Parameter (Englisch) DRKS	Parameter ClinicalTrials.gov	Modifications
Title	Official Title oder Brief Title	
Trial Acronym	Acronym	
Brief Summary in Lay Language	Brief Summary	
Brief Summary in Scientific Language	Detailed Description	
Keywords	Keywords provided by Additional relevant MeSH terms	
Date of Registration in Partner Registry or other Primary Registry	First Received Date	
Investigator Sponsored/Initiated Trial (IST/IIT)	/	performed semi-automatically on the basis of the sponsor name
Universal Trial Number (UTN)	Other Study ID Numbers	number is copied manually from "Other Study ID Numbers" into this field
EudraCT Number	Other Study ID Numbers	number is copied manually from "Other Study ID Numbers" into this field
Primary Registry ID	ClinicalTrials.gov Identifier: NCT Number	
BfArM No.	Other Study ID Numbers	number is copied manually from "Other Study ID Numbers" into this field
PEI No.	Other Study ID Numbers	number is copied manually from "Other Study ID Numbers" into this field
Other Secondary ID	Other Study ID Numbers	may have to be copied into the correct field
Health Condition or Problem studied	Condition	additional coding with ICD-10, performed bei DRKS staff
Other Health Condition or Problem studied	Condition	additional coding with ICD-10, performed bei DRKS staff
Intervention Arm 1	Intervention	
Intervention Arm 2	Intervention	
Further Intervention Arms	Intervention	
Study Type	Study Type	
Study Type Non-interventional	Study Type	(filled in only in case of observational trial)
Allocation	Intervention Model und Allocation	is composed of two parameters
Blinding	Masking	
Control	Intervention Model und Arms	is composed of two parameters
Purpose	Primary Purpose	
Assignment	Intervention Model	
Phase	Study Phase	
Primary Outcome	Current Primary Outcome Measures	
Secondary Outcome	Current Secondary Outcome Measures	
Countries of Recruitment	Location Countries	
Location	Locations	if Country = Germany
(Anticipated or Actual) Date of First Enrollment (dd.mm.yyyy)	Start Date (mm.yyyy)	day is added (end of month)

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Target Sample Size	Estimated Enrollment oder Enrollment	if the planned sample size is not available the field is filled with the actual sample size
Study Closing (LPLV) (dd.mm.yyyy)	Completion Date (mm.yyyy)	day is added (first day of the month)
Inclusion Criterion Gender	Gender	
Inclusion Criterion Minimum Age	Ages	conversion from minutes to days and from hours to days
Inclusion Criterion Maximum Age	Ages	conversion from minutes to days and from hours to days
Inclusion Criteria	Eligibility Criteria	Eligibility Criteria are devided into two parameters
Exclusion Criteria	Eligibility Criteria	Eligibility Criteria are devided into two parameters
Recruitment Status	Recruitment Status	
If "Other" Reason for Recruiting Stop please specify.	Recruitment Status (why stopped)	
Total Number of Participants (all Sites worldwide) after Recruiting complete	Enrollment	
Primary Sponsor	Sponsor	
Contact for Scientific Queries	<overall_official> oder <overall_contact>	
Contact for Public Queries	<overall_contact> oder <overall_official>	
Collaborator, Other Address	collaborators	
Trial Publications. Results and other Documents	Publications, Additional Information	