The YODA Project Research Proposal Due Diligence Assessment

	Part 1: General Information	
YODA Project (Protocol) ID:	2021-4649	
Date:	15 April 2021	
Product Name:	Abiraterone acetate	
Therapeutic Area:	Oncology	
Product Class:	CYP17 inhibitor	
Condition(s) Studied:	Prostate Cancer	
Protocol Number(s) and Title(s):	NCT01715285 212082PCR3011 - A Randomized, Double-blind, Comparative Study of Abiraterone Acetate Plus Low-Dose Prednisone Plus Androgen Deprivation Therapy (ADT) Versus ADT Alone in Newly Diagnosed Subjects With High-Risk, Metastatic Hormone-naive Prostate Cancer (mHNPC) NCT00638690 COU-AA-301 - A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study of Abiraterone Acetate (CB7630) Plus Prednisone in Patients With Metastatic Castration-Resistant Prostate Cancer Who Have Failed Docetaxel-Based Chemotherapy	
Part 2: Data Availability		
Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments:		Yes
Data Holder has sharable electronic clinical trial data or data can be converted to electronic format.		Yes
Comments: De-identification and redaction of clinical trial data in accordance with current HIPAA Yes and EU criteria allows protection of participant privacy and confidentiality. Comments:		
The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments:		Yes
Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments:		
Part 3: Data Availability Summary		
Based on the responses to the above Data Availability questions, the requested clinical trial data are available for a data sharing request.		Yes
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. A similar analysis is underway or completed/pending disclosure by Janssen.		Yes
A similar analysis is underway or completed/pending disclosure by Janssen. No Comments:		