

SHIONOGIは、以下の研究テーマ・ニーズに関するシーズやパートナーを求めています

SHONOGI Wish List





[Infection (1/3)]

テーマ Theme	ニーズ゛ "Needs"	アイデア・コンセプト Idea/Concept	創薬標的 DrugTarget	シーズ化合物 Seeds Compound	評価系·創薬技術 EvaluationSyetem· DrugDiscoveryTechnology	診断技術・BM Diagnosis・ Biomarker
A cure for HIV and HBV that replaces chronic treatment	How to make a cure possible and how to confirm it					
Drug discovery for pandemics caused by RNA viruses	Viral drug target functions and molecules that are expected to act broadly against RNA viruses					
Research on host response to infectious diseases	Establishment of treatments to prevent the severity and death of acute respiratory viral infections: Acquisition of intervention methods, drug discovery targets, and evaluation techniques					
	By intervening with the immune system against chronic infections, it is possible to improve the severity of the infection or cure it completely.					
Mycobacterial infection	 Non-tuberculous mycobacterial drug discovery improving infection mortality and quality of life: Target functions/molecules, and new modalities that can control them. Associated with these are rapid screening technologies, non-clinical evaluation systems that can be bridged with clinical practice, rapid diagnosis of target pathogen identification and drug susceptibility, methods for monitoring infection status, and innovative manufacturing technologies. Establishment of methods to prevent recurrence (relapse/reinfection) 					
	Establishment of a diagnostic method that can determine early termination of treatment Search for indicators that allow patients to feel the effects					
Intractable microbial infections	Drug discovery aiming for zero deaths due to infection: Target functions/molecules, and new modalities that can control them. Associated with these are rapid screening technologies, non-clinical evaluation systems that can be bridged with clinical practice, rapid diagnosis of target pathogenic microorganisms and drug susceptibility, methods for monitoring infection status, and innovative manufacturing technologies.					







[Infection (2/3)]

テーマ Theme

Respiratory diseases

MRSA infection

LAP (Long-acting parenteral formulation) drug discovery

Oral long-term drug

analysis Nucleic acid analog drug discovery research crystal analog Drug discovery method aimed at covalent Fundan binders

drug di Drug discovery technology and screening Fundan method for targeting undraggable targets attackir Prediction systems and other drug discovery Ames a assets aimed at avoiding Ames acid and



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New assets that can safely remove senescent cells, ideas for drug discovery, and technical infrastructure for evaluating senescent cell removal (It is known that senescent cells accumulate due to infection, and it is thought that this may be the starting point for the disease becoming more severe and difficult to treat. Therefore, we will build a foundation and consider applying it to new drug discovery to treat infections. Consider expanding the scope of development to respiratory diseases.)					
Drug discovery ideas, technology bases, and assets that restore/improve functions of degraded MP and T cells					
Drug discovery targets and assets that can be expected to restore tissue and improve respiratory function in respiratory diseases The Information & Method for undestanding the unmet medical needs for MRSA. (Especially the information about beta-lactam)					
New fundamental technology and evaluation method for LAP					
Fundamental technologies and evaluation methods to realize oral long-lasting drugs					
h Mechanism analysis method, co-crystal structure analysis method (technology that easily obtains the crystal structure of the ternary complex of nucleic acid analog triphosphate + RNA + RdRp), etc.					
Fundamental technologies and evaluation methods for drug discovery aimed at covalent binders					
Fundamental technologies and evaluation methods for attacking undraggable targets					
ryAmes avoidance measures and prediction tools in nucleic acid analog drug discovery, etc.					





[Infection (3/3)]

テーマ Theme

	Cell construction technology that allows /arious virus assays	Improving performin current si tasks and the cell lin (CPE assa
ſ		Shorter ir extrapola



アイデア Idea

ng task efficiency and reducing time when ing various virus assays. Measures to address the situation where it is difficult to simply increase d horizontally compare activity values because line changes for each virus.

say if possible. Techniques other than MucilAir.)

in-vivo testing and more accurate clinical ation



ア・コンセプト /Concept	創薬標的 DrugTarget	シーズ化合物 Seeds Compound	評価系·創薬技術 EvaluationSyetem· DrugDiscoveryTechnology	診断技術・BM Diagnosis・ Biomarker



[Central Nervous System (1/2)]

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	Mouse MRI measurement and analysis technology				
Biomarker research: fMRI	Preparation of IT/analysis environment for constructing fMRI-based symptoms/neural network database				
Biomarker research: patient stratification	Biomarkers to identify patient segments that are particularly likely to benefit from drugs to improve cognitive function				
Biomarker research: Others	Biomarker analysis technology to detect the mechanism of action on drug discovery targets and predict pathological changes				
Drug discovery for the treatment of core symptoms of dementia	Drug targets or their tool compounds that can be effective even when dementia-related proteins accumulate in the brain. for example: 1) Targets that can enhance memory through transcription and translation 2) Targets that can enhance memory through energy metabolism				
	Efficacy evaluation system that can reflect brain activity (non-clinical)				
	Measurement technology for cell-specific biomarkers using extracellular vesicles (EV)				
Drug discovery for the treatment of behavioral and psychiatric symptoms of	Novel drug target expected to be effective in treating BPSD				
dementia (BPSD)	Pathological model and evaluation system enabling drug efficacy evaluation for BPSD				





[Central Nervous System (2/2)]



Drug discovery for the treatment of attention deficit hyperactivity disorder (ADHD)

Dependency

Autism spectrum disorder (ASD)

New inte mechani

Evaluatio efficacy

Evaluatio

Interme is highly

Translati

Model co

Ideas, co ADHD d to chang

Elucidati targets

A simple than exist discrimir

Novel dr

Novel dr

ニーズ "Needs"	アイデア・コンセプト Idea/Concept	シーズ化合物 Seeds Compound	評価系·創薬技術 EvaluationSyetem· DrugDiscoveryTechnology	診断技術・BM Diagnosis・ Biomarker
tervention points and targets different from existing nisms]			
tion system that can quantitatively determine y (non-clinical)				
tion system with higher human extrapolation				
ediate phenotype (BioMarker) related to ADHD that ly extrapolable to humans				
itional medicinal biomakers				
construction and expansion of evaluation system				
concepts, and drug targets for the next generation drug and exploration of medical needs in response nges in the external environment				
tion of dependence mechanisms and new drug				
le and highly extrapolable evaluation model other kisting evaluation methods, such as CPP, ination, or self-administration				
drug targets for relapse prevention				
drug targets based on ASD onset mechanism				





[New Therapeutic Area (1/2)]



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	High throughput and high resolution inner ear tissue analysis technology Cutting-edge nervous system and inner ear tissue immunostaining analysis technology					
	High throughput and high resolution inner ear tissue analysis technology Automated technology for inner ear whole mount tissue immunostaining					
Sensorineural Hearing Loss: Non-clinical evaluation systems	High throughput and high resolution inner ear tissue analysis technology Image analysis technology using machine learning					
	Hearing loss non-clinical model, evaluation system Nonclinical in vivo model of sudden hearing loss					
	Hearing loss non-clinical model, evaluation system Non-clinical hearing loss model with less variation in pathogenesis					
	Hearing loss non-clinical model, evaluation system Adult mouse ex vivo inner ear tissue evaluation system					
	Inner ear administration technique Intracochlear administration technique to mice and rats					
Sensorineural Hearing Loss: Non-clinical evaluation systems, DDS techniques	Inner ear administration technique Continuous injection technology into the inner ear using a pump for mice and rats					
Sensorineural Hearing Loss: Non-clinical evaluation systems	How to assess hearing quality System for evaluating auditory nerve function					
	Biomarker that can detect hair cell shedding					
Sensorineural Hearing Loss: Research on pathophysiology, diagnosis and	Biomarker that can detect auditory neuropathy					
stratification of hearing-impaired patients	Biomarker that can detect inner ear inflammation and oxidative stress					



[New Therapeutic Area (2/2)]

テーマ Theme

Drug discovery research for sleep apnea syndrome Non-clinical /Clinical evaluation systems	High-thr system One of reduced negative upper ai with EM controllin it would measure
Drug discovery research for sleep apnea syndrome Non-clinical evaluation systems	Arousal One of lowered which av concentre esophag How to a Loop gat Respire accordine sleep, bu OSAS is breathine function
Drug discovery research for sleep apnea syndrome Understanding the unmet medical needs for OSAS	Ideas fo for RWD analysis A techno
Allosteric modulator or biased ligand screening technology	The way betweer binding Biased li avoidanc
Simultaneous evaluation technology for arousal threshold, loop gain, and genioglossus muscle	A metho by treat (some c



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hroughput genioglossus muscle activity evaluation					
٦					
of the important pathological traits of OSAS is the					
d responsiveness on the genioglossus muscle to					
ve pressure caused by respiratory effort resulting					
airway obstruction. How to evaluate this feature					
MG(ElectroMyoGraphy) is needed. Since the					
lling nerves change depending on the sleep stage,					
d be better if there was a system for simultaneous					
rement with EEG(ElectroEncephaloGraphy).					
I threshold evaluation system					
of the important pathological traits of OSAS is the darousal threshold that is the phenomenon in					
awakening occurs when blood CO2 and O2					
itrations rise and negative pressure in the					
agus exceeds a certain threshold during sleep.					
assess this function in vivo is needed.					
ain evaluation system					
piration is controlled by chemical regulation					
ing to the blood CO2 or O2 concentration during					
but one of the important pathological traits of					
is that this control function becomes excessive and					
ing becomes unstable. How to assess this					
n in vivo is needed.					
for exploring and validating OSAS needs. Proposals					
D obtained from OSAS patients and ideas for its					
IS.					
nology that can screen for allosteric modulators.					
ay to obtain compounds with high selectivity					
en receptor subtypes that have very similar					
g sites.					
ligand screening technology that may lead to					
nce of side effects by selecting 2nd signals					
nod that can evaluate additive/synergistic effects					
ating each OSAS phenotypic trait at the same time					
combinations may be possible, but not all).					





[Vaccine (1/2)]

テーマ Theme

Themes related to predicting the emergence of new pathogens and mutant of pathoc strains

Antigen Design

Particularization of vaccine antigens

Vaccine antigen evaluation technology



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 Technologies related to predicting the emergence of nepathogens and mutant strains In silico analysis method for predicting the emergence of pathogenic mutant strains that enable immune evase and/or enhanced infectivity, and prediction method for next pandemic infectious disease species Technologies related to search and design of vaccine antigens New antigens that are vaccine candidates for infection diseases, including viruses and bacteria Technologies related to search and design of vaccine antigens In silico techniques for searching for New antigens that are vaccine candidates for infectious diseases, including viruses and bacteria Technologies related to particularization of vaccine antigens New particle technology such as VLPs and nanoparti to enhance the immunogenicity of antigens 	e ion us at			
Technologies related to particularization of vaccine antigens New microparticulation technology capable of loadin multiple antigens and enhancing cross-immunogenicity Technologies related to vaccine antigen evaluation Systems and evaluation systems that can predict the vivo immunity induction ability of vaccine antigens in v Technologies related to vaccine antigen evaluation A small animal model that extrapolates highly to humans in evaluating the immunogenicity of vaccines Technologies related to vaccine antigen evaluation Protein engineering technology for optimal vaccine antigen design	in			





[Vaccine (2/2)]

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Antigen Design	Technologies related to search and design of vaccine antigens Technology related to antigen design or technologies or screening method to enhance cross- immunogenicity or to expand the spectrum to pathogens of vaccines including both of Bcell and Tcell reactivity. Technologies related to new administration routes of					
New administration routes of vaccines	vaccines New technologies related to new delivery systems that can effectively inducer mucosal immunity by nasal or oral administration routes					
	Technologies related to new administration routes of vaccines Innovative ideas and concepts related to immunity induced by new administration routes					
	Vaccine adjuvants New targets for creating adjuvants that can enhance the persistence of antibody induction					
	Vaccine adjuvants New technologies for creating adjuvants that can enhance the persistence of antibody induction					
Vaccine adjuvants	Vaccine adjuvants New targets for creating novel adjuvants that induce mucosal immunity in the nasal cavity and intestinal tract					
	Vaccine adjuvants New technologies for creating novel adjuvants that induce mucosal immunity in the nasal cavity and intestinal tract					
	Vaccine adjuvants Pre-clinical evaluation system (animal model, in vitro evaluation system) that can predict clinical side effects caused by adjuvants (local pain, fever, etc.)					







[Drug Discovery Technology (1/2)]

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Polyclonal Antibody Drug Development	Polyclonal Antibody Manufacturing Technology					
	An efficient method for evaluating the three-dimensional structure of RNA (including highly accurate prediction method)					
Research on virtual screening of small molecule compounds against RNA targets	High through put in silico screening technology for RNA					
High through put in silico screening technology	virtual screening methods that can screen a large-scale virtual library with high accuracy and speed					
Research on protein design by in silico technology	Protein design methods utilizing state-of-the-art AI technologies					
	Technologies to identify targets that are the main cause of disease and would be beneficial to degrade (especially those related to infection diseases, excludes cancer and tumors)					
	Technologies that can comprehensively analyze on-target and off-target					
	Efficient evaluation method for degraders (cell-free if possible, including computational methods)					
	Identification of intracellular PPI targets for treatment of SHIONOGI's focus diseases including infectious diseases					
Intra-cellular drug delivery research or technologiesfor medium sized compounds	An efficient method for transferring peptide drug molecules into cells					
Research on docking of peptides to target	Technology for highly accurate docking (SBDD) and simulation (LBDD) of peptide drug molecules. In particular, calculation methods or indicators to measure the degree of conformational locking of peptide drug molecules					
	An automated system in chemistry which can operate throughout the day					





(Drug Discovery Technolog

テーマ Theme	
In vitro evaluation method for ocular toxicity	An in vitr identify tl
Research on ototoxicity	An in vitr and regen evaluation Brainsten
System toxicology research	In silico t estimate networks
Imaging technology for organ toxicity	A technol live anima substitute evaluation prioritizat
Technology that can identify drug targets with in silico/AI	Developm distinguis on the inp and mach biochemic

ニーズ "Needs"

ro model that can evaluate ocular toxicity and the mechanism of ocular toxicity

ro evaluation system that can evaluate damage eneration of inner ear hair cells and an in vivo on system for ototoxicity using ABR (Auditory) m Response), etc.

toxicity mechanism analysis system that can the toxicity mechanism from molecular s, expression changes, etc.

plogy for predicting toxicity target organs using nal image diagnosis (CT, MRI, etc.) as a te for pathological examination or tests or preon for prioritizationpre-evaluation for ation

ment of an alert system that roughly shes whether it is adverse or non-adverse based nput of biochemical data; ie., a system using AI chine learning technology based on non-clinical nical parameter data (quantitative values) from toxicity studies conducted in-house



アイデフ

Idea

第価系・創薬技術 診断技術・BM EvaluationSystem・ DrugDiscoveryTechnology Biomarker
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[CMC R&D Technology (1/2)]						
テーマ Theme	ニーズ "Needs"	アイデア・コンセプト Idea/Concept	評価系・創薬技術 EvaluationSyetem・ DrugDiscoveryTechnology	診断技術・BM Diagnosis・Biomarker	その他 others	
Crystal and powder properties of drug substances (small molecule)	Evaluation and control technology for crystal/powder properties of drug substances (small molecule)					
Innovative medium component for cell culture	To increase and improve cell culture performance for biologics production.					
LAP (Long-acting parenteral formulation) formulation tecnology	 To avoid subcutaneous/intramuscular irritation To increase dose volume in subcutaneous/intramuscular 					
	An inexpensive LAP formulation to be developed					
_AP formulation human PK prediction technology	To predict the human PK of LAP preparations in a short period of time.					
Oral administration long-acting formulatio technology	nTo release the drug for NLT one week with an oral formulation.					
Long-acting formulation technology for mproved absorption	To sustained release a drug improved absorption without reducing AUC and Ctrough.					
Stability improvement technology	To store products that are stored in the refrigerator at room temperature.					
Child-safe preservatives	To develop a suspension that does not allow microorganisms to grow for a long period of time.					
orally disintegrating tablets (ODTs) formulation tecnology	Long-term patented technology for orally disintegrating tablets (ODTs) Estimated patent term: Exclusivity period over 2040					





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The	eme

ICMC R8	D Technology	(2/2)			
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Characterization technology of lipid particl and emulsion	Understanding of lipid particle and emulsion e charcterization such as existence mode of the components, the component release profile in biological environment, particle solidness and membrane fluidity.				
Device technology for injectable formulation to administrate 2 types liquid with one shot	Enhancement usability without complicated preparation				
Nebulizer that can be used for pulmonary administration of drugs at home	Co-development of a nebulizer suitable for home administration				
GMP manufacturing technology for ligand- modified mRNA-LNP (ligand addition technology after LNP formation)	To increase manufacturing feasibility in preparation for the launch of ligand-modified LNP formulations.				



