The YODA Project Research Proposal Due Diligence Assessment

Part 1: General Information		
YODA Project (Protocol) ID:	2016-0912	
Date:	13 May 2016	
Product Name:	THERMOCOOL® SMARTTOUCH™ Catheter	
Therapeutic Area:	CV	
Product Class:	Cardiovascular devices	
Condition(s) Studied:	Atrial Fibrillation	
Protocol Number(s) and	NCT01385202-Smart-AF-THERMOCOOL® SMARTTOUCH™ Catheter	
Title(s):	for the Treatment of Symptomatic Paroxysmal Atrial Fibrillation	
Part 2: Data Availability		
	Question:	Response:
Data Holder has authority to p	rovide clinical trial data or development	Yes
partner has agreed to share cli	nical trial data.	_
Comments: N/A		
Data Holder has shareable electronic clinical trial data or data can be		Yes
converted to electronic format Comments: N/A		
De-identification and redaction of clinical trial data in accordance with current Yes		
HIPAA and EU criteria allows protection of participant privacy and		163
confidentiality.		
Comments: N/A		
The product and relevant indication studied has either been approved by		Yes
regulators in the US and EU, or terminated from development.		
Comments: N/A		
Data Holder has completed the clinical trial and trial has been completed for a		Yes
period of at least 18 months (or results published in peer-reviewed		
biomedical literature). Comments: N/A		
Part 3: Data Availability Summary		
Based on the responses to the above Data Availability questions, the		Yes
requested clinical trial data car	n be made available for data sharing.	
Part 4: Proposal Review		
Question:		Response:
Summary-level CSR data is appropriate for the proposed analysis.		No
Participant-level data is appropriate for the proposed analysis.		Yes
A similar analysis is underway or completed/pending disclosure by Janssen.		No
Comments:		