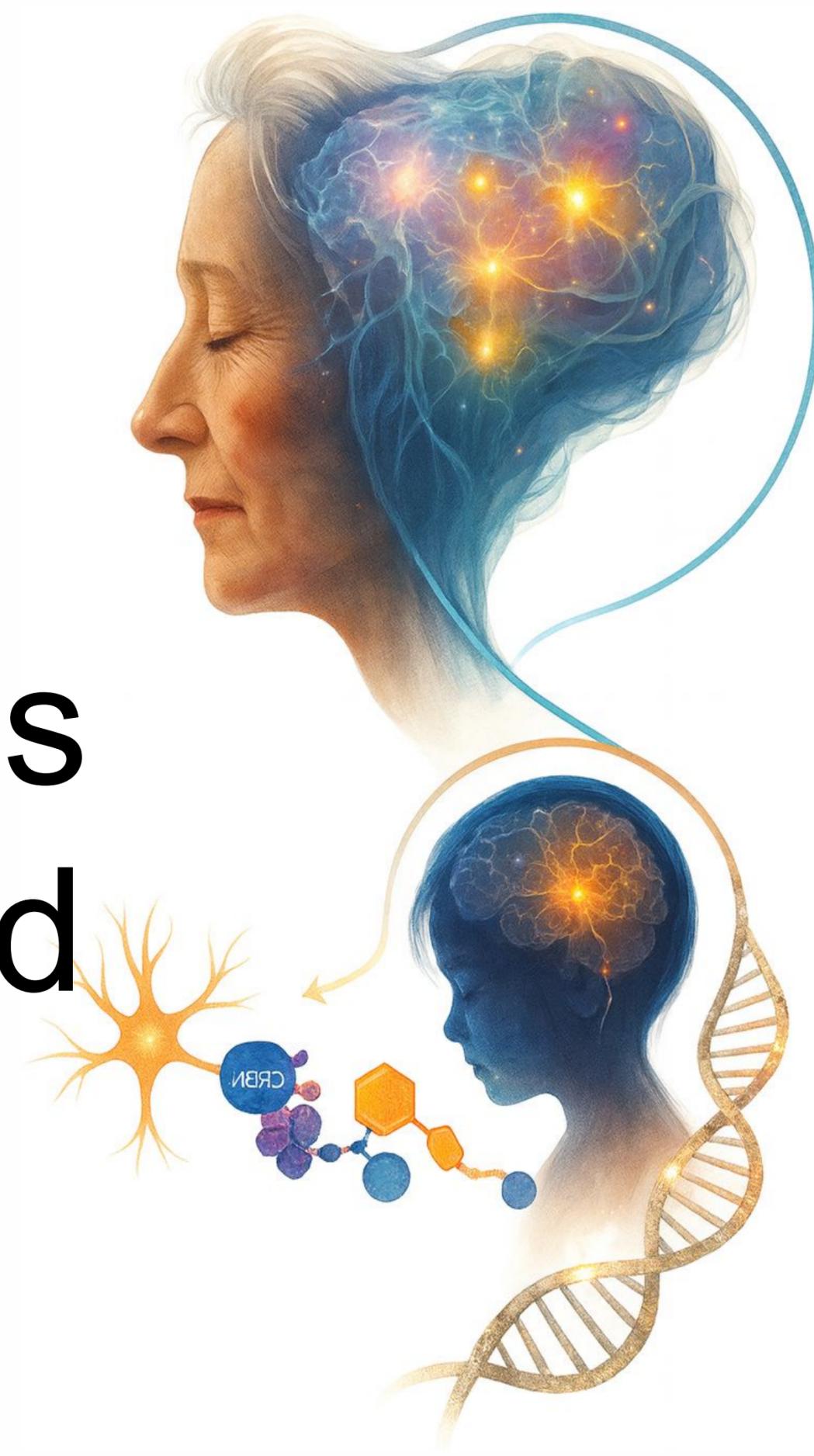




LigronBio

⌘ BREAKTHROUGH
H RESEARCH

Reversing Alzheimer's via Gene Therapy and Molecular Glues



Founder &

Nataraj

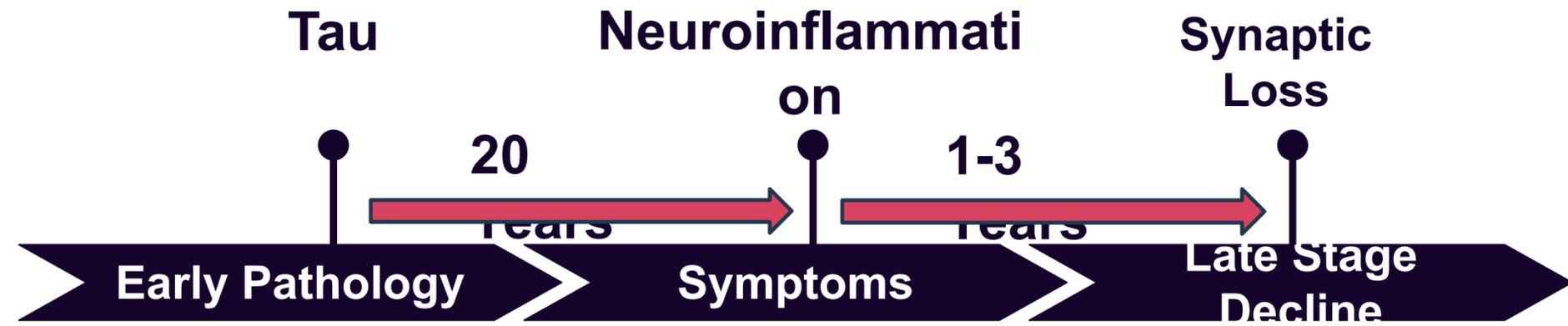


Alzheimer's Therapy : 9.2T / 2050

The Challenge

- Key Pathways (Amyloid Beta, Tau & Synaptic Loss) hard to Drug
- BBB Limits Drug Delivery
- Single Target Drugs Fail to Stop Progression

Why Current Therapies Fall Short



50M+

Current Cases

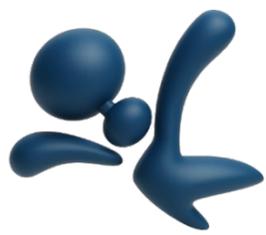
150M+

2050

50%

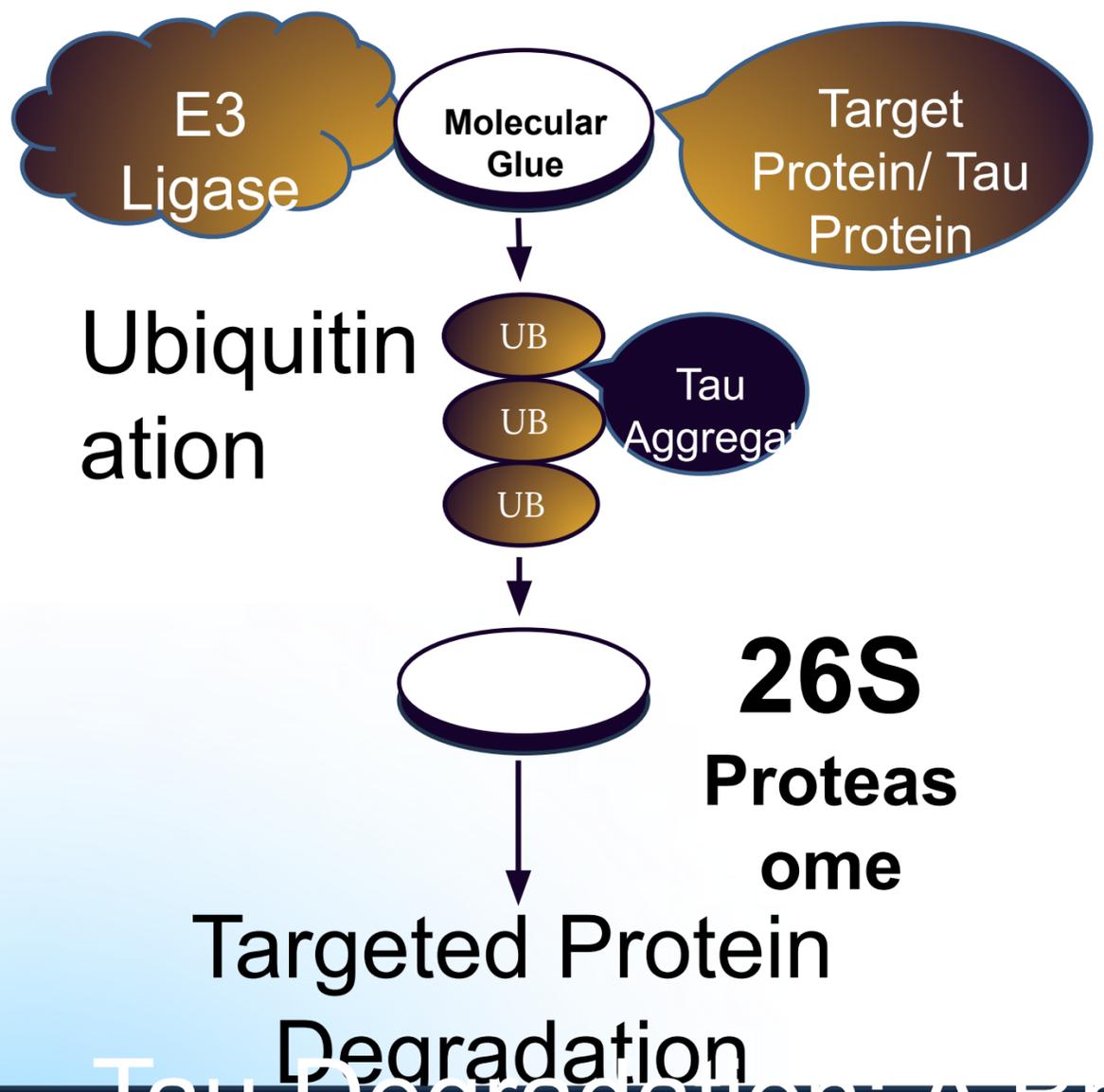
Genetic Inheritance / 2050

Too little = No Drug efficacy / Too much = Toxicity



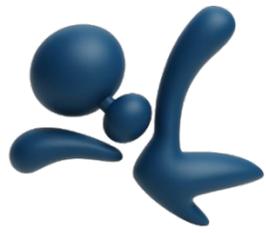
Solution : Glu-NcAA TPD via Molecular Glues with Gene Therapy

How Molecular Glues Works: They Create A Bridge Between Disease-causing Proteins And The Cell's Natural Disposal Machinery (E3 Ligase), Triggering Protein Ubiquitination And Degradation Through The 26S Proteasome.



-  Unlock Treatment for Undruggable Proteins
-  Clearance Of Toxic Proteins
-  Prevent Tau Protein Phosphorylation
-  Prevent Neuroinflammation
-  Restore Memory Function

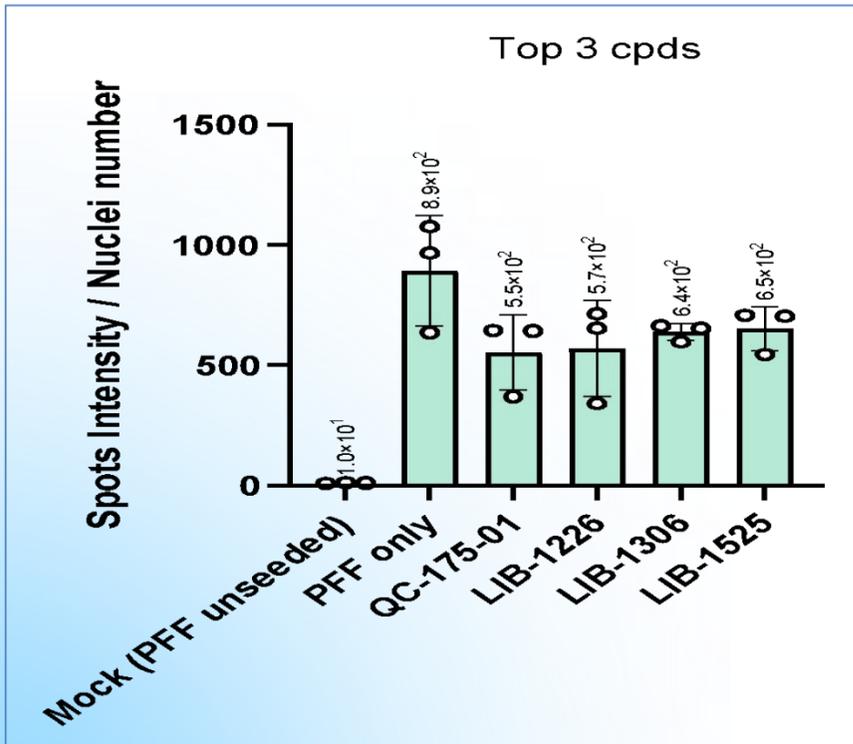
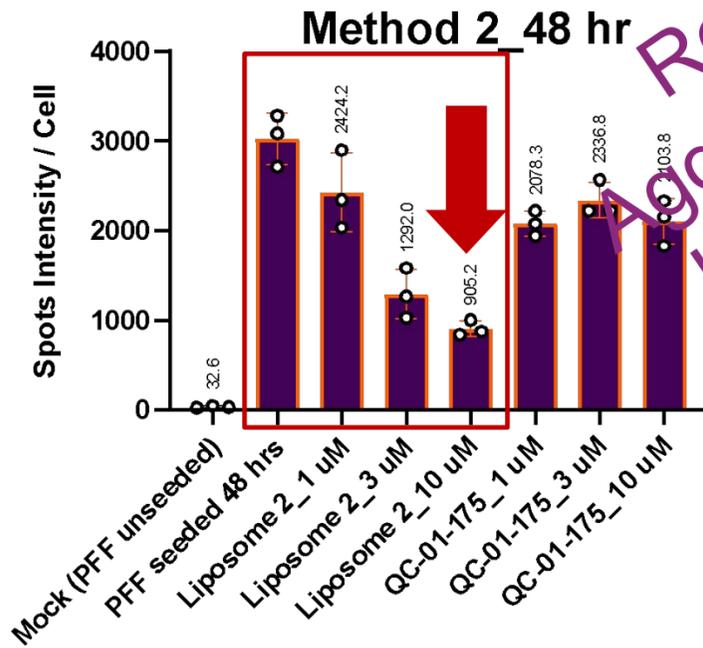
Tau Degradation: A Breakthrough Pathway to Reverse Cognitive Decline



Reverses Neurodegeneration

DC₅₀: ≈ 57%

Reduced Aggregated Tau Intensity/Cell



- ❑ Novel Tau-Targeted Therapeutic Platform
 - Selectively Degrades Pathological Tau Protein Aggregates
- ❑ Scalable Platform
 - Small-molecule, Manufacturable, and Cost-efficient
- ❑ De-risked Path
 - Clinically and Regulatorily Validated Modality
- ❑ Protected IP
 - Strong Patent Coverage Extending Through 2026

Degrading Toxic Aggregates, Restoring Neurons

Human Tau-K18-GFP

HEK293T



Discovery Pipeline

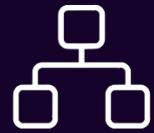
Precision-Engineered Protein Degradation Targeting Multiple Neurodegenerative Pathways With Enhanced Selectivity Profiles



Glu-NcA

Aβ Peptide-
Tau Aggregate Degradation for AD
Therapy

• Targeting 2028 For IND Filing



LIB-1226

c-RAF Protein
Degradation For



AD

LIB-067

TDP-43 Protein Degradation
for Early-Onset of AD & ALS



Discover

Additional targeted protein degradation
candidate in FTD



HumanAI™ & L-Tag Assay : Making

the 'Undruggable' Druggable

(Provisional Patent)

AI-Trimatrix™ Analyzer™

(Patent)

Diagnostic Kit

(Patent Pending)

Therapeutics

Data

Integration

Proteome Mapping

Validation in Live cells

Human clinical Trials

1

Input

2

3

Output

Molecular Glue discovery

4

5

Invitro & In Vivo Studies

6

Disease Therapy

Amino acid/Peptide

Generati

Molecular Glue Optimization

AI-driven discovery

Rapidly identifies and optimizes molecular glues without requiring 3D protein structures



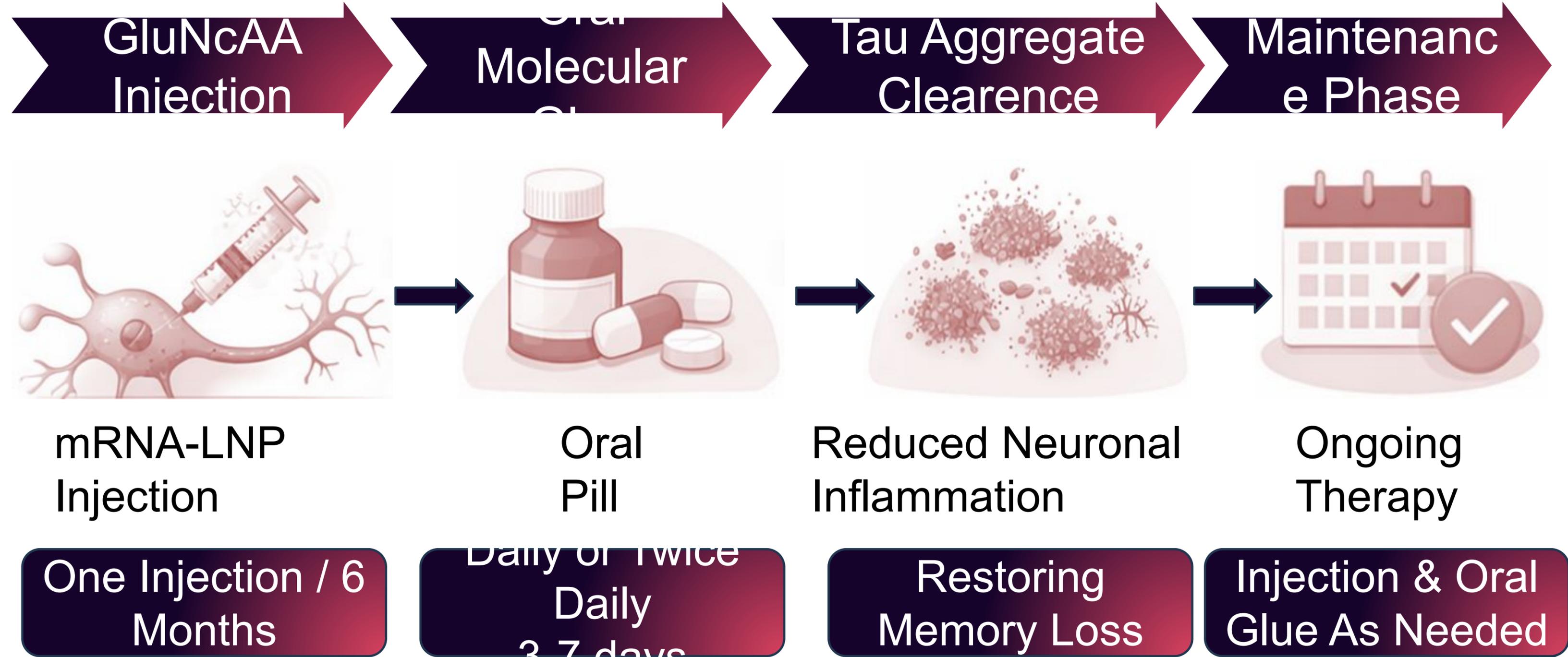
Membrane Penetration

Our compounds demonstrate excellent Membrane penetration and metabolic stability





GluNcAA Alzheimer's Treatment





Glues

One Injection every 4-6 weeks Drives Sustained Protein Clearance, Slowing Cognitive Decline And Restoring Memory

Once Or Twice Daily For 3-7 Days Following The Glu-NcAA Injection



**Safe
ty**

□ Precision Targeting

- Right Protein, Right Time, Right Brain Exposure
- Eliminating Infusion Burden and Immune Side Effects



**Efficac
y**

□ Therapeutic Index

- >100× Vs. Traditional Inhibitors
- Prolonged Action : 24 - 48+ Hours Post-dose



**Conveni
ence**

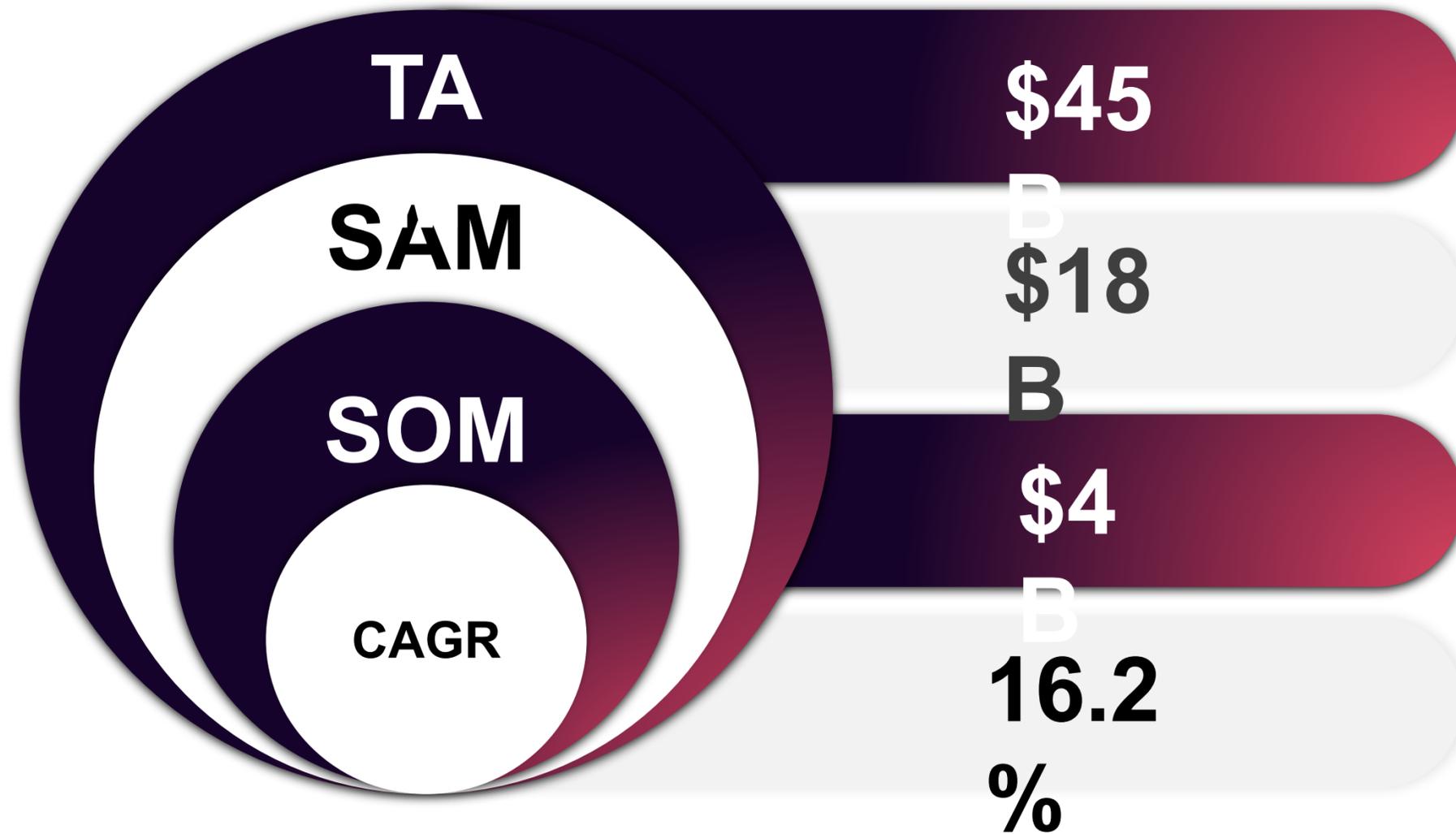
□ Precision Therapy

- One Pill. Right Dose. Right Target. Right Time.
- New Standard of Care

Safety + Superior Efficacy +



Market Share by 2035



64M /
2030
Revenue
Projection

Revlimid Molecular Glue set as a



B2B

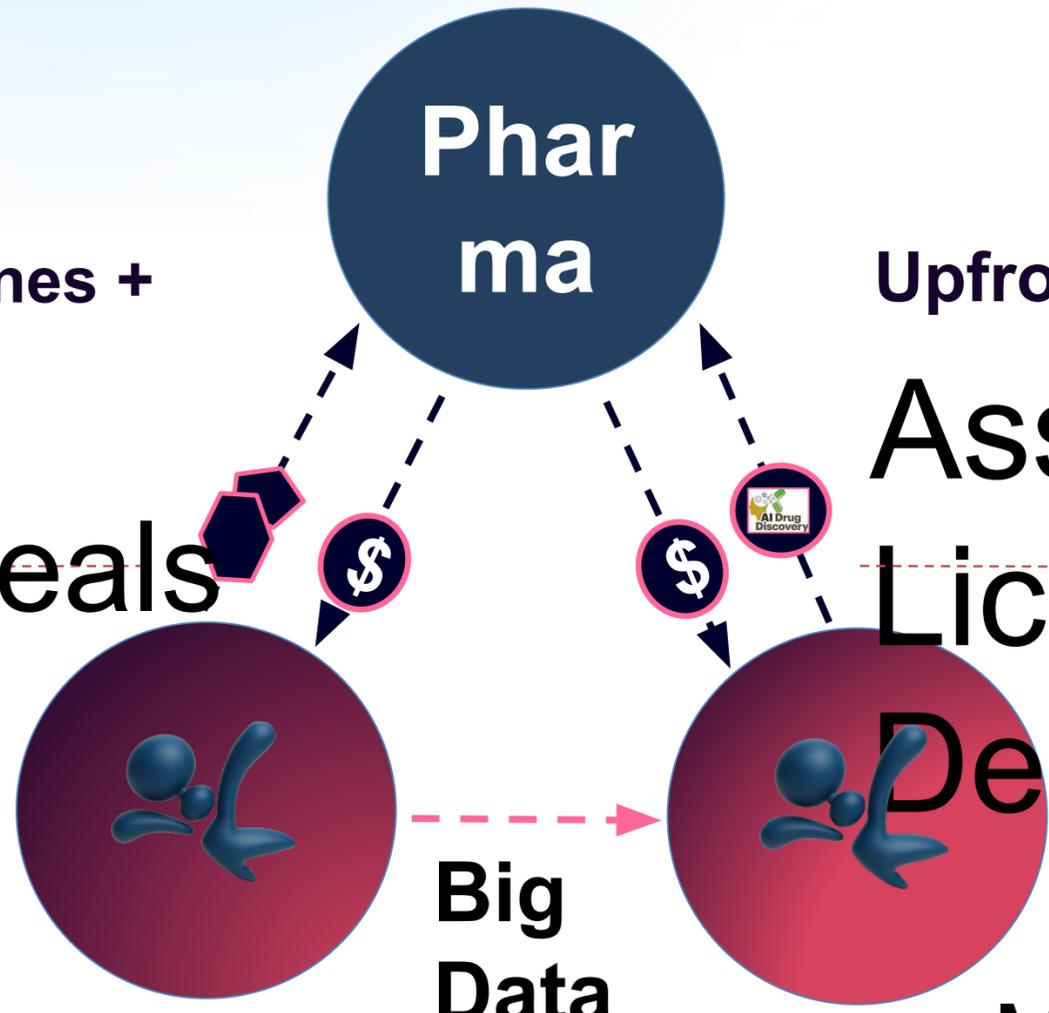
Business Model

Upfront fees + Milestones + Royalties

Discovery Partnership Deals

Upfront fees + Milestones + Royalties

Asset Licensing Deals



- Human AI - TriMatrix™ Analyzer™ Platform +
- Diagnostic Kit (L-Tag Assay)
 - Biomarker discovery and drug efficacy prediction
 - Therapeutic Candidate Optimization

Molecular Glue

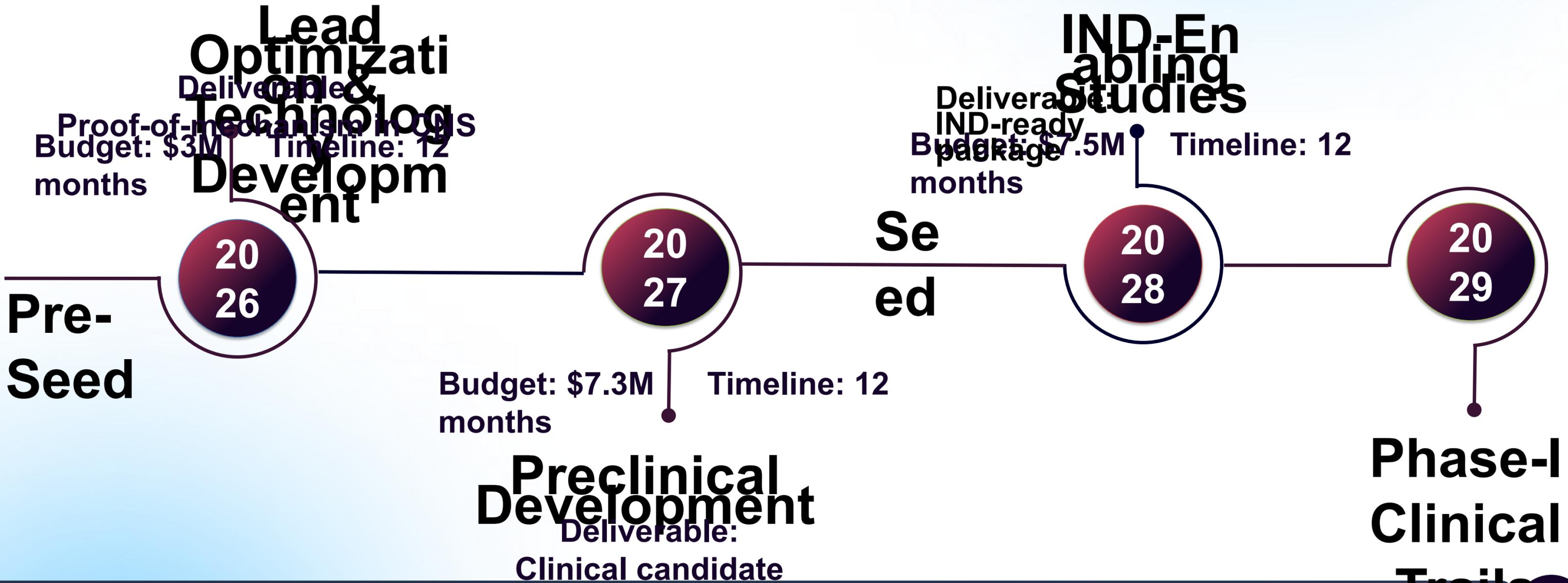
Alzheimer's, ALS, Parkinsons & Oncology

Therapeutic Candidates



Business & Clinical Development Strategy

Our strategic roadmap accelerates novel therapeutics from discovery through clinical trials, leveraging proprietary platforms to address critical unmet needs in neurodegenerative disease



Total Investment: \$17.8M Over 36 Months Positions Lead Asset For Phase 1 Clinical Trials, With Potential For Accelerated Regulatory Pathways In Orphan Indications



We are Raising \$3M Pre Seed

(18 Months) on a SAFE

Key Milestones Advancing Our Glu-NCAA Technology Through Proof-of-concept Validation

Q1

Glu-NCAA Tech & MO Development
35% Budget
\$1,05M

\$1,05M

Q3

Preclinical Validation Phase
23% Budget
\$400,000

\$400,000

Q2

Medicinal Chemistry Optimization
16.7% Budget
\$500,000

\$500,000

Q4

InVivo Target Assessment & POC
9% Budget
\$550,000

\$550,000

Operational Costs Budget : \$300,000 (10%)

IP Protection &

R

20-50X

Safe : 20% Discount

Investor Returns

- Equity Appreciation
- Licensing Deals
- Acquisition & Partnerships

\$115K : Boot Strap
\$100K : Angel Investor



Next-Gen Therapeutics

Computational
Chemistry



**Nataraj Pagadala,
PhD**
Fellow, Carnegie Mellon University
President & CEO, UNIVERSITY OF ALBERTA

Learning &
Memory



Prof. Albert La
SUC Irvine Center for the
Neurobiology of Learning & Memory

Drug Metabolism &
Pharmacokinetics



**Prof. Oliver
Bouvier**
VANDERBILT
UNIVERSITY

Animal Models



**Christopher M. Butt,
Ph.D.**
VANDERBILT
UNIVERSITY

Clinician



**Dr. Venu
Prasad**

Immunology



**Amir Landi,
PhD**
Principal, Medical Discovery &
Translational Research
Leader
UNIVERSITY OF
ALBERTA

Clinical Trials



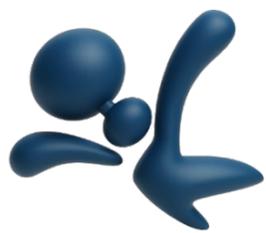
**Claudia Shojaei,
CRSM**
Principal, Clinical Trial
Operations

Fractional CFO



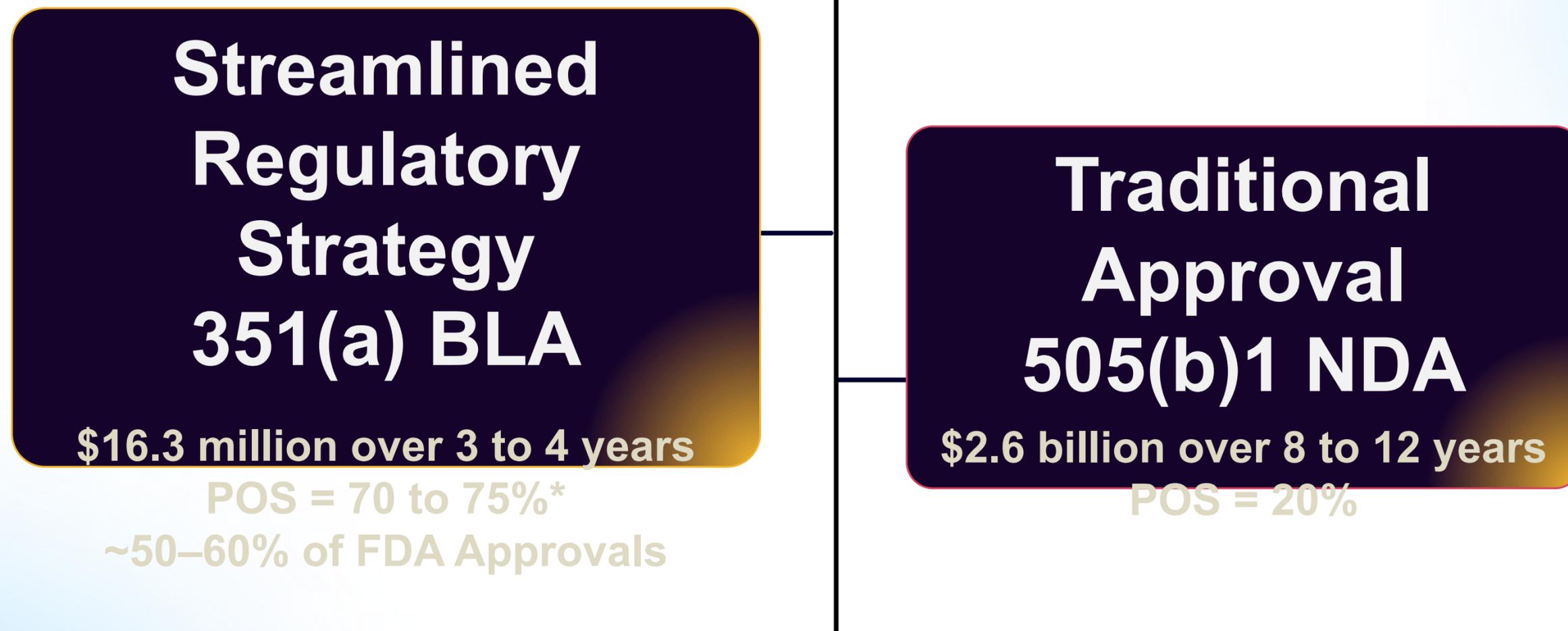
Ashish Sinha
CEO @
StartupFin.co

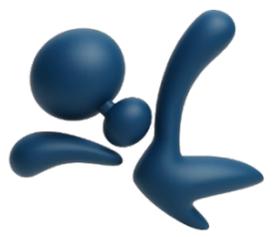




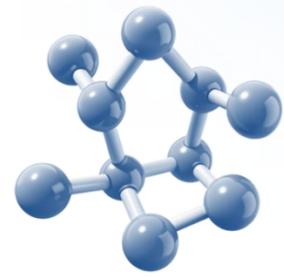
Regulatory

FDA-Regulated; Marketing Authorization
Required Prior To Licensing



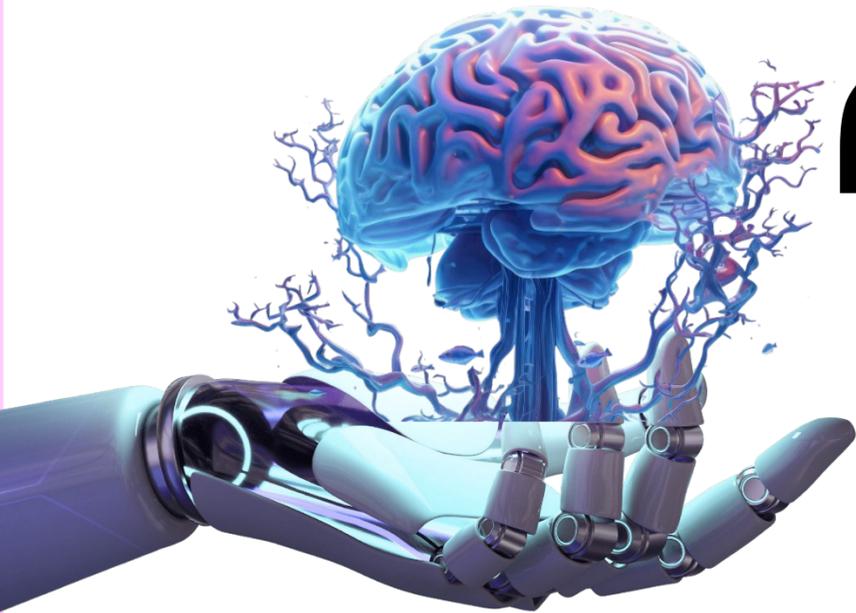


Health Crisis Than COVID-19



Molecular Glues

The New Revolution in
Alzheimer's Treatment



“We Are Fighting To
Protect Future
Generations From
Alzheimer's

Let's

LigronBio Inc.
Aquillius Innovation Hub,
10918 Technology Pl San Diego,
CA 92127
Nataraj
Pagadala,
PhD Founder &
CEO
npagadala@ligr
onbio.com
<https://www.ligronbio.com>