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FACILITY NAME & ADDRESS

Facility Name	Facility Type	Facility Address
National Hospital Organization Okayama Medical Center	Hospital or Medical Center	1711-1 Tamasu kita-ku, Okayama, Okayama, Japan, 701-1192

FACILITY CONTACTS

Primary FPM?	Name	Email Address	Roles
Yes	Hamada, Yuuki	hamada.yuuki.rh@mail.hosp.go.jp	Facility Profile Manager, Delegation Manager
No	Nishiyama, Atsuko	nishiyama.atsuko.vd@mail.hosp.go.jp	Facility Profile Manager

THERAPEUTIC AREAS & PATIENT POPULATION

Therapeutic Area(s)	
Therapeutic Area	**Sub Therapeutic Area**
Bacterial Infections and Mycoses	
Cardiovascular Diseases	
Congenital, Hereditary, and Neonatal Diseases and Abnormalities	
Digestive System Diseases	
Endocrine System Diseases	
Hemic and Lymphatic Diseases	
Respiratory Tract Diseases	Bronchial Diseases
Respiratory Tract Diseases	Laryngeal Diseases
Respiratory Tract Diseases	Lung Diseases
Respiratory Tract Diseases	Respiration Disorders
Respiratory Tract Diseases	Respiratory Hypersensitivity
Respiratory Tract Diseases	Respiratory System Abnormalities
Respiratory Tract Diseases	Respiratory Tract Infections
Respiratory Tract Diseases	Thoracic Diseases
Respiratory Tract Diseases	Tracheal Diseases
Other Areas of Expertise	
Study Phase Capabilities	
Phase I, Phase II, Phase III, Phase IV	

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Other Facility Details	
Do you have Affiliated Research Sites or Satellite Sites/Clinics? A Satellite Site is a secondary location where the investigator sees clinical trial subjects, usually this is the same investigator who sees subjects at the primary site location.	No
What study types does your Facility have experience with?	Industry, Investigator Initiated
Is your Facility affiliated with a government agency or part of a government funded health service?	Yes

Patient Population	
Patient Population Demographics	Pediatrics - Less than or equal to 17, Adults - Ages 18-64, Geriatrics - Greater than or equal to 65
Patient Population Comments	

IRB/ERB/ETHICS COMMITTEE

General Questions	
What is the average time (in days) to start a study once you have received the regulatory package?	30-60
Does your Facility perform IRB/ERB/Ethics Committee submissions?	Yes
Does your Facility have a Facility or group to perform IRB/ERB/Ethics Committee submissions?	Yes
Department Contact Name	Clinical trial
Department Contact Phone Number	086-294-9519
Department Contact Email Address	504-chiken-crc@mail.hosp.go.jp
Is your Facility able to initiate study activities prior to IRB/ERB/Ethics Committee protocol approval?	Yes
What types of IRB/ERB/Ethics Committee does your Facility use?	Central Acting as Local, Local
Does your institution and/or local regulation mandate the distribution of safety reports [e.g., Development SafetyUpdate Report (DSUR), suspected unexpected serious adverse reaction (SUSAR)] to a local Review only IRB/ERB/Ethics Committee?	Yes
Are there any other steps that the Sponsor should be aware of for your IRB/ERB/Ethics Committee review and submission?	No

LOCAL IRB/ERB/ETHICS COMMITTEE

Local IRB/ERB/Ethics Committee: National Hospital Organization Okayama Medical Center	
IRB/ERB/Ethics Committee Name	National Hospital Organization Okayama Medical Center
Address	1711-1 TAmasu kita-ku, Okayama, Okayama, Japan, 701-1192
Registration#	**Registering Body**
No Records	
What is the meeting frequency of your Local IRB/ERB/Ethics Committee?	Monthly
How long before IRB/ERB/Ethics review is the Submission Packet required?	Greater than 2 weeks
Does the IRB/ERB/Ethics Committee require payment prior to release of final approval documents?	No
Does the IRB/ERB/Ethics Committee require contract/budget approval prior to release of final approval documents?	No

LOCAL IRB/ERB/ETHICS COMMITTEE ATTACHMENTS		
Document Type	**Document Name**	**Document Description**
No Records		

OTHER REVIEW BOARDS

Does your Facility have Other Review Boards that need to approve the study prior to IRB/ ERB/Ethics Committee submission? For example, scientific, radiation safety committees, or others.	No

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Local Lab

Is your Facility using a Local Lab?	Yes
Local Lab: National Hospital Organization Okayama Medical Center Clinical Laboratory Department of	
Lab Name	National Hospital Organization Okayama Medical Center Clinical Laboratory Department of
Lab Contact First Name	
Lab Contact Last Name	
Address	1711-1,tamasu,kita-ku, Okayama, Okayama, Japan, 701-1192
Phone Number	086-294-9911
Fax Number	
Email Address	
Local Lab Accreditation	Others
Other Local Lab Accreditation	Japanese Association of Medical Technologists,Okayama Prefecture Medical
Additional Questions	
Does your Facility have a SOP/written procedure for documenting bio-specimen (Sample) processing steps/chain of custody?	
What is the system or tool that the site currently has or utilizes to document Bio-specimen (Sample) Processing Steps/ Chain of Custody?	
Please indicate tissue collection and processing capabilities at your site?	
Does your Facility has established processes to oversee staff compliance with study-specific lab manual instructions for bio-specimen processing?	
What are your Facility’s capabilities for tissue collection and/or processing (embedding)?	
Are LOINC codes available for the Local Lab? (If Yes, you can upload the relevant LOINC list as an attachment in Lab Documentation)	

Attachments

Document Type	Document Name	Document Description
No Records		

CONSENT & TRAINING

Consent	
Does your Facility have a written SOP/Policy/Procedure for: Informed Consent?	No
Does your Facility have a written SOP/Policy/Procedure for: Minor Assent for Pediatric Populations?	No
Does your Facility have a written SOP/Policy/Procedure for: Other Vulnerable Populations?	No
Will your Facility require language translations for consents?	Yes
Select the required languages	Japanese
If located in the US, has your Facility used or are you able to use the informed consent short form?	Not Applicable
What types of eConsent platforms does your Facility use? (Select all that apply)	

Training	
Does your Facility have a training program for the research staff?	Yes
Does the course content include GCP?	Yes
Does your Facility use an external program to conduct research training?	Yes
Please provide program course name.	eAPRIN
Do you have a process or program in place to retrain research staff when a protocol is amended?	No
Does the study staff that prepares or transports dangerous goods have training that meets the IATA International Air Transport Association (US) or other countries hazardous training requirements for shipping dangerous goods?	No

FACILITY & EQUIPMENT

Facility Capabilities	
Can your Facility support patient visits on weekends?	No
Can your Facility support in-patient admissions for research studies?	Yes
Does your study staff have sufficient English knowledge to understand communications in English?	No
Does your Facility have access to translators and translation support for study conduct (e.g. consent, study specific instruction)?	No
Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Is the lab kit storage space able to support early phase studies which may require an increased number of kits?	Yes
Does your Facility have the ability to collect and store PK/PD specimens?	Yes
Does your Facility have the ability to collect PK/PD samples beyond normal business hours?	No
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Equipment	
Identify the Diagnostic Equipment available at or near the Facility to support Research studies?	Computerized Tomography Scan, Dual-Energy X-ray Absorptiometry or Bone Densitometry, Magnetic Resonance Imaging, Fluoroscopy, X-Radiation, Magnetic Resonance Angiography, Magnetic Resonance Spectroscopy, Mammography, Nuclear Medicine (e.g.Bone scan,Thyroid scan,Thallium cardiac stress test), Electrocardiogram
General Equipment	
Does your Facility have an SOP or process that ensures routine calibration and maintenance of general equipment? Examples of general equipment include: scale, pulse oximeter, stadiometer, sphymomanomer, etc.?	No
Does your Facility have the necessary equipment to treat medical emergencies (ie. code cart)?	Yes
Identify the equipment available at the Facility to support Research studies?	Refrigerated Centrifuge, Centrifuge, Refrigerator (2 to 8 Degrees C), Freezer (-20 to -30 Degrees C), Freezer (-70 to -80 Degrees C)
Equipment Capabilities: Refrigerator (2 to 8 Degrees C)	
Do you have the ability to generate a temperature monitoring log for this equipment?	Yes
Does this equipment provide Min/Max Temperature Monitoring?	Yes
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	By Minute
Does this equipment have back-up power?	Yes
Does this equipment have a temperature alarm?	Yes
Do you have an SOP which supports calibration of this equipment?	No

Equipment Capabilities: Freezer (-20 to -30 Degrees C)	
Do you have the ability to generate a temperature monitoring log for this equipment?	Yes
Does this equipment provide Min/Max Temperature Monitoring?	Yes
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	By Minute
Does this equipment have back-up power?	Yes
Does this equipment have a temperature alarm?	Yes
Do you have an SOP which supports calibration of this equipment?	No
Equipment Capabilities: Refrigerator (-70 to -80 Degrees C)	
Do you have the ability to generate a temperature monitoring log for this equipment?	Yes
Does this equipment provide Min/Max Temperature Monitoring?	No
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	
Does this equipment have back-up power?	Yes
Does this equipment have a temperature alarm?	Yes
Do you have an SOP which supports calibration of this equipment?	No
Computer Capabilities	
Does your Facility have computers which are dedicated to research studies?	Yes
Does your Facility have antivirus software on all these electronic devices?	
Does your Facility have an SOP or process that documents both physical and firewall security measures that are used for electronic systems?	
Does your Facility perform periodic system reviews to ensure they remain in a validated state?	
Does your Facility have an SOP or process that documents user management processes that are used for electronic systems?	
What type of computer operating system(s) does your institution use to support studies?	Windows (Windows XP, Windows 7, Windows 8, etc.)
What type of internet access does your Facility have?	Cable or DSL
What type of internet browser(s) does your institution use to support studies? (Check all that apply)	
Is access to the internet secured against unauthorized use?	
Does your Facility limit or prohibit access and use of external web-based tools or sites for clinical research? (e.g. web portals to submit documents to sponsors or CROs)	Yes
Does your institution allow the download of information from an external portal ?(e.g. eDC system)	
Does the Facility have access to local IT support?	I don't Know
Does your Facility prohibit the use of an external USB device (e.g. to download and send data from a temperature monitoring device)?	No
Business Continuity Plan	
Does your Facility have Business Continuity Plan (BCP) to protect essential business operations which describes how those processes will be performed during a crisis at your Facility?	Yes
Does your Facility's BCP include a process for dealing with Cybersecurity incidents?	

Attach Your BCP or SOP

Document Type	Document Name	Document Description
No Records		

INVESTIGATIONAL PRODUCT & CONTROLLED SUBSTANCES

Investigational Product Shipping Details					
IP Recipient Name	Address	Phone Number	Fax Number	Email Address	Site Specific Requirements
National Hospital Organization Okayama Medical Center	1711-1,Tamasu,Kita-ku, Pharmacy Department, Atsuko Nishiyama, Okayama, Okayama, Japan, 701-1192	086-294-9911	086-294-9508	nishiyama.atsuko.vd@mail.hosp.go.jp	

Investigational Product Storage Location				
IP Storage Location Name	Address	Phone Number	Fax Number	Email Address
Pharmacy	1711-1,Tamasu,Kita-ku, Pharmacy Department, Atsuko Nishiyama, Okayama, Okayama, Japan, 701-1192	086-294-9911	086-294-9508	nishiyama.atsuko.vd@mail.hosp.go.jp

Investigational Product Storage Equipment	
Identify the Investigational Product Storage Equipment at your Facility	Refrigerator (2 to 8 Degrees C), Freezer (-20 to -30 Degrees C)
Equipment Capabilities: Refrigerator (2 to 8 Degrees C)	
Do you have the ability to generate a temperature monitoring log for this equipment?	Yes
Does this equipment provide Min/Max Temperature Monitoring?	Yes
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	By Minute
Does this equipment have back-up power?	Yes
Does this equipment have a temperature alarm?	Yes
Do you have an SOP which supports calibration of this equipment?	No
Equipment Capabilities: Freezer (-20 to -30 Degrees C)	
Do you have the ability to generate a temperature monitoring log for this equipment?	Yes
Does this equipment provide Min/Max Temperature Monitoring?	Yes
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	By Minute
Does this equipment have back-up power?	Yes
Does this equipment have a temperature alarm?	Yes
Do you have an SOP which supports calibration of this equipment?	No
Investigational Product Storage And Handling	
Is the Investigational Product Storage Room secured with controlled access?	Yes
Do you have the ability to generate a temperature monitoring log for this Investigational Product Storage Room?	Yes
Does the Investigational Product Storage Room provide Min/Max temperature monitoring?	Yes
Does the Investigational Product Storage Room have back-up power?	Yes
Does the Investigational Product Storage Room have a temperature alarm?	Yes
Do you have an SOP which supports calibration of the temperature monitoring equipment?	No
Does your Facility have the ability to manage on-site or off-site destruction of Investigational Product?	Yes
Does your Facility have a written SOP/Policy/Procedure for destruction of Investigational Product?	No
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Not Applicable

Does your Facility have a written SOP/Policy/Procedure to ensure that Investigational Product is appropriately maintained during transportation to Satellite Site(s)?	Not Applicable
Describe additional Investigational Product Storage And Handling Capabilities	
Preparation and Administration Of Investigational Product	
Identify the Investigational Product preparation capabilities at your Facility	Extemporaneous Preparation, Vertical laminar flow hood (chemo/hazardous drugs)
Is your Facility capable of administering infusions?	Yes
Is your Facility adequately staffed to support studies with both blinded and un-blinded Investigational Product?	Yes
Controlled Substances	
Does the Facility have the required licenses or registrations to receive, store, dispense and return controlled substances as required by local law?	Yes
Is the storage area for controlled substances securely constructed with restricted access in accordance with local law?	Yes
Does the Facility have the ability to handle radio-labelled Investigational Product?	No
Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	Not Applicable

Attachments

Document Type	Document Name	Document Description
No Records		

SOURCE DOCUMENTATION & REMOTE MONITORING

Source Documents	
What type of source documents will be used?	Paper, Electronic
Do all of the Facility's systems/technology used to store and manage source data conform with international system standards (e.g. ICH/GCP/EU)?	
Does your Facility have secure storage for patient records?	Yes
Does your Facility have patient record archiving on-site?	Yes
Does your Facility have an SOP or process that documents how to limit access and prevent the premature destruction of the Facility's files?	
What type of investigator site file/regulatory binder used (select all that apply)	Paper, Electronic
What investigator site file (eISF) / eRegulatory system do you use?	Others: Agatha
Are monitors able to access eISF/eReg while off-site?	Yes
Do the eISF/eReg systems conform with international system standards (e.g. ICH/GCP/EU)?	
Please list any access limitations/ requirements for eISF/eReg	Access is possible by the administrator creating individual accounts.Issuance of ID and password by the administrator
Electronic Medical Records (EMR) / Electronic Health Records (EHR)	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
What EMR/EHR system do you use?	In-house system
For Facilities with satellite sites, where is the monitor required to access source documents?	Main Facility Only
Please list any access limitations/requirements for the Electronic Medical Records.	ID,Password
Do the EHR/EMR systems conform with international system standards (e.g. ICH/GCP/EU)?	
Do you work with a vendor that can electronically exchange data for clinical research from the EHR/EMR?	No

Do you have institutional approval to export data from the EHR/EMR for the clinical research?	Yes
Are monitors able to access EHR/EMR while off site?	No
Does your Facility require Sponsor representative to sign any local form (paper or electronic) for access, or any other purpose?	Yes
Provide details of information requested	Viewing the electronic medical records requires logging in with an ID and password. Additionally, the granted permissions are limited to viewing only. Please note that access from outside the facility is not permitted.
Does your Facility utilize any additional electronic systems to capture source data outside of the EMR/EHR?	
Does your Facility utilize any additional electronic systems or paper-based systems to access patient medical history?	
Do the systems used for clinical trial data entry and management have audit trail functionality?	

Bring Your Own Device	
Can your Facility support the use of bring your own device (BYOD) technologies to conduct Clinical Trials?	

Monitoring	
Check all equipment that will be available to Monitors:	Copy Machines, Internet Access
What Electronic Data Capture (EDC) systems has your staff used for clinical trials?	Oracle Inform, Medidata Rave, Oracle RDC Remote Data Capture, Others
Describe Other EDC Systems	DATA TRAK,My Trials,iMedidata
Does your site/institution and/or local regulations allow remote source data verification of study participant data to support remote monitoring?	No

Attachments

Document Type	Document Name	Document Description
No Records		

ADDITIONAL LOCATIONS

Additional Locations

Add any addresses you wish to be available in the Study Site Profile. These addresses will be available for selection in the following sections of the Study Site Profile - Additional Study Locations - These addresses can be added to your FDA Form 1572, if applicable.

SIP Registered Locations

Location Name	Address
No Records	

SIP Non-Registered Locations

Location Name	Contact Name	Address	Phone Number	Fax Number	E-mail Address
No Records					

ADDITIONAL INFORMATION & ATTACHMENTS

Additional Information
Please provide additional information not captured in other sections of the Facility Profile that you feel is important for Sponsors to know about your site. Please reference the section name if applicable.

Facility Attachments		
Document Type	**Document Name**	**Document Description**
No Records		

ORGANIZATION AFFILIATIONS

Organization Affiliations			
The Organization (s) that requested Affiliation with your Facility are listed below with Affiliation Status			
Organization Name and Address	**Organization Affiliation Type**	**Organization Affiliation Status**	**Status Date**
No Records			

ASSOCIATED SITE USERS

Associated Site Users

☐ Once checked, this checkbox will enable the Approval/Rejection workflow for this Facility. Any site user requesting to associate with this Facility would require to send the affiliation requests and only once Approved, this Facility will be shown on User's Profile.

Site User Association Requests				
Name	**E-mail Address**	**Request Affiliation Date**	**Affiliation Status change Date**	**Affiliation Status**
Yamamoto,Nozomi	nozomi-yamamoto.ce@cmicgroup.com	03-Feb-2023		Pending

Associated/Confirmed Site Users				
Name	**User Role**	**Request Affiliation Date**	**Affiliation Status change Date**	**Affiliation Status**
Tomihama,Itsuko	Clinical Research User	06-Oct-2025	06-Oct-2025	Confirmed
Uemoto,Akemi	Clinical Research User	22-Oct-2025	22-Oct-2025	Confirmed
Miyasaka,Hitomi	Clinical Research User	16-Oct-2025	16-Oct-2025	Confirmed
takahashi,chiho	Clinical Research User	09-Dec-2022	09-Dec-2022	Confirmed
Okada,Hirofumi	Investigator	05-Dec-2022	05-Dec-2022	Confirmed
Yokohama,Fumi	Investigator	05-Dec-2022	05-Dec-2022	Confirmed
Kobashi,Miki	Clinical Research User	09-Dec-2022	09-Dec-2022	Confirmed
Hayashi,Kazuna	Investigator	13-Dec-2022	13-Dec-2022	Confirmed
Sogabe,Rie	Clinical Research User	09-Dec-2022	09-Dec-2022	Confirmed
Yamamoto,Sayaka	Clinical Research User	09-Dec-2022	09-Dec-2022	Confirmed

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Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Kanezawa,Misaki	Investigator	13-Dec-2022	13-Dec-2022	Confirmed
SURUGA,KAZUKI	Investigator	01-Dec-2022	01-Dec-2022	Confirmed
shimokawahara,hiroto	Investigator	01-Dec-2022	01-Dec-2022	Confirmed
Ogawa,Aiko	Investigator	05-Dec-2022	05-Dec-2022	Confirmed
Yamamoto,Nozomi	Clinical Research User	03-Feb-2023	03-Feb-2023	Confirmed
katsube,Sho	Investigator	10-Apr-2023	10-Apr-2023	Confirmed
Nishiyama,Midori	Clinical Research User	07-Apr-2023	07-Apr-2023	Confirmed
yamaguchi,yuta	Investigator	11-Apr-2023	11-Apr-2023	Confirmed
Nishizaki,Mari	Investigator	13-Apr-2023	13-Apr-2023	Confirmed
Kimura,Tomonari	Investigator	27-Apr-2023	27-Apr-2023	Confirmed
Sato,Kimi	Investigator	28-Apr-2023	28-Apr-2023	Confirmed
Honda,Akira	Investigator	27-Apr-2023	27-Apr-2023	Confirmed
Karaki,Yumi	Clinical Research User	26-Apr-2023	26-Apr-2023	Confirmed
kawara,yuko	Clinical Research User	01-May-2023	01-May-2023	Confirmed
Yamaguchi,Tomoko	Investigator	15-Jan-2019	15-Jan-2019	Confirmed
Sunami,Kazutaka	Investigator	17-Mar-2020	17-Mar-2020	Confirmed
Akaishi,Makiko	Clinical Research User	21-Apr-2020	21-Apr-2020	Confirmed
Korai,Mutsuko	Clinical Research User	11-Aug-2020	11-Aug-2020	Confirmed
Okada,Rieko	Clinical Research User	18-Feb-2021	18-Feb-2021	Confirmed
Fujii,Yumi	Clinical Research User	27-Dec-2021	27-Dec-2021	Confirmed
Makita,Masanori	Investigator	27-Dec-2021	27-Dec-2021	Confirmed
yoshioka,takanori	Investigator	17-Dec-2021	17-Dec-2021	Confirmed
Sando,YASUHISA	Investigator	21-Dec-2021	21-Dec-2021	Confirmed
Nishiyama,Atsuko	Clinical Research User	06-Apr-2022	06-Apr-2022	Confirmed
Kaneko,Tamami	Clinical Research User	06-May-2022	06-May-2022	Confirmed
Ueno,Anna	Clinical Research User	11-May-2022	11-May-2022	Confirmed
Masumoto,Fumi	Clinical Research User	07-Jun-2022	07-Jun-2022	Confirmed
Tsuno,Saeko	Clinical Research User	11-Nov-2022	11-Nov-2022	Confirmed

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Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
tabuchi,isao	Investigator	29-Nov-2022	29-Nov-2022	Confirmed
Fukuda,Yoshitake	Investigator	26-Nov-2022	26-Nov-2022	Confirmed
Shigetoshi,Masataka	Investigator	15-Nov-2022	15-Nov-2022	Confirmed
Kobashi,Soichiro	Investigator	18-Nov-2022	18-Nov-2022	Confirmed
Watanabe,Atsuyuki	Investigator	14-Sep-2022	14-Sep-2022	Confirmed
Matsubara,Hiromi	Investigator	20-Oct-2022	20-Oct-2022	Confirmed
Mizuno,Aiko	Clinical Research User	03-Oct-2024	03-Oct-2024	Confirmed
Zaizen,Kaori	Clinical Research User	07-Oct-2024	07-Oct-2024	Confirmed
Watanabe,Hiromi	Investigator	10-Mar-2025	10-Mar-2025	Confirmed
Sato,Ken	Investigator	04-Mar-2025	04-Mar-2025	Confirmed
Ichikawa,Takeru	Investigator	17-Mar-2025	17-Mar-2025	Confirmed
Daido,Yusuke	Investigator	16-Apr-2025	16-Apr-2025	Confirmed
Matsumoto,Shoichiro	Investigator	23-Mar-2025	23-Mar-2025	Confirmed
Matsuoka,Suzuka	Investigator	12-Mar-2025	12-Mar-2025	Confirmed
Kudo,Kenichiro	Investigator	10-Mar-2025	10-Mar-2025	Confirmed
Hamada,Yuuki	Clinical Research User	25-Mar-2025	25-Mar-2025	Confirmed
Takigawa,Yuki	Investigator	21-Mar-2025	21-Mar-2025	Confirmed
Wakabayashi,Aiko	Clinical Research User	07-Nov-2023	07-Nov-2023	Confirmed
Iwamoto,Keiko	Clinical Research User	31-Aug-2023	31-Aug-2023	Confirmed
Furujo,Mahoko	Investigator	19-Sep-2023	19-Sep-2023	Confirmed
Shiraha,Keisuke	Investigator	13-May-2025	13-May-2025	Confirmed
Goda,Mayu	Investigator	19-May-2025	19-May-2025	Confirmed
Nishimura,Jun	Investigator	12-May-2025	12-May-2025	Confirmed
Murakami,Michiko	Clinical Research User	18-Jan-2024	18-Jan-2024	Confirmed
Shimizu,JUNYA	Clinical Research User	19-Jan-2024	19-Jan-2024	Confirmed
Kageyama,Misao	Investigator	14-Feb-2024	14-Feb-2024	Confirmed
Tamai,Kei	Clinical Research User	18-Jan-2024	18-Jan-2024	Confirmed
takao,miyabi	Clinical Research User	29-Feb-2024	29-Feb-2024	Confirmed

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Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Otsuki,Minako	Clinical Research User	17-Jan-2024	17-Jan-2024	Confirmed
Maitani,Sachiko	Clinical Research User	17-Jan-2024	17-Jan-2024	Confirmed
Fujiwara,Shintaro	Investigator	03-Feb-2024	03-Feb-2024	Confirmed
Honda,Momo	Clinical Research User	04-Mar-2024	04-Mar-2024	Confirmed
Igarashi,Jun	Clinical Research User	04-Mar-2024	04-Mar-2024	Confirmed
KAMIYA,YUSAKU	Clinical Research User	14-Feb-2024	14-Feb-2024	Confirmed
Nakamura,Makoto	Clinical Research User	06-Feb-2024	06-Feb-2024	Confirmed
Sano,Nagako	Clinical Research User	01-Mar-2024	01-Mar-2024	Confirmed
Saito,Takashi	Investigator	05-Feb-2024	05-Feb-2024	Confirmed
Higuchi,Yousuke	Clinical Research User	18-Jan-2024	18-Jan-2024	Confirmed
Watanabe,Keiko	Clinical Research User	04-Mar-2024	04-Mar-2024	Confirmed
Yasutake,Miyu	Clinical Research User	28-Nov-2023	28-Nov-2023	Confirmed
UENO,MASAYA	Investigator	22-Dec-2023	22-Dec-2023	Confirmed
Takeuchi,Akihito	Clinical Research User	09-Jan-2024	09-Jan-2024	Confirmed
Kondo,Akira	Investigator	27-Dec-2023	27-Dec-2023	Confirmed
Uenaga,Noriko	Clinical Research User	28-Dec-2023	28-Dec-2023	Confirmed
matsuda,masayuki	Investigator	05-Jan-2024	05-Jan-2024	Confirmed
Tanita,Noriko	Clinical Research User	29-Dec-2023	29-Dec-2023	Confirmed
Eda,Mutsumi	Clinical Research User	05-Dec-2023	05-Dec-2023	Confirmed
Kuronuma,Keiichiro	Investigator	22-Jul-2024	22-Jul-2024	Confirmed
Honda,Hiroyuki	Investigator	29-Jul-2024	29-Jul-2024	Confirmed
Hasegawa,Kou	Investigator	30-Jul-2024	30-Jul-2024	Confirmed
Ota,Kosuke	Investigator	29-Mar-2024	29-Mar-2024	Confirmed
Hattori,Mariko	Clinical Research User	15-Apr-2024	15-Apr-2024	Confirmed
Fujiwara,Keiichi	Investigator	23-Apr-2024	23-Apr-2024	Confirmed
Nakamura,Makoto	Investigator	28-Apr-2024	28-Apr-2024	Confirmed
Ikeda,Misa	Clinical Research User	15-Apr-2024	15-Apr-2024	Confirmed
Chikama,Shunsuke	Investigator	30-Apr-2024	30-Apr-2024	Confirmed

Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Ando,Tsubasa	Investigator	04-May-2024	04-May-2024	Confirmed
Minokami,Fumiko	Clinical Research User	27-Aug-2025	27-Aug-2025	Confirmed
Kishigami,Daiki	Clinical Research User	02-Sep-2025	02-Sep-2025	Confirmed
Hamaguchi,Yasuhito	Investigator	28-May-2024	28-May-2024	Confirmed
Fujiwara,Masumi	Clinical Research User	25-May-2024	25-May-2024	Confirmed
Iwai,Fumi	Clinical Research User	11-Dec-2025	11-Dec-2025	Confirmed
Yamaguchi,Mao	Clinical Research User	02-Feb-2026	02-Feb-2026	Confirmed
Tatsukami,Taiko	Investigator	06-Apr-2026	06-Apr-2026	Confirmed
Erika,Araki	Clinical Research User	30-Apr-2026	30-Apr-2026	Confirmed
tabuchi,haruno	Clinical Research User	01-Jun-2026	01-Jun-2026	Confirmed
Nagahisa,Tomomi	Clinical Research User	22-May-2026	22-May-2026	Confirmed
Sasano,Yuto	Investigator	30-May-2026	30-May-2026	Confirmed
Inoue,Yusuke	Investigator	03-Apr-2026	03-Apr-2026	Confirmed
Tomoya,Osedo	Investigator	02-Jun-2026	02-Jun-2026	Confirmed