

Real Life™

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Grants Working Group Recommendations CLIN

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CIRM
CALIFORNIA'S STEM CELL AGENCY

OUR MISSION

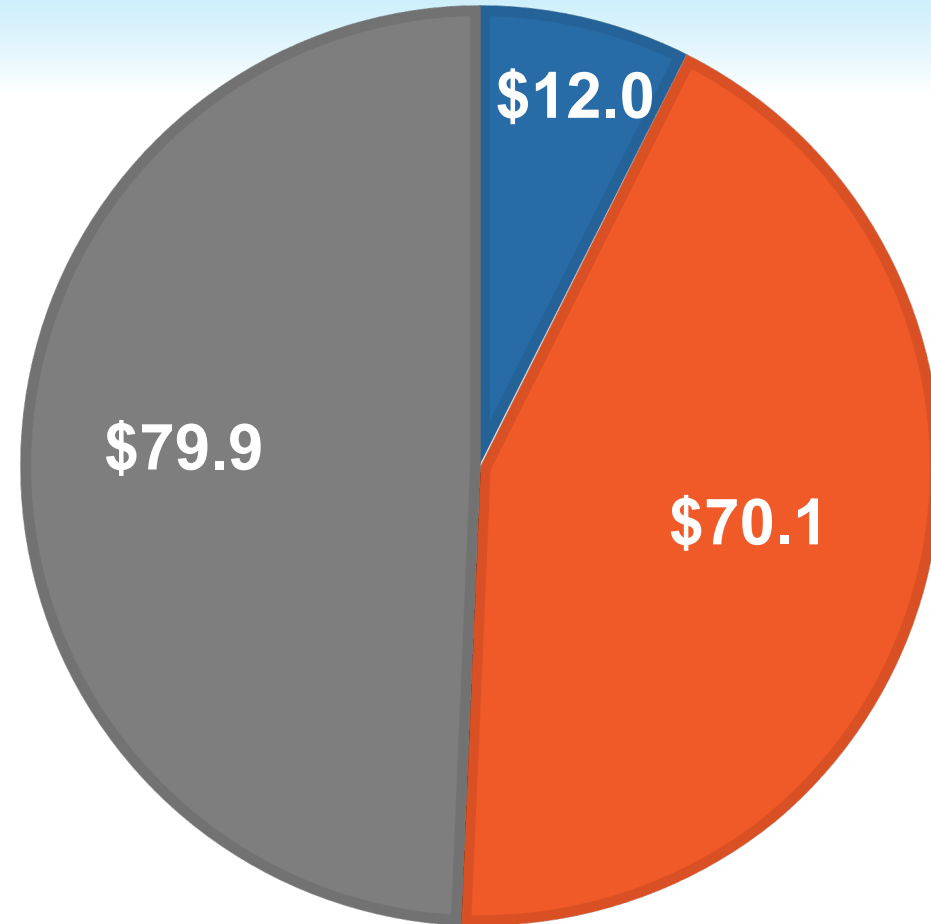
Accelerating world class science
to deliver transformative
regenerative medicine treatments
in an equitable manner to a
diverse California and world



Annual Allocation: \$162 million

- Amount Requested Today
- Approved Awards
- Unused Balance

Amounts are shown in millions



- **Score of “1”**

Exceptional merit and warrants funding.

May have minor recommendations and adjustments that do not require further review by the GWG

- **Score of “2”**

Needs improvement and does not warrant funding at this time but could be resubmitted to address areas for improvement.

GWG should provide recommendations that are achievable (i.e., “fixable changes”) or request clarification/information on key concerns.

- **Score of “3”**

*Sufficiently flawed that it does not warrant funding and the same project should not be resubmitted **for at least 6 months.***

Applications are scored by all scientific members of the GWG with no conflict.

1. Does the project hold the necessary significance and potential for impact? (i.e., what value does it offer; is it worth doing?)
2. Is the rationale sound? (i.e., does it make sense?)
3. Is the project well planned and designed?
4. Is the project feasible? (i.e., can they do it?)
5. Does the project address the needs of underserved communities?

Scientific GWG
Member



Scientific evaluation (disease area expert,
regulatory, CMC, product development)
Provides scientific score on all applications

Patient Advocate
or Nurse GWG
Member



DEI evaluation, patient perspective on significance
and potential impact, oversight on process
Provides DEI score on all applications
Provides a suggested scientific score

Scientific
Specialist
(non-voting)



Scientific evaluation (specialized expertise as
needed)
Provides initial but not final scientific score

Title	Phase 1, open label, dose escalation study of oncolytic virus (OV)-loaded cytokine induced killer (CIK) cells in patients with advanced solid tumors
Therapy	Cytokine-induced killer cells with oncolytic virus that target cancer cells
Indication	Advanced refractory solid tumors
Goal	Completion of phase 1 clinical trial to assess safety and tolerability
Funds Requested	\$7,999,689 (co-funding: \$0)

Maximum funds allowable for this category: \$8,000,000

Clinical Background: Refractory solid tumors such as colorectal, ovarian, breast, and osteosarcoma that fail to respond to standard treatments represent a significant unmet medical need.

Value Proposition of Proposed Therapy: The standard of care varies by tumor type but may involve chemotherapy, radiation, resection, and/or available drugs. If successful, the proposed therapy would provide a safe and effective therapeutic option for patients with solid tumors where approaches such as CAR-T have been less successful.

Why a stem cell or gene therapy project: Hematopoietic progenitor cells are used to manufacture the therapy.

Application/ Award	Project Stage	Project End Date	Indication	Candidate	Mechanism of Action
TRAN	Pre-IND	Jul 2022	Ovarian cancer	Neural stem cells loaded with oncolytic virus	Neural stem cells target solid tumor cells to deliver oncolytic virus.

Applicant has not previously received a CIRM award.

GWG Recommendation: Exceptional merit and warrants funding

Scientific Score	GWG Votes
1	14
2	1
3	0

DEI Score: 8 (scale 1-10)

CIRM Team Recommendation: Fund (concur with GWG recommendation)

CIRM Award Amount: \$ 7,999,689*

*Final award shall not exceed this amount and may be reduced contingent on CIRM's final assessment of allowable costs and activities.

Title	Hematopoietic Stem Cell Gene Therapy for X-linked Chronic Granulomatous Disease (XCGD)
Therapy	Autologous gene-corrected hematopoietic stem cells
Indication	X-linked Chronic Granulomatous Disease (XCGD)
Goal	Completion of IND-enabling studies and IND submission
Funds Requested	\$3,999,959 (co-funding: \$0)

Maximum funds allowable for this category: \$4,000,000

Clinical Background: X-linked Chronic Granulomatous Disease (X-CGD) is a rare immune disorder that prevents white blood cells from killing foreign invaders. This results in severe, recurrent infections that can impact quality and length of a patient's life. X-CGD is usually diagnosed before age 5, but without treatment, children die before age 10.

Value Proposition of Proposed Therapy: The current standard of care involves ongoing antibacterial and antifungal prophylaxis and allogeneic hematopoietic stem cell transplant. If successful, the therapy offers patients the potential for immune restoration and cure.

Why a stem cell or gene therapy project: The therapeutic candidate is composed of gene-modified hematopoietic stem cells.

Application/ Award	Project Stage	Project End Date	Indication	Candidate	Mechanism of Action
CLIN2	Phase 1/2 clinical trial	Dec 2022	XCGD	Autologous gene-corrected CD34+ cells	Lentiviral vector correction of gene defect in patient CD34+ cells.

Applicant has not previously received a CIRM award.

GWG Recommendation: Exceptional merit and warrants funding

Scientific Score	GWG Votes
1	8
2	7
3	0

DEI Score: 5.5 (scale 1-10)

CIRM Team Recommendation: Fund (concur with GWG recommendation)

CIRM Award Amount: \$ 3,999,959*

*Final award shall not exceed this amount and may be reduced contingent on CIRM's final assessment of allowable costs and activities.