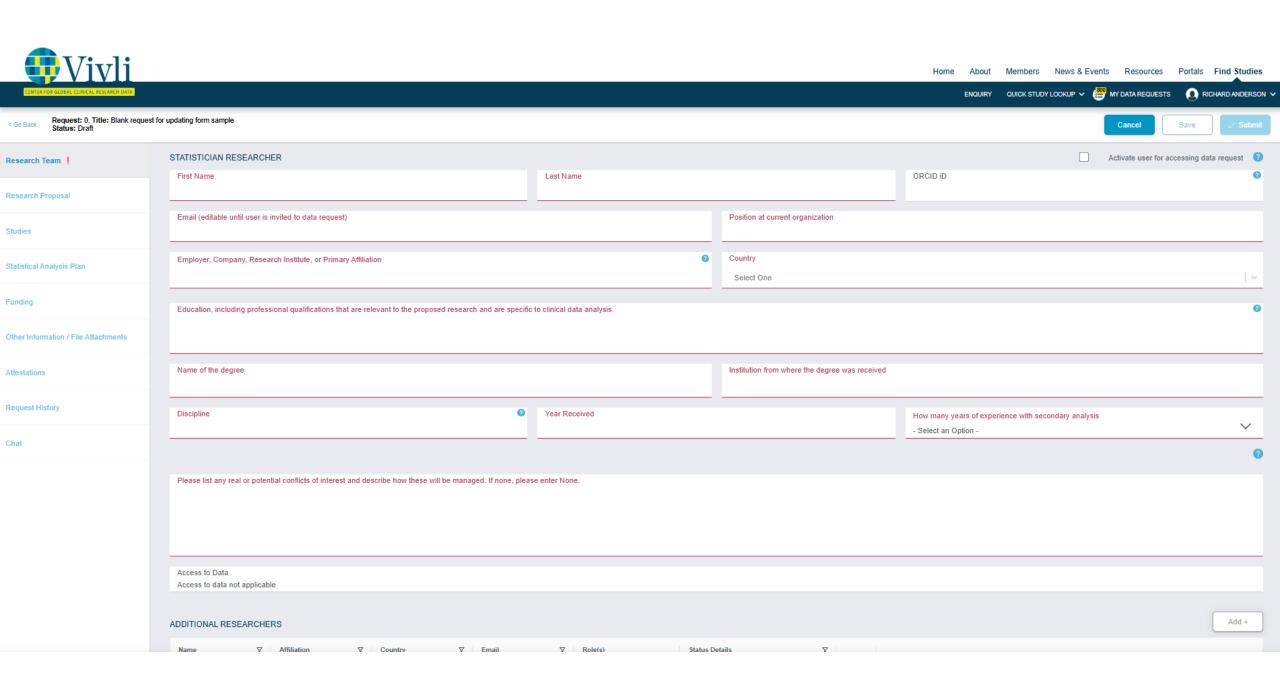


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Request: 0, Title: Blank request for updating form sample Status: Draft Cancel Research Team ! LEAD RESEARCHER Activate user for accessing data request Lead Researcher is also Statistician Researcher ORCID iD First Name Last Name Research Proposal Email (editable until user is invited to data request) Position at current organization Studies 0 Statistical Analysis Plan Employer, Company, Research Institute, or Primary Affiliation Country Select One Funding Education, including professional qualifications that are relevant to the proposed research and are specific to clinical data analysis. Other Information / File Attachments Attestations Institution from where the degree was received Name of the degree Request History Year Received Discipline How many years of experience with secondary analysis - Select an Option -Chat 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





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ENQUIRY QUICK STUDY LOOKUP V 🐸 MY DATA REQUESTS 🜘 RICHARD ANDERSON 🗸 Request: 0, Title: Blank request for updating form sample < Go Back Cancel Status: Draft Research Team GENERAL Title of the Proposed Research (Overall brief description, including the condition, about the proposed research question that is understandable by a general audience) Research Proposal Blank request for updating form sample Studies Lay Summary of the Research Request Statistical Analysis Plan Funding What are the specific aims/objectives of the proposed research, including the specific hypotheses to be evaluated? Other Information / File Attachments Attestations Request History PURPOSE OF ANALYSIS ② Purpose of Analysis is required. OUTCOME(S) Outcome(s) is required. Chat Inform Patient Care Decisions New research question to examine treatment effectiveness on secondary endpoints and/or within subgroup populations Inform Patient Care Decisions New research question to examine treatment safety Algorithms / Code A.I. algorithm Research that confirms or validates previously conducted research on treatment effectiveness Algorithm for predicting treatment response Code Research that confirms or validates previously conducted research on treatment safety Machine Learning Clinical Guidelines Preliminary research to be used as part of a grant proposal Clinical guidelines Summary-level data meta-analysis Designing Future Trials, Trial Protocol Clinical trial design Participant-level data meta-analysis Clinical trial patient selection / recruitment

Optimization of clinical trial parameters

(non-generative)

Generative Models / Large Language Models (LLMs) - ex: GPT, Generative Adversarial Networks (GANs), Diffusion models

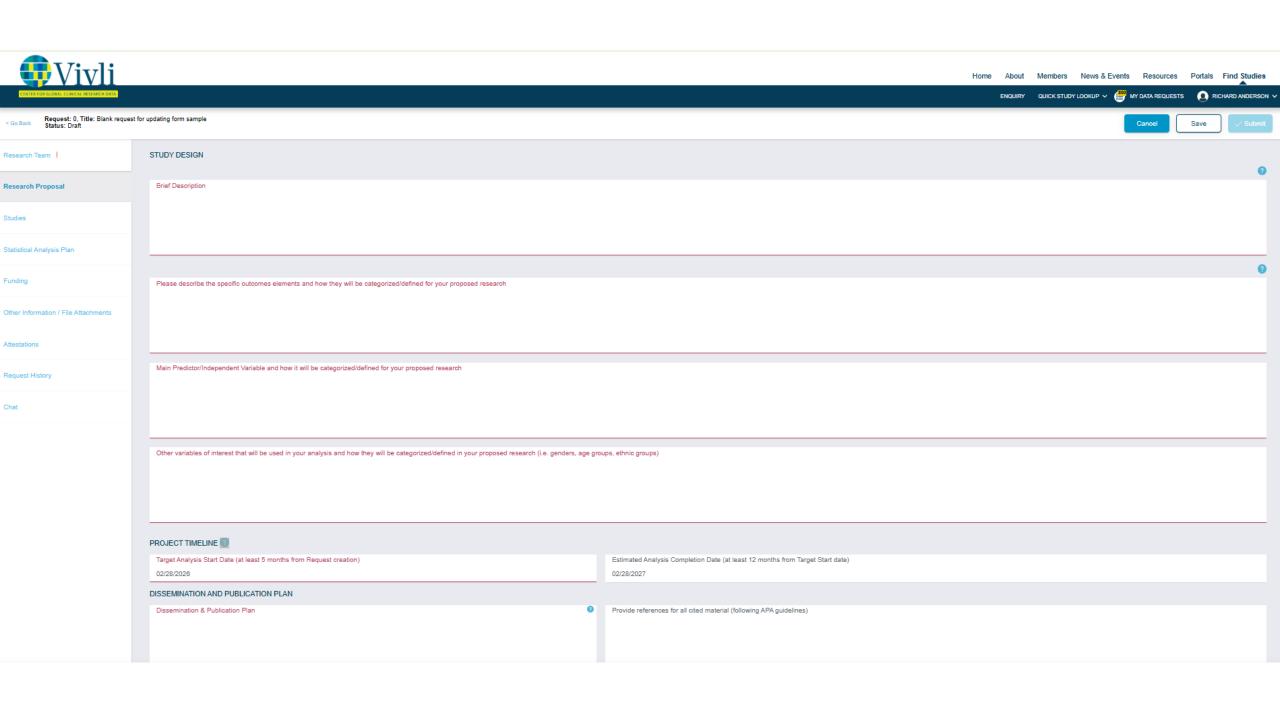




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Request: 0, Title: Blank request for updating form sample

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|--------------------------------------|---|--------|------|-------|
| Research Team | | | | |
| Research Proposal | Response required. Will your analysis include the use of existing trained models? * | | | |
| TO OCCUPANT TO POOL | | | | |
| Studies | ○ No | | | |
| | Yes | | | |
| Statistical Analysis Plan | Response required. | | | |
| Funding | Will you be exporting a model from the Vivli research environment? * | | | |
| Other Information / File Attachments | ○ No | | | |
| Other information / File Attachments | ○ Yes | | | |
| Attestations | | | | |
| | Response required. | | | |
| Request History | Please select how the data will be used * | | | |
| Chat | To validate (to confirm model performance) - Data is used to confirm that a model is performing as expected | | | |
| Citat | To train (to adjust existing model algorithim parameters) – Data is used to adjust algorithms parameters | | | |
| | To create a model – Data used to build a new custom model | | | |
| | | | | |
| | Response required. | | | |
| | Will the model form the basis of a commercial product? * | | | |
| | ○ No | | | |
| | Yes | | | |
| | | | | |



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Remove X

Request: 0, Title: Blank request for updating form sample Status: Draft

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Data already on platform

Research Team

Research Proposal

Studies

Funding

Statistical Analysis Plan

Other Information / File Attachments

Request History

Attestations

Chat

REQUESTED STUDIES 2



VIVLI-LISTED AND PROVISIONED STUDIES

A double-blind, randomized, stratified, placebo-controlled study of Ro 64-0796 (also known as GS4104) in children with influenza.

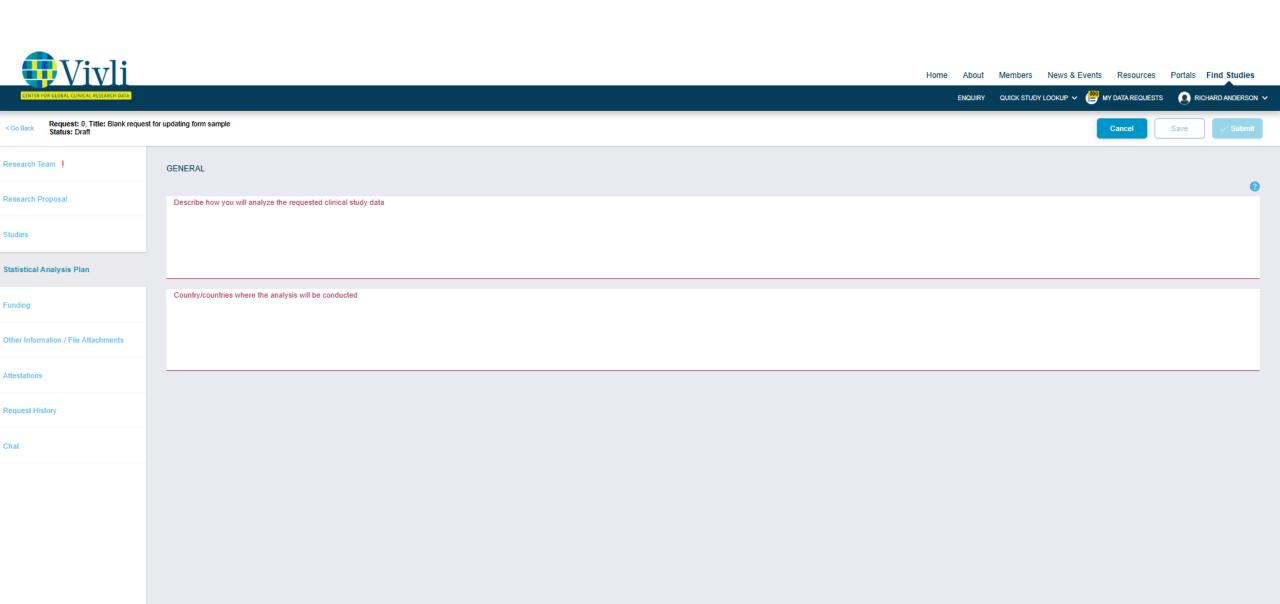
Study ID: WV15758 Sponsor ID: WV15758
Data Request ID:
Data Contributor: Roche IRP/Approver: Wellcome Trust

VIVLI-LISTED STUDIES PROVISIONED BY EXTERNAL PROVIDERS

No Studies Found

STUDIES, DATA OR TOOLS NOT LISTED ON VIVLI Add +

No Studies Found





<Go Back Request: 0, Title: Data Request Example Status: Draft

Research Team ! Research Proposal Studies

Statistical Analysis Plan

Funding

Other Information / File Attachments

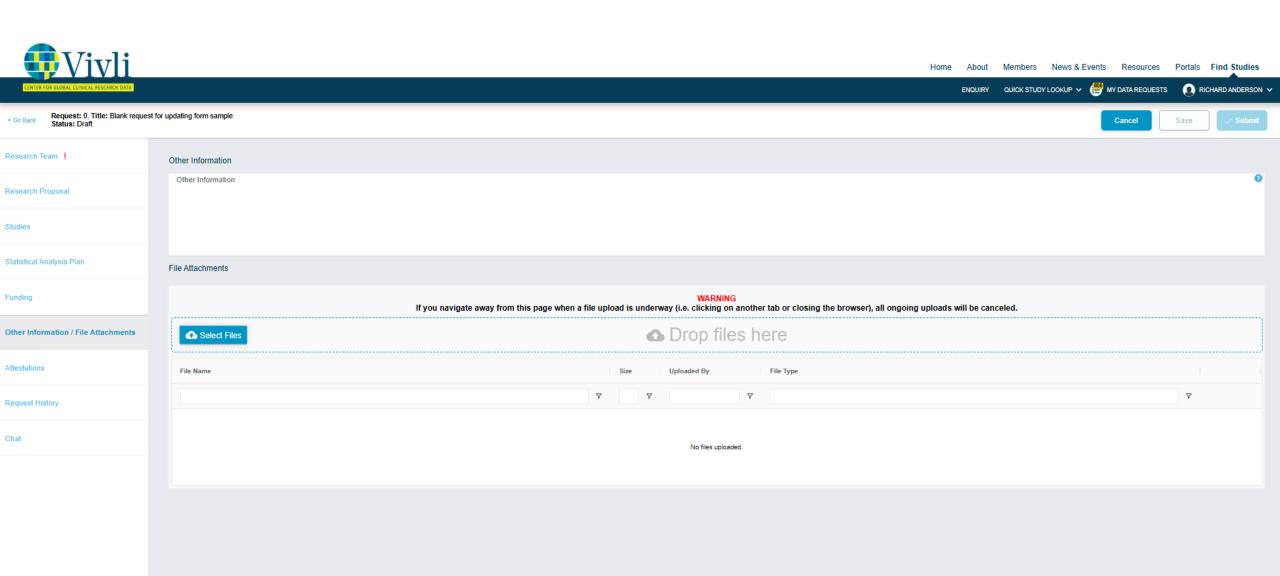
○ Yes ● No

Attestations

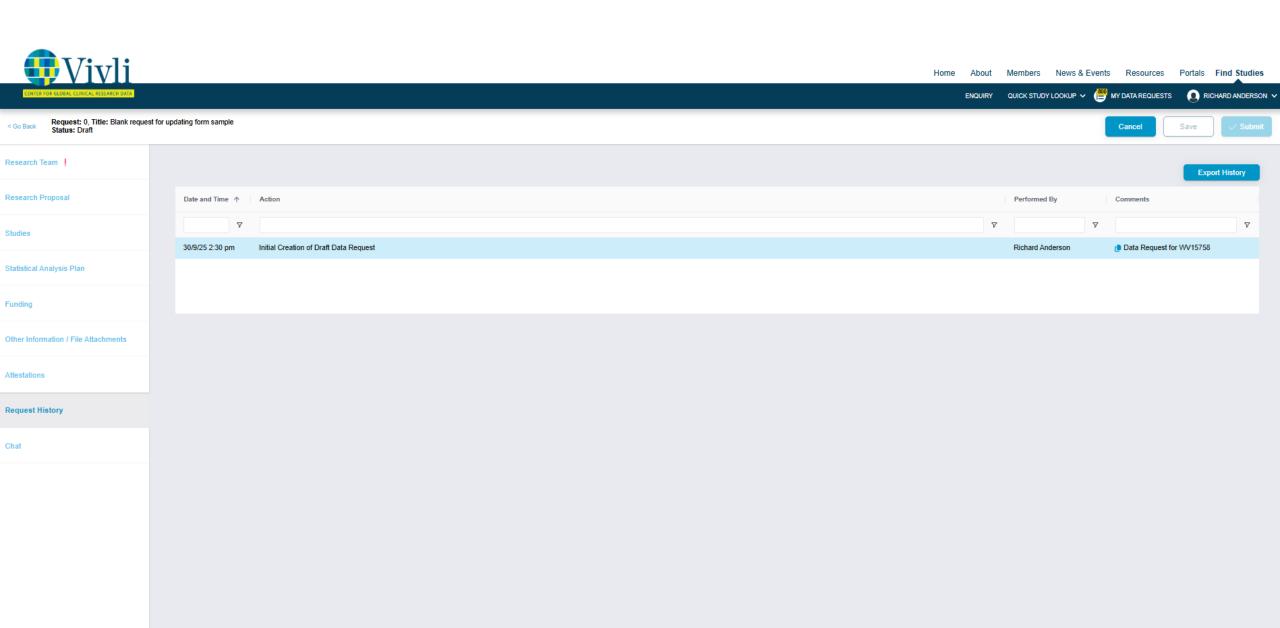
Request History

Chat

| GENERAL | |
|------------|---|
| Is the pro | posed research being funded by research grants from government agencies? |
| ○ Yes | ® No |
| Is the pro | posed research being funded by employers through employment contracts? No |
| Is the pro | posed research being funded by additional contracts or consultancies? No |
| Is the pro | posed research being funded by commercial organizations? |



| CENTER FOR GLOBAL CLINICAL RESEARCH DATA | ENQUIRY QUICK STUDY LOOKUP V (29) MY DATA REQUESTS (20) RICHARD ANDERSO |
|---|---|
| Go Back Request: 0, Title: Blank reques Status: Draft | t for updating form sample |
| search Team ! | Certify Complete and Accurate |
| search Proposal | 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. |
| udies | Certify the information provided is complete and accurate. |
| atistical Analysis Plan | 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 Vivii, and as such, the agreement is non-negotiable. The DUA |
| nding | 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). |
| her Information / File Attachments | If you have any questions regarding the DUA, please contact a Vivli admin at support@vivli.org. |
| testations | |
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Cancel

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Save

Request: 0, Title: Blank request for updating form sample Status: Draft

Research Team Research Proposal Studies Statistical Analysis Plan Funding Other Information / File Attachments Attestations Request History Chat

Open Chat Communicate with Vivli staff, the Data Contributors, Independent Review Panel and the Research Team. Uploaded Files WARNING If you navigate away from this page when a file upload is underway (i.e. clicking on another tab or closing the browser), all ongoing uploads will be canceled. ♠ Drop files here Select Files File Name Uploaded By Uploaded Date ∇ ∇ ∇ No files uploaded.