INTERMACS Eligibility

Please submit the data below to confirm eligibility for INTERMACS inclusion. Only enter information on consented patients with FDA approved durable (potential for patient discharge) devices. FDA approved temporary devices should be included only if they are implanted simultaneously with a durable device or subsequent to a durable device.

There is one exception to this rule, data entry for patients receiving the Berlin Heart EXCOR device. Authorized Berlin Heart medical centers should proceed to enter EXCOR recipient data in INTERMACS.

Device type:*	LVAD
Device brand:*	
Device brand (RVAD):*	
Specify:*	
Specify RVAD:*	
Implant date:*	mm/dd/yyyy

Reminders:

1. Obtain consent pre-implant. If this is not possible, obtain consent immediately post-implant.
2. Refer to Appendix K of the Protocol for the current list of devices and their FDA approval status.
3. Please remember to complete and submit any device tracking forms that accompany the implanted devices.