

**SOW-8: PREPARE IUCLID ENDPOINT  
(ROBUST) STUDY SUMMARIES**

- 8.1 *Contractor* shall prepare study summaries and robust study summaries, as required by the *REACH Regulation*, of all relevant and available data to fulfill the requirements of Article 10 (a)(iv), (vi), (vii) and (ix) for a joint submission for each substance listed by the *Buyer*.
- 8.2 *Contractor* shall prepare summaries for [the data and information identified under SOW-5.] [or] [the following data and information:
- *list available data & information here and identify the source]*
- 8.3 *Contractor* shall obtain copies of any study reports to be summarized directly from the data holder, along with the Universal Unique Identifier (UUID) / legal entity object of the data holder. *Contractor* shall import the data holder's UUID into the IUCLID 5 software before the endpoint (robust) summaries are prepared, revised, and/or updated.
- 8.4 *Contractor's* study summaries and robust study summaries shall be in the IUCLID 5 format (IUCLID Sections 4, 5, 6, and 7) and prepared in the [English] language.
- 8.5 *Contractor's* study summaries and robust study summaries shall be consistent

# SAMPLE

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- 8.8 *Contractor* shall determine the reliability of existing data and information as described in the *REACH Guidance* document titled: *Guidance on Information Requirements and Chemical Safety Assessment, Chapter R.4. Evaluation of Available Information*.
- 8.9 *Contractor* shall [not] include a peer review or technical review step as these reviews will [not] be performed separately by the *Buyer*.