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**Lead Researcher**

- **First Name**
- **Last Name**
- **ORCID ID**
- **Position**
- **Email**
- **Company/Research Institute**
- **Country**

**Education**

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

**Ethical Considerations**

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

**Data Access/Use Agreement**

- Admin Approval Based on Approved DUA
- DUA Approval Not Applicable
Data Request Example

**STATISTICIAN RESEARCHER**

- **First Name**: [Input]
- **Last Name**: [Input]
- **ORCID ID**: [Input]
- **Email**: [Input]
- **Employer, Company, Research Institute**: [Input]
- **Position**: [Input]
- **County**: [Select]
- **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.
- **Conflicts of Interest**: Please list any real or potential conflicts of interest and describe how these will be managed. If none, please enter None.

**VII Access Administration**

- **DUA Approval Not Applicable**

**ADDITIONAL RESEARCHERS**

[Add +]
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?

Purposes of Analysis as 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

Outcomes as required:
- Algorithms / Code
  - AI algorithms
  - Algorithm for predicting treatment response
  - Code
  - Machine Learning
- Clinical Guidelines
  - Clinical guidelines
  - Overview from Title: First Sentence
STUDY DESIGN

Brief Description

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

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).
Main Predicted 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 (e.g., genders, age groups, ethnic groups)

PROJECT TIMELINE

- Target Analysis Start Date: 09/16/2023
- Estimated Analysis Completion Date: 09/16/2023

DISSEMINATION AND PUBLICATION PLAN

- Dissemination & Publication Plan: Provide references for all cited material (following APA guidelines)
Requested Studies

Vivli Listed and Provisioned Studies

A Multicenter, Open-Label Conversion of Valproate Monotherapy to Lamotrigine Monotherapy in Patients With Epilepsy
Study ID: NCT01042914
Sponsor ID: LAM40113
Data Source: Vivli
Data Type: Clinical Trial - Phase II/III
IRB Approval: Yes
Institution: Weill Cornell

Vivli Listed Studies Provisioned by External Providers
No Studies Found

Studies, Data, or Tools Not Listed on Vivli
No Studies Found
Data Request Example

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
Data Request Example

Other Information

File Attachments

NO FILES IN PACKAGE

Select Files

Drop files here
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 Requesters 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.
Communicate with stakeholders involved in this data request.