

Funding Area Preferences

Rosa Canet-Avilés, Ph.D.
CSO, CIRM
ICOC Meeting
January 29, 2026



Presentation Overview

1 Context & Objectives

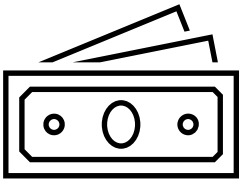
2 Guiding Principles

3 PDEV & CLIN2

4 DISC4

5 Closing

Proposition 14 & Where We Are Now



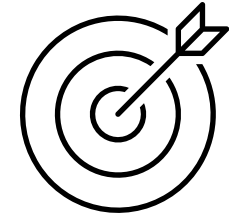
Finite Funds

- Finite remaining runway
- CIRM cannot fund everything



Expanded Mandates

- CNS
- Access & affordability

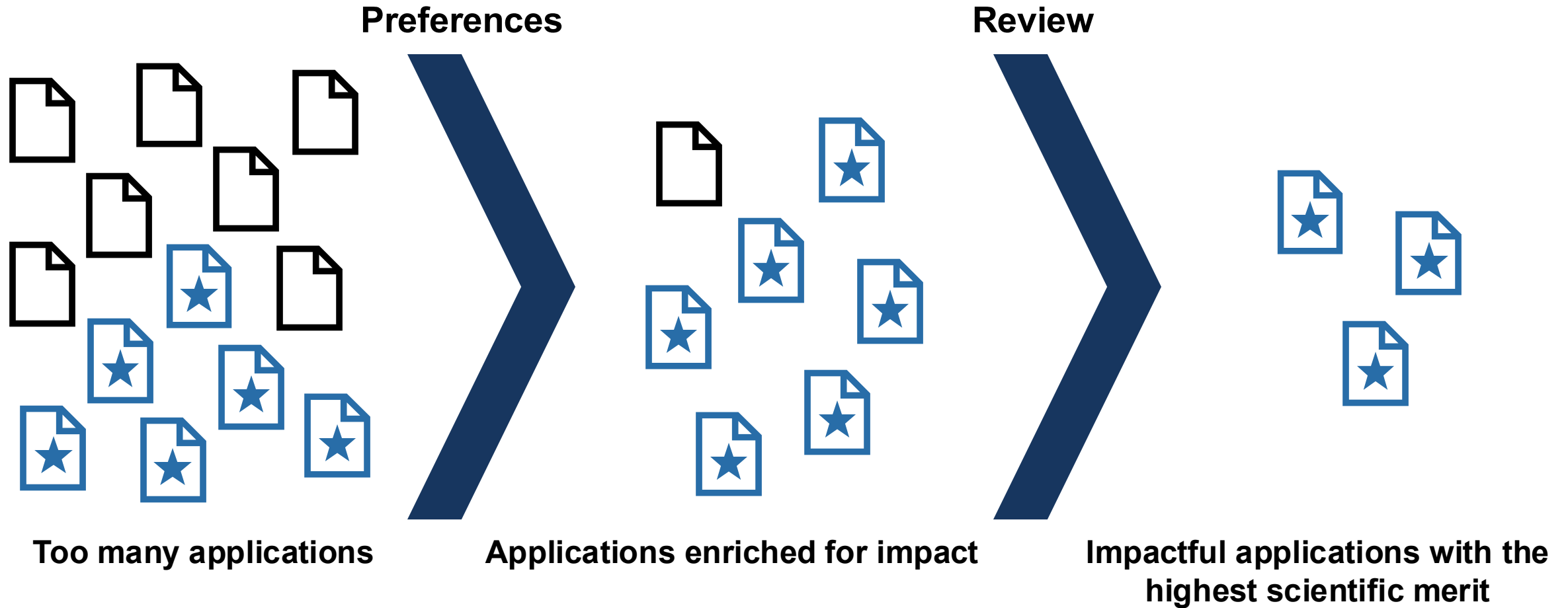


Goal

- Translate public investment into **therapies reaching patients** within CIRM's lifetime

Challenge: CIRM receives **more applications than it can fund**

Preferences Help Achieve Therapies in CIRM's Lifetime



What We Have Heard

- Anxiety that proposals were triaged without scientific review
- CIRM is not funding progression events
- Perception of modality bias
- Desire for clarity on qualification scoring, preferences, and intent
- Misunderstanding of what preferences do / don't do
- Need for clearer communication & broader reach

It is critical for CIRM to **clarify how actions taken** so far will lead to **therapies for patients** and achieving its mission

Today's Objectives

- Review **rationale for preferences**
- **Share learnings** from first funding cycles
- **Identify portfolio analyses needed** to refine preferences

Presentation Overview

1 Context & Objectives

2 Guiding Principles

3 PDEV & CLIN2

4 DISC4

5 Closing

Guiding Principles for Preference Setting

Guiding Principles

1. Offer potential for transformative clinical impact
2. Address bottlenecks to access, affordability, & translational feasibility
3. Fill critical funding gaps & advance CIRM's statutory mandates
4. Can realistically achieve key regulatory and development path within CIRM's finite runway
5. Address diseases affecting Californians
6. Diversify CIRM's active award portfolio

PDEV Pre-Submission Rubric

Preference	Points
At least one of the following: <ul style="list-style-type: none">• PSC-derived therapies• In vivo gene therapies• Diseases of the CNS	3
• Non-viral nucleic acid delivery	1
• Pre-IND or INTERACT meeting conducted	1
• Progression from DISC2 or TRAN1	1
• Targeting disease area under-represented in CIRM active awards portfolio	1
• Novelty of therapeutic approach compared to CIRM active awards portfolio	0-2

CLIN2 Qualification Rubric

Preference	Points
• PSC-derived therapies	1
• Diseases of the CNS	1
• In vivo genetic therapy	1
• Non-viral genetic therapy	1
• Accelerated regulatory designation (RMAT, Breakthrough, Fast Track)	1
• Progression from earlier stage CIRM award	1
• California Organization	1
• Pivotal trial	2

Presentation Overview

1 Context & Objectives

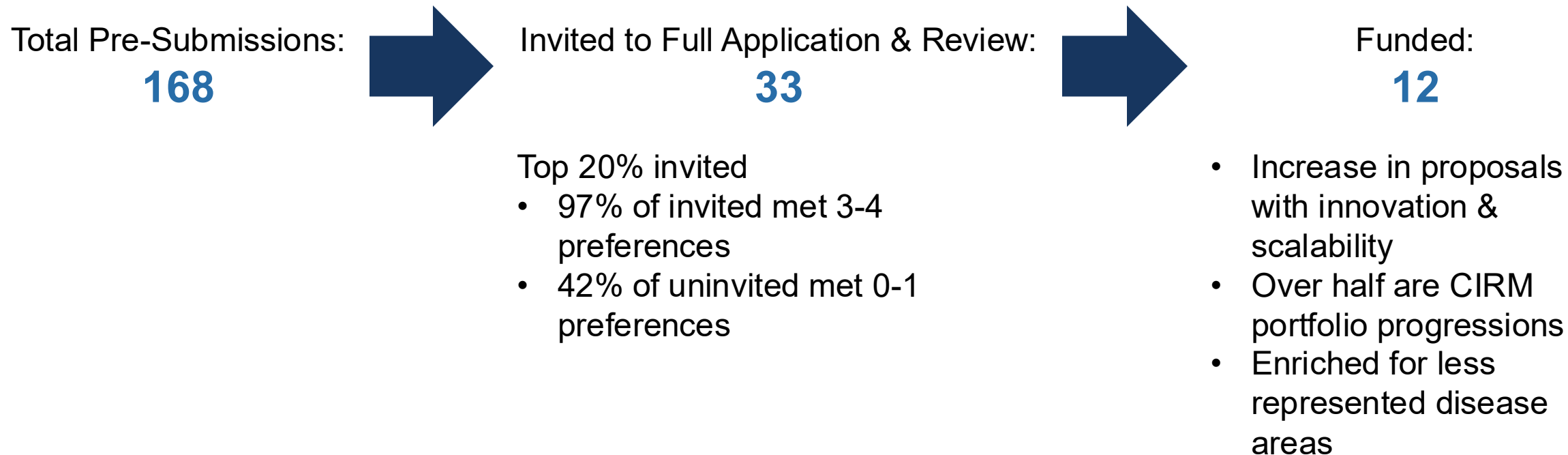
2 Guiding Principles

3 PDEV & CLIN2

4 DISC4

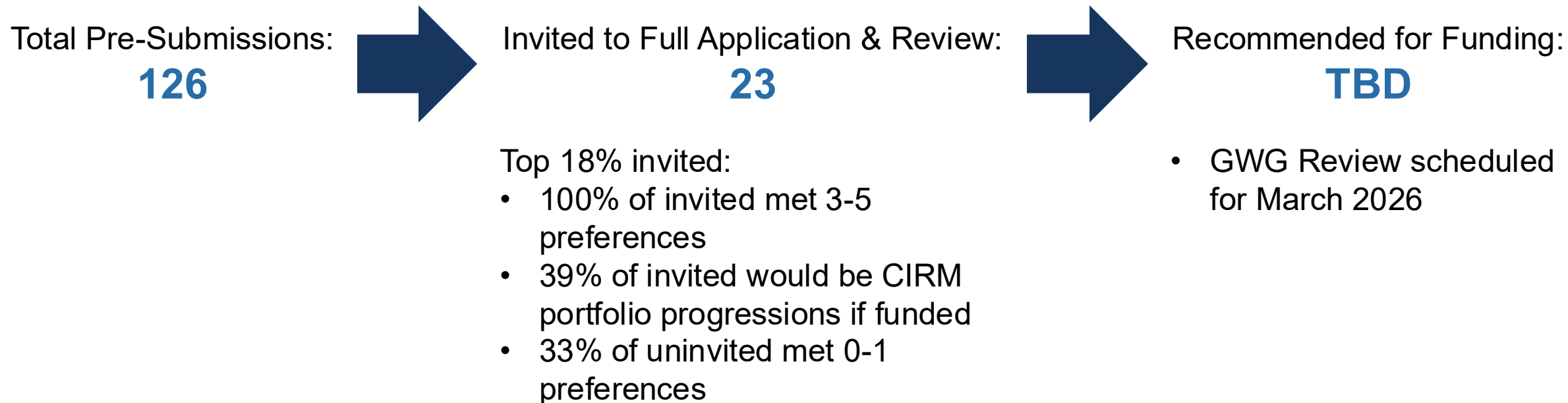
5 Closing

PDEV Cycle 1 Results



Awarded programs **reflect preferences** and CIRM's preclinical **portfolio is now better balanced** across modality, disease, & Prop 14 CNS priorities

PDEV Cycle 2 Results



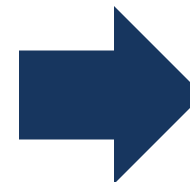
23 invited applications represent broader disease areas and include more cancer & immunology

CLIN2 Cycle 1 Results

Total Applications:
23



Advanced to Full Review:
7



Funded:
4

- Ranked by preference points

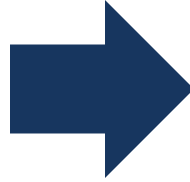
- All 7 had 3-4 preference points
 - All targeted CNS
 - 6 were pluripotent stem cell or in vivo genetic therapy
 - 4 had advanced designations

- All use next-generation or platform modalities
- All have feasible delivery paths consistent with A&A strategy
- 3 are CIRM portfolio progressions

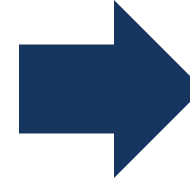
The system worked directionally as intended, but **scoring weights need refinement**

CLIN2 Cycle 2 Results

Total Applications:
21



Advanced to Full Review:
7



Funded:
TBD

- Ranked by preference points
- All 7 advancing had 3-4 preference points
 - 4 target CNS
 - 5 are CIRM portfolio progressions
 - 5 are in vivo genetic therapies or PSC-derived therapies
- GWG Review scheduled for January 27, 2026

Cycle 2 results are pending GWG Review & ICOC/ARS approval

PDEV & CLIN2 Cycle 1 | Summary

- Awards in both PDEV and CLIN2 reflect the innovation, readiness, Access & Affordability, and CNS criteria in the Program Announcements
- Applications advancing were largely CIRM progressions that also met multiple preferences
- The resulting portfolio now includes more candidates with feasible, scalable modalities that strengthen Access & Affordability potential

Caveats

- These are preliminary signals
- Portfolio-level impact cannot be assessed yet

Presentation Overview

1 Context & Objectives

2 Guiding Principles

3 PDEV & CLIN2

4 DISC4

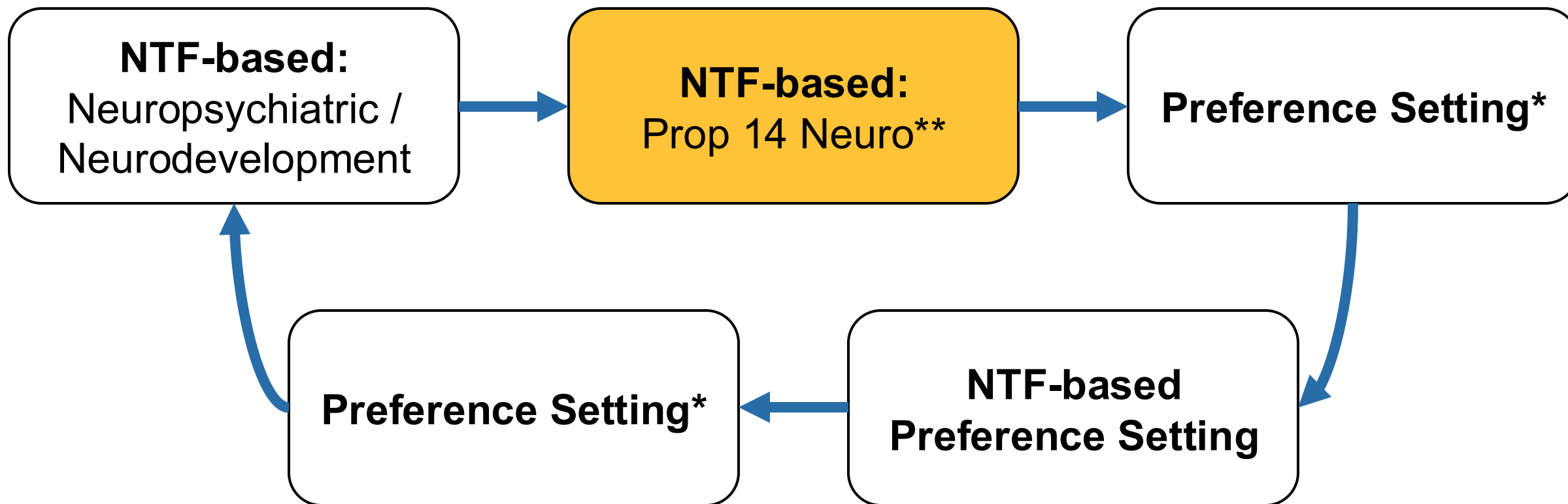
5 Closing

What We Set Last Year

- CIRM staff proposed alternating preference cycles for DISC4, with Neuro every other year
- The Board approved Neurological Disease as the preference for the FY25/26 DISC4 cycle; the program was open to all comers

DISC4 Preferences | Alternating Neuro Cycles

Enabling NTF Prioritization while allowing other diseases to use this structure



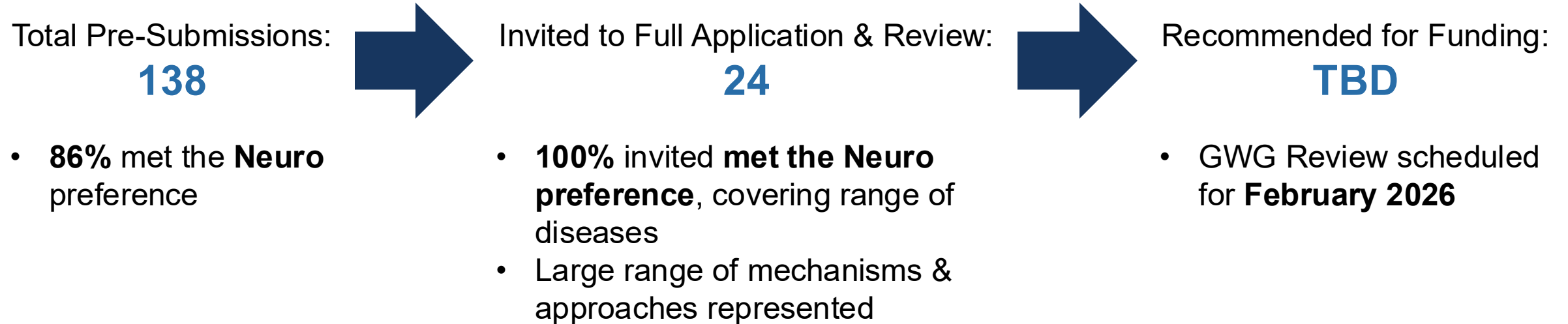
*All cycles will be open to all-comers with alternating preferences determined by either NTF or by the Board each year based on portfolio analyses

** FY25/26, ICOC selected broad Neuro preference– In first round, Neuro preference had a weight of 36% overall

DISC4 FY25/26 Preference Scoring

Criteria	Description (listed in program announcement)	Weight (%)
Preference Topic: Neuro	Does the project align with the cycle-specific preference topic?	36%
Relevance to human disease biology	Does the project hold strong relevance for understanding or addressing human diseases?	24%
Cross disciplinary and systems biology	Does the project integrate cross-disciplinary approaches?	24%
Stem cell or genetic research innovations	Does the project incorporate innovative approaches particularly in stem-cell or genetic research?	16%

DISC4 Cycle 1 | Pre-submission Results



Preferences enriched the pool: all invited applications aligned with the Neuro preference while maintaining scientific diversity

Presentation Overview

1 Context & Objectives

2 Guiding Principles

3 PDEV & CLIN2

4 DISC4

5 Closing

What We Learned

- **Initial signals from preference setting**

- Applications that advanced to GWG review met multiple preferences
- High percentage of CNS projects funded for CLIN2
- Wide range of disease areas and modalities funded for PDEV
- High percentages of funded projects are CIRM progressions (>50% PDEV, 75% CLIN2)

January presentation **tees up the questions** for **Board**; **March** brings the **portfolio** analysis and any potential preference refinements based on Board guidance

Planned Analyses for March ICOC

- Definition & examples of what we categorize as innovation, progressions, and disease areas
- Active portfolio & application cycles broken down by disease area, modality, progression status, and how CIRM is driving innovation

Does the ICOC request **any other portfolio analyses** to **inform preference setting**?