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Data Request Example

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LEAD RESEARCHER

Invite user to access data request

Lead Researcher is also Statistician Researcher

First Name

Last Name

ORCID iD

Emai...

Position

Employer, Company, Research Institute...

Country

- Select an Option -

Education, including the degree, discipline and institution where the degree was granted, and professional qualifications that are relevant to the proposed research and are specific to clinical data analysis.

Character Count: 0/1000

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

VM Access Admin Approval Based on Approved DUA

DUA Approval Not Applicable

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STATISTICIAN RESEARCHER

Invite user to access data request

First Name

Last Name

ORCID iD

Emai...

Position

Employer, Company, Research Institute...

Country

- Select an Option -

Education, including the degree, discipline and institution where the degree was granted, and professional qualifications that are relevant to the proposed research and are specific to clinical data analysis.

Character Count: 0/1000

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

VM Access Admin Approval Based on Approved DUA

DUA Approval Not Applicable

ADDITIONAL RESEARCHERS

Add +

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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)

Lay Summary of the Research Request

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

Purpose of Analysis is required.

Outcome(s) is required.

PURPOSE OF ANALYSIS

- [ ] 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

OUTCOME(S)

Inform Patient Care Decisions

- [ ] Inform Patient Care Decisions

Algorithms / Code

- [ ] A.I. algorithm
- [ ] Algorithm for predicting treatment response

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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

08/15/2023

Estimated Analysis Completion Date

08/16/2023

DISSEMINATION AND PUBLICATION PLAN

Dissemination & Publication Plan

Provide references for all cited material (following APA guidelines)

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REQUESTED STUDIES

VIVLI-LISTED AND PROVISIONED STUDIES

A Multicenter, Open-Label Conversion of Valproate Monotherapy to Lamotrigine Monotherapy in Patients With Epilepsy

Study ID: NCT00043914 Sponsor ID: LAM40013
Data Request ID:
Data Contributor: GlaxoSmithKline IRP/Approver: Wellcome Trust

Data to be loaded after approval

Remove

VIVLI-LISTED STUDIES PROVISIONED BY EXTERNAL PROVIDERS

No Studies Found

STUDIES, DATA OR TOOLS NOT LISTED ON VIVLI

Add +

No Studies Found

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GENERAL

Describe how you will analyze the requested clinical study data

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

- The reasoning behind/criteria used for selecting a specific study (ie. 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 Cox

Country/countries where the analysis will be conducted

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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

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Other Information

Other Information

File Attachments

NO FILES IN PACKAGE

Select Files

Drop files here

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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.

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Open Chat

Communicate with stakeholders involved in this data request.

NO FILES IN PACKAGE

Select Files

Drop files here

Send