

CSL 2024 Research Acceleration Initiative

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Research Innovation

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Overview of CSL



CSL Research
Acceleration Initiative



Benefits of collaborating with CSL



CSL's core Therapeutic Areas



Areas of interest for collaboration



Questions



Overview of CSL







Top 25 Biotech Companies of 2024

Rank	Company	Ticker Symbol	Market Cap (US\$ Billion)
1	Novo Nordisk	NOVO-B (CPH)	430.96
2	Thermo Fisher Scientific	TMO (NASD)	189.20
3	Amgen	AMGN (NASD)	143.98
4	Gilead Sciences Inc	GILD (NASD)	98.41
5	Regeneron Pharmaceuticals	REGN (NASD)	91.51
6	Vertex Pharmaceuticals	VRTX (NASD)	90.24
7	CSL Ltd	CSL (ASX)	84.82
8	Chugai Pharmaceutical	4519 (TOKYO SE)	60.86
9	Daiichi Sankyo	4568 (TOKYO SE)	54.33
10	Seagan	SGEN (NASD)	41.31
11	Jiangsu Hengrui Medicine Co Ltd	600276 (SHSE)	40.59
12	Samsung Biologics	207940 (KRX KE)	38.31
13	Agilent Technologies	A (NYSE)	37.16
14	Sun Pharmaceutical Industries	SUNPHARMA (NSE)	35.54
15	Biogen	BIIB (NASD)	34.68
16	WuXi App Tec	603259 (SSEC)	31.46
17	Moderna	MRAN (NASD)	30.61
18	Lonza	LONN (SWX)	27.77
19	Argenx	ARGX (NASD ENX)	26.79
20	BioNTech	BNTX (NASD)	24.79

CSL's Key Global R&D Locations





Research Acceleration Initiative



CSL's Research Acceleration Initiative

Objective: to build relationships with entrepreneurial researchers and fastrack discovery of innovative medicines that address unmet needs

Why? Early collaborations with high quality academic partners are key to building a sustainable pipeline

CSL's RAI provides a differentiated approach to partnering:

- ✓ Up to USD \$400,000 funding over 2 years
- ✓ CSL scientific champion assigned to each project.
- ✓ Focused on early-stage projects
- ✓ Simple and fast 300-word initial application
- ✓ Clear and transparent timelines



WHY COLLABORATE WITH CSL?



Global capabilities on your doorstep.



Work with one of the world's leading biotech companies.



Funding for successful proposals.



Access to commercial R&D, clinical, intellectual property, marketing and manufacturing expertise.



Accelerate translation of your research to deliver new therapies to patients.

CSL Research Acceleration Initiative

Seeking Expressions of Interest from Research Organizations

CSL is a leading global biotech company that develops and delivers innovative biotherapies to help people living with life-threatening medical conditions live full lives.

CSL's Research Acceleration Initiative aims to fast-track discovery of innovative biotherapies through partnerships between CSL and global research organizations. These partnerships provide funding and access to industry experts for scientists working on novel biotherapeutic strategies in CSL's therapeutic areas.

Expressions of interest are sought from Business
Development / Commercialization representatives across
global research organizations that wish to participate in
the 2024 CSL Research Acceleration Initiative.

The 2024 Research Acceleration Initiative will focus on innovative research projects that address unmet medical needs and are aligned with CSL's **Therapeutic Areas** and scientific **Platforms**:



To register your research organisation please email RAI@csl.com.au by 15th December 2023

CSL has invested in 30+ RAI partnerships since 2019

"We had a **stellar experience participating** in the CSL RAI process. The information material, informational webinars, and access to the program team for Q&A was well received by our faculty..."

RAI 2023 participant University of Pittsburgh

"Peerless experience – timely, transparent, actionable communication."

RAI 2023 participant University of British Columbia

"Well-designed, easy and clear process. **Highly engaged and highly responsive to all questions** and provided well
contemplated and customised feedback."

RAI 2023 participant University of Toronto "It has been a great pleasure to collaborate with our colleagues at CSL. The Research Acceleration Initiative (RAI) is an **outstanding platform that helps bridge the academic world with industry**."

RAI 2021 awardee Justus-Liebig-Universität Giessen "CSL has proven to be an exceptional collaborator,

fostering a scientifically focused partnership marked by open scientific exchange and generosity. Their extensive research expertise has consistently enriched our collaborative efforts making the interaction with CSL an indispensable asset to our joint projects"

> RAI 2021 awardee Klinikum der Universität München (KUM)

> > "...the webinar session was very useful because it **clearly indicated which areas CSL was interested in funding**, thereby allowing me to focus my thoughts on them."

> > > RAI 2022 awardee Nanyang Technological University

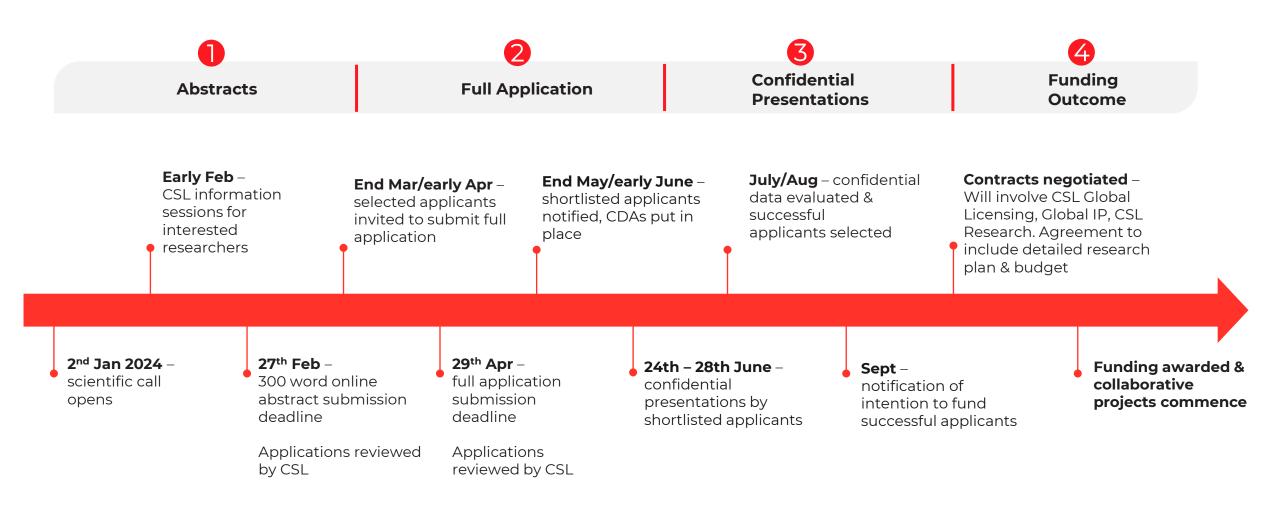
"...the opportunity to address **feedback**from CSL and to refine the project was
particularly valuable"

RAI 2023 participant The University of Adelaide

"The types of projects CSL were looking for was made very clear, the process of submitting an **application was easy** and did not require excessive time or effort."

RAI 2023 participant Auckland UniServices Ltd

CSL 2024 Research Acceleration Initiative Process



No obligation for registered organizations to submit applications

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Agreement Guidance



Separate collaboration agreements will be negotiated for each project which reflect the nature of the project, nature of funding and support, and the contributions of both parties



Under these negotiated agreements, CSL will be granted certain rights of interest to the program results for further R&D and/or commercialization



© Collaboration agreements will typically include the following terms (although CSL may propose other conditions depending on the nature of the project):

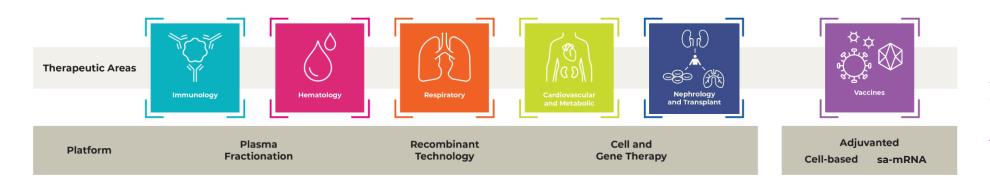
- Research organization will generally own results arising under the project
 - CSL would typically own any results which relate to proprietary CSL products or materials contributed to the project or may seek joint-ownership of results to which it has made a significant contribution (e.g. protein or antibody discovery and engineering activities).
 - The RAI is designed to accelerate the translation of novel discoveries made by research scientists for proposals outside this scope, we may propose that projects be progressed outside the RAI
- CSL will be granted an exclusive option to negotiate an exclusive, worldwide licence
- CSL supports publication of research outcomes

Further details on agreement terms can be provided on request

Eligibility

To be eligible to apply, researchers/clinicians must satisfy the following 2 conditions:

- 1. Be employed by a research organization registered to participate in the 2024 Research Acceleration Initiative
- 2. Submit a 300-word online abstract that is aligned with CSL's Therapeutic Areas and scientific Platforms:



Specific indications of focus within each TA are provided on slides 28-35.

Abstract submission via online portal

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Step 1/2 - Lead Investigator Information Applications for the 2024 CSL Research Acceleration Initiative open 2nd January 2024 and close 27th February 2024. Applications received outside these dates (before or after) will not be reviewed. 1 Fields with * are mandatory		Step 2/2 - Describe your opportunity and confirm submission Modality										
		Please describe and categorize your opportunity.						Recombinant protein (including antibodies)	Plasma protein	Gene therapy		
		• Fields with * are mandatory						Cell therapy	RNA therapeutic (non-vaccine)	Vaccine (including RNA vaccines)		
		Proposal Title *						Peptide	Small molecule	Other modality		
First Name *	Salutation							Opportunity Type *				
		Primary Focus Area *						Novel therapeutic candidate	Novel therapeutic target	Drug target discovery		
Last Name •	Job Title *	(8)	\Diamond			G _P D	\$\frac{1}{4} \psi \qquad \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq \qqqq \qqq		Vaccine – novel target or candidate	Vaccine – immune mechanisms	RNA platform development	
		(/(35/\)				∞ 86 €	4 6		Patient stratification	Biomarker	New use for CSL product or pipeline candidate	
Organization (start typing to search) *	Organization (if not listed)	Cardiovascular and Metabolic	Hematology	Immunology	Respiratory	Nephrology and Transplant	Vaccines Ce	ene Therapy	Other			
€ Email *	Confirm Email *	Plasma Protein Research	Oral Delivery						Project Description (max. 300 words) * Tips for a competitive abstract: - Abstracts should focus on a defined project (as opposed to a general overview of the applicant's research interests) - If you have multiple projects of potential interest, please submit multiple abstracts - Where possibilize/spiciatio, please describe the treate, disease area, high level mechanism of action (1-2 sentences), and key supporting data (1-2 sentences)			
Address		Secondary Focus Area (Optional, do no select if not applicable)							- An example abstract is as follows: "We have discovered a novel target expressed on X cells. We have generated data in X assay(s) and/or X model(s). We have shown the mechanism of action is modated via X pathway(s), Inhibition of this target could be used to treat X indication(s). This novel strategy could address an important unmet need for patients and be superior to			
City	Zip/Postcode	Cardiovascular and Metabolic	Hematology	Immunology	Respiratory	Nephrology and Transplant	A CONTRACTOR OF THE CONTRACTOR	ene Therapy	standard of care and other therapeutics in development for reasons X, X and X.* - Please do not include any confidential information			
Country or Territory *	Geographical region *								I have read the privacy policy and agree with it. Read more * I hereby confirm that my submission does not contain any confidential information. *			
Are you an existing collaborator, or have you previously collaborated with CSL (including CSL Behring, CSL Seqirus or CSL Vifor)?		Plasma Protein Research	Oral Delivery						I'm not a robot			
○ Yes ○ No		Disease Areas/Indicat	tions *						Privacy - Tarms	BACK SUBMIT		

Submission T&Cs on last slide Driven by Our Promise

What makes for a competitive proposal?

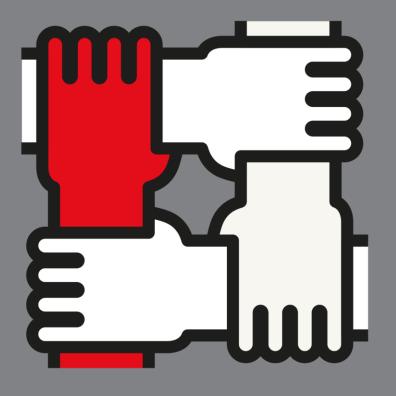
- ✓ Aligned with our focus areas and modalities (slides 28-35)
- ✓ Project is clearly defined (as opposed to a general overview of the applicant's research interests)
- ✓ Focused on a novel target or therapeutic candidate
- ✓ Clear differentiation of approach from competitors and current standard of care
- ✓ Research team has capacity and expertise to complete the bulk of the experimental work over the course of the program (with CSL guidance and support)
- ✓ If third party IP is required, ensure your research organization has secured all necessary rights to grant CSL an exclusive option to negotiate an exclusive, worldwide licence

Examples of activities funded in previous RAI partnerships

- ✓ Human target validation and translational studies using patient samples
- ✓ Mechanism of action studies for therapeutic candidates
- ✓ Benchmarking to provide proof-of-concept for the differentiation of novel therapeutics to standard-of- care or competing therapeutics in development
- ✓ Target validation using genetic knock-out/knock-in or tool compounds in preclinical disease models
- ✓ Characterization of therapeutic candidates (e.g. affinity, potency, selectivity, and developability)

CSL

Benefits of collaborating with CSL



Benefits of CSL's Research Acceleration Initiative



Collaborate
with one of the world's
leading biotech
companies



Publish with CSL 200+ publications with our collaborators since 2020



Funding of up to \$400,000 USD over 2 years



Access expertise
CSL scientific champion
assigned to provide you
with industry guidance



Recognition

Awardees may use title

"CSL Research Acceleration
Initiative Fellow"



Accelerate
the translation of
your research into
new therapies

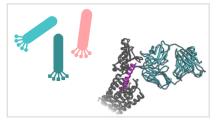


Access global capabilities in R&D, clinical, intellectual property, manufacturing and commercial



Demonstrate impact of your research to funding bodies via industry collaboration

Capabilities from Discovery to Patients



Antibody Discovery and Protein Engineering



In vitro pharmacology



Animal Models of Disease



Toxicology & Product Development



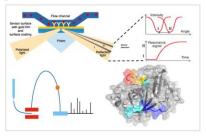
Patients

R&D CAPABILITIES

CLINICAL CAPABILITIES



Protein production and purification



Analytical Biochemistry



Translational Medicine & Data Science



Phase I-III/Launch Manufacturing

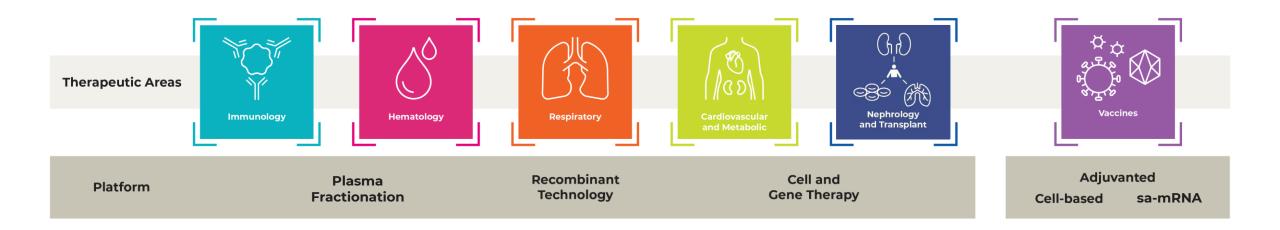
CSL

CSL's core Therapeutic Areas

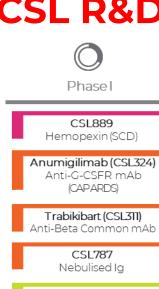


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CSL's Core Therapeutic Areas & Platforms



CSL R&D Portfolio – FY24



Anti-Beta Common mAb

C1 Inhibitor (AIS)

CSL040 Complement R1 Inhibitor

> SQ009 Cell-based Influenza (H2N3) Vaccine

SO012 sa-mRNA Influenza (H5N1) Vaccine

ARCT2138 sa-mRNA Quadrivalent Influenza Vaccine

LASN01 Anti-IL-11R mAb (IPF,TED)



Phase II

Anumigilimab (CSL324) Anti-G-CSFR mAb(HS)

VAMIFEPORT Ferroportin inhibitor (SCD)

CSL301 α2AP mAb (PE)

> Garadacimab Anti-FXIIa mAb (ILD/IPF)

Clazakizumab Anti-IL-6 mAb (ESKD)

Mavrilimumab Anti-GM-CSFR mAb (GCA, COVID)

Eblasakimab (CSL334) Anti-IL-13R mAb(AD)

Phase III

HIZENTRA® (DM)

KCENTRA® 4F-PCC (Trauma)

> CSL112 apoA-I(AMI)

SNF472 Calcification inhibitor (CUA-ESKD)

FILSPARI™(Sparsentan) Dual ET_A & AT₁ antagonist (FSGS)

Clazakizumab Anti-IL-6 mAb(ca-AbMR)

CSL964 Alpha 1Antitrypsin (Treatment of a GvHD)

> CSL964 Alpha 1 Antitrypsin (Prevention of aGvHD)

SQ036 (aQIVc) Adjuvanted Cell-based Ouadrivalent Influenza Vaccine

Registration/Post-Registration

HAEGARDA® (HAE)

HIZENTRA® (SCIa) 20% Liquid

PRIVIGEN® (IVIg) 10% Liquid

Garadacimab Anti-FXIIa mAb (HAE)

> AFSTYLA® rFVIII (Haem A)

IDELVION® rFIX-FP (Haem B)

HEMGENIX® (Haem B)

ZEMAIRA®/RESPREEZA® Alpha 1 Antitrypsin

■FILSPARI™(Sparsentan) Dual ET_A & AT₁ antagonist (IaAN)

■■KORSUVA®/KAPRUVIA® KOR agonist (CKD-aP)

RAYALDEE® Oral ext. release calcifediol (SHPT)

TAVNEOS® Oral C5a receptor inhibitor (AAV)

VFI PHORO® Sucroferric oxyhydroxide (Serum P control in CKD)

VELTASSA® Oral potassium binder (HK)

AUDENZ™

Adjuvanted Cell-based Monovalent Influenza A (H5N1) Vaccine

FOCLIVIA®/AFLUNOV® Adjuvanted Egg-based Influenza A (H5N1) Vaccine

FLUAD® Trivalent Adjuvanted Egg-based

Influenza Vaccine

FLUAD® Quadrivalent Adjuvanted Egg-based Influenza Vaccine

FLUCELVAX® **Ouadrivalent** Cell-based Influenza Vaccine

ARCTI54

sa-mRNA Vaccine (COVID)

INJFCTAFFR® (Ferric carboxymaltose) (HF-ID)

Immunology

Vaccines

Haematology

Respiratory

Cardiovascular & Metabolic

Nephrology & Transplant

Product and pipeline highlights



Privigen® (10% intravenous Ig) Primary immunodeficiencies (PID), Secondary Immune Deficiency (SID)*, Chronic inflammatory demyelinating polyneuropathy (CIDP)

Hizentra® (20% subcutaneous Ig) PID, CIDP, SID* Dermatomyositis (DM). Ph III

Haegarda® (C1 Esterase Inhibitor) Hereditary angioedema

Garadacimab (Anti-FXIIa mAb) Hereditary angioedema, Ph III

CSL324 (Anti-G-CSFR mAb) Hidradenitis suppurativa (HS), Ph I



Idelvion® (Recombinant FIX-FP) Hemophilia B

Hemgenix® (AAV FIX gene therapy) Hemophilia B

Afstyla® (Recombinant FVIII) Hemophilia A

Kcentra® (Prothrombin complex concentrate) Urgent warfarin reversal

Vamifeport (Oral ferroportin inhibitor) Sickle cell disease, PhIIa

CSL889 (Hemopexin) Sickle cell disease, Ph I

CSL888 (Haptoglobin) Sub-arachnoid hemorrhage, preclinical development



ZEMAIRA®/RESPREEZA® (Alpha 1 Antitrypsin)

Garadacimab (Anti-FXIIa mAb) Idiopathic Pulmonary Fibrosis, Ph IIa

CSL311 (Anti-β-common mAb) Airways inflammation, Ph I

CSL787 (Nebulised Ig) Respiratory infections, Ph I



CSL112 (ApoA-1) Acute coronary syndrome. Ph III

CSL300 (Anti-IL-6 mAb) End stage kidney disease Ph IIb



FLUAD Ouadrivalent Adjuvanted Influenza Vaccine

FLUCELVAX Ouadrivalent Cell-based Influenza Vaccine

Adjuvanted Cell Culture Influenza Vaccine (aQIVc), Ph II

sa-mRNA Influenza Vaccine, PC

ARCT-154 COVID-19 Vaccine, Ph III



CSL964 (Alpha 1 Antitrypsin) Graft versus host disease, Ph

Clazakizumab (Anti-IL-6 mAb) Antibody mediated rejection, Ph III

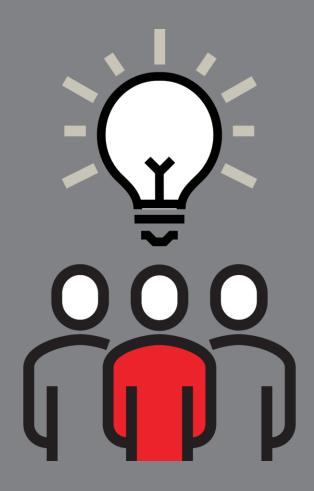
CSL040 (Novel Complement Inhibitor), Ph I



Areas of interest for collaboration

Areas <u>not</u> of interest

- Oncology (including hematological malignancies)
- Medical devices or diagnostics
- Small molecule approaches







Novel targets or best-in-class biologic therapeutics addressing:

- . B cell and plasma cell depletion or inhibition
- 2. T cell modulation, immune checkpoint agonism or co-stimulatory antagonism, Regulatory T cell stimulation or Tolerance
- 3. Modulation of cytokines, chemokines and immune-super family members (e.g., TNF, IL-1, other), particularly approaches enabling multi-pathway inhibition
- 4. Depletion/modulation of innate immune effector cells

Autoimmune diseases:

Inflammatory Idiopathic Myopathies including Dermatomyositis, Primary Sjögren's Syndrome, Small Fiber Neuropathy, ANCA-Associated Vasculitis and Autoimmune Hepatitis

Not of interest:

Target discovery campaigns or platforms, intracellular targets, complement inhibition





Acute hemorrhage control and hemorrhagic stroke

- Novel biologic therapies to treat and prevent acute hemorrhage (e.g. intracerebral hemorrhage (ICH), reversal of anticoagulation/anti-platelet associated bleeding)
- 2. Novel biologic targets and therapies for the treatment of secondary brain injury in subarachnoid hemorrhage and ICH
- 3. Omics approaches for patient stratification and drug discovery

Acute thrombotic conditions (macro- and micro-circulation)

- Novel biologic therapies for targeted fibrinolysis/thrombolysis in acute thrombosis (ischemic stroke, pulmonary embolism)
- 2. Novel biologic therapies to treat and prevent microvascular thrombosis and endotheliopathies (e.g. TMAs, APS and DIC).

Benign hematology adjacencies*

- 1. Novel biologic therapies for the treatment of anemias
- 2. Novel biologic therapies to treat bone marrow disorders





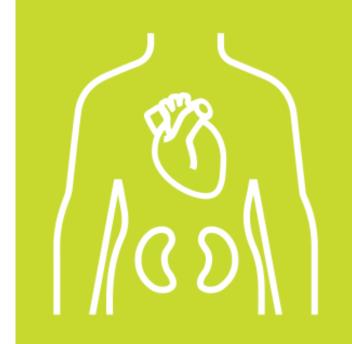
Idiopathic pulmonary fibrosis (IPF), pulmonary sarcoidosis and progressive pulmonary fibrosis (PPF)

- Novel biologic therapies or target proposals derived from translational or biobank cohorts
- 2. Therapies targeted at reversing remodelling of fibrotic lung tissue
- 3. Multiomics-based approaches to target discovery

Community acquired pneumonia (CAP)-associated complications

(Acute Respiratory Distress Syndrome (ARDS), Sepsis, Acute kidney injury)

- Novel biologic therapies or target proposals derived from translational or biobank cohorts
- 2. In Silico approaches for patient stratification to delineate CAP patients at risk for ARDS/Sepsis/AKI



Cardiovascular and Metabolic



Atherosclerotic plaque stabilization in high-risk patient groups

Novel targets or biologic therapies to prevent atherosclerotic plaque rupture/erosion and Major Adverse Cardiovascular Events (MACE)

Rare lipid disorders

Novel targets or biologic therapies (including gene therapies) for rare /severe lipid disorders e.g. homozygous familial hypercholesterolemia, hypertriglyceridemia

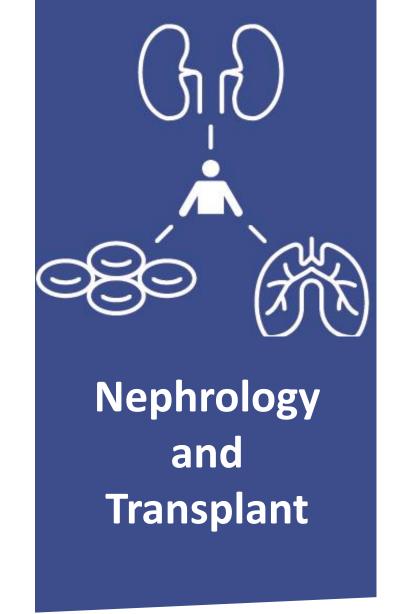
Myocarditis

Novel targets or biologic therapies for immune checkpoint inhibitor myocarditis

Biomarker approaches for patient stratification

Inflammatory cardiomyopathies

Novel targets or biologic therapies for inflammatory cardiomyopathies Biomarker approaches for patient stratification





Acute and chronic solid organ transplant rejection (kidney/lung) therapies

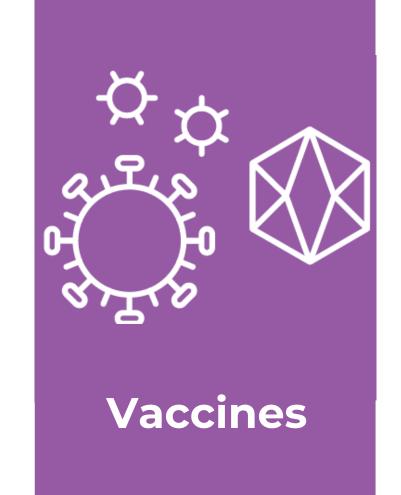
Novel biologic therapies or targets to prevent or treat acute and chronic solid organ transplant rejection of the kidney and lung

Chronic graft versus host disease (GvHD)

Novel biologic therapies for the treatment and prevention of chronic GvHD

Tolerance for organ transplant rejection

Novel biologic therapies for the induction of tolerance to prevent or treat organ transplant rejection





Respiratory vaccines

- New antigenic targets (epitopes or combinations)
- 2. Methods (e.g. Al/machine learning) to predict respiratory viral evolution/pathogenicity to inform vaccine development

New vaccine targets

Development of novel targets/approaches for any disease

RNA delivery and therapeutics

- RNA delivery, enhanced stability, route of administration and/or expression strategies
- 2. mRNA-encoded protein therapies encompassing cellular targeting technologies

Immune mechanisms

Understanding innate and adaptive responses to vaccines



Cell & Gene Therapy



Gene editing / genomics

- 1. Improve insertional editing efficiencies in vivo
- Genetic elements enhancing regulation of cells of the immune system (e.g. promoters and enhancers)

In vivo Delivery

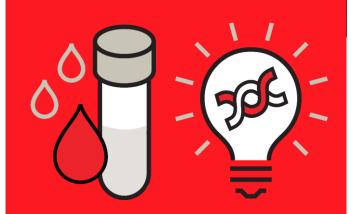
- 1. Delivering nucleic acid templates for insertional gene editing
- 2. Targeting moiety for HSCs

GT safety

Technologies that minimize SAEs from insertional gene editing

Areas not of interest

- Oncology (including hematological malignancies)
- Ex vivo cell therapy



Plasma Protein Research



Novel plasma therapeutic candidates

- 1. Seeking plasma candidates aligned with CSL's therapeutic areas
- 2. CSL can provide native human plasma proteins (≥ µg/L plasma concentration) for preclinical proof-of-concept studies

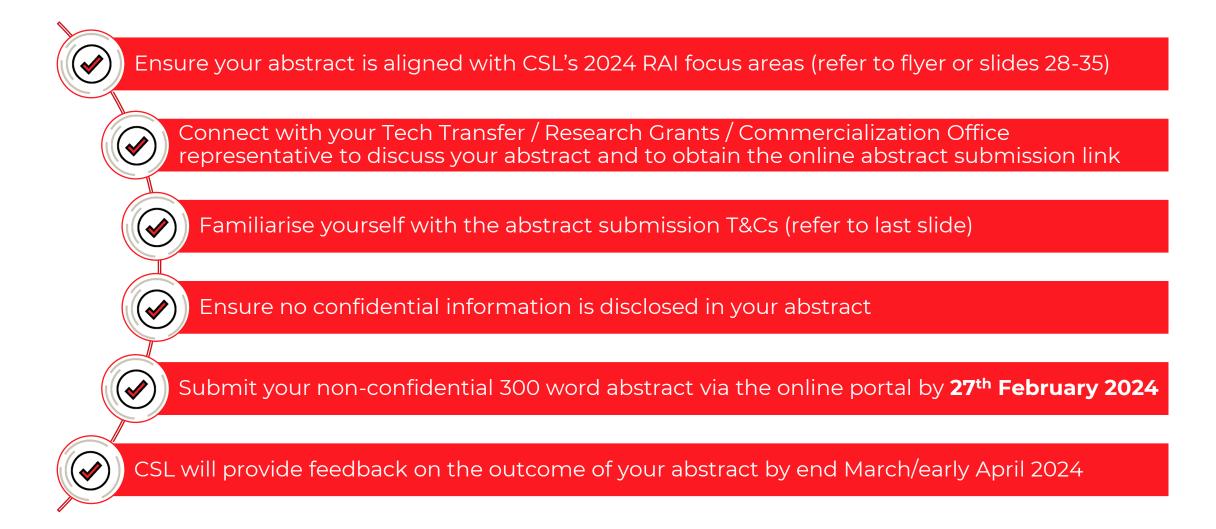
Novel association of plasma protein function with disease

- 1. Based on healthy and patient clinical data sets aligned with CSL's therapeutic areas, or
- 2. Access to patient data sets with corresponding clinical data to enable association studies to be performed

Novel methods for plasma protein purification

Protein purification systems capable of targeted purification from plasma with high purity at research scale (methods translatable to manufacturing scale will be prioritized)

Checklist for 2024 Research Acceleration Initiative



SEVEN NEW CSL RESEARCH



Dr Laurent Martinez

Institute of Cardiovascular and Metabolic Diseases (I2MC), IHU HealthAge, INSERM / University of Toulouse, France

Prof. Delphine Borgel

INSERM - APHP - Université Paris SACLAY, France

Prof. Denis Vivien

INSERM / Caen Normandie University Hospital, France

Research Director Benoit Salomon

INSERM / University of Toulouse, France

Assoc Prof. Tan Meng How

Nanyang Technological University, Singapore

Prof. Elisa Laurenti

University of Cambridge, United Kingdom

Prof. Leon Schulte

Philipps-Universität Marburg, Germany







CSL

Questions



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THANK YOU

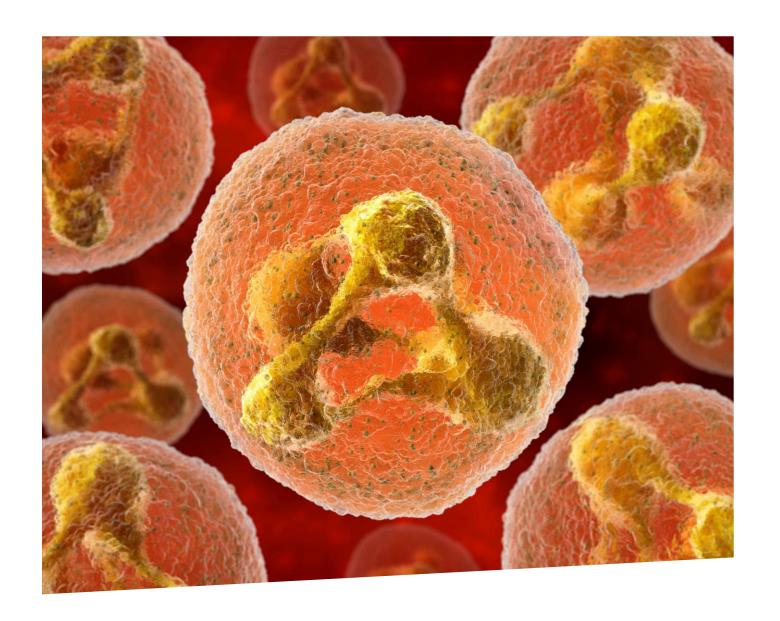
Dr Marthe D'Ombrain Executive Director & Head, Global Research Innovation

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RAI@csl.com.au



Terms and Conditions for Research Acceleration Initiative Portal ("RAI Portal")

- 1. This RAI Portal is an online portal operated by CSL Innovation Pty Ltd ("CSL") for the purpose of allowing individuals to submit scientific proposals for consideration by CSL for its Research Acceleration Initiative program. By using this website and the RAI Portal, and by providing your submission and personal information to CSL, you are agreeing to abide by these terms and conditions.
- 2. You acknowledge and agree that CSL has no obligations of confidentiality or non-use in relation to the submission provided. You warrant that your submission does not contain confidential information of any kind. Further, you acknowledge that notwithstanding the existence of any confidentiality agreements previously entered into between you and CSL, the terms of such agreements will not apply with respect to any information submitted by you through the RAI Portal.
- 3. You further represent and warrant that:
- a. you have the right and authorisation (including where relevant after consultation with all relevant commercialisation or technology transfer offices) to submit an application to the RAI Portal and to accept the terms and conditions set out herein;
- b. you are an employee or are otherwise affiliated with a registered organisation authorised by CSL to submit an application to the RAI Portal; and
- c. to the best of your knowledge and without making any further enquiries, the information provided in your submission (and CSL's use of that information in connection with the Research Acceleration Initiative program) shall not infringe on the intellectual property rights of any third party, including your current or former employer, university, public research institute or other registered organisation.
- 4. CSL may disclose personal information collected in connection with your use of this website or the RAI Portal to your employer, university, public research institute or other registered organisation (if applicable) as at the time your application was submitted, solely for the purpose of reviewing and determining your application. CSL will ensure that any personal information collected, used or disclosed in connection with your use of this website or the RAI Portal is handled in accordance with all relevant privacy legislation and with CSL's privacy policy, a copy of which is available at https://www.csl.com/privacy-policy.
- 5. CSL is under no obligation to respond to any individual application submitted to the RAI Portal, and may in its sole discretion choose not to progress an application further for any reason without any further communication with you.