## Screening Log - Pedimacs 11/07/2024

Blank View

Implant Date	
	of implant support device (MCSD) which is FDA approved 2 (The device does not need to be the first implant for the patient)
Device type	○ LVAD ○ RVAD ○ Both (LVAD + RVAD in the same OR visit) ○ Total Artificial Heart
Device brand	Berlin Heart EXCOR (paracorporeal) HeartWare HVAD HeartMate II LVAS HeartMate III HeartMate IP HeartMate VE HeartMate XVE Micromed DeBakey VAD - Child Novacor PC Novacor PCq Thoratec IVAD Thoratec IVAD Abiomed AB5000 Abiomed BVS 5000 Thoratec Centrimag (Levitronix) Thoratec Pedimag TandemHeart Biomedicus Maquet Rotaflow Sorin Revolution Abiomed Impella 2.5 Abiomed Impella CP Abiomed Impella RP Abiomed Impella 5.5 Other, Specify

Specify brand

	12/05/2024
Device brand (RVAD)	
Specify brand (RVAD)	
Age Range	0 to 2 3 to 4 5 to 9 10 to 12 13 to 15 16 to 18
Race	<ul> <li>American Indian or Alaska Native</li> <li>Asian</li> <li>African-American or Black</li> <li>Hawaiian or other Pacific Islander</li> <li>White</li> <li>Unknown / Undisclosed</li> <li>Other / none of the above</li> </ul>
Ethnicity: Hispanic or Latino	<ul><li>Yes</li><li>No</li><li>Unknown</li></ul>
Gender	<ul><li>Male</li><li>Female</li><li>Unknown</li></ul>
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?	<ul><li>Yes</li><li>No</li><li>Unknown</li></ul>
What is the name of the study?	
Is this an industry sponsored post approval study?	<ul><li>Yes</li><li>No</li><li>Unknown</li></ul>