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Adverse Events and Device Deficiency Sheet
Study ID: H-34 Q1 2022

Adverse Events																																																
Patient ID	Report type	Initial Report Date	Report Date	Date of Onset	Date of Study Team Aware	Time team became aware (24 hr clock)	Title of clinical study	Protocol ID	Name of the healthcare facility	Name of the Principal Investigator	Name of the medical device involved	Components involved	ID number of study subject	Age	Gender	Attribution within the clinical investigation	Adverse Event (disease, if known, or signs/symptoms)	Event narrative	Admission date (dd/mm/yyyy)	Discharge date (dd/mm/yyyy)	Classification of the adverse event	Outcome of the event	Relevant concomitant therapy and date of administration	Detail all possible and suspected causes including relevant medical history	Current medical status	Severity	Causality: relationship to study procedure	Expectedness	Action taken	Detailed treatment	Was the device permanently removed?	Attachments: Relevant Medical History	Attachments: Relevant Medical Records	Attachments: Imaging Laboratory Results	Attachments: Other	Submitter	Name of submitter	Notes	Status	Initial / Follow-up	communicated to PMS	If Yes, date	Date of reply	Device relationship to PMS dept	Relationship ID PMS Dept	Complaint	Further evaluation performed	Final decision/communication relationship and expediency
101	9	Initial & Final	07/12/2021	07/12/2021	01/12/2021	08:00	DELTA Revision study	H-34	Samodzielny Publiczny Szpital Kliniczny im. Prof. Adama Grucy CMKP	Jerzy Bialecki	Delta Revision Cup	Delta Revision Cup, Hemispherical module, Bone screws, Insert	101 - 9	60	Male	Treatment Group	Cup loosening	Early postoperative cup loosening.	19-Nov-2021	19-Nov-2021	Medical or surgical intervention required to prevent any of the above	Ongoing	None	Procedure done successfully in first operation with Delta Revision implantation. Too proximal cup implantation and not sufficient cup hook stabilization at the time of surgery. There could be wrong position of stable stem, various position in femur.	Severe	Not Related	Not Related	N/A	Not pharmacological	Revision surgery with cup and stem removal and Reimplantation Trident Multihole cup and stem replacement system.	Yes	None	None	Available	None	Investigator	Pawel Bartosz	N/A	Valid	Initial	Yes	09/12/2021	10/12/2021					
101	10	Initial	19/01/2022	19/01/2022	14/01/2022	08:51	DELTA Revision study	H-34	Samodzielny Publiczny Szpital Kliniczny im. Prof. Adama Grucy CMKP	Jerzy Bialecki	N/A	N/A	101 - 10	76	Male	Treatment Group	Dyspnoea, fluid collection in lungs	Chronic heart failure exacerbation	30-Dec-2021	30-Dec-2021	Hospitalization or prolongation of hospitalization	Ongoing	Furosemid 40mg 2xday from 14.01.2022.	Chronic heart failure	Good	Moderate	Not Related	Not Related	N/A	Pharmacological		No	None	None	None	None	Investigator	Pawel Bartosz	N/A	Draft	Initial	No						
101	5	Initial	03/02/2022	03/02/2022	20/10/2021	00:00	DELTA Revision study	H-34	Samodzielny Publiczny Szpital Kliniczny im. Prof. Adama Grucy CMKP	Jerzy Bialecki	Revision Hip	Stem17mm/20 Dmm	101 - 5	76	Female	Treatment Group	Intraoperative fracture of femur shaft	The femur shaft was weakened because of previous use of metal loop. 25 years ago ThM of both hips, 2011- revision of right hipEcirc;	18-Oct-2021	27-Oct-2021	Hospitalization or prolongation of hospitalization	Ongoing	N/A	On the first control visit 22 Nov 2021 the femur shaft was still weakened. The patient was able to walk with 2 crutches.	Severe	Not Related	Not Related	Anticipated	Other	Intraoperative reduction and fixation of the fracture	No	None	None	None	None	None	Investigator	Julia Macias	N/A	Draft	Initial	No						

Role: _____ Date: _____ Signature: _____

Written by: Lisa Claffarin AJCL

Approved by:
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