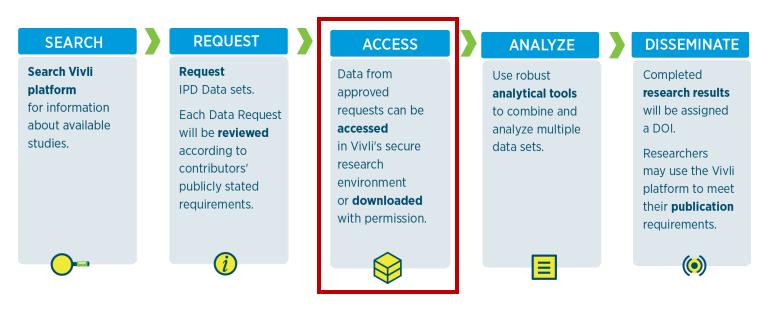


# How to Access Data for Analysis and Publication Process

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# 1.0 Data Access Introduction

Your team's data request has been approved and your Data Use Agreement has been executed. Now that one of the studies from your request has been uploaded, your research team is able to begin analyzing the data. Depending on the access options available on your requested study(ies), you can either download the data (<u>See section 12.0</u> for more information) or you can access the data via a secure, cloud-based, isolated workspace known as a research environment.

- The Research Environment is a where users have access to various tools to analyze their data and conduct their research, including <u>R</u>, <u>Python</u>, <u>Jupityr</u>, the Microsoft Office suite, STATA, and <u>SAS (Academic-license only)</u> depending on the type of the Research Environment selected. A <u>complete list of the software</u> included and versions in the Research Environment can be found on the Vivli website resources page.
- The Vivli Research Environment can also accept your own preferred analytical tools if you can provide the license key for the tool, or if it is an open-source tool. Your research team can load R packages from the CRAN repository yourself R packages in other locations (such as Github) or Python packages can be added by the Vivli Team

# 3.0 Getting Started

- Once one of your requested studies data packages has been uploaded, you will be notified via email that your data is available.
- At this point, you may initiate the Secure Research Environment. Please note that only the "Owner" (normally the originator) of the data request can start the Research Environment.
- Note, dependent on when you initiated your environment, your background may appear either white (initiated before Summer 2024) or blue (initiated after Fall 2024). The software and tools are the same in both environments.

Older research environment desktop



Updated research environment desktop



#### 3.1 Locating the Data Request and seeing how many studies are available for analysis

- 1. Click on My Data Requests in the top right corner of the screen:
  - Figure 2 My Data Requests

<b>Viv</b>	li	Home About	t Mombers	News & Events Re	sources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARC	NBOA		>	• 🗸 😬 MY DATA REQUES	rs 🗕 data requester 🗸
A Dashboard	Welcome, Data Requester!	Organization Memberships			
	This is your view of Vivli at a glance. Here you can view your organizational memberships and roles, any pending requests that require your approval, as well as any studies awailing Data Package upload from your organization. You can also generate metrics for data requests involving your organization's studies. If you have any questions, please contact Vivli Support. Thanks! Data Requests Awaiting My Approval No Data Requests Awaiting Approval				
	Studies Awaiting Data Package Upload Only data contributors are authorized to upload IPD data				
	Only data commonions are admonized to upload in D data				

2. The request will appear under **Active**:

Vivli			Home	About	Members	News & Events	Resources	Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA				QUICK	STUDY LOOKUP	✓ <sup>166</sup> MY DATA REG	QUESTS	DATA REQUESTER 🗸
My Data Requests (166)							2549	
Draft     Active     Not Approved	Withdrawn	Archived						
ASCENDING MULTIPLE-DOSE SAFETY, TOLERANCE, F	PHARMACOKINE	ETIC, AND F						
Vivli ID: 00002549								
Status: At least one Data Package Provided and Available								

Figure 3 - Fulfilled Requests

3. If the request has been approved, the Data Use Agreement validated and the requested data from at least one requested study is available, the request will appear under "Active", with a status of "At least one Data Package Provided and Available"

CENTER FOR GLO	DBAL CLINICAL RESEARCH DATA				Home	About	Members STUDY LOOKUP	News & Events	Resources	DATA REQUESTE
My Da	ta Reques	ts (166)						e	254	
Oraft	Active	Not Approved	Withdrawn	Archived						
ASCENDI		E SAFETY, TOLERAN		INETIC. AND F						
Vivli ID: 0000		,								
	least one Data Packag	e Provided and Available								
Status: At										
Status: At										
Status: At										

Figure 4 – At least one Data Package Provided

To determine **which** studies have been loaded, open the request and click on the Studies tab:

Vivl <sup>2</sup>	Home About Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DA	QUICK STUDY LOOKUP 🗸 👹 MY DATA REQUESTS 👔 DATA REQUESTER
< Go Back Ascending	g Multiple-dose Safety, Tolerance, Pharmacokinetic, and Pharmacodyna
Studies	
Attachments	VIVLI-LISTED AND PROVISIONED STUDIES
Request History	Ascending Multiple-dose Safety, Tolerance, Pharmacokinetic, and Pharmacodynam Pi: Sponsor: BMS Study ID: AEGR-CV145-002 IRP/Approver: Novelion Therapeutics Awaiting Data Package upload by Data Contributory Data Contributor: Novelion Therapeutics IPD Uploaded: No
Signed Agreements	
Safety Concerns	A Patient Preference Evaluation Study of Fluticasone Furoate Nasal Spray and PI: Sponsor: GlaxoSmithKline Study ID: NCT02397915 IRP/Approver: Wellcome Trust Data Request Sponsor ID: 201474 Data Contributor: GlaxoSmithKline IPD Uploaded: No Awaiting Data Package upload by Data Contributor
Chat	An Open Label Study of the Efficacy and Safety of Re-treatments With Rituxima
Research Team	Pt: Spensor: Hoffmann La Rodie Sudy ID: NOT2097745 IRP/Approver: Wellcome Trust Data Package Provided to Requestor Data Request ID: 00002549 Sponsor ID: WA17531 Data Contributor: Roche IPD Upleaded: No
Research Environment	An Open Label, Non-comparative Study To Evaluate Parasitological Clearance Ra
Request Details/Print View	PI: Sponsor: Plizer Study ID: NCT01103713 IRP/Approver: Plizer Inc. Data Request ID: 00002549 Awaiting Data Package upload by Data Contributor Sponsor ID: A0661201 Data Contributor: Plizer Inc. IPD Uploaded: No
	VIVLI-LISTED STUDIES PROVISIONED BY EXTERNAL PROVIDERS
How To Guide Privacy Co	okie Policy EEA Disclosure Policy Contact Us e Copyright 2017 - 202

*Figure 5 – Data Package Provided to Requestor* 

If you start the research environment before all of the data is available, then as additional data is made available, it will not automatically appear in the research environment– you will have to request that the new data be added to your research environment– see section "6.2 How to add additional data to your research environment as it becomes available"

### 3.2 Starting the Secure Research Environment

Once you have opened your request, click on the **Research Environment** tab on the left-hand side of your screen to begin initiating the environment. Please read and acknowledge the Vivli Terms of Use for the Research Environment. Click on **Sign Now**:

<b>Wivli</b>	Home About Members News & Events Resources	Find Studies				
CENTER FOR GLOBAL CLINICAL RESEARCH DATA	QUICK STUDY LOOKUP 🗸 📛 MY DATA REQUESTS	DATA REQUESTER 🗸				
< Go Back Ascending	Multiple-dose Safety, Tolerance, Pharmacokinetic, and Pharmacodyna	Print				
Studies						
Attachments						
Request History						
Signed Agreements	Data Package available for 1 of 4 studies in the Data Request					
Safety Concerns						
Chat	YOU MUST E-SIGN THE VIVLI TERMS OF USE AGREEMENT					
Research Team	TO CONTINUE					
Research Environment	Sign Now					
Request Details/Print View						
How To Guido Brivooy Coo	kie Policy EEA Disclosure Policy Contact Us	@ Consider 0017 - 0001 18-5				
How to Guide Privacy Coo	Reforcy LLA Disclosure Forcy Contact us	© Copyright 2017 - 2021 Vivli				

Figure 7 – Start Signing Process

4. The following pop-up window will appear:

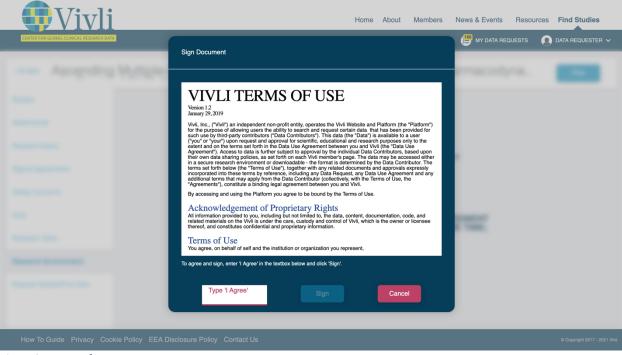


Figure 8 - Terms of Use Pop-Up

5. Type I Agree in the dialogue box and click Sign.

Each Team member accessing the Research Environment must Sign the Terms of Use when they first access the Research Environment.

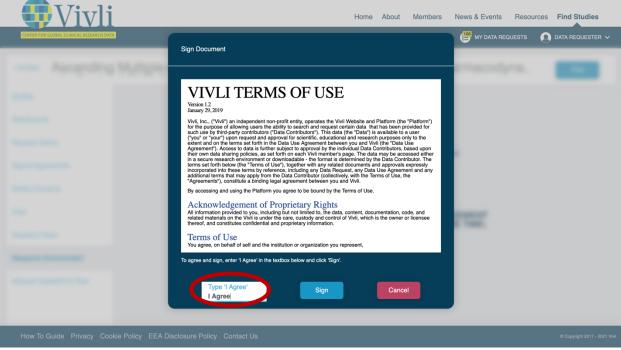


Figure 9 - Terms of Use Read, Acknowledge and Sign

#### 3.3 Secure Research Environment Options

Studies	Data Package available for 1 of 1 studies in the Data Request							
Attachments	Information on using the Research Environment is available here.							
Request History	RESEARCH ENVIRONMENT DETAILS							
Signed Agreements	Advanced Options:	Standard Environment	Premium Environment					
Safety Concerns	Initial Cost	No charge for 365 days	No charge for 90 days					
Chat	After Initial Period	\$12/Day after 365 days, 2 concurrent logins	\$25/Day after 90 days, 2 concurrent logins					
Research Team	Machine Size	2CPUx7GB	4CPUx14GB					
lesearch Environment	Jupyter Notebook	•	*					
	Python, R	v	*					
Public Disclosures	STATA							
Request Details/Print View	Academic license for SAS. Alternative pricing applies for industry users of SAS, email support@witing for details.							
		Start Standard Environment	Start Premium Environment					

1. After you agree to the Terms of Use, the following options window will appear:

2.. Most research teams find that the standard environment meets their needs. If you need a larger environment, the Premium will normally suffice.

2b. If you need a dramatically larger environment, choose Advanced Options for additional environment size types. Please note that if you choose a larger size, you cannot move to a smaller size, but you can start with a smaller size and upgrade to a larger size research environment. In addition, the no charge period will change based upon the larger machine size that is used and will take into account the no charge time already provided. For example, if you move from a standard to a premium, after 90 days, you will not be entitled to a further 90 days of no charge.

Data Package available for 1 of 1 studies in the Data Request								
Once the machine is started, the request cannot be sent back for revision								
Before starting the Research Environment, read the short introduction here.								
RESEARCH ENVIRONMENT DETAILS								
Advanced Options: 🗹	Premium Environment	Large Environment	Compute Optimized Environment	Memory Optimized Environment	Compute Optimized			
Initial Cost	No charge for 90 days	\$65/Day	\$125/Day	\$275/Day	\$220/D			
After Initial Period	\$25/Day after 90 days, 2 concurrent logins	\$65/Day	\$125/Day	\$275/Day	\$220/D			
Machine Size	4CPUx14GB	16CPUx64GB	32CPUx64GB	64CPUx128GB	64CPUx5(			
Jupyter Notebook	*	*	*	*	*			
Python, R	~	*	*	*	*			
STATA								
Academic license for SAS. Alternative pricing applies for industry users of SAS, email <u>support@vivil.org</u> for details.								
	Start Premium Environment	Start Large Environment >	Start Optimized Environment	Start Memory Optimized Environment >	Start Memory Optimized (			

4. Select the research environment appropriate for your needs and check the boxes as desired for the software you require and click the button "Start Environment".

As you consider which environment to choose, if you don't want to use SAS, but if the data is provided as SAS data, the system includes three R-studio packages that can help:

- Foreign
- Haven
- SAS7bdat

When you click the "Start..." button, your selection is final, and the provisioning will begin. For more information, contact <u>support@vivli.org</u>.

#### 3.4 Initiating the Secure Research Environment

1. After selecting the size of the environment, the data will be provisioned into the Secure Research Environment. While provisioning is taking place, the following screen will appear:

Uivli	Home About Members News & Events Resources Find Studies							
CENTER FOR GLOBAL CLINICAL RESEARCH DATA	QUICK STUDY LOOKUP 🗸 📑 MY DATA REQUESTS 💽 VIVLI INFO 🗸							
Coo Back Efficacy of erythromycin in prevention of anthrax.								
Studies	Data Package available for 1 of 3 studies in the Data Request							
Attachments	For more information on starting and using the Research Environment click here to view it in another tab or download,							
Request History	Your Research Environment is Provisioning							
Signed Agreements								
Safety Concerns								
Chat								
Research Team								
Research Environment	0							
Request Details/Print View	This is a long-running operation and may take a while							
	If you think something has gone wrong, you can always <u>Retry Provisioning</u>							
-	kie Policy EEA Disclosure Policy Contact Us e Capyright 2017 - 2019 Vide							

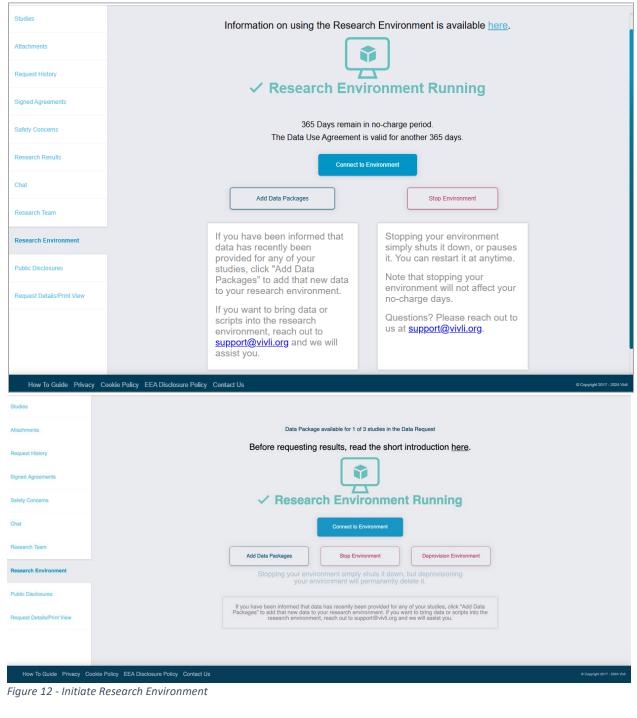
Figure 11 - Provisioning notification

2. While the system is provisioning, you can exit this screen and allow the provisioning to proceed in the background. You will receive an email when the process is complete. This process typically takes 20-25 minutes.

# 4.0 Access to the Environment

#### 4.1 Accessing the Secure Research Environment

1. Once the provisioning is complete, you will see the screen below:



2. Then click on the box "Connect to Environment"

2. Connect to Environment

3. After clicking on "Connect to environment," the research environment will load in a new browser tab.



4. The secure research environment home screen opens automatically:

Figure 16 - Secure Research Environment Home Screen

• Note that the first time you connect to the Research Environment, the system will be slower than normal as the system does some first-time configuration.

#### 4.2 Working in the Research Environment

#### 4.2.1 Finding your data and using space

#### 1. Open Disk V:

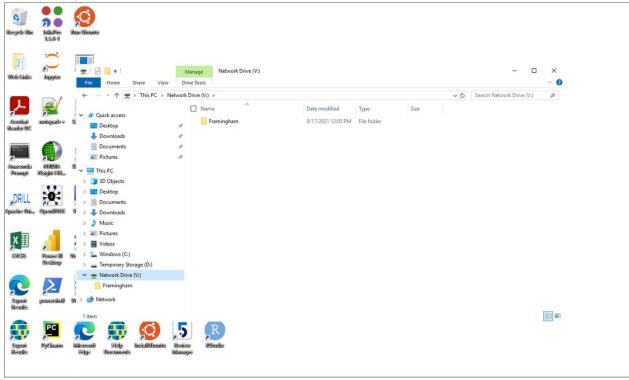


Figure 24 - Open source data

In the Network Drive (drive V: ) you will find one folder for each study that has been provided –the folder name will be the sponsor-assigned ID. Unlisted studies will appear with the ID you used to request the study. Within the ID, some characters that are not allowed for filenames will be replaced by an underscore character, including ∧:\*?\*<>|

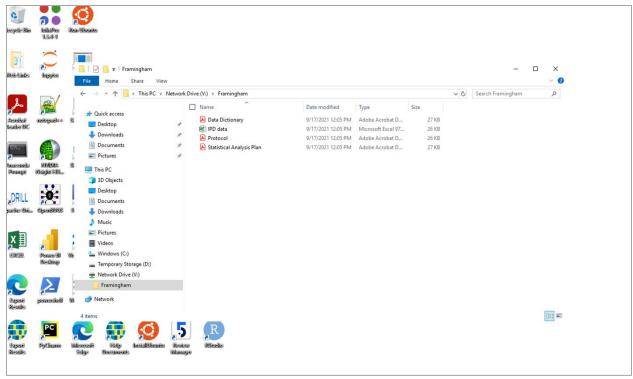


Figure 25 - Open source data – files

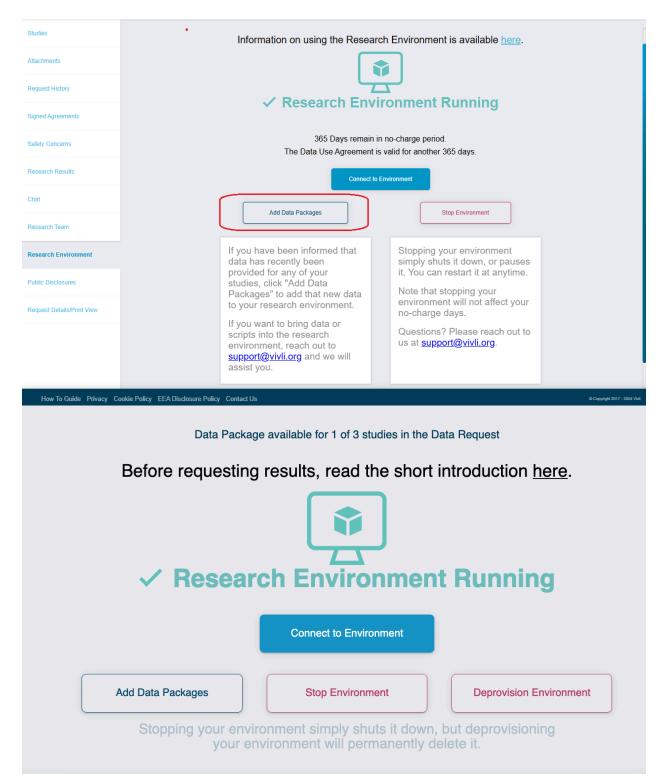
# Do not place any important data onto the temporary storage drive D:- when the machine is stopped, even temporarily, the temporary storage will be deleted.

If the data has been provided in the form of a zip file, you can open the file as though it were a folder, and read the files in the zip file, but you will not be able to modify them in-place. To modify the files, you will need to copy the file(s) to a new, un-zipped folder. Alternatively, if you drag the icon representing the zip file onto another folder, 7-zip will offer to extract all of the files.

Additional network space will expand as you add files, up to a terabyte. We recommend that you not put anything but transient data onto C: drive. Data on the V: drive will be placed into long-term storage after you have completed your analysis. If you have questions about the data and what has been provided, use the Chat function within the Vivli platform or email <u>support@vivli.org</u>; Please direct questions about the source data to the data contributors. Responses to questions about the source data is at the discretion of the data contributor.

#### 4.2.2 If you started before all of the data was available

If you have decided to start before \*all\* of the data is available, then when additional data is provided by the contributor, it won't be added to your research environment automatically. <u>See Section 6.0</u> for instructions on how to load newly provided data.



#### 4.2.3 Additional Tips for working in the Research Environment

From the research environment tab, you can review the (1) remaining no-charge period left in the environment and (2) the number of valid days remaining for the Data Use Agreement (DUA). –

- The no-charge period refers to how many days remain before billing will start for the environment selected (refer to Section 11.0 Paying for the Research Environment).

- The DUA is valid for one year from the date it is signed and a request for extension will be required to continue analysis beyond this date (refer to Section 10.0 Extensions to the Data Use Agreement via the Data Request Progress Report).

_ • Vivli	Home About Members News & Events Resources Portals Find Studies
CENTRATION REGISTAL CONCERNMENTATION DATA	ENQUIRY QUICK STUDY LOOKUP 🗸 📛 MY DATA REQUESTS 🚺 RICHARD ANDERSON
Co Back Request: 49046, Title: Stan Test Upbaci part the second Status: All Data Packages Provided and Available	
Studies	Information on using the Research Environment is available here.
Attachments	
Request History	✓ Research Environment Running
Signed Agreements	• Resource Environment Running
Safety Concerns	365 Days remain in no-charge period. The Data Use Agreement is valid for another 58 days.
Chat	Connect to Environment
Research Team	Add Data Packages Stop Environment
Research Environment	
Public Disclosures	If you have been informed that Stopping your environment data has recently been simply shuts it down, or pauses provided for any of your it. You can restart it at anytime.
Request Details/Print View	studies, click "Ádd Ďata Packages" to add that new data to vour research environment. environment will not affect your
	If you want to bring data or scripts into the research environment, reach out to support@vivli.org. Use the second of the second

#### 4.3 Disconnecting from the research environment

• When you are done working with the research environment, unless you are leaving a longrunning analysis running, we recommend that you Sign Out – this will free up memory and will allow other members of the research team to use the research environment.

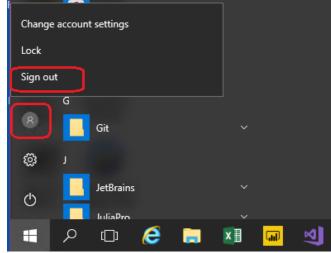


Figure 17 – How to Sign-out

• There is also an option to Sign Out by selecting the SignOff icon from the desktop.



• Only two team members can use the Secure Research Environment at a time. If a third member tries to log on, they will be given the option of disconnecting a team member:

	Windows sign-in X Select a user to disconnect so that you can sign in. There are too many users signed in → vm489948563\vivliuser5712 Active	
	→ vm489948563\vivliuser9582 Active Cancel	
C	mey contract os	e copyrgite cont - conte vite
🖬 🔿 📄 🥭 🥥 Administr 🌀 Types of D 💿 Vivli -	Goo 🝥 🛛 Goston Int 🖉 Document 🧲 Nauvya_Sr	

Figure 18 - too many users signed on

Figure 19 - disconnecting user

• The user in the research environment will receive the following message:

S connectio	n (15) - 40.122	2.72.85 - Rem	note Desktop Connection							- 🗆 X
Racycle Bin	RetBrains PyCharm C	R-Sarvar-A.								
ana	JuliaPro - Comman	Rsturlie		re View	3be-40d1b54c23ce IA (E:) ≥ Source Data ≥ c6f0a300-b9-	144-4841-93be-40d1b54c23ce		~ U	- C X ~ 2 Search c6f0a300-b944-4841-9 /2	
Links	kino iter Juliaite.	SAS 9.4 (Firefish)	<ul> <li>Quick access</li> <li>Desktop</li> <li>Downloads</li> <li>Documents</li> </ul>		IG Data Dictionary Documentation ig Protocol	Date modified         Type           9/26/2018 4:58 PM         PDF File           9/26/2018 4:58 PM         PDF File	Size 119 KB 179 KB			
Analytics Labs	Jupyter Notebo	SAS Study 3.71	<ul> <li>Pictures</li> <li>Screenshots</li> <li>This PC</li> <li>Network</li> </ul>	* =	Remote Desktop Do you want to allow vm48 Click OK to disconnect you					
	Alterescult Azere Po	State5: 1 (64-50)			No action will disconnect y	your session in 30 seconds.	ок	Cancel		
	NVIDIA Noight HU.	Visual Obje Tagging Te								
Ewince	Power H Deskiop		6 items 1 item selecte	d 1.35 MB					() () () () () () () () () () () () () (	· · · · · · · · · · · · · · · · · · ·
How	To Guide	Privacy	Cookie Policy EE	A Disclosu	re Policy Contact Us					© Copyright 2017 - 2018 Vivli
		_	user notific		Sticky XII Excel WII Word	📗 Untitle 💿 (•) Metr 📢	🕽 Vivli 🧕 Skype f (	🖻 Inbox 🖳 connec	🩋 Vivli-Ze 🔨 ា 🛋 🌈	d× 🗗 11:28 AM 📮

• The user in the research environment must click **OK** to disconnect and allow their team member access. Save your work regularly, and especially before disconnecting.

#### 4.4 Team Access to Secure Research Environment

- All members of the research team working in the research environment must have a signed Data Use Agreement (DUA). If a research team member is part of the same institution as the Principal Investigator, they are covered under the Principal Investigator's DUA. If a team member is from another institution, they must sign a DUA before accessing the research environment.
- The data request administrator may add team members to access the data request, which also grants access to the Research Environment.
- To add a user, the user must first create an account using the "Sign-up" link on the initial page. For information on creating an account, see the <u>User Quick Start Guide</u>
- Once the Research team member creates a Vivli Account, the data request administrator may activate the team members to access the data request and the research environment.

#### Figure 21 - change user notification

• When the research team is activated to access the data request, the request will appear on their "My Data Requests" screen and they will be given access to the data request. Once Vivli has validated that they are covered by a signed Data Use Agreement, they will be given

access to the research environment.

Uivl	i	Home About Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH C		ENQUIRY QUICK STUDY LOOKUP 🗸 👹 MY DATA REQUESTS 💽 DATA REQUESTER 🗸
< Go Back Request: 48056, Title Status: At least one D	: Liz - testing study added to request in process ata Package Provided and Available	Print
Studies	RESEARCHERS	Add +
Attachments Request History	Liz Test (Lead Researcher / statistician / data request administrator)	Emotive Team Member Activate Member for Access to Data Request
Signed Agreements	Jacob Wiley (ADDITIONAL RESEARCHER)	
Safety Concerns		
Research Results		
Research Team		
Research Environment		
Public Disclosures		
Request Details/Print View		
How To Guide Privacy Co	aokie Policy EEA Disclosure Policy Contact Us	@ Copyright 2017 - 2024 Vv4

• To remove a team member who is no longer involved in the project, click on the three dots to the right of their name, and click on "Remove Team Member"

Studies Attachments	RESEARCHERS	Remove Team Member
Request History	Liz Test (LEAD RESEARCHER / STATISTICIAN / DATA REQUEST ADMINISTRATOR)	Ac Transfer Administrator Role to Member
Signed Agreements	Jacob Wiley (ADDITIONAL RESEARCHER)	Access Granted DUA Approval Required
Safety Concerns		
Research Results		
Chat		
Research Team		
New 25 Environment		
Public Disclosures		
Request Details/Print View		

Figure 22 - change user notification

• Reminder: you are unable to change the Principal Investigator or the Lead Statistician. If you need to change the statistician or Principal Investigator, contact <a href="mailto:support@vivli.org">support@vivli.org</a>.

#### 4.5 Adding additional software, tools and data during your analysis

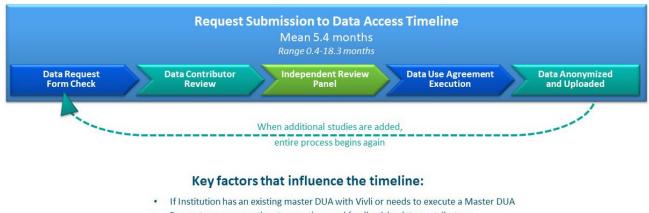
During the course of your analysis, if you find that you need to load R scripts from the CRAN repository, you can do that yourself using the following steps:

- To install a CRAN package in R, use the install.packages() function. This simple command downloads the package from a specified repository (by default, CRAN) and installs it on your machine.
- If your research environment was provisioned before June 1<sup>st</sup>, 2023, please reach out to the Vivli Technical Team via <a href="mailto:support@vivli.org">support@vivli.org</a> and we will complete a one-time update to point your machine to the latest replica of CRAN.

If you need any other R or Python scripts not pre-installed in the Vivli Research Environment, reach out to Vivli via chat or <u>support@vivli.org</u> and let them know the names of the additional scripts that you require and they will add them to your research environment. Any scripts that you have written or github packages can be emailed to <u>support@vivli.org</u> and they will be added to the research environment; please include the request number of your environment to help Vivli locate the correct environment.

If you have additional software, please email Vivli at <u>support@vivli.org</u> with the name of the software and any license key, if required. If this software was not included in your request, Vivli will need to reach out to the data contributors involved and ask for their agreement to include this software or data.

If you wish to add additional study data that is available on Vivli after your request is approved, you will need to submit a new request. Please note that this process can take an additional 2-5 months for review, approval and uploading of the additional data. Vivli strongly recommends that all studies needed for analysis are included before submitting your data request.



- Requesters response time to questions and feedback by data contributors
- Number of studies being requested

# 5.0 Stopping and Starting the Research Environment

#### 5.1 Stopping or Pausing the research environment

- Stopping the research environment is like powering off your laptop: nothing is lost, and the environment can be restarted quickly (less than 5 minutes).
- If you expect to leave the research environment idle for several days or longer, we recommend that you stop the Research Environment.
- When you are ready to use the Research Environment again, you may restart it and the environment will be restarted with all the saved data as you left it.

To Stop the Research Environment:

- 1. Login to the Vivli Platform
- 2. Navigate to the Research Environment, and click on the button "Stop Environment":

Studies	Information on using the Research Environment is available here.			
Attachments				
Request History				
Signed Agreements	Research Environment Running			
Safety Concerns	365 Days remain in no-charge period. The Data Use Agreement is valid for another 365 days.			
Research Results	Connect to Environment			
Chat				
Research Team	Add Data Packages Stop Environment			
Research Environment	If you have been informed that data has recently been simply shuts it down, or pauses			
Public Disclosures	provided for any of your it. You can restart it at anytime. studies, click "Add Data Packages" to add that new data Note that stopping your			
Request Details/Print View	to your research environment. environment will not affect your no-charge days.			
	If you want to bring data or scripts into the research environment, reach out to support@vivli.org and we will assist you.       Questions? Please reach out to us at support@vivli.org.			
How To Guide Privacy C	iookie Policy EEA Disclosure Policy Contact Us ecopyriget 2017-2004 VAR			
Studies	Data Package available for 1 of 3 studies in the Data Request			
Attachments	Before requesting results, read the short introduction here.			
Request History				
Signed Agreements				
Safety Concerns Research Results	✓ Research Environment Running			
Chat	Connect to Environment			
Research Team	Add Data Packages Stop Environment Deprovision Environment			
Research Environment	Stopping your environment simply shuts it down, but deprovisioning			
Public Disclosures	your environment will permanently delete it.			
Request Details/Print View	If you have been informed that data has recently been provided for any of your studies, click "Add Data Packages" to add that new data to your research environment. If you want to bring data or scripts into the research environment, reach out to support@vivil.org and we will assist you.			

*Figure 31 – Stop Research Environment* 

This will stop the secure research environment.

# 5.2 Restarting the research environment

To restart the research environment:

- 1. Login to the Vivli Platform
- 2. Navigate to the research environment; the start button will be where the Stop button was located:

	Data Packane available for 3 o	of 5 studies in the Data Permet			
Data Package available for 3 of 5 studies in the Data Request					
		onment Stopped			
		n no-charge period. s valid for another 351 days.			
	Add Data Packages	Start Research Environment			
	If you have been informed that data has recently been provided for any of your studies, click "Add Data Packages" to add that new data to your research environment. If you want to bring data or scripts into the research environment, reach out to <u>support@vivil.org</u> and we will assist you.	Stopping your environment simply shuts it down, or pauses it. You can restart it at anytime. Note that stopping your environment will not affect your no-charge days. Questions? Please reach out to us at <u>support@vivli.org</u> .			
udies					
achments		Data Package available for 1 of 3 studies in the Data Request			
quest History		equesting results, read the short introduction here.			
gned Agreements					
esearch Results	Research Environment Stopped				
nat	Add Data Packa	ages Start Research Environment Deprovision Environment			
esearch Team	Startin	ng your environment will make it so you can connect, but			
esearch Environment		rövisioning your environment will permanently delete it.			
ublic Disclosures	If you have been informed that data has recently been provided for any of your studies, click "Add Data Packages" to add that new data to your research environment. If you want to bring data or scripts into the research environment, reach out to support@vivil.org and we will assist you.				
Request Details/Print View		and a set of the set o			
How To Guide Privacy Cookie	Policy EEA Disclosure Policy Contact Us	e Capyrie			

3. Click on **Start Research Environment** – the platform will tell you that the research environment is starting:

• Vivl	Home About Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH	ENQUIRY QUICK STUDY LOOKUP 🗸 🔮 MY DATA REQUESTS  🐧 DATA REQUESTER 🗸
< Go Back Request: 48056, Titl Status: At least one I	: Liz - testing study added to request in process sta Package Provided and Available Print
Studies	
Attachments	
Request History	Your Research Environment is Starting
Signed Agreements	
Safety Concerns	
Research Results	
Chat	
Research Team	٠
Research Environment	This is a long-running operation and may take a while
Public Disclosures	This is a long-running operation and may take a write
Request Details/Print View	If you think something has gone wrong, you can always Retry Starting
How to Guide Privacy 0	ookie Policy EEA Disclosure Policy Contact Us e Copyright 2017-2024 VMI

Figure 33 - Research Environment restarting

4. After approximately 5 minutes, the system will display the "Running" screen:

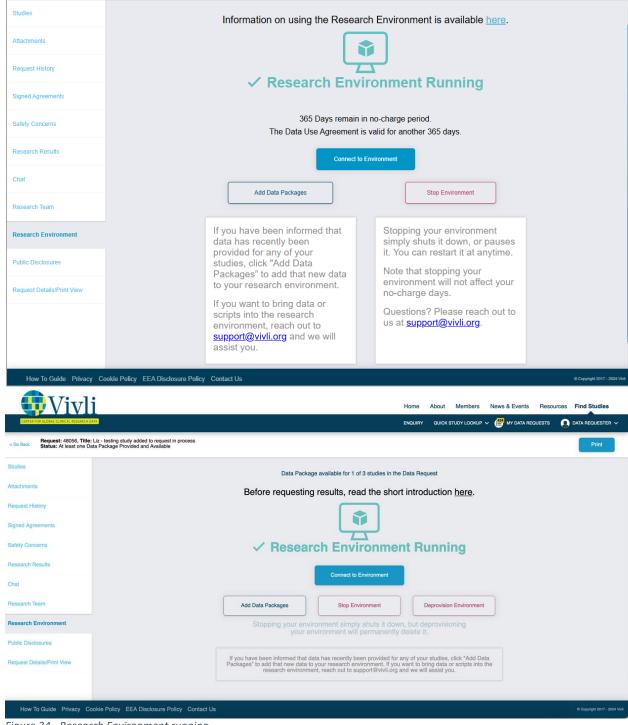


Figure 34 - Research Environment running

After restarting the machine, the first login may take a few minutes.

## 6.0 Adding Data Packages to the Research Environment

If you started the Research Environment before all the data was provided, then as additional data packages are loaded into the system, they will not be placed into your Research Environment automatically. You will however receive an email notification that additional data has been provided. When you are ready, you can ask the system to add the additional data packages to your Research Environment.

#### 6.1 How will you know when data is available?

As additional data packages are loaded into the system and made available, you will receive an email notification. Within the Vivli Platform, you can look at the "Active" tab, select the request you are interested in:

	QUICK STUDY LOOKUP	MY DATA REQUESTS	(1) DATA REQUESTER
ly Data Requests (166)			2549
Draft Octove Not Approved Withdrawn Archived			
ASCENDING MULTIPLE-DOSE SAFETY, TOLERANCE, PHARMAGOKINETIC, AND PHARMAGODYNAMIC STUDY OF BMS-201038 IN HEALTHY			
Vivil Ib detodste Status: Al least one Data Package Provided and Available			

Figure 35 – Available data packages

To determine *which* studies have been uploaded, click on the studies tab, and this page will provide a status of each study:

entributor
, ,
, ,
intributor
or >
ontributor >
m

Figure 36 – Available data packages

#### 6.2 How to add additional data to your research environment as it becomes available

Once you have determined that you are ready to ask the platform to load the additional data into your research environment, open the data request, select the Research Environment tab, and click on the "Add Data Packages" button:

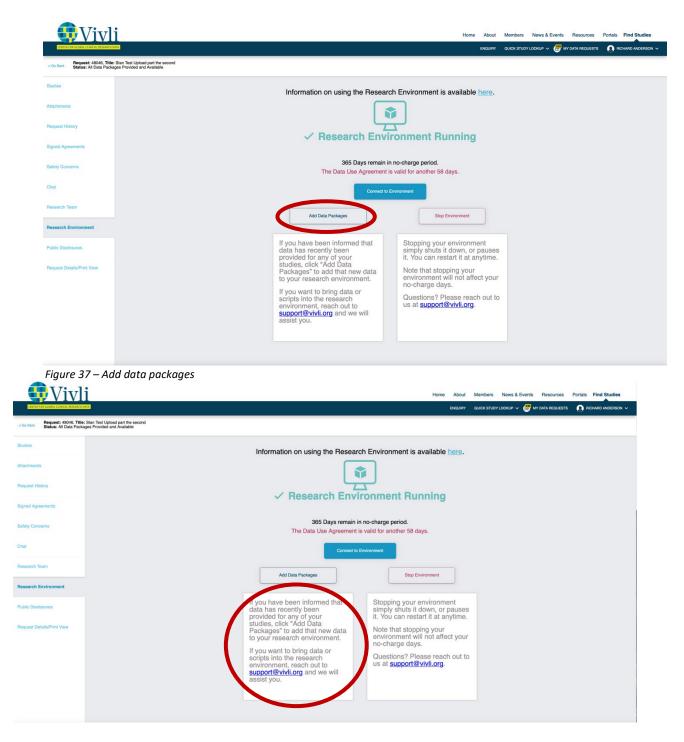


Figure 38 – Research Environment Adding data packages

#### 6.3 Refreshing previously loaded data

The Vivli platform determines what new data should be loaded by looking for an existing folder with the appropriate name: if it finds a folder in the Research Environment with the original folder name for the study, it will leave that existing folder alone. This means that:

- If you have started to make changes to previously loaded data (e.g. for harmonization), your changes will **not** be overwritten.
- If you would like a fresh copy of the data for any reason, you can simply rename the existing folder, and the system will load a fresh copy of the data package.
- If you rename a data folder for any other reason, the system will load a fresh copy of the data using the original folder name. If the extra copy is redundant, delete it.

# 7.0 Safety Concerns

If you discover any information regarding the safety or risks of a product related to their requested data, you must inform Vivli and the Data Contributor of this discovery within 24 hours, per the terms of the Data Use Agreement.

The steps for reporting safety concerns are as follows:

- 1. Login to the Vivli Platform.
- 2. Click on the "Safety Concerns" tab on the dashboard:

#### The following screen will appear:

Vivli	Home About Members News & Events Resources Find Studies
CENTER FOR CLOTAR CURICAL RESEARCH DATA	QUICK STLDY LOOKUP V 📛 MY DATA REQUESTER V
< Go Back Ascending	Multiple-dose Safety, Tolerance, Pharmacokinetic, and Pharmacodynamic Study of BMS-201038 in Healthy Volunteers
Studies	Supply your contact information and safety concern description below, then click 'Submit Safety Concern' to continue.
Attachments	Name Data Requester
Request History	Email Address Phone Number Datarequester vivil@gmail.com
Signed Agreements	Describe the Safety Concern
Safety Concerns	Submit Sately Concern
Research Results	
Chat	Previously Submitted Safety Concerns
Research Team	
Research Environment	
Request Details/Print View	
How To Guide Privacy Cook	kle Policy EEA Disclosure Policy Contact Us s copyon 2017-2021

Figure 39 - Safety Concerns tab

3. Complete the form and click on **Submit Safety Concerns**:

Vivli	Home About Members News & Events Resources Fin	nd Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DAT	SINA QUICK STUDY LOOKUP 🗸 📛 MY DATA REQUESTS 💽 DATA	A REQUESTER 🗸
< Go Back Ascending	ng Multiple-dose Safety, Tolerance, Pharmacokinetic, and Pharmacodynamic Study of BMS-201038 in Healthy Volunteers	Print
Studies	Supply your contact information and safety concern description below, then click 'Submit Safety Concern' to continue.	
Attachments	Name Data Requester	
Request History	Email Address Phone Number Datarequester.vivl@gmail.com \$55-555	
Signed Agreements	Describe the Safety Concern Safety concern	
Safety Concerns	Submit Safety Concern	_
Research Results		
Chat	Previously Submitted Safety Concerns	
Research Team		
Research Environment		
Request Details/Print View		
How To Guide Privacy Coo	Cookle Policy EEA Disclosure Policy Contact Us	Copyright 2017 - 2021 Vivi

Figure 401 - Submit safety concerns

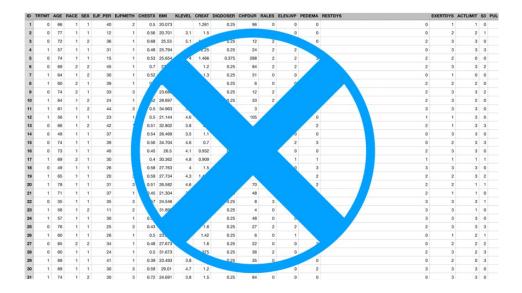
# 8.0 Exporting Research Results

If you have preliminary or final results from your analysis, you may request the ability to export summary results from the Secure Research Environment. Any custom scripts you may have developed during the research can also be exported with the results. A shortcut to submit this request is on the desktop.



When submitting your request to export results, please note that Individual Participant Data (IPD) requested from this study will need to remain in the Vivli secure research environment. As such, there should be no attempt to try to remove IDP data or re-identify individual participants in the study you are requesting. Per your signed Data Use Agreement, any results derived from your analysis in the Vivli Research Environment, which will be used in subsequent presentation or publication, should be obtained by submitting a request for removal of this data.

Results exports should not include any individual participant data in the results. Acceptable results export is limited to summary representation of data (e.g. means, standard deviations, counts), derived data (e.g. slopes, clin pharm parameters) data, analytical and representations of results (e.g. graph), scripts or programs that you developed in the Research Environment. The following graphic is an example of a request to export results spreadsheet, including IPD, which would not be approved.



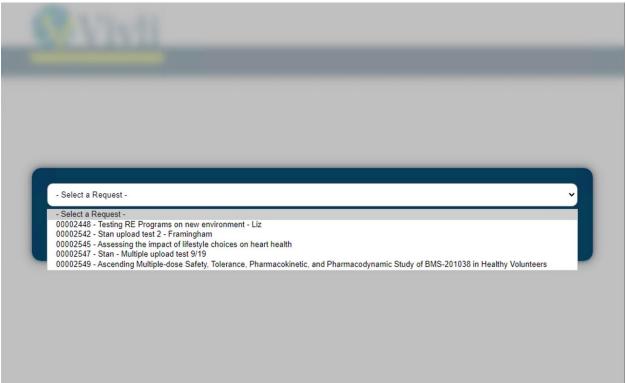
Please note that all results should be submitted through the results export option.

#### 8.1 Request to Export Results

- 1. From the Vivli Research Environment, double click the Export Results icon on the Desktop.
- 2. Enter the Vivli User credentials you use to log onto the Vivli Platform your username will be your email address.

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C To be most productive with Microsoft E	https://vivliqa.b2clogin.com/vivliqa.onmicrosoft.com/b2c_1a_signi				
	<image/> <image/> <image/> <image/> <form><text><text><text></text></text></text></form>				

3. From the dropdown menu, select your Vivli request. Please note, if you have more than one Vivli request which has a secure research environment, each request will appear in the menu.



4. From the results export menu, select the help text to open the Vivli Research Environment guide. Section 8.0 Exporting Research Results has detailed instructions for submitting this request.

testenquirvrequest		
Before requesting results, read the short introduction here.		Create New Results Request
	No Result Requests Found	
b		0 to 0 of 0 K < Page 0 of 0 > >
		@ Copyright 2017 - 2024 Vivii

5. Create a new results request.

Vivli
CENTER FOR GLOBAL CLINICAL RESEARCH DATA
Assessing the impact of lifestyle choices on heart health
Create New Results Request
No Result Requests Found
0 to 0 of 0 K < Page 0 of 0 > >I
@ Copyright 2017 - 2021 Vivli

Create New Result Request?
Yes No

6. Drag and drop, or select files, to copy your files into the export folder. If your results are organized into more than one folder, we recommend that you place the files into a single zip file before placing them in the results folder. 7-zip is included on the research

environment for that purpose. Similarly, if you have more than 5-10 files, place them into a single zip file.

CENTER FOR GLOBAL CLINICAL RESEARCH DATA					
Assessing the impa	ct of lifestyle of	choices or	n heart health		
Requesting expor	t of research r	esults			
To request export of research results, use Drag and Drop to drop the requested files onto the "Drop f					
here" control, then click sul approved, you will connect individual participant data i	omit when they are co to the main Vivli Plat	omplete. When	n the results have be	en reviewed an	hd
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Select Files	🙆 Dro	p files h	nere		
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CENTER FOR BLOBAL CLINICAL RESEARCH DATA Assessing the impa	ct of lifestyle c	hoices on	heart health	© Copyright 2	2017 - 2
Assessing the impa			heart health	© Copyright 2	2017 - 2
Assessing the impa Requesting export	of research re	esults			
Assessing the impa	of research re ch results, use Drag a mit when they are co to the main Vivli Platfe	esults and Drop to dro mplete. When	op the requested files the results have bee	s onto the "Drop n reviewed and	
Assessing the impa Requesting export To request export of researchere" control, then click sub approved, you will connect	of research re ch results, use Drag a mit when they are co to the main Vivli Platfe	esults and Drop to dro mplete. When	op the requested files the results have bee	s onto the "Drop n reviewed and	
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Assessing the impart Requesting export To request export of researchere" control, then click sub approved, you will connect individual participant data is Select Files	of research re ch results, use Drag a mit when they are co to the main Vivli Platfe	esults mplete. When form to downloa	op the requested files the results have bee	s onto the "Drop n reviewed and	
Assessing the impart Requesting export To request export of researchere" control, then click sub approved, you will connect individual participant data is Select Files UPLOADED FILES	of research re ch results, use Drag a mit when they are co to the main Vivli Platfe not allowed.	esults mplete. When form to downloa	op the requested files the results have bee ad the results. Remin	s onto the "Drop n reviewed and ider - exporting	
Assessing the impart Requesting export To request export of researchere" control, then click sub approved, you will connect individual participant data is Select Files UPLOADED FILES	of research re ch results, use Drag a mit when they are co to the main Vivli Platfe not allowed. Size 0.02	esults mplete. When form to downloa	op the requested files the results have bee ad the results. Remin	s onto the "Drop n reviewed and ider - exporting	

7. Submit files for review.

Submit Files?	
Are you sure all files have been uploaded? This action cannot	be undone.
Yes No	
Vivli	
CENTER FOR GLOBAL CLINICAL RESEARCH DATA	
Assessing the impact of lifestyle choices on	
Assessing the impact of lifestyle choices on heart health	(Request by Data Requester on 2021- 10-01)
	Create New Results Request
2021-10-01 Data Requester	Result Request in Progress
	1 to 1 of 1 K < Page 1 of 1 > >1
Results not yet available for this request.	

Please note, the processing time for a results export request is 5-7 business days. You will receive an email notification when a decision is recorded on the platform and can also monitor by checking the Research Results tab on the platform.

## 8.2 Downloading Approved Results

Once the request to export results is approved, the data requestor may download their results.

### 8.2.1 Steps, Exporting Results:

- 1. Login to the Vivli Platform
- 2. Navigate to My Data Requests and select the request:

Vivli	Home About Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA	QUICK STUDY LOOKUP 🗸 进 MY DATA REQUESTS 🛛 🗕 DATA REQUESTER 🗸
My Data Requests (166)	2545
Draft     Active     Not Approved     Withdrawn     Archived	
ASSESSING THE IMPACT OF LIFESTYLE CHOICES ON HEART HEALTH I 1 STUDY	
Vivil ID: 00002545 Status: All Data Packages Provided and Available	

3. Click on Research Results:

Vivli		Home	About Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA			QUICK STUDY LOOKUP 🗸 🏥 MY DATA REQUESTS 🗕 DATA REQUESTER
< Go Back Assessing th	ne impact of lifestyle choices on	heart health	Print
Studies			
Attachments	Vivli Platform password, Use Drag and Drop to drop t	esearch environment, click on the Export Results icon on the deskt he requested files onto the "Drop files here" control, then click subn he results. Reminder - exporting individual participant data is not al	nit when they are complete. When the results have been reviewed
Request History	2021-10-01	Data Requester	Result Retrieval Succeeded
Signed Agreements			
Safety Concerns			
Research Results			1 to 1 of 1 IC < Page 1 of 1 > >I
Chat	Result Request Status: Result Retrieval Succeeded UPLOADED FILES		
Research Team	Filename	Size	Jploaded By
Research Environment	result export.txt	< 1 kB [	Data Requester
Request Details/Print View			
How To Guide Privacy Cookie	Policy EEA Disclosure Policy Contact Us		@ Copyright 2017 - 202

If you have made more than one request, click on the row corresponding to your most recent request – this will be on the top of the list. The row you have selected will be shown in dark blue.

4. Here, you will see if the request to export results has been approved:

Vivli		Home About	Members News & Events Resources Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA		QUICK S	STUDY LOOKUP 🗸 🥶 MY DATA REQUESTS 🛛 💽 DATA REQUESTER 🥆
< Go Back Assessing	the impact of lifestyle choices on hea	rt health	Print
Studies			
Attachments	To request export of research results, connect to the resear Vivli Platform password. Use Drag and Drop to drop the req and approved, you will return back here to download the res	ch environment, click on the Export Results icon on the desktop. In the uested files onto the "Drop files here" control, then click submit when t sults. Reminder - exporting individual participant data is not allowed.	e browser window, log in with your email address and they are complete. When the results have been reviewed
Request History	2021-10-01	Data Requester	Result Request in Progress
Signed Agreements	2021-10-01	Data Requester	Result Retrieval Succeeded
Safety Concerns			
Research Results			1 to 2 of 2 IC < Page 1 of 1 > >I
Chat	Results not yet available for this request.		
Research Team			
Research Environment			
Request Details/Print View			

5. If the request to export results has been approved, click on **Download**:

Uivli		Home At	bout Members News & Events Resources Find Stur
CENTER FOR GLOBAL CLINICAL RESEARCH DATA		C. C	QUICK STUDY LOOKUP 🗸 🚰 MY DATA REQUESTS 💽 DATA REQUE
Go Back Assessing	the impact of lifestyle choices on	heart health	Print
tudies			
tachments	Vivli Platform password. Use Drag and Drop to drop	e research environment, click on the Export Results icon on the desktop the requested files onto the "Drop files here" control, then click submit d the results. Reminder - exporting individual participant data is not allow	when they are complete. When the results have been reviewed
equest History	2021-10-01	Data Requester	Result Retrieval Succeeded
igned Agreements	2021-10-01	Data Requester	Result Retrieval Succeeded
afety Concerns			1 to 2 of 2 i< < Page 1 of 1 > >i
esearch Results			
hat	Result Request Status: Result Retrieval Succeeded	I Contraction of the second	
	UPLOADED FILES		
esearch Team	Filename	Size Upk	paded By
esearch Environment	result export.txt	< 1 kB Data	a Requester
equest Details/Print View			

If a review of the results identifies the presence of Individual Participant Data (IPD) in the result files, the Result Requests screen will display a message indicating that, and you will need to remove the IPD from the results files before requesting subsequent export of results again.

6. Once the *final* results are exported, please reach out to the Vivli Team via chat and we will provide next steps for deprovisioning the research environment and progression to publication.

## 9.0 Publication Review

## 9.1 Publication Follow up by Vivli

The Vivli administrator sends periodic reminders (at 3 months, 9 months and 1 year) to the researchers to get an update on the status of any potential publications from the time final results from the research environment are downloaded or a year after the downloadable data package was made available to the Data Requestor.

### 9.2 Publication Reviews by Data Contributors

- Once your researcher team has completed your analysis and you are ready to submit your findings for dissemination, either through a learned forum such a publication or conference abstract, requestors must submit to Vivli a copy of any Publication materials at least 30 days prior to submission. Please submit this publication using the Chat function.
  - Please submit 2 files or less for review. If there are additional files, please combine the documents into no more than 2 files (e.g. 1 file containing the final draft of your publication and 1 file containing your attachments such as supplemental figures, graphs, etc.).
- The following acknowledgement should be added to the dissemination of findings. Also, please use the following language in your acknowledgment section:

This [publication or presentation, as applicable] is based on research using data from data contributors \*Data Contributor(s) Name\* that has been made available through Vivli, Inc. Vivli has not contributed to or approved, and is not in any way responsible for, the contents of this publication.

- During this 30-day review period, data contributors may provide you with non-binding comments regarding the scientific content. They may also possibly request the deletion of any confidential information (confidential information as defined in the signed DUA).
- Once the 30-day publication review period is up or once all data contributors have commented on the manuscript, whichever comes first, the research team may submit their manuscript. The Vivli team will follow up monthly to check on the status.
- If you are submitting an abstract to a conference or learned forum and it is accepted as a poster or presentation, the new material (poster, presentation, etc.) must be submitted for the 30-day review period using the Chat function.

### 9.3 Publication Notification by Data Requestor

• Once your team has been notified that your publication has been accepted by the journal and the DOI for your citation made available, please let the Vivli team know via chat.

- The Vivli team will then update your data request on the Vivli website by including the statistical analysis plan and the DOI to the publication.
- The Vivli team will also add the citation to your data request on the Vivli platform. The citation will be linked to the study(s) involved in your request as well and will be viewable when viewing study details on the Vivli platform.
- If this is your final analysis for the research project, please let Vivli know and if so, we will move your analysis to long-term storage. See <u>section 12.0 Stopping the Research Environment</u>. Vivli team will send you a survey about your experience on Vivli platform.
- If you have more publications linked to this research project, please follow step 9.2 Publication Reviews by Data Contributors for each subsequent publication.

### 9.4 Public disclosure not possible

If for whatever reason, you are unable to publish your results, you must provide a summary report of your findings and Vivli will publish this report on its website. For an example see: <a href="https://vivli.org/identification-of-biomarkers-associated-with-specific-sleep-variables/">https://vivli.org/identification-of-biomarkers-associated-with-specific-sleep-variables/</a>. Vivli will then de-provision your research environment and your project will be complete.

Alternatively, you may fulfill the obligations under the Vivli DUA and submit to a pre-print server such as <u>https://www.medrxiv.org/</u>. This preprint server was started by the BMJ and Yale and is a free distribution server for preprints of articles covering all aspects of research. Once posted you would receive a DOI so it would be citable and discoverable. You can cite this on your CV and continue to submit to other journals if you like.

- Once your publication is posted on a pre-print server, please let the Vivli Team know via chat.
- The Vivli Team will follow up to check the status of publication in a peer-review journal.

### 9.4.1 Reasons why analysis may not be completed

If the analysis is not completed, this fact and the reason that it was not completed, should be stated as a summary of the research.

- The analysis may not be performed or completed due to the following reasons:
  - Technical e.g. datasets can't be combined, the research questions can't be answered, data mapping isn't possible
  - $\circ$  Scientific e.g. the studies can't be combined due to design differences or endpoint differences

 Logistical - e.g. the Data Requestor loses funding or key research personnel According to Vivli policy, if a user fails to meet their DUA obligations and does not respond with reasons why their analysis has not been completed, they are unable to submit a subsequent request from Vivli.

# 10.0 Extensions to the Data Use Agreement via the Data Request Progress Report

- Access to the data listed in the approved research proposal is valid for one year from when the Data Use Agreement (DUA) is executed.
- 90 days prior to the 1-year DUA expiration, the Vivli team will reach out to the Data Requestors for an update on the data request and analysis by sending the lead investigator thethe Data Request Progress Report.
- The Lead Researcher fills out the Data Request Progress Report to request an extension. If additional team members from other institutions have a signed DUA, they will acknowledge the completed Data Request Progress Report to extend their access as well.
- Vivli team will follow up on the Data Request Progress Report 1 week prior to the DUA expiration.
- Based on the response, Vivli makes the decision regarding extending access to the data in 1year intervals. These extensions are only granted due to extenuating circumstances. Vivli will respond in 10 business days with a decision that has been reached and upload the document via Signed Agreements on the Vivli platform.
- If no response is received before the end of the Data Use Agreement, Vivli team stops the Research Environment and remove the team's access to the research environment on the day of DUA expiration.
- Note: For any requests for downloadable data, Vivli team will request evidence of data destruction as per the data security addendum.
- Note that an extension to the Data Use Agreement (a legal agreement) is entirely separate from the No Charge period on the Research environment and does not extend the No Charge period.

### 10.1 Failure to meet DUA obligations

According to Vivli policy, if a user fails to meet their DUA obligations, they are unable to submit a subsequent request from Vivli.

# 11.0 Paying for a Research Environment

The cost and length of no charge periods for the Secure Research Environment access are available on the Vivli <u>website</u>. Please note these charges are subject to change.

Note that the no charge period (i.e., the length of time where you will not be charged) begins on the date you first provision the Research Environment, which usually is later than the date that the Data Use Agreement is signed. Per Vivli policy, the no charge period for the Vivli Research Environment begins at initiation and is not impacted by the days you actually use the environment; billing is not impacted by pausing. Renewing the Data Use Agreement does not extend the no charge period.

As you near the end of your no charge period, Vivli will send you notices via email to the email addresses of the research team members on the data request. In these notices we will remind you of the end date of your no charge period. **We will ask you to provide payment via a credit card once your no charge period has ended.** Charges will be automatically completed on the

same day of the month, approximately every 30 days, as when payment is first submitted for the next month's usage. A constant 30 days per month will be charged, and your subscription will be cancelled and no further charges made as soon as you submit a public disclosure for courtesy review and confirm with the Vivli team you wish to stop using your research environment.

If no payment method is arranged, access to the Research Environment will be revoked.

Please note that you will be responsible for all charges once your no charge usage period has ended. Once you have submitted a public disclosure for courtesy review and confirmed with the Vivli team that you wish to stop your secure research environment then this will stop the accrual of additional fees. See <u>section 12.0</u> for further information about stopping the research environment.

- At the end of your no charge period, the Vivli team will email you with the appropriate link to sign up for subscription billing for the secure Research Environment within 1 week of your no charge period end date. The subject line will be "Action Required: Vivli Research Environment ALERT – Payment Request to Maintain Access to data request"
- 2. Upon opening the link, you will be directed to the PayPal page where you can login with your PayPal account or enter the credit/debit card number to be used for billing.

with a p-	Pay with PayPal
with a Pa	yPal account, you're eligible for Purchas Protection and Rewards.
Email or	mobile number
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orgot pas	sword?
	Log In
	or
	Pay with Debit or Credit Card

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3. After entering the credit card details, click the Agree & Subscribe button. Please note that you will be charged every 30 days on approximately the same date each month going forward.

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4. Lastly, you should receive confirmation from PayPal that you have successfully signed up for your subscription for the Research Environment and also a receipt of your first payment to the email address shared with PayPal.

If you have any questions about this process, please contact Vivli at support@vivli.org.

## 12.0 Pausing the Research Environment

- When you have submitted your publication for review and are awaiting feedback, you may reach out to the Vivli Team in chat to request a "pause" of your research environment. This will preserve any remaining no charge period days for subsequent analysis from peer review feedback. If you are actively paying for your environment, requesting to "pause" will stop the accrual of additional fees.
- Once your research environment has been stopped, this will stop the accrual of additional fees until a request to re-access has been submitted.
- Important: Before requesting to "pause" the environment, please copy any data that you
  might need later access to onto the Network Drive (Drive V:\). Once disabled, access to
  content in the Research Environment will be terminated. Note that long-term archive will
  ONLY save the contents of drive V. Data saved in any other drive will be deleted: so please
  place any essential data to Drive V. Remember to check your remote desktop and
  documents folders for any data you wish to be archived.
- If you need access to data while your request is "paused", simply reach out to the Vivli Team in chat or via <u>support@vivli.org</u> and will provide next steps for resuming your analysis.

## 13.0 Deprovisioning the Research Environment

- When you have completed your research, exported your final results and published your findings, the Vivli team will provide next steps, as needed, to deprovision your research environment and prepare data for long term archive. Please reach out when you have saved all data to which you may need future re-access to the V:drive then reach out to the Vivli Team via <a href="mailto:support@vivli.org">support@vivli.org</a> to request deprovisioning.
- Important: Before requesting to deprovision the environment, please copy any data that you might need later access to onto the Network Drive (Drive V:\). Once disabled, access to content in the Research Environment will be terminated. Note that long-term archive will ONLY save the contents of drive V. Data saved in any other drive will be deleted: so please place any essential data to Drive V.
- Shortly after your research environment has been marked for deprovisioning, Vivli will place a copy of the contents of the data disk into long-term archive.
- If you need access to data held in the Vivli long-term archive, please send a request to the Vivli Team via <u>support@vivli.org</u> including the Data Request number and Project Name. Processing this request takes 5-7 days.
- If you need access to long-term storage data for a new research project, to address a different question and/or if you are requesting additional data, you will need to create and submit a new data request on the Vivli platform. In the Narrative, reference the original data request, including the request number.

If you accidentally deprovision a research environment, contact Vivli at <u>support@vivli.org</u>. Vivli will re-provision the environment, including any results files.

## 14.0 Downloadable data

Some Data Contributors will allow you to download their data directly from the Vivli Platform using the following process:

1. Log in and open your approved data request:

- 🖶 Vivli		Home About	Members	News & Events	Resources	Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA			QUICK STUD	rlookup 🗸 🔮 n	IY DATA REQUESTS	🗶 IDASIM 🗸
«Go Back Hepatitis	Project					Print
Studies	REQUESTED STUDY TYPES ()					
Attachments	VIVLI-LISTED AND PROVISIONED STUDIES					
Request History	A Phase II Open Label, Dose-Escalation Study to Determine the Safety, Tolerability and Efficacy of Microsomal Triglyceride Transfer (MTP) Inhibitor BMS 201038 in Patients With Homozygous Familial Hypercholeterolemia PL Sponsor Agenton Pharmaceutcale, inc. Subj ID NC10156806 IRPRepriver Venstat Data Reguest ID 00001280 Sponsor ID UP1001	Protein	Data F	ackage Provided t	o Requestor	>
Signed Agreements	VIVLI-LISTED STUDIES PROVISIONED BY EXTERNAL PROVIDERS					
Safety Concerns	No Studies Found STUDIES PROVIDED ON VIVLI PARTNER PLATFORMS (NOT LISTED ON VIVLI) OR OTHER DATA					
Chat						
Research Team						
Research Environment						
Request Details/Print View						
How To Guide Privacy Cool	kie Policy EEA Disclosure Policy Contact Us					@ Copyright 2017 - 2018 Vivi

Figure 26 - Approved Data Request

2. Click on the study to get to the **Study details** screen:

Vivli					Home Al	oout Members	News & Events	Resources	Find Studies
CENTER FOR GLOBAL CLINICAL RESEARCH DATA						QUICK STUD	r lookup 🗸 🔮	MY DATA REQUESTS	s  🔕 ida sin
Phase II Open Label, Do omozygous Familial Hyp			Safety, Tolerability and Efficad	y of Microsomal Triglyceride Tra	nsfer Protein (M	FP) Inhibitor E	8MS-201038 ir	n Patients W	(ith
Study Details Administrativ	ve Details Dow	nload Data Package							
Phase Phase 2				Condition or Disease Homozygous Familial Hypercholester	rolemia				
Brief Summary									
The primary objective of this stur objectives of this study included	the evaluation of the	pharmacodynamics of lomitap	de based on: - Percent change in low-	given as an initial low dose and then escala density lipoprotein cholesterol (LDL-C), total previous dose phase(s) Changes in othe	l cholesterol (TC), trigl	ycerides, and very	low density lipop	rotein cholester	ol (VLDL-C)
Ages Eligible For Study		Sexes Eligible For S	hulu	Accepts Healthy Volunteers		Actual Enrollr	nent		
13 Years and older		All	wwy	No		6			
Study Start Date 06/2003			Study Completion Date 02/2004		Recruitment Sta Completed	tus			
Locations United States (1)									

Figure 27 - Study details screen

3. Click on Download Data Package:

		QUICK STUDY LOOKUP 🗸 🎒 MY DATA REQUESTS
		ity and Efficacy of Microsomal Triglyceride Transfer Protein (MTP) Inhibitor BMS-201038 in Patients With
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Figure 28 - Download Data Package

4. This will take you to the Download screen:

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5. Depending on your browser, a pop-up will appear:

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Figure 30 - Downloadable data file selection pop-up

- 6. Select the target file, choose whether you would like to Save or Open the files and click **Ok**.
- 7. Repeat Step 6 for any additional files.

## Tips for downloading files from Vivli

When files are available for direct download from the Vivli platform, the usual approach to downloading files is to open the approved data request, open the studies tab, click on the "card" for the study, go the download data package tab and click "Download" for each individual file that you need to download.

If the size of the file is more than 1000MB, you should take a few extra precautions before starting the download

- In your computer settings, set Power Plan to sleep "Never" when plugged in. (sleeping will interrupt the download)
- Once you start the download, leave the computer running and the browser open.
- The progress of the download is shown in different places depending on the browser (for example for Chrome it is shown in the lower left, and for Firefox it is show in the upper right - note that for Firefox, the progress pop-up is taken down after a few minutes - you can check back on progress by clicking on the download icon on the upper right)
- If the download takes more than 30 minutes, the Vivli Platform will automatically log you out, but the download will continue until it is complete.
- Do not download more than one file at a time

The time the data will take to download can vary a great deal based on your available bandwidth, the size of the file(s), the time of day, what else is happening on your local network (e.g. Zoom calls can use up bandwidth) and even how full or fragmented your disk is. To provide a \*very\* general guideline, in home environments, download time can vary from 1 minute per gigabyte (for a network with 400 Mbps download speed) to 15 minutes or more. When downloading using a business address or a high-end fiber network, it may be faster, sometimes much faster.

Other download tips:

- If it is practical, often downloading is faster in the evening or overnight, as you are competing with less traffic on the internet.
- For large downloads, before starting the upload, it can be useful to reboot your computer this can free up some memory and reset some elements of the operating system.

Finally, if the total volume of data to download is much greater than 25 Gb or so, or if you have problems with large downloads (such as network glitches, other errors, or the download just takes too long) reach out to Vivli at support@vivli.org - we can provide instructions for using a Microsoft upload/download manager that takes a few extra steps, but is faster and more reliable.

If the files have extensions of the form .001, .002, then this was a large zip file that was split into pieces by 7-zip for ease of upload and download. Make sure you download all of the numbered files into the same folder. If you don't already have 7-zip, download a copy of 7-zip from <u>https://www.7-zip.org/download.html</u>. Right-click on the .001 file and drag to a new folder, and from the pop-up menu, choose 7-zip -> Extract here. The utility 7-zip knows enough to combine the pieces logically together as part of the unzip operation.

# 15.0 Feedback and Support

If you need technical help with your research environment, please email Vivli at <u>support@vivli.org</u> and we will assist you as soon as possible. Please provide as much information as you can to the problem, including the research proposal number, the time the problem and any messages you received from the platform occurred as this will make it easier for Vivli to diagnose and fix the issue.

If you ever have thoughts on how to improve the system or processes, please email <u>support@vivli.org</u> or reach out to Vivli via chat. When you have completed your project, Vivli will ask you to complete a short survey to help us continue to improve.