	○ Yes ○ No ○ Unknown