

Abeta active immunotherapy agreement with Takeda

NASDAQ: ACIU | Investor Presentation - May 2024



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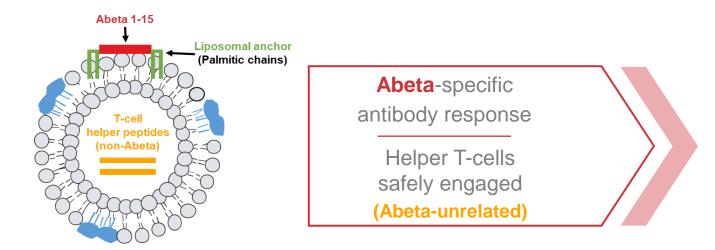
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# Unique Abeta targeted active immunotherapy – ACI-24.060

Best-in-class SupraAntigen®-based active immunotherapy in Phase 2

- Generates antibody responses against toxic forms of Abeta (oligomeric and pyroGlu-Abeta)
- FDA Fast Track supports disruptive potential in Alzheimer's disease treatment and prevention



#### **Unprecedented Performance**

Immunogenicity	<b>~</b>
Target & conformation specificity	<b>~</b>
Avidity increase over time	<b>~</b>
Sustainable response	<b>~</b>
Boostable response	~



Initial Phase 2 ABATE 6-month AD data on Abeta plaque reduction measured via PET imaging



Phase 2 ABATE 12-month AD data<sup>1</sup> on Abeta plaque reduction measured via PET imaging

(1) 12-month data will also include safety and immunogenicity results

### Landmark deal for ACI-24.060 in Alzheimer's disease

Supports promise of active immunization for neurodegenerative diseases



■ Today's deal with Takeda covers AC Immune's unique, class-leading Abeta targeted active immunotherapy ACI-24.060



- Deal terms:
  - \$100 million upfront payment received for exclusive option to license global rights
  - Option exercise fee in the low-to-mid nine-figure range linked to ABATE clinical data
  - Up to approximately \$2.1 billion in potential payments including option exercise fee and development, commercial and sales-based milestones
  - Royalties in the mid-to-high teens on global sales



■ Combines AC Immune's leadership in developing products for NDDs¹ with Takeda's clinical development expertise and history of driving neuroscience innovation

(1) Neurodegenerative diseases

# ABATE: Biomarker-based Phase 1b/2 study of ACI-24.060 in AD¹ and DS²

### Placebo-controlled Phase 1b/2 Study Overview

#### **Trial Schematic**

### **Adaptive Study Design**

Both

- Interim analyses of safety/tolerability & immunogenicity
- Biomarker analyses including Abeta PET<sup>3</sup> and others

AD

- Up to 4 different doses and/or dose regimens
- Expansion of one cohort to assess effect on Abeta PET

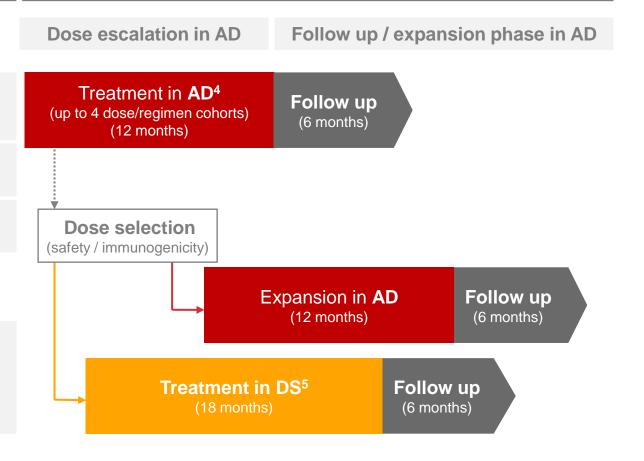
DS

 Initiation using selected dose identified in AD (based on safety/tolerability and immunogenicity)

#### **Outcome measures**

Soth

- Safety/tolerability
- Pharmacodynamics: Serum anti-Abeta antibody titers
- Abeta-PET imaging
- Exploratory biomarkers and clinical endpoints



(1) Alzheimer's disease; (2) Down syndrome-related AD; (3) Positron emission tomography; (4) AD participants must between 50 – 85 years of age and have prodromal AD with Clinical Dementia Rating Global Score of 0.5 and Abeta pathology confirmed by PET scan; (5) Cohort comprised of non-demented people living with DS (age 35 – 50 years) and Abeta pathology confirmed by PET scan



## AC Immune technology platforms driving validating pharma deals

Strategy: optimize value to risk ratio and retain significant upside

**SupraAntigen® Morphomer**<sup>®</sup> **Platform** ACI-24.060 ACI-35.030 Mor Tau Tau PET **Program Partner** Janssen Takeda Molecular Imaging

■ An integrated approach to Central Nervous System (CNS)-specific therapies



# AC Immune strong track record in deals<sup>1</sup> with leading pharma co.s

Strategy: optimize value to risk ratio and retain significant upside

Program	Phase	Total value <sup>2</sup>	Upfront <sup>2</sup>	Milestones received <sup>2</sup>	Royalties	Partner
ACI-24.060 (anti-Abeta active immunotherapy)	Phase 1b/2	>USD 2,100	USD 100		Mid-to-high teens	Takeda
ACI-35.030 (anti-pTau active immunotherapy)	Phase 2b	CHF 500	CHF 26	CHF 20	Low-double digits to mid-teens	Janssen PRABUSCHICA COUNTRIES OF SCHOOLS SPECIAL COUNTRIES
Tau Morphomer® drugs	Phase 1 <sup>6</sup>	CHF 1,860	CHF 80 +USD 50 <sup>7</sup>	CHF 40	Low-double digits to mid-teens	Lilly
PI-2620 (Tau PET <sup>4</sup> tracer)	Phase 3 <sup>5</sup>	EUR 160	EUR 0.5	EUR 7	Mid-single digits to low-teens	Life Molecular Imaging
Crenezumab (anti-Abeta antibody)	Phase 2	USD 65 <sup>3</sup>	USD 25	USD 40		*
Semorinemab (anti-Tau antibody)	Phase 2	CHF 59 <sup>3</sup>	CHF 17	CHF 42		*
Total (millions)8		CHF ~4,750	CHF 255.2 9	CHF 147.4		



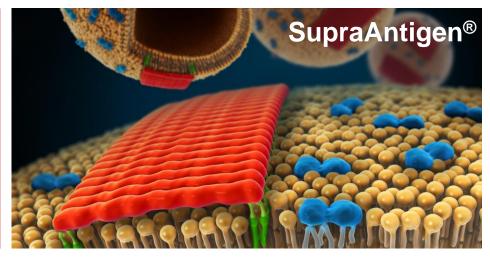
<sup>(1)</sup> Disclosure limited due to confidentiality agreements with collaboration partners; (2) In millions; (3) Total payments received from partner until termination of agreement; (4) Positron emission tomography; (5) In Alzheimer's disease; (6) Phase 1 completed; (7) Equity investment; (8) Converted to CHF on date of receipt; (9) Excludes convertible note agreement of USD 50 million; \* previously licensed to Genentech (a member of the Roche Group)

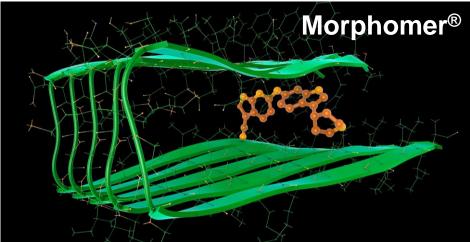


# AC Immune technology platforms driving validating pharma deals

Strategy: optimize value to risk ratio and retain significant upside

#### **Platform**





# Unpartnered Programs

- a-syn active immunotherapy
- Anti-TDP-43 mAb<sup>1</sup>
- Anti-NLRP3-ASC mAb

- Mor-a-syn
- Mor-TDP-43 PET
- Mor-NLRP3-ASC

■ Considerable additional potential value in our unpartnered clinical and preclinical programs

### Conclusions

AC Immune's leadership in precision medicine



Significant shareholder value is created through **class-leading active immunotherapies** in neurodegeneration



- Today's Takeda deal reinforces AC Immune's track record in deal execution
- >USD 4.5 billion in potential milestones from partnerships



- Clinically validated technology platforms continuously deliver high-valued assets
- Multiple unpartnered assets against key targets for development in-house and with potential partners



Cash reserves of CHF 104.8 million as of March 31, 2024, and upfront payment provides 3-year cash runway<sup>1</sup>

# AC Immune: Pioneering science and precision medicine

Shifting the treatment paradigm for neurodegenerative disease towards precision medicine and disease prevention

