

Alzinova

We will make it possible for Alzheimer's patients to live an independent and active life

Company presentation | December | 2023



Forward looking statements

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Imagine if there was a vaccine against Alzheimer's disease



Alzinova is developing the first specific vaccine against Alzheimer's disease



Swedish biopharma company | Based in Gothenburg Founded 2011



High unmet medical need for Alzheimer's disease

huge market opportunity



Innovative science & proprietary technology enables development of disease-modifying treatments for Alzheimer's



Pipeline - Vaccine ALZ-101 - phase 1b

- Antibody ALZ-201 - preclinical



Nasdaq First North listed as ALZ Shareholders ~4 500, Market cap ~150 MSEK



Alzheimer's disease

- Slowly progressing disease
- Caused by toxic formations in the brain called oligomers consists of Aβ-42 peptide
- Destroys brain structures



Need for targeted treatment that can neutralize the toxic oligomers – stopping the disease progression



Alzheimer's disease – huge unmet medical need

Dementia

>55 million people suffer from dementia

>150 million people in 2050

60-70% are Alzheimer's patients

Impacts quality of life

Patients, families, care givers

US \$1.3 trillion

(US \$2.8 trillion by 2030)

Annual cost

World Alzheimer's Report 2015, 2018, 2019, 2021 | Alzheimer's Disease Facts and Figures. Alzheimer's & Dementia 2018;14(3):367-429) | The Lancet Feb 2021



Unmet medical need



Cognitive and physical decline, leads to increased dependency on caregiving and later full-time care.



Caregiver

Unpaid family and friend caregivers can experience physical, financial & emotional strains.



Elderly care requires institutionalized housing, managing health issues, and more healthcare staff.



Aging populations and high incidence pose healthcare challenges, costing 1.3 billion each year.



Dementia affects 1 in 3 seniors, surpassing breast and prostate cancer deaths combined. Alzheimer's deaths doubled from 2000 to 2019.



>100 million people will suffer from Alzheimer's in 2050



Enormous market potential

Substantial market potential for Alzinova

Alzheimer's therapeutics market > US \$13 billion by 2028

2 out of 7 predicted blockbusters by 2026 are new anti-Alzheimer's drugs with US \$1.7 - 4.5 billion projected annual revenue

Market growth **driven** by introduction of novel therapeutics

Global Data 2020 I Drugs to watch report 2022



Potential breakthrough in Alzheimer's

High unmet medical & market potential for disease-modifying treatments of Alzheimer's disease



Short-term goal:

- ✓ First clinical trial targeting oligomers with a therapeutic vaccine Q3 2021
- ✓ Interim data safe and well tolerated, immune response Q2 2023
- ✓ Top-line data Positive results, met primary endpoint and immune response Q4 2023



Mid-term goal:

- > ALZ-101 Enter Phase 2 in 2024
- > ALZ-201 "Phase 1 ready" TBD



Long-term goal:

- Bring to market in partnership with Big Pharma
- Provide effective treatment, halting and preventing Alzheimer's



Alzinova's
business model –
develop, document,
prepare, partner &
bring to market



Develop breakthrough disease-modifying treatments based on proprietary AβCC technology



Document safety and tolerability in patients with Alzheimer's disease



Prepare (in parallel) for phase 2 to optimize the candidates value



Partner with big pharma on "phase 2-ready" candidates



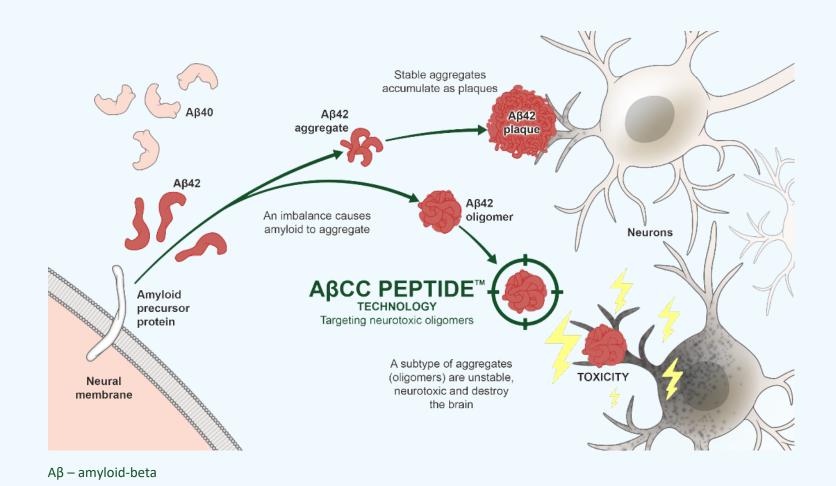
Income through upfront, milestone and royalty payments



Our candidates for treating Alzheimer's disease

Alzinova's **AβCC** peptide[™] technology enables high precision, with antibodies specifically targeting the toxic oligomers





Two differentiated, oligomerspecific products based on the same proprietary technology

- **Highly specific disease modifying treatments**, directed towards the toxic accumulations of amyloid beta in the brain
- Opportunity to become effective and well tolerated treatments to halt and prevent Alzheimer's
- "Best in Class" potential



ALZ-101 represent a potential breakthrough in treating Alzheimer's Disease



- The body generates specific antibodies against toxic oligomers
- High patient compliance (few booster doses)
- Long-acting
- Preventive treatment possible
- Large potential patient population



ALZ-201 a complement to the vaccine ALZ-101 in treating Alzheimer's Disease



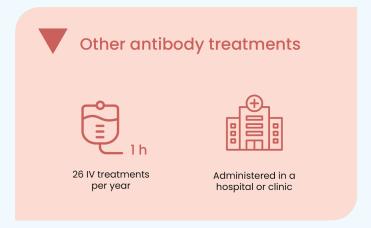
- Unique binding profile for targeting toxic oligomers
- Several potential uses for the antibody:
 - complement to the vaccine ALZ-101 for patients who, for various reasons, need higher levels of antibodies
 - stand-alone treatment against Alzheimer's
- Lower doses (vs. non-specific antibody)



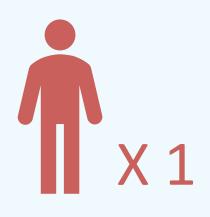
Improving quality of care – for more patients





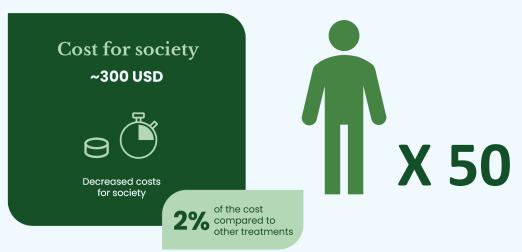












Based on Swedish healthcare system, and that the two treatments have equivalent clinical efficacy, total treatment duration and drug cost.

Sources: SCB Statistics Sweden, Swedish regional healthcare cost



Alzinova's product pipeline offers complementary ways to fight Alzheimer's Disease

Candidates		Discovery	Pre-clinical	Clinical	NDA/MAA	Next Major Milestone
ALZ-101 ¹⁾	therapeutic vaccine				top-line data mber 2023	Q1 2024, Full analysis of part A clinical phase 1b
ALZ-201 ²⁾	monoclonal antibody					TBD "Phase 1 ready"
New innovative discovery projects						Evaluating possiblities for expanding the portfolio

¹⁾ Phase 1b study in patients with Alzheimer's

²⁾ Pre-clinical development for phase 1b



What if there was a vaccine for Alzheimer's?

Alzinova

- ✓ Targeted treatment **oligomer specific**
- **✓** Vaccine
- ✓ Antibody as a complement to the vaccine
- **✓** Fast, effective and uncomplicated
- **✓** Likelihood for good efficacy
- Can start treatment early

Other actors

- Target plaques
- Non-specific treatments
- Often complicated drug treatments
- Unlikely to be sufficiently effective



ALZ-101 – favorable profile with "best-in-class" opportunity

	ALZ-101	Donanemab	Lecanemab
Companies	Alzinova	Eli Lilly	BioArctic/Eisai/Biogen
Primary target -> good efficacy	Aβ oligomers +++ (potential)	plaque (P-Glu3-Aβ) +	Aβ monomers/fibrils/oligomers +
Compliance	+++	+	+
Low cost*	+++	++	+
Low societal burden**	+++	+	+
Patient coverage	+++	+	+
Low risk of complications related to therapy (ARIA-E, ADA, immunogenicity etc.)	+++	+	+

^{*}treatment cost per patient

^{**} cost for the society, patients and families

ALZ-101 & ALZ-201 are patented through two patent families



We maintain an active patent strategy, for the future commercial products.

Stable amyloid-beta monomers and oligomers (ALZ-101)						
Region/Country	Patent number	Status	Expiry date			
Australia	2009236699	Granted	2029-04-14			
Europe ⁽¹⁾	2262526	Granted	2029-04-14			
Japan	5817060	Granted	2029-04-14			
Japan	6128450	Granted	2029-04-14			
India	292362	Granted	2029-04-14			
Canada	2721156	Granted	2029-04-14			
China	ZL200980122412.X	Granted	2029-04-14			
USA	9,688,734	Granted	2030-01-02			
USA	10,023,622	Granted	2029-04-14			
USA	10,138,281	Granted	2029-04-14			
Anti oligomer antibodies (ALZ-201)						
Region/Country	Patent number	Status	Expiry date			
Europe ⁽²⁾	2683738	Granted	2032-03-07			
USA	9062102	Granted	2032-07-30			

⁽¹⁾ Belgium, Switzerland, Lichtenstein, Czech Republic, Germany, Denmark, Spain, Finland, France, United Kingdom, Croatia, Ireland, Italy, Norway, Holland, Poland, Sweden.

⁽²⁾ Switzerland, Germany, France, United Kingdom, Ireland, Luxembourg, Monaco, Holland, Sweden.

Strong data supports clinical development of ALZ-101

Comprehensive pre-clinical results

- ✓ ALZ-101 is well tolerated in non-clinical safety and toxicity studies (monkeys and rabbits)
- ✓ Efficacy data in three different Alzheimer's models, all showing promising results
 - Transgenic mice
 - Aβ-immunodepletion of human AD brain extracts; and assayed in zebrafish embryos

Robust and validated process for manufacturing of drug substance and product

- ✓ GMP manufacturing of vaccine Phase 1b
- ✓ Scale up manufacturing of vaccine for Phase 2





Alzinova's ALZ-101 Phase 1b-study

ALZ-101 Phase 1b clinical study in patients with Alzheimer's disease



Study: Double-blind, randomized, placebo-controlled study on safety, tolerability and immunogenicity.

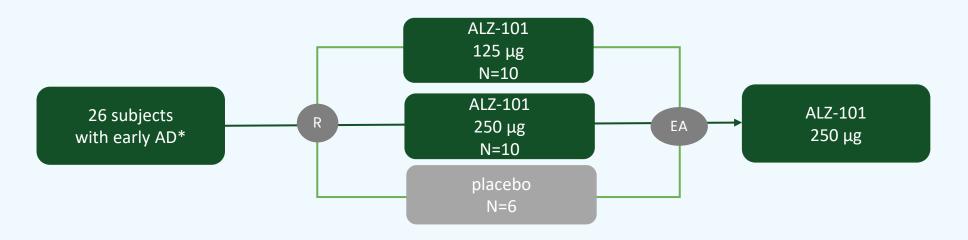
Part A - ALZ-101 (125 and 250 μg) or placebo, given at weeks 0, 4, 8 and 16, if not eligible for Part B, follow-up for additional 48 weeks.

Part B - ALZ-101 (250 μg), given at week 0, 4, 8 and 16 and for additional 48 weeks

Patient population: Patients between 50-80 years, MCI due to AD or mild AD*, CSF pattern consistent with amyloid plaque load and indicative of AD pathology.

Endpoints:

Primary; Safety, tolerability - **Secondary:** Immunogenicity (Aβ-specific) - **Exploratory:** Biomarkers (CSF & blood) and cognition



^{*}according to National Institute of Aging – Alzheimer's Association (NIA-AA)
MCI – Mild Cognitive Impairment; AD – Alzheimer's Disease; CSF – Cerebrospinal fluid; Aβ – amyloid-beta

Strong data in Phase 1b Study

Interim analysis - good safety and tolerability with a clear immune response

- 26 subjects included
- treated with at least 3 doses of ALZ-101 (125 and 250 μg) or placebo
- increasing antibody levels with number of doses given

Top-line results

- The study met the primary endpoint of safety and tolerability after 20 weeks.
- A high immune responder rate
- Antibody levels increasing with the number of doses given

Extension of phase 1b study

- All patients receive ALZ-101 (250 μg)
- Long-term data for safety & tolerability
- Biomarker & cognitive data
- Advantages ahead of Phase 2



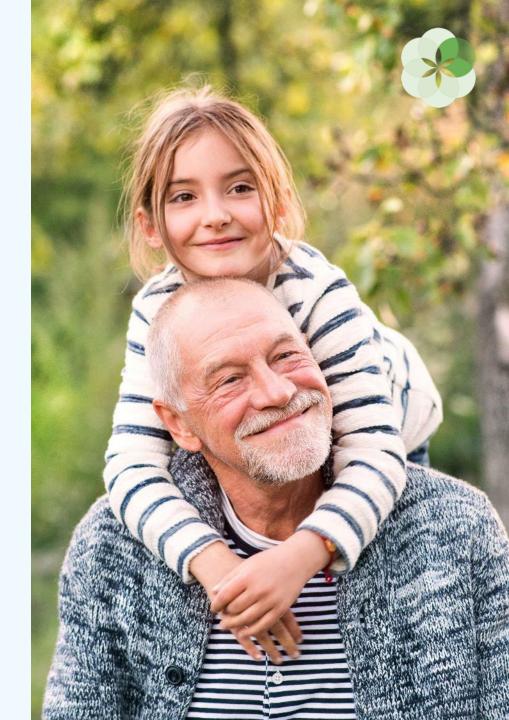
Phase 2 preparations

Interactions with Regulatory Authorities – FDA and EMA

Comprehensive preparation work

- Scalable manufacturing process
- Study design and protocol development
- CRO and site selection
- Fast track application based on Phase 1b data
- Intellectual property strategy work
- Manufacturing of clinical trial material

Strategic partnership discussions





Corporate & Finance

Experienced management team



Kristina Torfgård, PhD
Chief Executive Officer
30 years experience from leading roles in pharma and biopharma industry



Anders Sandberg, PhD
Chief Scientific Officer
Co-Founder
>20 years experience in protein research



Håkan Skogström
Chief Financial Officer
>25 years experience from leading finance positions in the industry



Kirsten Harting
Chief Medical Officer
>30 years experience in medicine,
clinical- and business development



Sebastian Hansson
Business Development Director
>15 years experience in IP/patent,
pharmaceutical R&D and production



Stefan Pierrou, PhD
Development Project Director
>25 years experience from project
leading and managerial roles in R&D



Ann-Sofie Sternås
Intellectual Property Officer
>30 years experience in IP/patent
in pharma and biopharma industry



Julian Aleksov Chairman

>25 years experience in finance and international development within the pharma industry



Anders Blom Member

>25 years experience in international finance and business development in pharma and medical device industry



Per-Göran Gillberg, PhD Member

35 years experience in drug development in pharma and biopharma industry. Adj prof neuroscience at Uppsala University



Clas Malmeström, MD, PhD Member

Chief physician MS-center, neurosurgery and lab of clinical immunology, Sahlgrenska University Hospital





Lena Degling Wikingsson, PhD
Member

25 years experience in drug development of biological drugs and vaccines



Carol Routledge, PhD Member

>30 years experience in drug development and acquisitions in pharma and biopharma industry



Anders Waas Member

>25 years experience in management, business & pharmaceutical development



Capitalization table, summary of shares and options



Shareholders, as per 29 September 2023:

Name	No. of shares	Capital, %
Maida Vale Capital AB	6,747,686	15.15%
Försäkrings AB Avanza Pension	3,223,681	7.24%
Nordnet Pensionsförsäkring AB	2,229,751	5.01%
Patrik Ahlvin	1,004,750	2.26%
Sara Gjertz	877,303	1.97%
Ålandsbanken, for owners	746,499	1.67%
MIVAC Development AB	711,787	1.60%
Özlem Erdogdu Gül	671,316	1.51%
MGC Capital Ltd	604,171	1.36%
Moll Invest AB	600,080	1.35%
Total 10 largest owners	17,417,024	39.12%
Total other owners	27,114,241	60.88%
Total all	44,531,265	100.00%

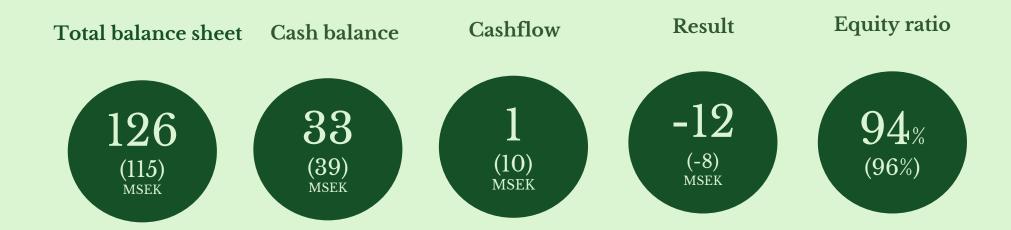
Company management and Board of Directors, direct and indirect holdings:

- CEO and other executive managers, 324,118 shares
- Board of Directors, 7,778,845 shares





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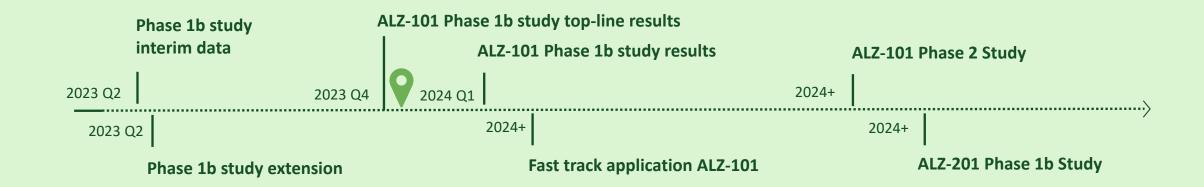


- Mainly invested in capitalized costs for R&D, focus on present ALZ-101 phase 1 study and preparing ALZ-101 for phase 2 study.
- Rights issue provided 30 MSEK during 2022, to use for finalising ALZ-101 phase 1 study and continue towards phase 2 readiness. Additional 25 MSEK (after cost) provided from rights issue in April 2023.
- Personnel costs slightly higher when building up the organisation.

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Roadmap & value inflection points



Cash to finance ALZ-101 through phase 1b and preparatory ahead of phase 2

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Investment highlights



Alzinova's goal is to enable Alzheimer's patients to live an independent and active life.



Alzinova's lead candidate, ALZ-101, is a therapeutic vaccine against Alzheimer's in clinical development with a phase 1b study in patients.



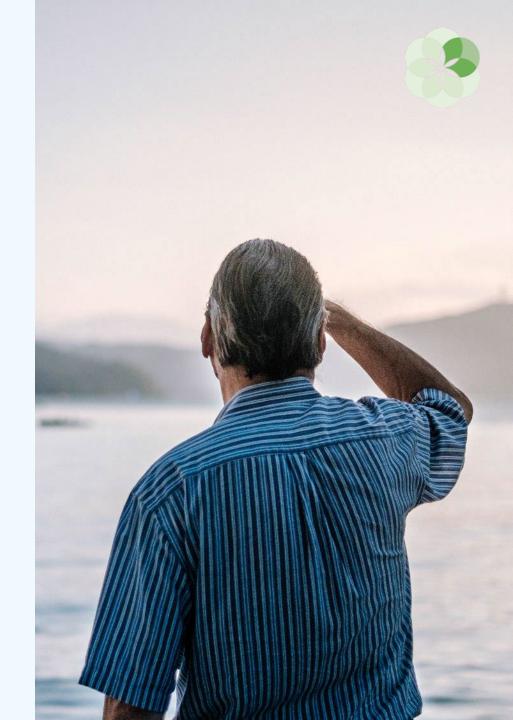
Based on the same technology, Alzinova is also developing a monoclonal antibody, ALZ-201, as a complementary treatment to fight Alzheimer's.



Data show that the unique specificity of Alzinova's vaccine (ALZ-101) and the monoclonal antibody (ALZ-201) provides "best-in-class" potential.



Preparatory activities underway for next clinical development phase, making Alzinova's candidates more attractive for strategic partnership.





Alzinova

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https://www.linkedin.com/company/alzinova-ab/