## The YODA Project Research Proposal Due Diligence Assessment

| Part 1: General Information   |  |           |
|---|--|-----------|
| YODA Project (Protocol) ID:   | 2024-0688  |           |
| Date:   | 16-July-2024   |           |
| Product Name:   | INVOKANA (Canagliflozin)   |           |
| Therapeutic Area:   | Cardiovascular & Metabolic Diseases  |           |
| Product Class:  | Diabetes Related – Other   |           |
| Condition(s) Studied:   | Diabetes Mellitus, Type 2  |           |
| Protocol Number(s) and Title(s):  | <ol> <li>NCT1032629 – 28431754DIA3008 – A Randomized,<br/>Multicenter, Double-blind, Parallel, Placebo-Controlled Study<br/>of the Effects of JNJ-28431754 on Cardiovascular Outcomes in<br/>Adult Subjects With Type 2 Diabetes Mellitus</li> <li>NCT1989754 – 28431754DIA4003 – A Randomized,<br/>Multicenter, Double-blind, Parallel, Placebo-Controlled Study<br/>of the Effects of Canagliflozin on Renal Endpoints in Adult<br/>Subjects With Type 2 Diabetes Mellitus,</li> </ol> |           |
| 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.  Comments:   |  | Yes       |
| De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality.                             |  | Yes       |
| The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development.  |  | Yes       |
| Comments:  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: |  | Yes       |
| 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.  Part 4: Proposal Review                             |  | Yes       |
| Question: Response  |  | Response: |
| Summary-level CSR data is appropriate for the proposed analysis.  |  | No No     |
| Participant-level data is appropriate for the proposed analysis.  |  | Yes       |
| A similar analysis is underway or completed/pending disclosure by Janssen.  |  | No        |
| Comments:   |  |           |