
Comprehensive Data Management & Streamlining Solution for Business- Critical Data (Data Stream) Request for Information

Background:

This Sources Sought Request for Information (RFI) is for information, planning and market research purposes only and shall not be construed as either a solicitation or obligation on the part of the Food and Drug Administration or its Centers. The purpose of this RFI is to help the FDA understand the technical capabilities for providing the full range of services described in the Statement of Work (SOW) attached at the end. FDA will use this market research information to assess the market's capability to successfully meet the Data Stream requirements.

FDA welcomes responses from all interested businesses. FDA does not intend to make a selection decision or award a contract on the basis of responses nor otherwise pay for the preparation of any information submitted or FDA's use of such information.

Acknowledgment of receipt of responses will not be made, no questions will be answered, nor will respondents be notified of the outcome of the FDA's review of the information received. Additionally, the FDA does not intend to hold discussions concerning this RFI with any interested parties. However, FDA reserves the right to contact vendors if additional information is required.

General Instructions:

A complete response to the RFI should include answers to all of the questions below.

Please note that a respondent's failure to provide a complete response to a question will be interpreted to mean that it does not understand the question and does not demonstrate the capability to provide services required. Responses should demonstrate capability, not merely affirm the respondent's capability (e.g.: The response must go beyond the statement that, "XYZ company can provide training support.").

The Face Page requested under Question 1 will not count toward the page limitation. The narrative response to Question 2 shall not exceed five (5) pages. Any pages exceeding the stated page limitation may not be reviewed. The RFI responses should be submitted via email to Jada Scott at jada.scott@fda.hhs.gov by 12:00 PM EST on October 19, 2026. E-Mail is the only acceptable method of delivery for this sources sought notice.

Questions:

1. On the FACE PAGE of your response, please provide the following:

RFI #	FDA-RFI-27-1251190	
RFI Title	Comprehensive Data Management & Streamlining Solution for Business-Critical Data (Data Stream)	
Company	Name	
	Website URL	
	Address	
	Phone	
	DUNS Number	
	Business Size Status	
	How product can be procured?	
Original Equipment Manufacturer	Name	
	DUNS Number	
	Business Size Status	
Contact	Name	
	Email Address	
	Address (if different from company address)	
	Phone Number	
	Cell Number (optional)	
	Fax Number (optional)	

2. The SOW describes the background, objective, requirements, deliverables and period of performance for the Data Stream requirement. Please briefly describe your proposed solutions to meet the requirements specified on the SOW.
3. Responses shall not contain any information that requires a non-disclosure agreement or has sensitive business information.

Attachments:

Draft Statement of Work