

**The YODA Project**  
**Research Proposal Due Diligence Assessment**

<b>Part 1: General Information</b>	
<b>YODA Project (Protocol) ID:</b>	2022-4992
<b>Date:</b>	8 July 2022
<b>Product Name:</b>	Risperidone/ Paliperidone
<b>Therapeutic Area:</b>	Neuroscience
<b>Product Class:</b>	Atypical antipsychotics
<b>Condition(s) Studied:</b>	Schizophrenia/ Schizoaffective Disorder
<b>Protocol Number(s) and Title(s):</b>	NCT00077714 R076477-SCH-304 NCT00078039 R076477-SCH-303 NCT00083668 R076477-SCH-305 NCT00085748 R076477-SCH-302 NCT00086320 R076477-SCH-301 NCT00105326 R076477-SCH-1010 NCT00309686 R076477-BIM-3003 NCT00309699 R076477-BIM-3002 NCT00334126 R076477SCH3015 NCT00397033 R076477SCA3001 NCT00412373 R076477SCA3002 NCT00460512 R076477SCH3017 NCT00488319 R076477PSZ3002 NCT00490971 R076477-BIM-3004 NCT00518323 R076477PSZ3001 NCT00524043 R076477-SCH-4012 NCT00566631 R076477SCH3018 NCT00645099 R076477SCH3020 NCT00645307 R076477-SCH-701 NCT00650793 R076477-SCH-703 NCT00752427 R076477-SCH-702 NCT01009047 R076477PSZ3003 NCT01258920 PALM-JPN-5 NCT01299389 PALM-JPN-4 NCT01662310 R076477-SCH-3041 NCT00299715 R076477-BIM-3001 NCT00074477 R092670-SCH-201 NCT00101634 R092670PSY3004 NCT00111189 R092670PSY3001 NCT00119756 R092670PSY3005 NCT00210548 R092670PSY3003 NCT00210717 R092670PSY3002 NCT00589914 R092670PSY3006 NCT00590577 R092670PSY3007 NCT00604279 R092670PSY3008 NCT01051531 R092670SCH3009 NCT01081769 R092670SCH3005 NCT01157351 R092670SCH3006 NCT01193153 R092670-SCA-3004

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NCT01281527	R092670SCH3010
NCT01515423	R092670PSY3011
NCT01527305	R092670SCH4009
NCT01529515	R092670PSY3012
NCT01606228	R076477SCH3033
NCT02713282	R092670SCH3015
NCT01624675	RIS-AUT-JPN-01
NCT N/A	RIS-INT-85
NCT00034749	RIS-USA-231
NCT00034762	RIS-USA-232
NCT00034775	RIS-USA-259
NCT00044681	RIS-INT-93
NCT00061802	RIS-SCP-402
NCT00076115	RIS-BIM-301
NCT00086112	RIS-ANX-301
NCT00088075	RIS-SCH-302
NCT00094926	RIS-BIP-302
NCT00095134	RIS-DEP-401
NCT00132678	RISBIM3003
NCT00216476	RISSCH3001
NCT00216528	RIS-KOR-66
NCT00216580	RIS-PSY-301
NCT00216632	RISSCH4026
NCT00216671	RISSCH4045
NCT00236353	RIS-USA-305
NCT00236379	RIS-USA-275
NCT00236444	RIS-INT-79
NCT00236457	RIS-INT-62
NCT00236470	RIS-INT-84
NCT00236587	RIS-USA-265
NCT00246246	RIS-BIP-301
NCT00249132	RIS-INT-3
NCT00249145	RIS-INT-24
NCT00249158	RIS-AUS-5
NCT00249223	RIS-INT-61
NCT00249236	RIS-IND-2
NCT00250354	RIS-CAN-19
NCT00250367	RIS-INT-46
NCT00253123	RIS-USA-63
NCT00253136	RIS-USA-121
NCT00253149	RIS-USA-102
NCT00253162	RIS-INT-69
NCT00257075	RIS-USA-239
NCT00261508	RIS-CAN-23
NCT00266552	RIS-USA-93
NCT00269919	RIS-KOR-64
NCT00297388	RIS-SCH-401
NCT00299702	RISSCH4060
NCT00369239	RISSCH4043
NCT00378092	RISSCH3024

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	NCT00391222	RISBMN3001
	NCT00495118	RIS-INT-80
	NCT00526877	RISSCH4119
	NCT00576732	RISAUT4002
	NCT00821600	RIS-SCH-1012
	NCT00992407	RISSCH4178
	NCT01050582	RISNAP4022
	NCT01855074	RISSCH4186
	NCT N/A	RIS-INT-83
	NCT N/A	RIS-USA-1
	NCT N/A	RIS-USA-150
	NCT N/A	RIS-USA-240
	NCT N/A	RIS-USA-72
	NCT N/A	RIS-BEL-14
	NCT N/A	RIS-USA-70
<b>Part 2: Data Availability</b>		
<b>Question:</b>		<b>Response:</b>
Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data.		Yes
Comments:	N/A	
Data Holder has sharable electronic clinical trial data or data can be converted to electronic format.		Yes
Comments:	N/A	
De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality.		Yes
Comments:	N/A	
The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development.		Yes
Comments:	N/A	
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).		Yes
Comments:	N/A	
<b>Part 3: Data Availability Summary</b>		
Based on the responses to the above Data Availability questions, the requested clinical trial data can be made available for data sharing.		Yes
<b>Part 4: Proposal Review</b>		
<b>Question:</b>		<b>Response:</b>
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:		