



CENTER FOR GLOBAL CLINICAL RESEARCH DATA

Vivli Data Request Form: Tips and Tricks

Platform Version 3.9

28 February 2026

What is the Vivli Data Request Form?

- The Vivli Data Request Form is used by data contributors and independent review panels to evaluate your proposal and make decisions on your data access for the studies requested.
- Each Vivli member describes the criteria and process for how they make decisions about the use of their completed clinical trial data.
- To ensure a timely review of the proposal, your data request form should be as detailed and complete as possible.

Research Team



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Research Team

LEAD RESEARCHER



Activate user for accessing data request



Lead Researcher is also Statistician Researcher

First Name

Last Name

ORCID ID

Email (editable until user is invited to data request)

Position at current organization

Employer, Company, Research Institute, or Primary Affiliation

Country

Select One

Education, including professional qualifications that are relevant to the proposed research and are specific to clinical data analysis.

PhD, Biostatistics, Boston University, DOIs for publications relevant to this project

Name of the degree

Details of highest degree, e.g., PhD, Master's

Institution from where the degree was received

Discipline

Year Received

How many years of experience with secondary analysis

- Select an Option -

Please list any real or potential conflicts of interest and describe how these will be managed. If none, please enter None.

Access to Data

Access to data not applicable

Research Team



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STATISTICIAN RESEARCHER



Activate user for accessing data request ?

First Name Last Name ORCID ID

Email (editable until user is invited to data request) Position at current organization

Employer, Company, Research Institute, or Primary Affiliation Country Select One

Education, including professional qualifications that are relevant to the proposed research and are specific to clinical data analysis.
List any skills or prior experience in performing relevant or similar statistical analyses as planned (i.e. previous publications)

Name of the degree Institution from where the degree was received

Discipline Year Received How many years of experience with secondary analysis - Select an Option -

Please list any real or potential conflicts of interest and describe how these will be managed. If none, please enter None.

Access to Data Access to data not applicable

ADDITIONAL RESEARCHERS

Add +

| Name | Affiliation | Country | Email | Role(s) | Status Details |
|------|-------------|---------|-------|---------|----------------|
|------|-------------|---------|-------|---------|----------------|

Research Proposal



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GENERAL

Title of the Proposed Research (Overall brief description, including the condition, about the proposed research question that is understandable by a general audience)

Data Request Example

Lay Summary of the Research Request

[Vivli Lay Summary Training Video - YouTube](#)

What are the specific aims/objectives of the proposed research, including the specific hypotheses to be evaluated?

PURPOSE OF ANALYSIS ?

Purpose of Analysis is required.

- New research question to examine treatment effectiveness on secondary endpoints and/or within subgroup populations
- New research question to examine treatment safety
- Research that confirms or validates previously conducted research on treatment effectiveness
- Research that confirms or validates previously conducted research on treatment safety
- Preliminary research to be used as part of a grant proposal
- Summary-level data meta-analysis
- Participant-level data meta-analysis

OUTCOME(S)

Outcome(s) is required.

- Inform Patient Care Decisions
 - Inform Patient Care Decisions
- Algorithms / Code
 - A.I. algorithm
 - Algorithm for predicting treatment response
 - Code
 - Machine Learning
- Clinical Guidelines
 - Clinical guidelines
- Designing Future Trials, Trial Protocol

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Artificial Intelligence / Machine Learning

Please answer the following questions related to use of Artificial Intelligence (AI) and Machine Learning (ML). Also, review the Vivli policy in this document: [Vivli AI/ML Model Requirements](#)

Please confirm if you will be using any Artificial Intelligence or Machine Learning in this request? *

- No
 Yes

Review Vivli's AI/ML policy <https://vivli.org/vivli-ai-ml-model-requirements>

Response required.

Please indicate which model types will be developed in your analysis (Note: you will comment on how these models support your analysis in the SAP section) *

- Machine learning models - Defined models similar in nature to standard statistical models (Lasso, Random Forests, Support Vector Regression, Decision Tree, K-Nearest Neighbors (KNN), Gradient Boosting, Kernel Machine, Boosting, Gaussian Mixture models (GMMs))
- Artificial Neural Networks (ANNs) and Similar Algorithms - ex: Artificial Neural Networks (ANNs), Convolutional Neural Networks (CNNs), Recurrent Neural Networks (RNNs), Transformer Models (non-generative)
- Generative Models / Large Language Models (LLMs) – ex: GPT, Generative Adversarial Networks (GANs), Diffusion models

Response required.

Will your analysis include the use of existing trained models? *

- No
 Yes

Response required.

Will you be exporting a model from the Vivli research environment? *

- No
 Yes

Research Proposal



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STUDY DESIGN

Brief Description

Please describe the specific outcomes elements and how they will be categorized/defined for your proposed research

Character Count: 0/1000

Main Predictor/Independent Variable and how it will be categorized/defined for your proposed research

Other variables of interest that will be used in your analysis and how they will be categorized/defined in your proposed research (i.e. genders, age groups, ethnic groups)

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Main Predictor/Independent Variable and how it will be categorized/defined for your proposed research

Other variables of interest that will be used in your analysis and how they will be categorized/defined in your proposed research (i.e. genders, age groups, ethnic groups)

PROJECT TIMELINE ?

Target Analysis Start Date

10/20/2025

Estimated Analysis Completion Date

10/20/2026

DISSEMINATION AND PUBLICATION PLAN

Dissemination & Publication Plan ?

Declare your intention to publish your results, and list of potential journals or conferences here

Provide references for all cited material (following APA guidelines)

Studies



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REQUESTED STUDIES ? ↓

VIVLI-LISTED AND PROVISIONED STUDIES

A Randomized, Double-blind, Single Migraine Attack, Placebo-controlled, Parallel-group Multicenter Study to Evaluate the Efficacy and Tolerability of Trexima (Sumatriptan S...

Study ID: NCT00329255 Sponsor ID: TRX105852
Data Request ID:
Data Contributor: GlaxoSmithKline IRP/Approver: Wellcome Trust

Data to be loaded after approval

Remove x

VIVLI-LISTED STUDIES PROVISIONED BY EXTERNAL PROVIDERS

No Studies Found

STUDIES, DATA OR TOOLS NOT LISTED ON VIVLI **Add +**

No Studies Found

Statistical Analysis Plan



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GENERAL

Describe how you will analyze the requested clinical study data

Country/countries where the analysis will be conducted

Describe how you will analyze the requested clinical study data, including:

- The reasoning behind/criteria used for selecting a specific study (i.e. search criteria)
- If your proposed research involves studies from other sources, provide a full list of studies requested from other sources so the Data Contributors can see the full scope of the proposed research. Also please propose a brief plan on how to combine the results.
- Include discussion of descriptive, bivariate and multivariable analyses
- Any other planned advanced analyses (such as propensity score methods, Kaplan-Meier or ...)

Funding



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GENERAL

Is the proposed research being funded by research grants from government agencies?

Yes No

Is the proposed research being funded by employers through employment contracts?

Yes No

Is the proposed research being funded by additional contracts or consultancies?

Yes No

Is the proposed research being funded by commercial organizations?

Yes No

Other Information / Attachments



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Select Files

Drop files here

File Name

Size

Uploaded By

File Type

| File Name | Size | Uploaded By | File Type |
|-----------|------|-------------|-----------|
| | | | |

No files uploaded.

Attestation



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Certify Complete and Accurate

Please check the box below to indicate that you as the Lead Researcher certify that the information provided is complete and accurate, and that you assume full responsibility for the research.

 I certify the information provided is complete and accurate.

Data Use Agreement

Please note that all Data Requestors wishing to receive access to data must execute the Data Use Agreement (DUA) before the data can be provided. The DUA is the product of extensive negotiation with the organizations that contribute data to Vivli, and as such, the agreement is non-negotiable. The DUA form must be completed and signed and is available [here](#).

You can either fill out the DUA form and sign it digitally, or print it out, sign it and scan it as PDF. Once the DUA has been signed by your organization, please upload it using the Signed Agreements tab of this data request (visible once the data request is submitted).

If you have any questions regarding the DUA, please contact a Vivli admin at support@vivli.org.

If the Submit button is not enabled, look in each of the tabs on the left for a field outlined in red which indicates that a required field needs to be completed

Chat



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Open Chat ?

Communicate with Vivli staff, the Data Contributors, Independent Review Panel and the Research Team.

Send

Uploaded Files

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Select Files

Drop files here

| File Name | Uploaded By | Uploaded Date |
|-----------|-------------|---------------|
| | | |

No files uploaded.

Insert questions here or email support@vivli.org