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FACILITY NAME & ADDRESS

Facility Name	Facility Type	Facility Address
National Hospital Organization Hokkaido Cancer Center		2-3-54 Kikusui 4-jo, Shiroishi-ku, Sapporo, Hokkaido, Japan, 003-0804

FACILITY CONTACTS

Primary FPM?	Name	Email Address	Roles
Yes	Suzuki, Hidetaka	suzuki.hidetaka.sg@mail.hosp.go.jp	Facility Profile Manager, Delegation Manager
No	Momoi, Yoshinori	momoi.yoshinori.yj@mail.hosp.go.jp	Facility Profile Manager, Delegation Manager
No	Sawano, Takako	ogawa.takako.ju@mail.hosp.go.jp	Facility Profile Manager, Delegation Manager
No	Shirooka, Yukari	shirooka.yukari.rw@mail.hosp.go.jp	Facility Profile Manager, Delegation Manager

THERAPEUTIC AREAS & PATIENT POPULATION

Therapeutic Area(s)	
Therapeutic Area	**Sub Therapeutic Area**
Oncology	Bladder
Oncology	Breast
Oncology	Carcinoma
Oncology	Cervical
Oncology	Colorectal
Oncology	Esophagael
Oncology	Gastric
Oncology	Gastrointestinal
Oncology	Genitourinary
Oncology	Head and Neck
Oncology	Hematologic Malignancies
Oncology	Hepatocellular Carcinoma
Oncology	Leukemia
Oncology	Lung
Oncology	Lymphoma
Oncology	Multiple Myeloma
Oncology	Ovarian
Oncology	Prostate

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Therapeutic Area	Sub Therapeutic Area
Oncology	Radiation Oncology
Oncology	Renal
Oncology	Sarcoma
Oncology	Uterine

Other Areas of Expertise

Study Phase Capabilities
Phase I, Phase II, Phase III, Phase IV

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, Academic
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?	Less than 30
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	National Hospital Organization Hokkaido Cancer Center Clinical Trial Management Office
Department Contact Phone Number	81118119111
Department Contact Email Address	100-mb08chk4@mail.hosp.go.jp
Is your Facility able to initiate study activities prior to IRB/ERB/Ethics Committee protocol approval?	No
What types of IRB/ERB/Ethics Committee does your Facility use?	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

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

Local Lab

Is your Facility using a Local Lab?	Yes

Local Lab: Clinical Laboratory	
Lab Name	Clinical Laboratory
Lab Contact First Name	Michio
Lab Contact Last Name	Sato
Address	4-2-3-54 kikusui shiroishi-ku, 2F, sapporo, Hokkaido, Japan, 003-0804
Phone Number	+81118119111
Fax Number	+81118119232
Email Address	100-ml080100c@mail.hosp.go.jp
Local Lab Accreditation	CAP, ISO, Others
Other Local Lab Accreditation	Japanese Medical AssociationJapanese Association of Medical Technologists
Additional Questions	
Does your Facility have a SOP/written procedure for documenting bio-specimen (Sample) processing steps/chain of custody?	Yes
Do your written procedures ensures that study-specific temperature bio-specimen storage requirements are known to responsible staff to ensure compliance?	Yes
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?	On site collection and Processing
Does your Facility has established processes to oversee staff compliance with study-specific lab manual instructions for bio-specimen processing?	No
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)	No

Attachments

Document Type	Document Name	Document Description
No Records		

CONSENT & TRAINING

Consent	
Does your Facility have a written SOP/Policy/Procedure for: Informed Consent?	Yes
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.	APRIN e-learning program (eAPRIN)
Do you have a process or program in place to retrain research staff when a protocol is amended?	Yes
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?	Yes
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)?	Yes
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?	Yes
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, Positron Emission Tomography Scan, X-Radiation, 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.?	Yes
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?	No
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	Daily
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?	Yes
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?	No
How frequently can temperature measurement occur? Check the most frequent measurement your equipment can support.	Daily
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.	Daily
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?	Yes

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, Wi-Fi
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)	No
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?	Yes
Does your Facility prohibit the use of an external USB device (e.g. to download and send data from a temperature monitoring device)?	Yes
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
clinical trial management office	4-2-3-54 kikusui shiroishi-ku, 3F, sapporo, Hokkaido, Japan, 003-0804	+81118119111	+81118119232	100-ml080100c@hosp.go.jp	

Investigational Product Storage Location

IP Storage Location Name	Address	Phone Number	Fax Number	Email Address
clinical trial management office	4-2-3-54 kikusui shiroishi-ku, 3F, sapporo, Hokkaido, Japan, 003-0804	+81118119111	+81118119232	100-ml080100c@hosp.go.jp

Investigational Product Storage Equipment	
Identify the Investigational Product Storage Equipment at your Facility	Refrigerator (2 to 8 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?	Yes
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?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of Investigational Product?	No
Does your Facility have a written SOP/Policy/Procedure for destruction of Investigational Product?	Not Applicable
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?	Not Applicable
Is the storage area for controlled substances securely constructed with restricted access in accordance with local law?	Not Applicable
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)?	

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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)	
Please list any access limitations/ requirements for eISF/eReg	
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?	
Please list any access limitations/requirements for the Electronic Medical Records.	
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?	
Are monitors able to access EHR/EMR while off site?	
Does your Facility require Sponsor representative to sign any local form (paper or electronic) for access, or any other purpose?	
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:	Phone, Fax, 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	Apices viedocmacro
Does your site/institution and/or local regulations allow remote source data verification of study participant data to support remote monitoring?	

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
No Records				

Associated/Confirmed Site Users				
Name	**User Role**	**Request Affiliation Date**	**Affiliation Status change Date**	**Affiliation Status**
Watanabe,Kenichi	Investigator	10-Jul-2022	10-Jul-2022	Confirmed
Shikishima,Karin	Investigator	15-Jun-2023	15-Jun-2023	Confirmed
Moriguchi,Takuya	Investigator	22-Jul-2025	22-Jul-2025	Confirmed
Mizugaki,Hidenori	Investigator	01-Mar-2023	01-Mar-2023	Confirmed
Fujimoto,Katsuya	Investigator	08-Mar-2023	08-Mar-2023	Confirmed
Kobayashi,Shinya	Clinical Research User	08-Mar-2023	08-Mar-2023	Confirmed
Kamada,Miku	Clinical Research User	12-May-2023	12-May-2023	Confirmed
Ito,Kazuyoshi	Clinical Research User	09-May-2023	09-May-2023	Confirmed
Ishihara,Chie	Clinical Research User	09-May-2023	09-May-2023	Confirmed
Seino,Madoka	Clinical Research User	18-May-2023	18-May-2023	Confirmed
Tashiro,Mayumi	Clinical Research User	12-May-2023	12-May-2023	Confirmed
Kadomasu,Shiori	Clinical Research User	10-May-2023	10-May-2023	Confirmed
Asahina,Hajime	Investigator	08-May-2023	08-May-2023	Confirmed
Hatakeyama,Yuki	Investigator	12-May-2023	12-May-2023	Confirmed
Takada,Shinya	Clinical Research User	08-May-2019	08-May-2019	Confirmed
Souma,Mikiko	Clinical Research User	12-Nov-2019	12-Nov-2019	Confirmed
Miura,Kiyofumi	Clinical Research User	12-Nov-2019	12-Nov-2019	Confirmed
Takamichi,Yuka	Clinical Research User	12-Nov-2019	12-Nov-2019	Confirmed
Todo,Yukiharu	Investigator	07-Jan-2020	07-Jan-2020	Confirmed
Echizen,Ayana	Clinical Research User	25-Nov-2019	25-Nov-2019	Confirmed
Uno,Asami	Clinical Research User	25-Nov-2019	25-Nov-2019	Confirmed
Kato,Hidenori	Investigator	12-Dec-2019	12-Dec-2019	Confirmed
Minobe,Shinichiro	Investigator	20-Dec-2019	20-Dec-2019	Confirmed
Tajima,Hiroe	Clinical Research User	25-Nov-2019	25-Nov-2019	Confirmed
Yamagishi,Kayo	Clinical Research User	13-Nov-2019	13-Nov-2019	Confirmed
Nitta,Masayo	Clinical Research User	25-Nov-2019	25-Nov-2019	Confirmed
Shirooka,Yukari	Clinical Research User	25-Nov-2019	25-Nov-2019	Confirmed
Nishiwaki,Satoko	Clinical Research User	18-Nov-2019	18-Nov-2019	Confirmed

Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Itagaki,Yoriko	Clinical Research User	12-Nov-2019	12-Nov-2019	Confirmed
Nakagawa,Yuko	Clinical Research User	12-Nov-2019	12-Nov-2019	Confirmed
Takahashi,Masato	Investigator	12-Dec-2019	12-Dec-2019	Confirmed
Minowa,Kaoru	Investigator	26-Dec-2019	26-Dec-2019	Confirmed
Fukai,Yuta	Clinical Research User	25-Jun-2020	25-Jun-2020	Confirmed
Sato,Yoshimi	Clinical Research User	24-Jul-2020	24-Jul-2020	Confirmed
Tsuruta,Tomohiko	Investigator	18-Sep-2020	18-Sep-2020	Confirmed
Yamada,Ryutaro	Investigator	18-Sep-2020	18-Sep-2020	Confirmed
Harada,Masao	Investigator	15-Oct-2020	15-Oct-2020	Confirmed
Oizumi,Satoshi	Investigator	15-Oct-2020	15-Oct-2020	Confirmed
Saito,Chiharu	Clinical Research User	10-Nov-2020	10-Nov-2020	Confirmed
Iguchi,Aki	Clinical Research User	10-Nov-2020	10-Nov-2020	Confirmed
YAMADA,NORIYUKI	Investigator	05-Nov-2020	05-Nov-2020	Confirmed
Yokouchi,Hiroshi	Investigator	05-Nov-2020	05-Nov-2020	Confirmed
Kudo,Risa	Investigator	24-Nov-2020	24-Nov-2020	Confirmed
Morioka,Yuuki	Investigator	10-Nov-2020	10-Nov-2020	Confirmed
Nishikawa,Noboru	Investigator	15-Dec-2020	15-Dec-2020	Confirmed
Nishiyama,Noriaki	Investigator	04-Dec-2020	04-Dec-2020	Confirmed
nishikawa,yukiko	Investigator	01-Dec-2020	01-Dec-2020	Confirmed
Fukumoto,Shin-ichi	Investigator	11-Dec-2020	11-Dec-2020	Confirmed
Harabayashi,Toru	Clinical Research User	18-Jan-2021	18-Jan-2021	Confirmed
Maruyama,Satoru	Investigator	18-Feb-2021	18-Feb-2021	Confirmed
Kikuchi,Minoru	Clinical Research User	05-Apr-2021	05-Apr-2021	Confirmed
SUZUKI,YUTARO	Investigator	29-Apr-2021	29-Apr-2021	Confirmed
Minatogawa,Hideki	Investigator	25-Jun-2021	25-Jun-2021	Confirmed
Shiroshita,Takashi	Clinical Research User	21-Sep-2021	21-Sep-2021	Confirmed
Sakakibara,Jun	Investigator	01-Sep-2021	01-Sep-2021	Confirmed
Yamamoto,Mitsugu	Investigator	03-Dec-2021	03-Dec-2021	Confirmed

Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Tomioka,Nobumoto	Investigator	21-Dec-2021	21-Dec-2021	Confirmed
Maeda,Hideki	Investigator	14-Dec-2021	14-Dec-2021	Confirmed
Sato,Keiichi	Clinical Research User	24-Mar-2022	24-Mar-2022	Confirmed
Sato,Yuma	Investigator	18-Apr-2022	18-Apr-2022	Confirmed
Kawamura,Nanami	Clinical Research User	11-Oct-2022	11-Oct-2022	Confirmed
Yamada,Natsue	Clinical Research User	16-Oct-2024	16-Oct-2024	Confirmed
Ozu,Shunsuke	Investigator	05-Dec-2024	05-Dec-2024	Confirmed
Suzuki,Hidetaka	Clinical Research User	03-Apr-2025	03-Apr-2025	Confirmed
Ohashi,Kasumi	Clinical Research User	13-Mar-2025	13-Mar-2025	Confirmed
sanada,kayo	Clinical Research User	15-Apr-2025	15-Apr-2025	Confirmed
Yamamoto,Koyori	Clinical Research User	12-Jun-2026	12-Jun-2026	Confirmed
Uegaki,Mai	Clinical Research User	12-Jun-2026	12-Jun-2026	Confirmed
Tanno,Maya	Clinical Research User	12-Jun-2026	12-Jun-2026	Confirmed
tokunaga,nao	Clinical Research User	30-Oct-2023	30-Oct-2023	Confirmed
Murooka,Hidetaka	Clinical Research User	01-Nov-2023	01-Nov-2023	Confirmed
Tachikawa,Hanae	Investigator	07-Nov-2023	07-Nov-2023	Confirmed
sakai,toshiya	Investigator	06-Nov-2023	06-Nov-2023	Confirmed
Koiso,Yukari	Clinical Research User	08-Nov-2023	08-Nov-2023	Confirmed
Furukawa,Shizuka	Clinical Research User	18-Jun-2025	18-Jun-2025	Confirmed
Kamada,Mai	Clinical Research User	18-Jun-2025	18-Jun-2025	Confirmed
Mizobuchi,Shohei	Investigator	22-May-2025	22-May-2025	Confirmed
Araki,Risa	Clinical Research User	12-Jun-2025	12-Jun-2025	Confirmed
Aramachi,Yukari	Investigator	11-May-2025	11-May-2025	Confirmed
Kamada,Ryohei	Investigator	21-May-2025	21-May-2025	Confirmed
Ueno,Miki	Clinical Research User	01-Mar-2024	01-Mar-2024	Confirmed
Nomoto,Keiko	Clinical Research User	04-Mar-2024	04-Mar-2024	Confirmed
ebata,ko	Investigator	11-Dec-2023	11-Dec-2023	Confirmed
Takahashi,Shogo	Investigator	04-Jan-2024	04-Jan-2024	Confirmed

Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Takada,Norikata	Clinical Research User	23-Aug-2024	23-Aug-2024	Confirmed
Abe,Sachiko	Clinical Research User	29-Aug-2024	29-Aug-2024	Confirmed
Ikenaga,Shigehiro	Clinical Research User	20-Sep-2024	20-Sep-2024	Confirmed
Saito,Haruna	Clinical Research User	25-Sep-2024	25-Sep-2024	Confirmed
Yamaguchi,Honoka	Clinical Research User	20-Sep-2024	20-Sep-2024	Confirmed
Kobayashi,Shiori	Clinical Research User	25-Sep-2024	25-Sep-2024	Confirmed
Sasaki,Mio	Clinical Research User	18-Sep-2024	18-Sep-2024	Confirmed
Sarashina,Takeyoshi	Investigator	20-Dec-2024	20-Dec-2024	Confirmed
Adachi,Chihiro	Clinical Research User	19-Dec-2024	19-Dec-2024	Confirmed
Kanazawa,Tomoko	Clinical Research User	15-Apr-2024	15-Apr-2024	Confirmed
Yoshida,Yukiko	Investigator	21-Apr-2024	21-Apr-2024	Confirmed
Anada,Yuri	Clinical Research User	15-Mar-2024	15-Mar-2024	Confirmed
Sawano,Takako	Clinical Research User	10-Apr-2024	10-Apr-2024	Confirmed
Saito,Masaaki	Clinical Research User	08-Mar-2024	08-Mar-2024	Confirmed
Koganezawa,Chinatsu	Investigator	16-Apr-2024	16-Apr-2024	Confirmed
Nakamura,Sai	Clinical Research User	07-Mar-2024	07-Mar-2024	Confirmed
Sato,Misako	Clinical Research User	15-Aug-2025	15-Aug-2025	Confirmed
Kataoka,Yuka	Clinical Research User	13-Aug-2025	13-Aug-2025	Confirmed
Shirai,Kuki	Clinical Research User	29-Aug-2025	29-Aug-2025	Confirmed
Yahaba,Yoshie	Clinical Research User	25-Sep-2025	25-Sep-2025	Confirmed
Aiba,Yukari	Clinical Research User	29-Aug-2025	29-Aug-2025	Confirmed
Momoi,Yoshinori	Clinical Research User	09-May-2024	09-May-2024	Confirmed
Shigehara,Mai	Clinical Research User	06-Jan-2026	06-Jan-2026	Confirmed
Ueda,Ayaka	Clinical Research User	17-Feb-2026	17-Feb-2026	Confirmed
Miyakawa,Minori	Clinical Research User	05-Feb-2026	05-Feb-2026	Confirmed
Kimura,Hirotoshi	Investigator	21-May-2026	21-May-2026	Confirmed
Ikezawa,Masafumi	Investigator	12-May-2026	12-May-2026	Confirmed
Harada,Shinpei	Investigator	07-May-2026	07-May-2026	Confirmed

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Name	User Role	Request Affiliation Date	Affiliation Status change Date	Affiliation Status
Mori,Hinata	Investigator	12-May-2026	12-May-2026	Confirmed
Hama,Miyuki	Clinical Research User	08-Apr-2026	08-Apr-2026	Confirmed
Uehara,Takashi	Investigator	09-Jun-2026	09-Jun-2026	Confirmed