



HANSA

BIOPHARMA

Investor Road Show
Presentation Q2 2022

Lund, July 19, 2022



Forward-looking statements

This presentation may contain certain forward-looking statements and forecasts based on our current expectations and beliefs regarding future events and are subject to significant uncertainties and risks since they relate to events and depend on circumstances that will occur in the future. Some of these forward-looking statements, by their nature, could have an impact on Hansa Biopharma's business, financial condition and results of operations [or that of its parent, affiliate, or subsidiary companies]. Terms such as "anticipates", "assumes", "believes", "can", "could", "estimates", "expects", "forecasts", "intends", "may", "might", "plans", "should", "projects", "will", "would" or, in each case, their negative, or other variations or comparable terminology are used to identify forward-looking statements. There are a number of factors that could cause actual results and developments to differ materially from those projected, whether expressly or impliedly, in a forward-looking statement or affect the extent to which a particular projection is realized. Such factors may include, but are not limited to, changes in implementation of Hansa Biopharma's strategy and its ability to further grow; risks and uncertainties associated with the development and/or approval of Hansa Biopharma's product candidates; ongoing clinical trials and expected trial results; the ability to commercialize imlifidase if approved; changes in legal or regulatory frameworks, requirements, or standards; technology changes and new products in Hansa Biopharma's potential market and industry; the ability to develop new products and enhance existing products; the impact of competition, changes in general economy and industry conditions and legislative, regulatory and political factors.

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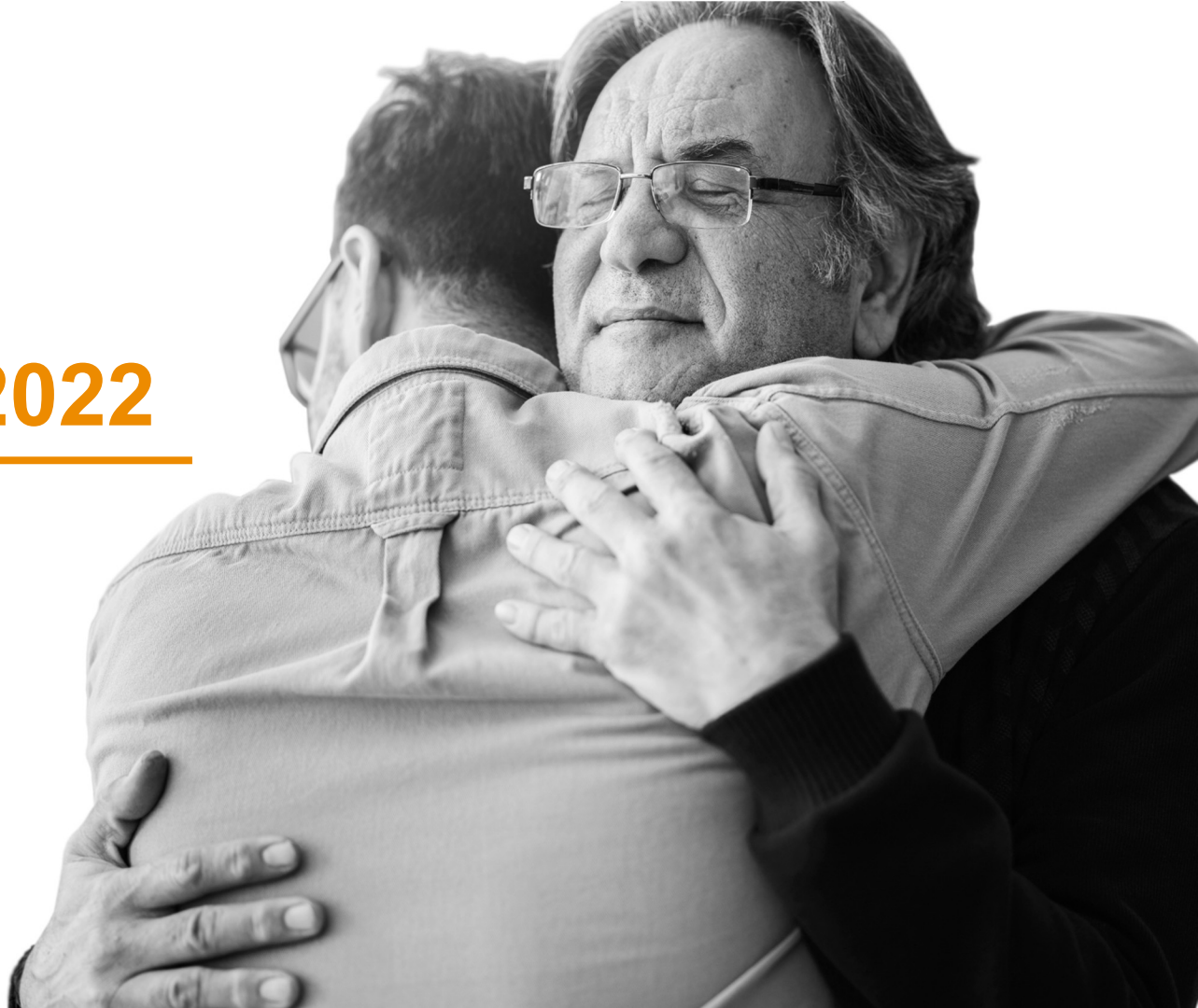
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Business update Q2'2022



Continued solid sales in Q2; Positive recommendation by NICE; Patient enrollment completed in AMR; Peter Nicklin appointed as new Chairman of the Board

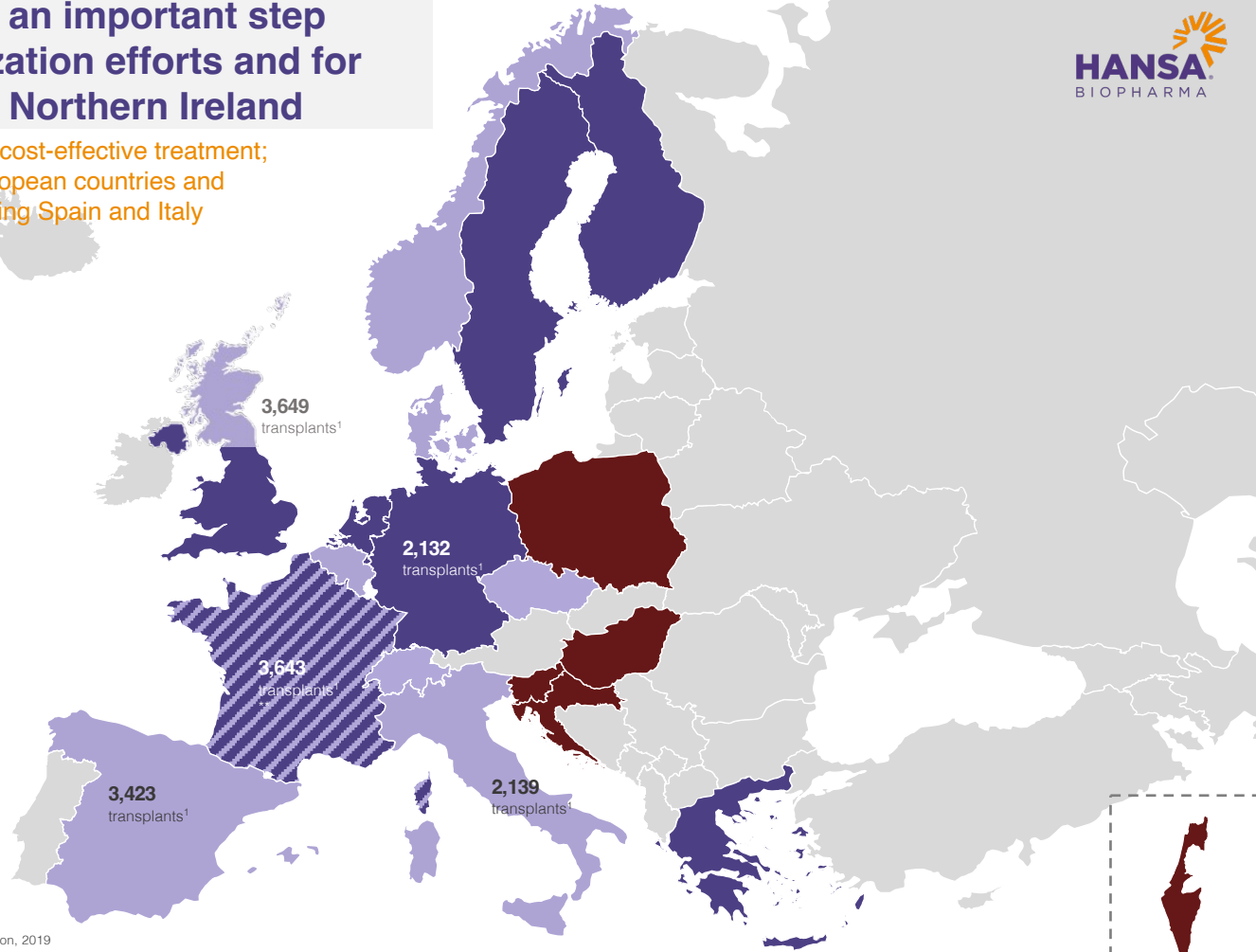
Highlights for the second quarter of 2022

- ✓ **Launch activities and market access efforts in EU progressing as planned**
 - Continued solid sales, with SEK 19.5m in product sales; Total revenue of SEK 26.4m
 - Market access obtained in England, Wales and Northern Ireland as NICE recommends Idefirix® for desensitization of highly sensitized patients
 - France grants Idefirix® ASMR 3 rating by the Transparency Commission (TC) of the French National Authority for Health (HAS)
 - Market access has now been secured in 7 countries and procedures are ongoing in 11 countries, including Spain and Italy
 - Temporary marketing authorization granted for Idefirix® in Switzerland
- ✓ **Clinical pipeline**
 - U.S. ConfIdoS Study in kidney transplantation: 22/64 patients enrolled
 - Anti-GBM: Expect to commence Phase 3 study later this year, as previously guided
 - AMR: Patient enrollment completed; First data read-out expected in H2'22
 - GBS: 18/30 patients enrolled in the GBS phase 2 study; Significant initiatives were implemented during H1 2022 to support the completion of enrollment in H2 2022
- ✓ **Annual General Meeting held on June 30, 2021**
 - All resolutions were approved by shareholders
 - Peter Nicklin appointed as new Chairman of the Board. Peter Nicklin brings significant experience from leading global teams in large and mid-size life science companies
- Events after the reporting period**
 - ✓ - First patient was treated in Hansa's European post approval efficacy study (PAES)
 - Concluded a USD 70 million non-dilutive financing transaction with NovaQuest Capital Management to support the continued development of Hansa's antibody-cleaving enzyme technology platform across multiple therapeutic areas while extending the expected cash runway through 2024.



Recommendation by NICE is an important step forward for our commercialization efforts and for patients in England, Wales & Northern Ireland

NICE considers Idefirix® to be a clinically- and cost-effective treatment; Market access has now been secured in 7 European countries and procedures are ongoing in 11 countries, including Spain and Italy



¹Annual kidney transplantations 2019 (pre-Corona)

²Transplantation data is from Global Observatory on Donation and Transplantation, 2019

^{**}Pricing & reimbursement obtained in France on an early access basis

First patient treated in post-authorization efficacy study (PAES) of Idefirix® (imlifidase) in highly sensitized kidney transplant patients

The study will provide further important insights regarding Idefirix® desensitization treatment of highly sensitized kidney transplant patients

An open-label Phase 3 study in 50 patients

- First patient was treated by Dr. Oriol Bestard, Chair of Nephrology and Kidney Transplantation at Vall d'Hebron University Hospital in Barcelona
- Study will enroll 50 highly sensitized patients with positive crossmatch against an available deceased donor across multiple countries and centers in Europe
- The study is an obligation under the conditional marketing authorization for Idefirix® granted by EMA in August 2020, in order to complete a full marketing authorization in the EU
- The aim will be to confirm the long-term efficacy and safety of Idefirix® with the primary objective to determine the one-year graft failure-free survival of the Idefirix® treated and transplanted patients.
- In addition, a total of 50-100 patients undergoing compatible kidney transplantation at the participating centers will be included and serve as a non-comparative concurrent reference cohort, with no formal comparison, to contextualize the one-year graft failure-free survival of the Idefirix® treated patients



Continuous progress in our ongoing clinical Programs

Enrollment status
July 19, 2022



Antibody Mediated Rejection Phase 2 study

- 30/30 patients enrolled in the AMR phase 2 study
- Completion of enrollment expected H1 2022*
- First data read out expected in H2 2022*



Guillain-Barré Syndrome Phase 2 study

- 18/30 patients enrolled in the GBS program
- Ten centers are active and open for recruitment
- Significant initiatives were implemented during H1 2022 to support the completion of enrollment in H2 2022*
- First data read out expected in H1 2023

Enrollment status
July 19, 2022



Anti-GBM Phase 3 study

- FDA has accepted Hansa's Investigational New Drug (IND) application to proceed with a Phase 3 study
- The planned study will commence this year targeting 50 patients across the U.S. and Europe*

- Patients enrolled
- Patients remaining



U.S. ConfldeS Phase 3 study

Randomized, controlled trial in highly sensitized kidney transplant patients across up to 15 centers

- 22/64 patients enrolled for randomization
- Ten centers are active and open for recruitment
- Completion of enrollment expected H2 2022*

- Patients enrolled
- Patients remaining

Broad clinical pipeline in transplantation and auto-immune diseases

Candidate/ Program	Indication	Research/ Preclinical	Phase 1	Phase 2	Phase 3	Marketing Authorization	Marketed	Next Anticipated Milestone
Imlifidase	EU: Kidney transplantation in highly sensitized patients ^{1,2}	Completed	Completed	Completed	Planned	Completed	Completed	EU: Additional agreements around reimbursement from H2'21
	US: Kidney transplantation in highly sensitized patients ^{1,2}	Completed	Completed	Completed	Ongoing			Completion of enrollment (64 patients) H2'22
	Anti-GBM antibody disease ³	Completed	Completed	Completed	Planned			Pivotal Phase 3 study expected to commence in 2022 (50 patients)
	Antibody mediated kidney transplant rejection (AMR)	Completed	Completed	Ongoing				First data readout H2'22
	Guillain-Barré syndrome (GBS)	Completed	Completed	Ongoing				Completion of enrollment (30 patients) H2 2022
	Pre-treatment ahead of gene therapy in Limb-Girdle (Partnered with Sarepta)	Ongoing						Preclinical phase
	Pre-treatment ahead of gene therapy in Duchenne (Partnered with Sarepta)	Ongoing						Preclinical phase
	Pre-treatment ahead of gene therapy in Pompe disease (Partnered with AskBio)	Ongoing						Preclinical phase
NiceR	Recurring treatment in autoimmune disease, transplantation and oncology	Ongoing						Completion of GLP toxicology studies in 2022
EnzE	Cancer immunotherapy	Ongoing						Research phase

¹ Results from the Phase 1 study have been published, Winstedt et al. (2015) PLOS ONE 10(7)

² Lorant et al American Journal of Transplantation and 03+04 studies (Jordan et al New England Journal of Medicine)

³ Investigator-initiated study by Mårten Segelmark, Professor at the universities in Linköping and Lund

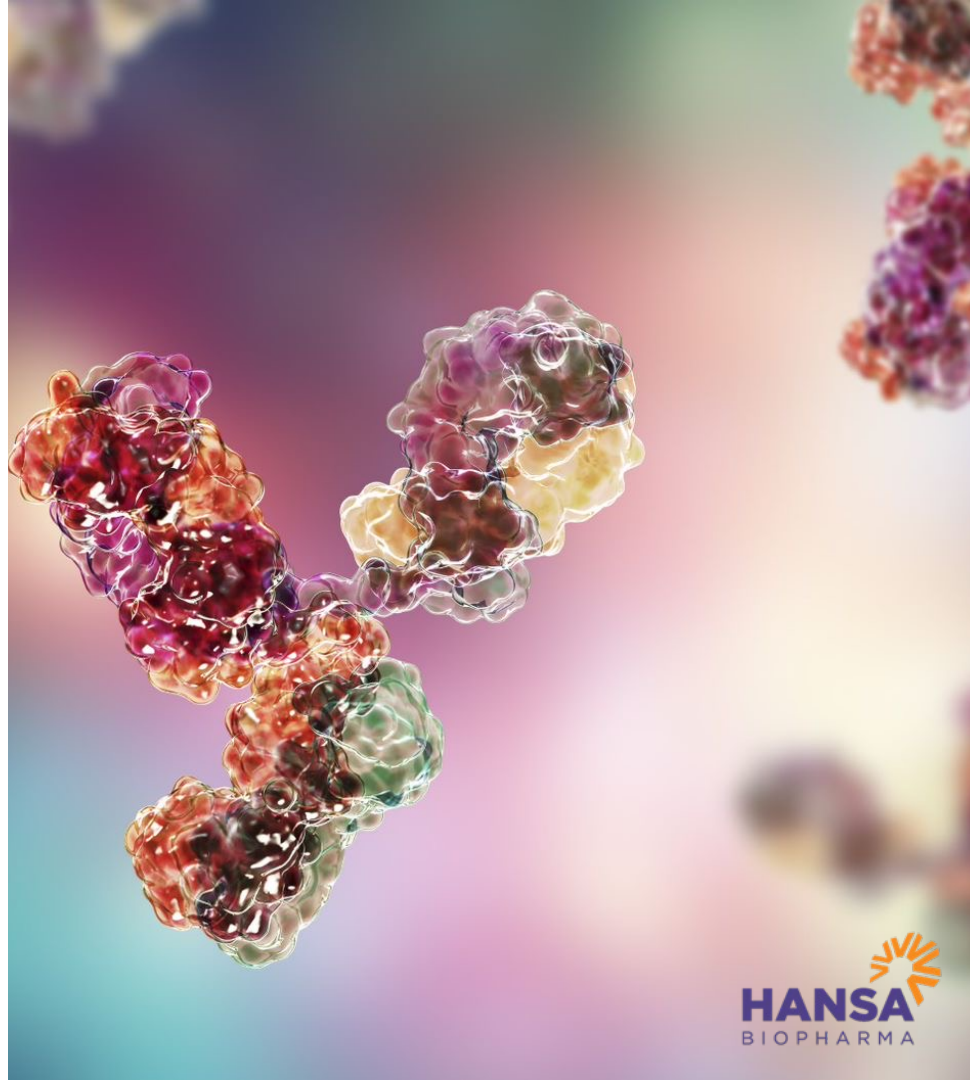
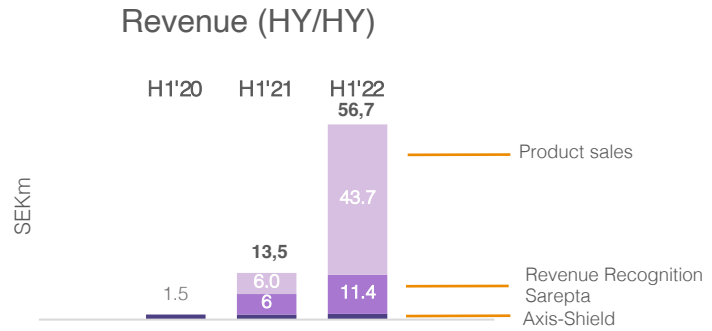
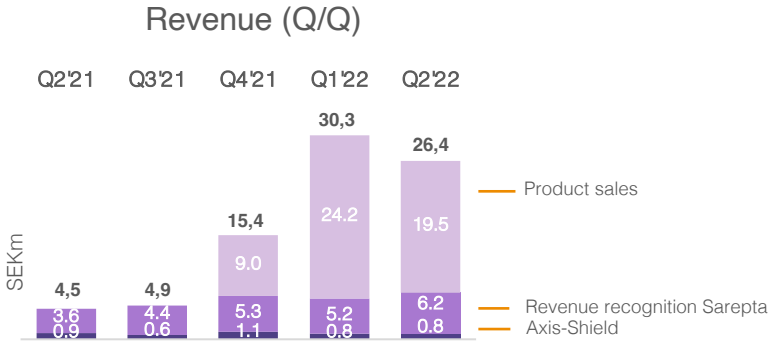
Completed

Ongoing

Planned

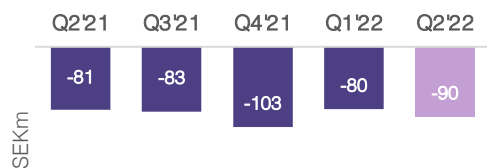
Post approval study running in parallel with commercial launch

Continued solid sales in Q2 with product sales of SEK 19.5m; Total H1-2022 revenue SEK 56.7m

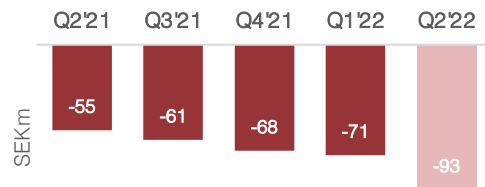


Continued investments in commercialization and our pipeline

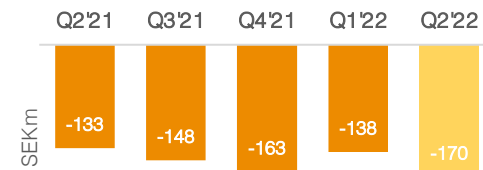
SG&A expenses (Q/Q)



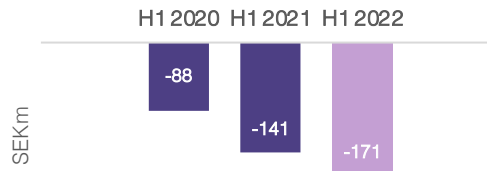
R&D expenses (Q/Q)



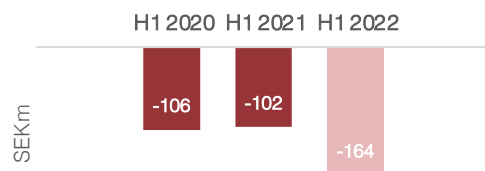
Net loss (Q/Q)



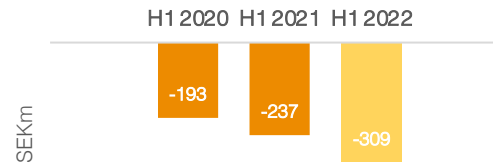
SG&A expenses (HY/HY)



R&D expenses (HY/HY)

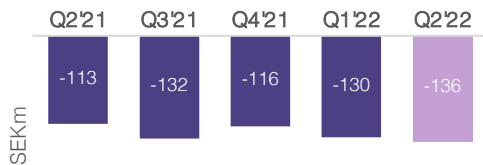


Net loss (HY/HY)

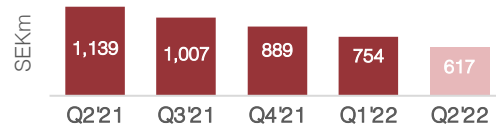


With recent financing transaction secured with NovaQuest; Hansa's cash runway has extended through 2024

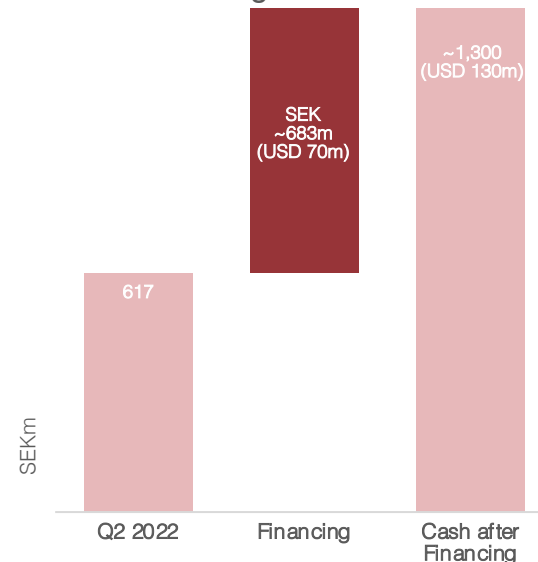
Operating cash flow (Q/Q)



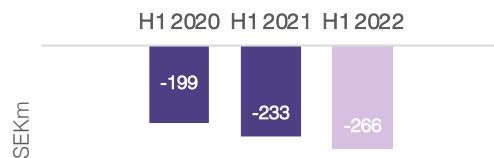
Cash & short-term investments (Q/Q)



Cash position post recent financing transaction



Operating cash flow (HY/HY)



Number of employees (Q/Q)

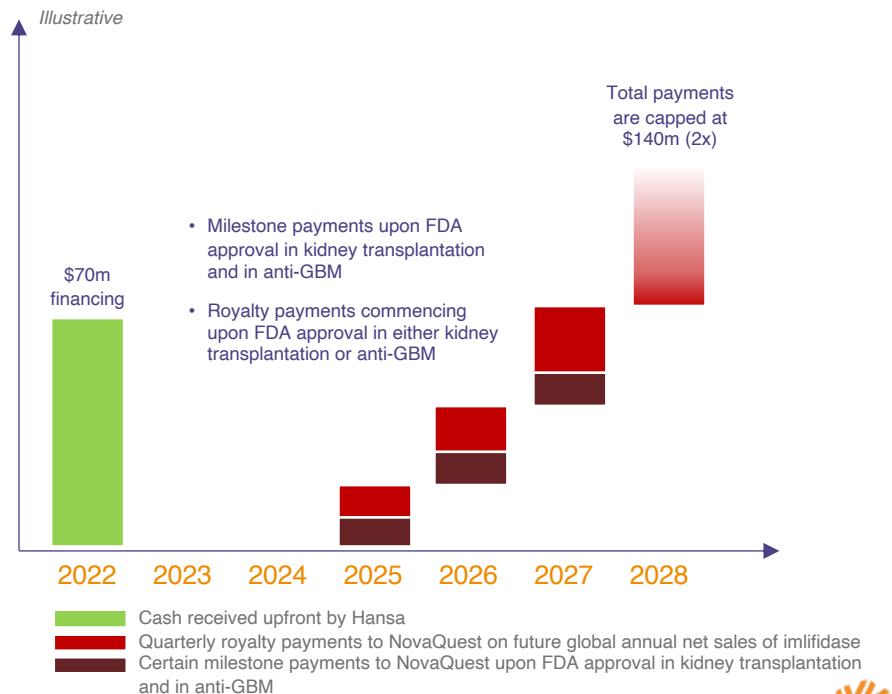


\$70 million non-dilutive financing transaction to support the continued development of Hansa's antibody-cleaving enzyme technology platform

Transaction extends cash runway through 2024 and helps bolster the ability to invest in the continued development of our unique antibody-cleaving enzyme technology platform across multiple therapeutic areas

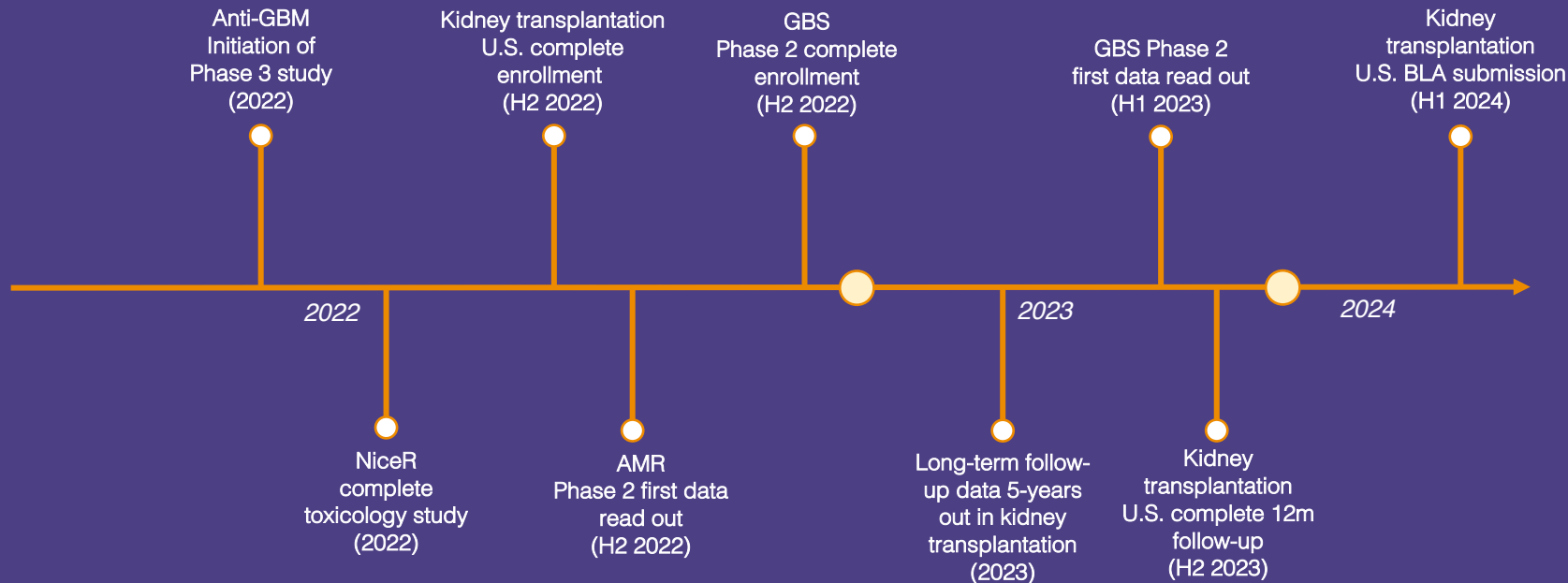
Proceeds from the transaction will chiefly be utilized to:

- Further strengthen Hansa's position in kidney transplantation through the continued support of ongoing European commercial launch activities for Idefirix (imlifidase) and execution for the U.S.
- Further fund ConfldeS trial of imlifidase, which is expected to support a potential Biologics License Application (BLA) submission to the U.S. Food and Drug Administration (FDA) under the accelerated approval pathway in the first half of 2024.
- Advance the global Phase 3 clinical trial of imlifidase in anti-GBM antibody disease, and
- Together with the existing cash, complete our ongoing Phase 2 programs in AMR and GBS and to advance Hansa's next generation of enzymes (NiceR) into clinical development



Upcoming milestones

Milestones subject to potential COVID-19 impact



Guidance assumes no persistent impact or further escalation of the COVID-19 pandemic potentially forcing trial centers to reprioritize patient recruitment or even shut down again.

Company overview



Successful track record...
Strong momentum...
Promising future...

A validated technology

VALIDATION ACROSS THREE AREAS

- ✓ Approval in kidney transplantations
- ✓ Proof of concept in autoimmune diseases
- ✓ Partnerships to explore gene therapy

Idefirix® is our first approved drug in Europe*

EUROPE KIDNEY TRANSPLANTS

For highly sensitized patients in Europe

Broad pipeline in transplantation and autoimmunity

PROGRAMS IN CLINICAL DEVELOPMENT

US kidney transplants
Anti-GBM
Guillain-Barré syndrome (GBS)
Antibody mediated kidney transplant rejection (AMR)

Established a high-performance organization

NEW COMPETENCIES ADDED

145 employees June 2022 (~3x in 3 years)
Highly qualified team with 20 years on average in life science
Purpose driven culture

With current cash position Hansa is financed through 2024

FINANCIALS

SEK 617 in Cash and short term investments (USD ~60m) June 2022
SEK ~1.3bn (USD ~130m) post NovaQuest financing transaction carried out July 2022

Created shareholder value and diversified our ownership base

MARKET CAPITALISATION (USD): ~220m

Listed on Nasdaq Stockholm
18,000 shareholders
Foreign ownership make up ~40% through leading international life science specialist funds



*Idefirix approved in EEA under conditional approval for kidney transplantation

**Actual patient has given consent to provide images

We are building a global leader in rare diseases

Today

We are launching our first commercially approved product for enablement of kidney transplantation in Europe*



Tomorrow

We envision a world where patients with rare immunologic diseases can lead long and healthy lives



* European Economic Area incl EU plus Iceland, Liechtenstein and Norway

Stock images

Our mission

We leverage our unique enzyme technology platform to develop innovative, lifesaving and life altering immunomodulating therapies, bring these to the patients with rare diseases who need them, and generate value to society at large.



Develop new therapies



Extend and improve human lives

Transplantation leads to **dramatically better quality of life and life expectancy than dialysis**

77% of transplanted patients are alive after 8 years vs 44% of patients on dialysis¹



Deliver value to society

Transplantation is a **cost-effective intervention vs. dialysis**

Idefix was named in EMA report as **Outstanding contribution to public health³**

USD 115bn, equivalent to 20% of the US Medicare budget, relates to kidney diseases²



Desensitization in **kidney transplant patients***



Exploring treatment options in **anti-GBM****



Exploring treatment options in **GBS****



Exploring treatment options in **AMR****

* Idefix approved in EEA under conditional approval for kidney transplantation
** Imifidase under investigation

¹ Orandi et al. N Engl J Med 2016;374:940-50
² <https://www.hhs.gov/about/news/2019/07/10>
³ https://www.ema.europa.eu/en/documents/report/human-medicines-highlights-2020_en.pdf

Hansa Biopharma's history

Pre-clinical proof-of-concept established

October 2007
39 SEKm valuation

Imlifidase GMP process development and toxicology studies initiated

2. Clinical development

After the first-in-man study in **2013**, Hansa has completed four Ph. II studies since **2014**. Additionally Hansa has ongoing and completed trials in Anti-GBM, AMR and GBS. In **2019** a MAA was submitted for imlifidase in kidney transplantation.

SAREPTA THERAPEUTICS

Hansa Biopharma enters the gene therapy space through partnership agreement with Sarepta Therapeutics to develop imlifidase as a pre-treatment to enable gene therapy in NAb+ patients

idefixir
(imlifidase)

The EU Commission grants conditional approval for Idefixir® in highly sensitized kidney transplant patients in Europe



1. Turning foe into friend

The therapeutic potential in using a bacterial enzyme with specificity for IgG-antibodies, to neutralize pathogenic antibodies was discovered around **2006**. The original enzyme, IdeS, has been developed by *Streptococcus pyogenes* over thousands of years and by transferring a majority of the IdeS coding nucleotide into harmless *E. coli*-bacteria, the IgG-cleaving part of the IdeS molecule can be expressed and purified, resulting in the Hansa Biopharma drug imlifidase, i.e. turning a former foe to a friend.

First-in-man study

First Ph. II study of imlifidase in kidney transplantation

Starts trading on NASDAQ Stockholm main board

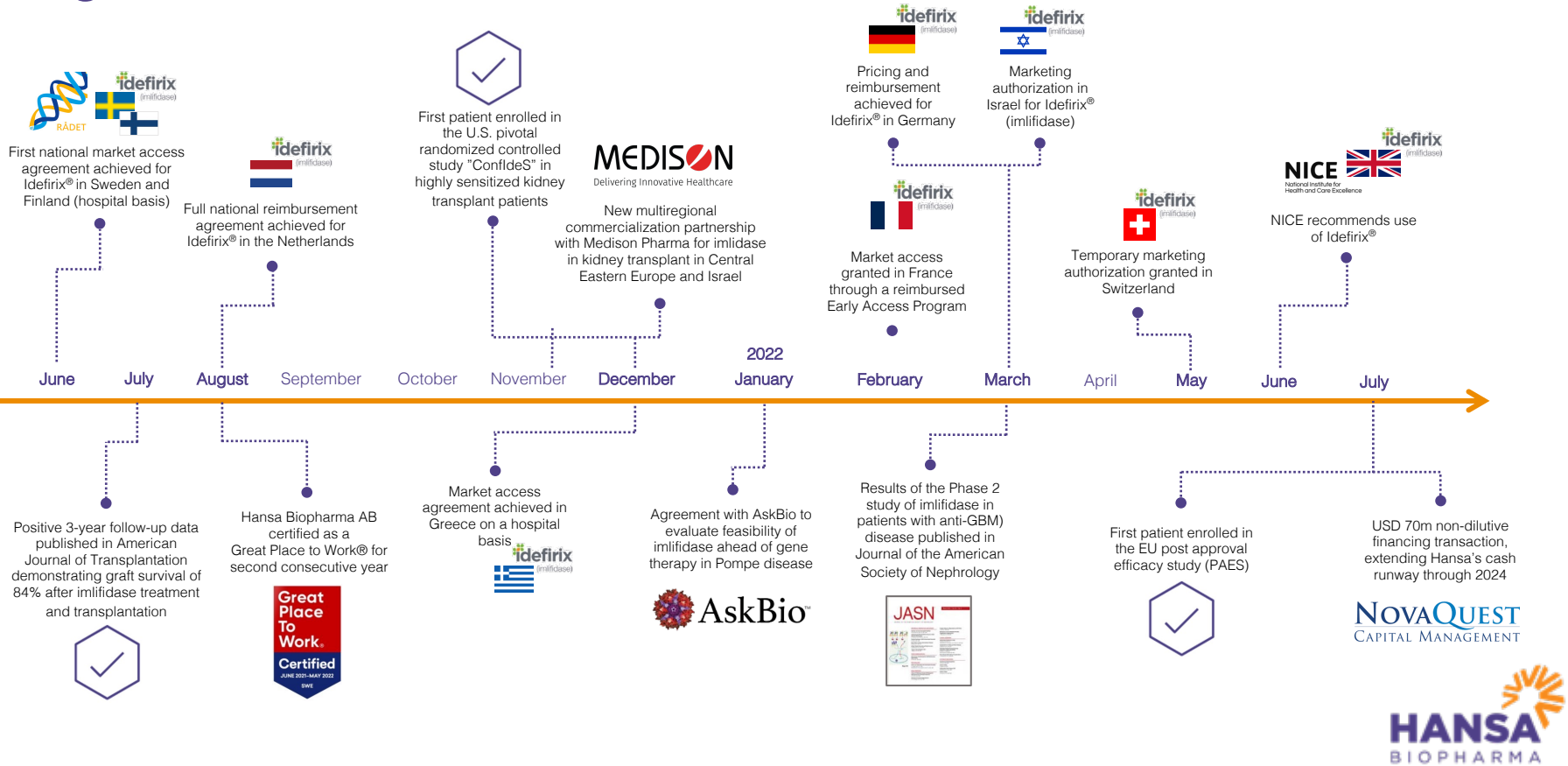
3. Commercialization

In august **2020**, Hansa received conditional approval for Idefixir (imlifidase) in kidney transplantation. Additionally, Hansa entered into the gene therapy field through a commercial partnership with Sarepta Therapeutics. Thus far, Hansa achieved market access in the first 7 European markets incl. UK, Germany and France (early access). Market Access procedures are ongoing in 11 additional countries

Hansa Biopharma enters into three collaborations with argenx (combination therapies); Medison (commercial) and AskBio (gene therapy)



Many milestones achieved during the last 12 months



Imlifidase

a novel approach to eliminate pathogenic IgG

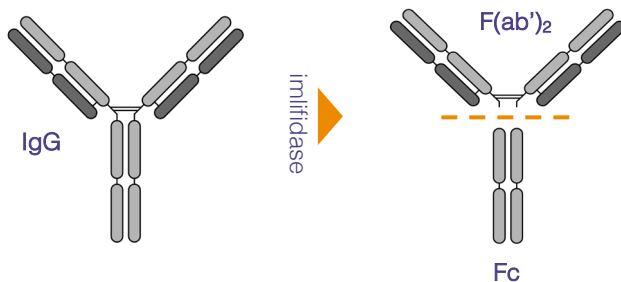
Origins from a bacteria *Streptococcus pyogenes*

- Species of Gram-positive, spherical bacteria in the genus *Streptococcus*
- Usually known from causing a strep throat infection



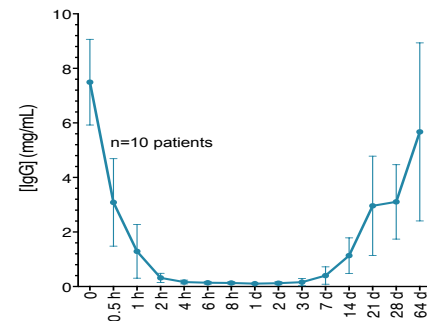
A unique IgG antibody-cleaving enzyme

- Interacts with Fc-part of IgG with extremely high specificity
- Cleaves IgG at the hinge region, generating one F(ab')₂ fragment and one homo-dimeric Fc-fragment



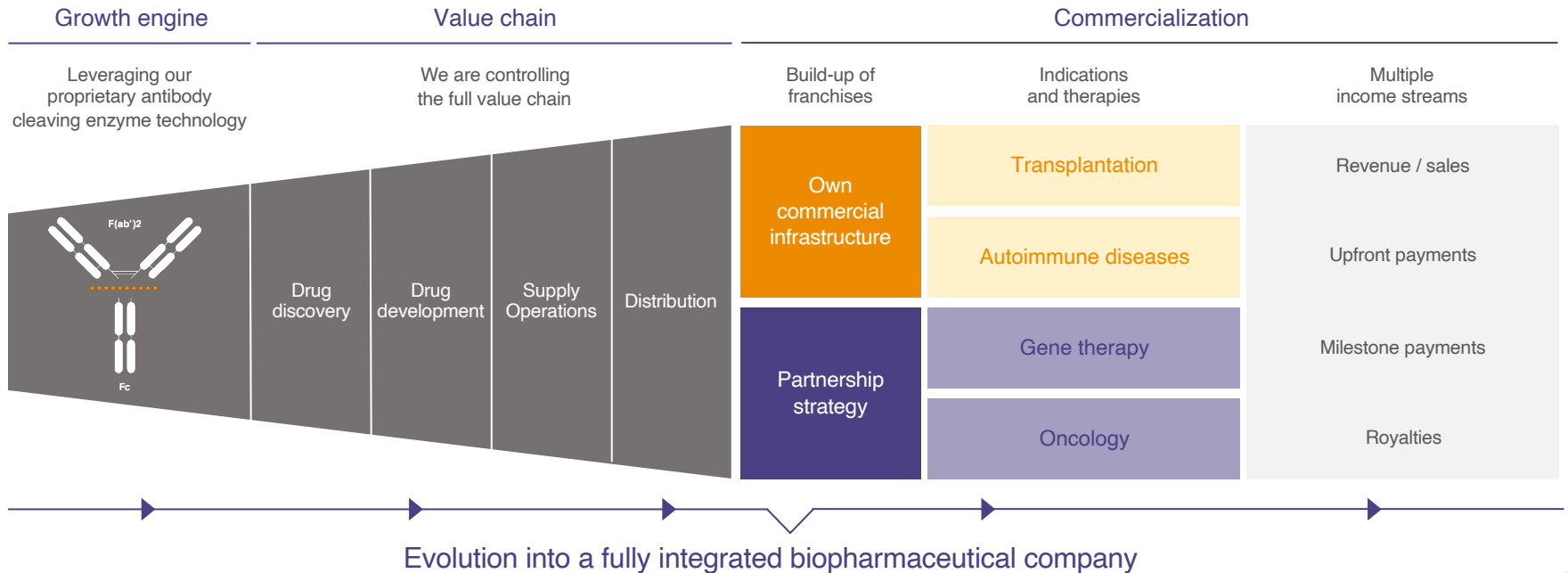
Inactivates IgG in 2-6 hours

- Rapid onset of action that inactivates IgG below detectable level in 2-6 hours
- IgG antibody-free window for approximately one week

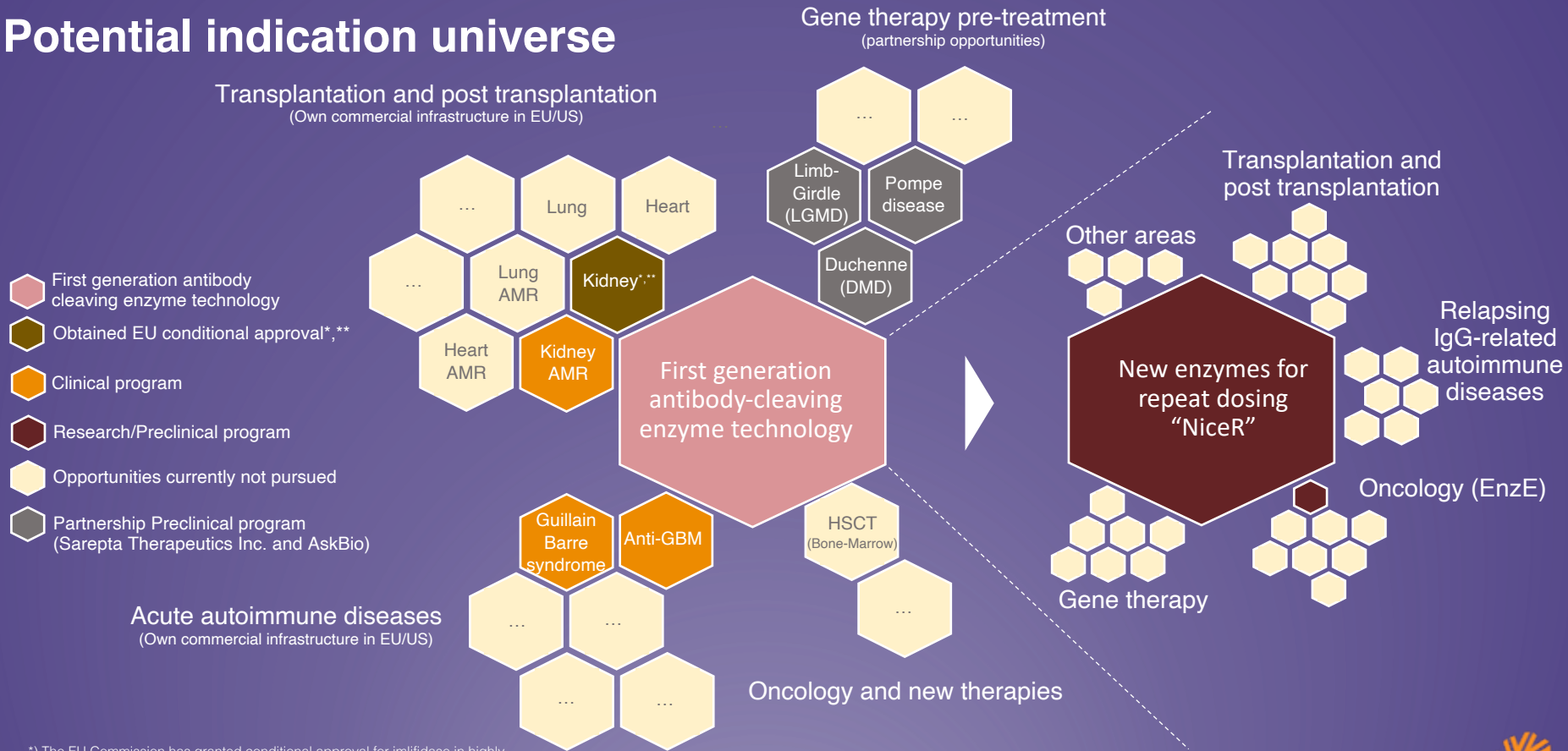


Our Business model

Leveraging our technology platform to develop new therapies targeting rare diseases with unmet medical need across a range of indications



Potential indication universe



*) The EU Commission has granted conditional approval for imlifidase in highly sensitized kidney transplant patients.

**) In the US a new study has commenced targeting a BLA filing by H1 2024

Our strategic priorities

Our mission is to become a global leader in rare diseases



- 1** Successfully commercialize Idefirix® in kidney transplantation in Europe, the U.S. and selected international markets
- 2** Advance our ongoing clinical programs in AMR and rare autoimmune diseases to regulatory approval
- 3** Develop imlifidase as pre-treatment to gene therapy, starting with our collaborations with Sarepta and AskBio
- 4** Develop our next generation IgG-cleaving enzymes to allow for recurring treatment
- 5** Successfully develop and market our products by pursuing a hybrid partnering model

Becoming a fully integrated commercial stage biopharmaceutical company

while expanding our technology and global footprint



Pre-clinical

Early-stage clinic

Late-stage clinic

Commercial stage

1

Creating a scientific platform

- Advanced imlifidase from preclinical models through to approval
- Initiated clinical studies in transplantation in EU and the US
- Built the R&D organization
- Validated through peer-reviewed publications (e.g. NEJM and AJT)

2

Preparing the company for commercial success

- Completion of four phase 2 studies in transplantation
- Development of GMP process
- Expanded the pipeline to post-transplantation and autoimmunity
- Established corporate and medical functions
- Expanding the footprint in EU and US

3

Building and capturing value in new indications and markets

- First drug approval in kidney transplantation in EU*
- Commercialisation
- Market Access secured in the U.K, Germany, France (early access), Sweden, Netherlands as well as Finland and Greece on individual hospital basis
- Expanding commercial teams and adding territory management
- Securing supply chain management
- Progressing pipeline and advancing our technology footprint

* Idefirix approved in EEA under conditional approval for kidney transplantation

Our culture is driven by people passionate about making changes



Purpose driven culture

Helping patients with rare diseases serves as a **strong purpose** for our colleagues to **go the extra mile**



Diverse and international

~45%

Internationals across ~30 nationalities

55/45

Male/female gender split in the leadership team



Skilled and experienced team

>50%

With relevant PhD in R&D

~20 years*

of life science experience on average from Big Pharma, Biotech and Academia

**covers Management, R&D, and Commercial functions*



Motivated workforce

For second consecutive year Hansa is certified as a “**Great Place to Work**” with **100%** participation rate in the survey



Experienced Board and Executive Committee

Extensive experience from the global healthcare industry

Executive Committee



Sören Tulstrup

President & CEO (2018)
+30 years in the Healthcare sector
Ex-CEO at Vifor Pharma
Ex-SVP at Shire Pharmaceuticals
Ex-CEO at Santaris Pharma
Shareholding: 26,541



Christian Kjellman

SVP & CSO/COO (2008)
+20 years in the Healthcare sector
Ex-Head of Research at Cartela
Ex-Senior Scientist at Biolvent,
MSc Chemical Biology, PhD in Tumour
Immunology from Lund University
Shareholding: 6,213



Donato Spota

SVP & CFO (2019)
+20 years in the Healthcare sector
Ex-CFO Basilea Pharmaceutica
Senior Finance roles at Roche
Shareholding: 6,673



Achim Kaufhold

SVP & CMO (2020)
+40 years in the Healthcare sector
Ex-CMO Basilea Pharmaceutica
Ex-CEO Affitech (merged with
Pharmexa A/S)
Ex-CMO Chiron (acquired by
Novartis)
Shareholding: 0



Henk D. van Troostwijk

SVP & CCO (2016)
+20 years in the Healthcare sector
Ex-GM at Raptor
Pharmaceuticals
Ex-BU Director at Genzyme
Europe
Shareholding: 2,564



Anne Säfström Lanner

SVP & CHRO (2019)
Ex-Head of HR European
Spallation Source
Ex-Head of HR Cellavision
Shareholding: 3,565



Peter Nicklin

Chairman (2022)
+30 years in the Healthcare sector
Chairman of Tunstall Healthcare,
Sciensus & Versantis
Held senior executive roles at Baxter,
Bayer, Novartis & Bristol-Myers Squibb
Shareholding: -



Hilary Malone

Board Member (2021)
COO at Valo Health (US).
Chief Regulatory Officer & Head of Global
Regulatory Affairs at Sanofi (2013-2019)
SVP & Head of Worldwide Regulatory
Strategy at Pfizer (2009-2011)
Shareholding: 0



Anders Gersel Pedersen

Board Member (2018)
+30 years in the Healthcare sector
Ex-EVP R&D H.Lundbeck
Chairman of Hansa Biopharma's
Scientific Committee
Shareholding: 2,500



Eva Nilsagård

Board Member (2019)
Board member of several companies,
e.g. Adolife, Bufab, Itras, Abriane
Ex-CFO of Vitrolife and Plasta
Chairman of Hansa Biopharma's
Audit Committee
Shareholding: 3,000



Mats Blom

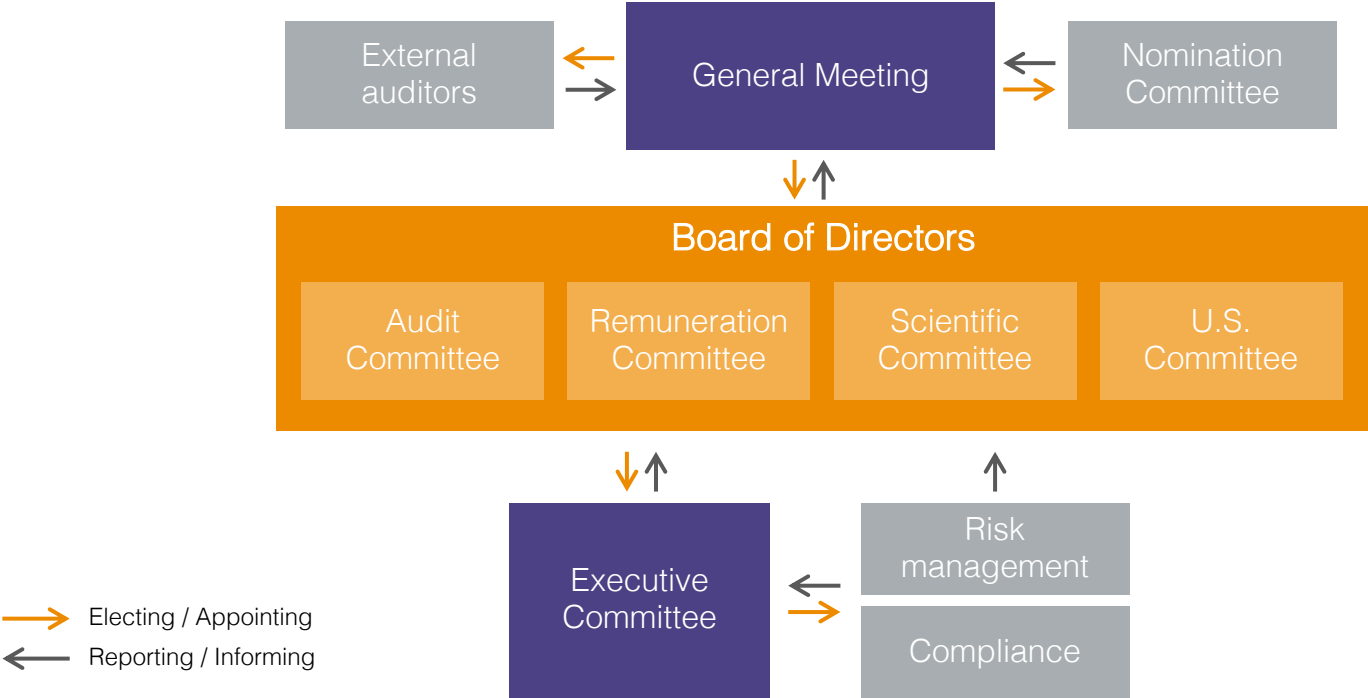
Board Member (2019)
CFO of NorthSea Therapeutics
Ex-CFO Zealand Pharma
Member of Hansa Biopharma's Audit
Committee
Shareholding: 1,000



Andreas Eggert

Board Member (2018)
Ex- SVP at H. Lundbeck A/S
Ex-VP Wyeth/Pfizer in the U.S.
Member of Hansa Biopharma's Audit
Committee and Remuneration Committee
Shareholding: 5,500

Hansa Biopharma's Governance Structure



Strong technology protection

through patents and orphan drug designations

Patent coverage out to 2035 in key markets

- Our lead product, imlifidase, is protected by six patent families including both granted patents and pending applications and cover the use of isolated imlifidase
- Patents cover use of isolated imlifidase at least in:

Medical use in IgG mediated medical conditions

Dosing in combination therapies incl. transplantation & oncology

Autoimmune diseases

Treatment of transplant rejection

Orphan drug designation

- Orphan drug designation is granted to drugs intended for rare diseases (affecting max 5 patients in 10,000 persons in EU or affecting less than 200,000 patients in the US)
- The designation provides development and commercial incentives, including ten years of market exclusivity in the EU and seven years in the U.S., protocol assistance on the development of the drug, including clinical studies and certain exemptions from or reductions in regulatory fees

EMA

Marketing Approval with orphan drug designation

- Conditional marketing approval for imlifidase, for the prevention of graft rejection following solid organ transplantation, was achieved in 2020

Orphan drug designation

- Imlifidase for the treatment of the rare and acute disease anti-GBM (2018)

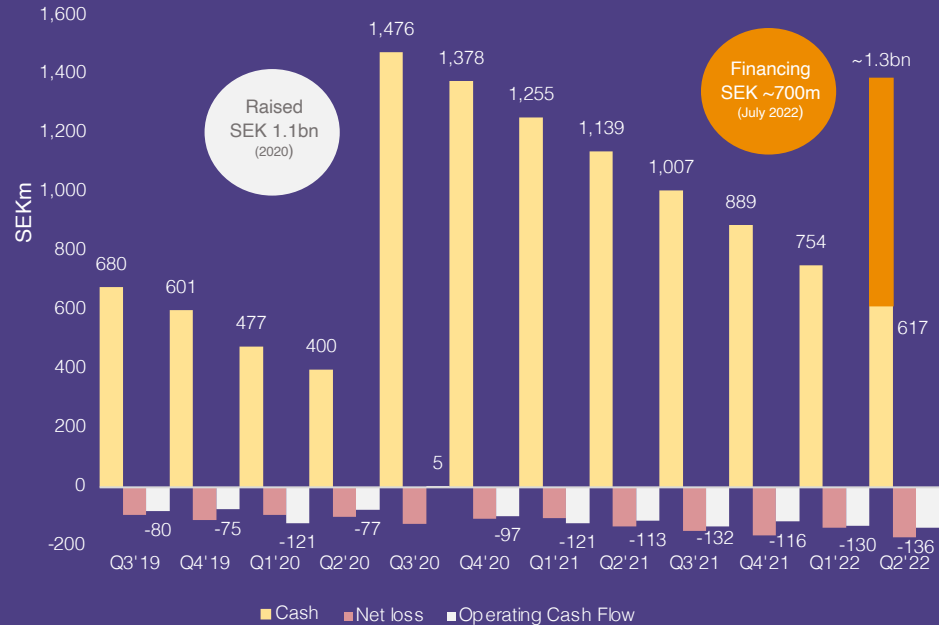
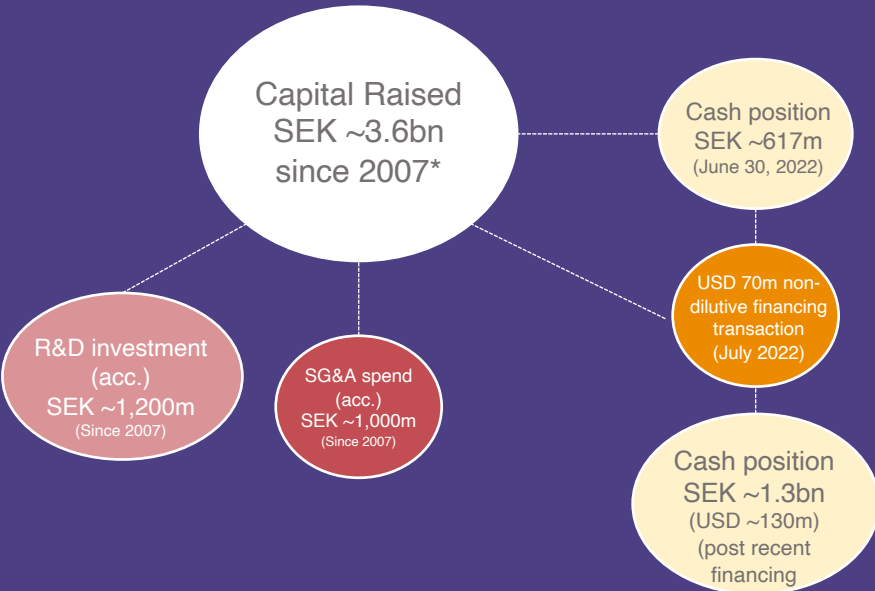
FDA

Orphan drug designations

- Imlifidase for the prevention of antibody-mediated organ rejection in solid organ transplantation (2015)
- Imlifidase for the treatment of Guillain-Barré Syndrome (2018)
- Imlifidase for the treatment of the rare and acute disease anti-GBM (2018)

Hansa Biopharma is financed into 2024

\$70 million non-dilutive financing transaction announced in July 2022 to support the continued development of Hansa's antibody-cleaving enzyme technology platform



*Including SEK ~750m from NovaQuest financing agreement & SEK ~100m upfront payments from Sarepta

Mid-term financial priorities

Our key financial priorities over the coming years will be focused on ensuring a successful European launch of Idefirix[®], while targeting mid-term product profitability

Hansa is fully financed into 2024 and we expect to use our current cash position to:

SEK ~1.3bn

(USD ~130m)

in cash and short-term investments
post recent financing
(July 2022)



Fund the launch and commercial expansion of Idefirix[®]
in kidney transplantation across Europe

Complete our EU post-approval commitments and patient enrollment in our ConfldeS study as well as advance our our long-term follow-up study to the five-year data readout in 2023

Advance our ongoing phase 2 programs in AMR and GBS and initiate a phase 3 clinical program in anti-GBM

Complete the preclinical program for our lead molecule from our next generation enzymes for repeat dosing (“NiceR”) and advance our initiatives in our other indications such as gene therapy and oncology

Fund working capital and general corporate purposes

An exciting journey ahead!

✔ This is just the beginning!

- ✔ Clinical validation
- ✔ External validation
- ✔ Regulatory validation
- ✔ Validated manufacturing
- ✔ Strong IPR
- ✔ Exciting pipeline
- ✔ Strong team

Key milestones to be achieved

- Expand Idefix[®] label in transplantation and in other solid organs
- Obtain regulatory approval in anti-GBM, GBS and AMR
- Demonstrate PoC in our next gen enzymes (NiceR)
- Expand partnerships in gene therapy and oncology
- Advance clinical studies with imlifidase as pre-treatment in Limb-Girdle, Duchenne and Pompe Disease therapies with Sarepta and AskBio
- Show PoC in new indications such as oncology
- Advance combination treatment into the clinic with argenx to potentially enable new therapeutics in transplantation and autoimmune diseases

Our future

Hansa Biopharma is a recognized global leader in rare diseases across multiple broad therapeutic areas with several market leading products and a highly valuable pipeline of late stage drug candidates



Stock images

Imlifidase in kidney transplantation



Idefix® (imlifidase) has received conditional approval in the European Union



Highly sensitized patients that are likely to be transplanted with a compatible donor

Highly sensitized patients unlikely to be transplanted under available KAS, including prioritization programs

Idefix® is indicated for

desensitization treatment of highly sensitized adult kidney transplant patients with positive crossmatch against an available deceased donor.

The use of Idefix® should be reserved for patients unlikely to be transplanted under the available kidney allocation system including prioritization programs for highly sensitized patients

Potential patients

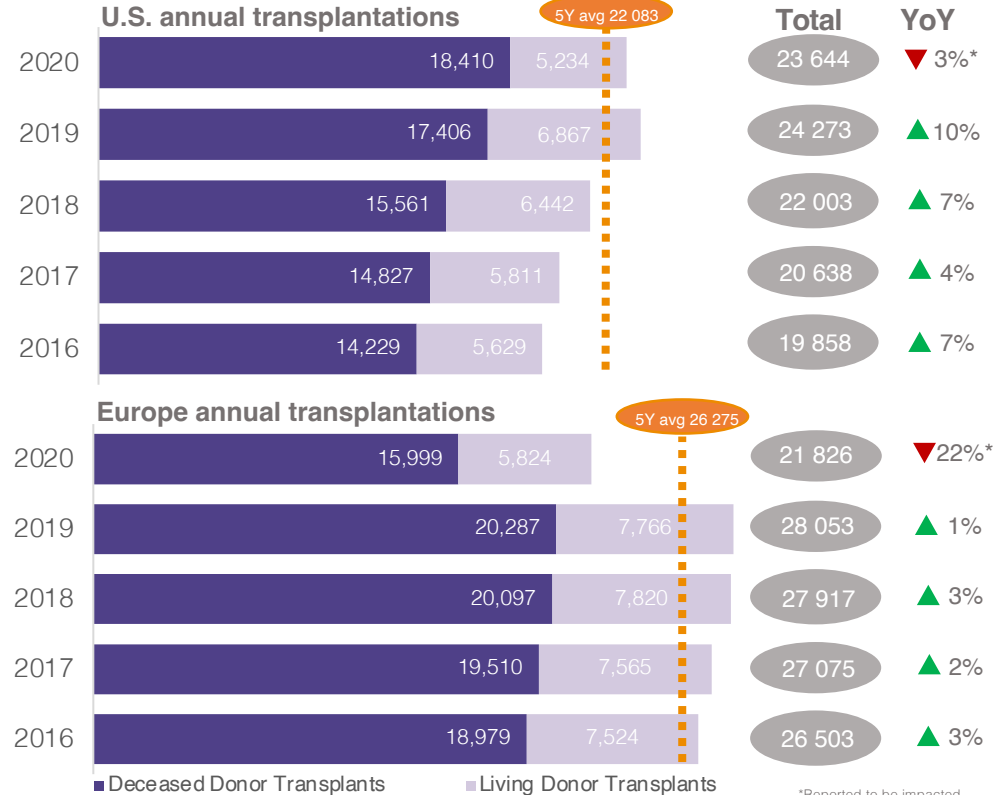
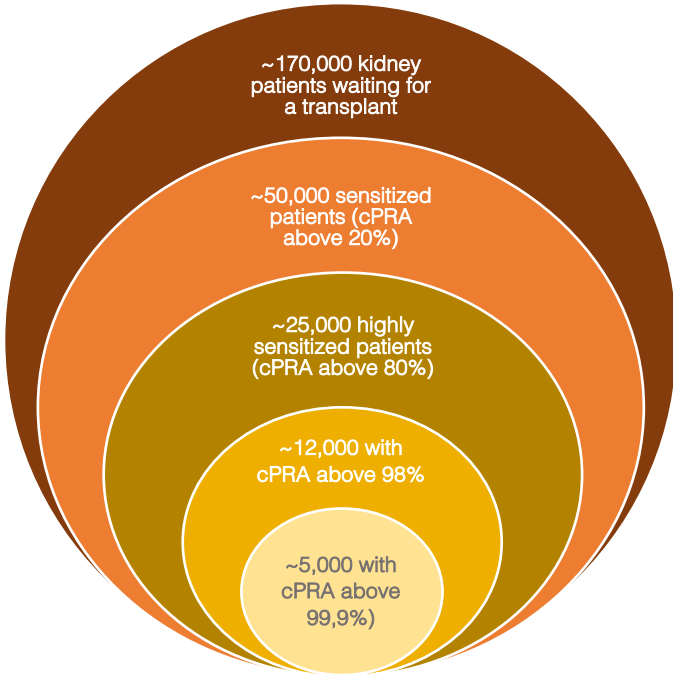


The kidney transplantation landscape in Europe and the U.S.

Up to 15% of patients waiting for a new kidney are highly sensitized

~50,000 transplants done annually in the U.S. and Europe

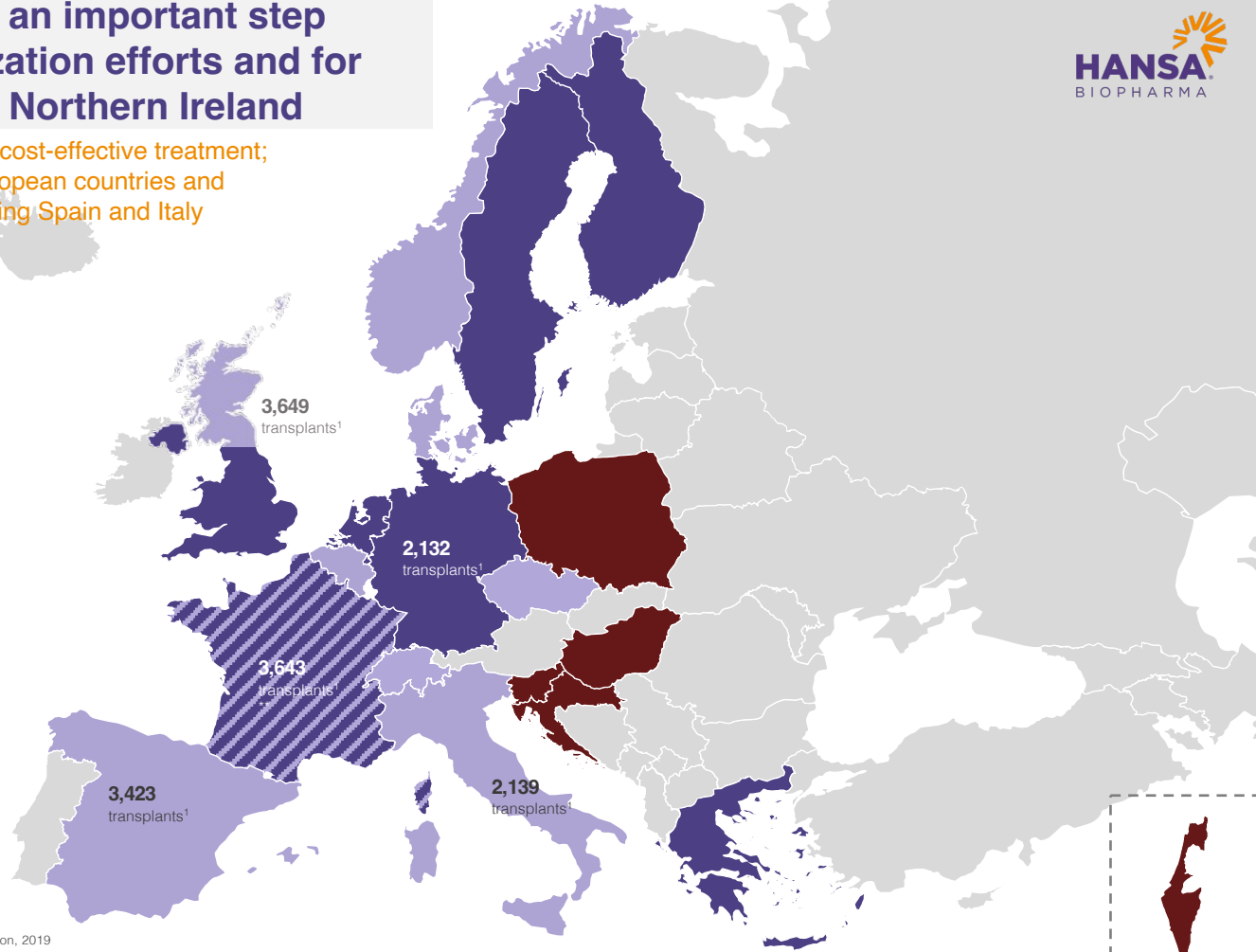
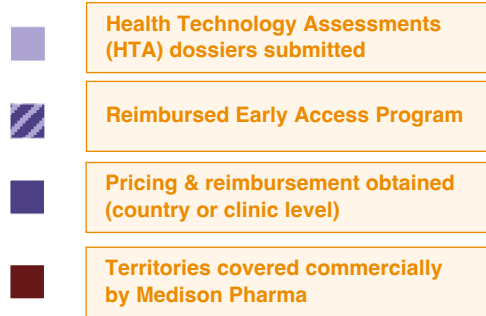
Breakdown of the kidney transplant waitlist in U.S. and EU



*Reported to be impacted by the COVID-19 pandemic

Recommendation by NICE is an important step forward for our commercialization efforts and for patients in England, Wales & Northern Ireland

NICE considers Idefirix® to be a clinically- and cost-effective treatment; Market access has now been secured in 7 European countries and procedures are ongoing in 11 countries, including Spain and Italy



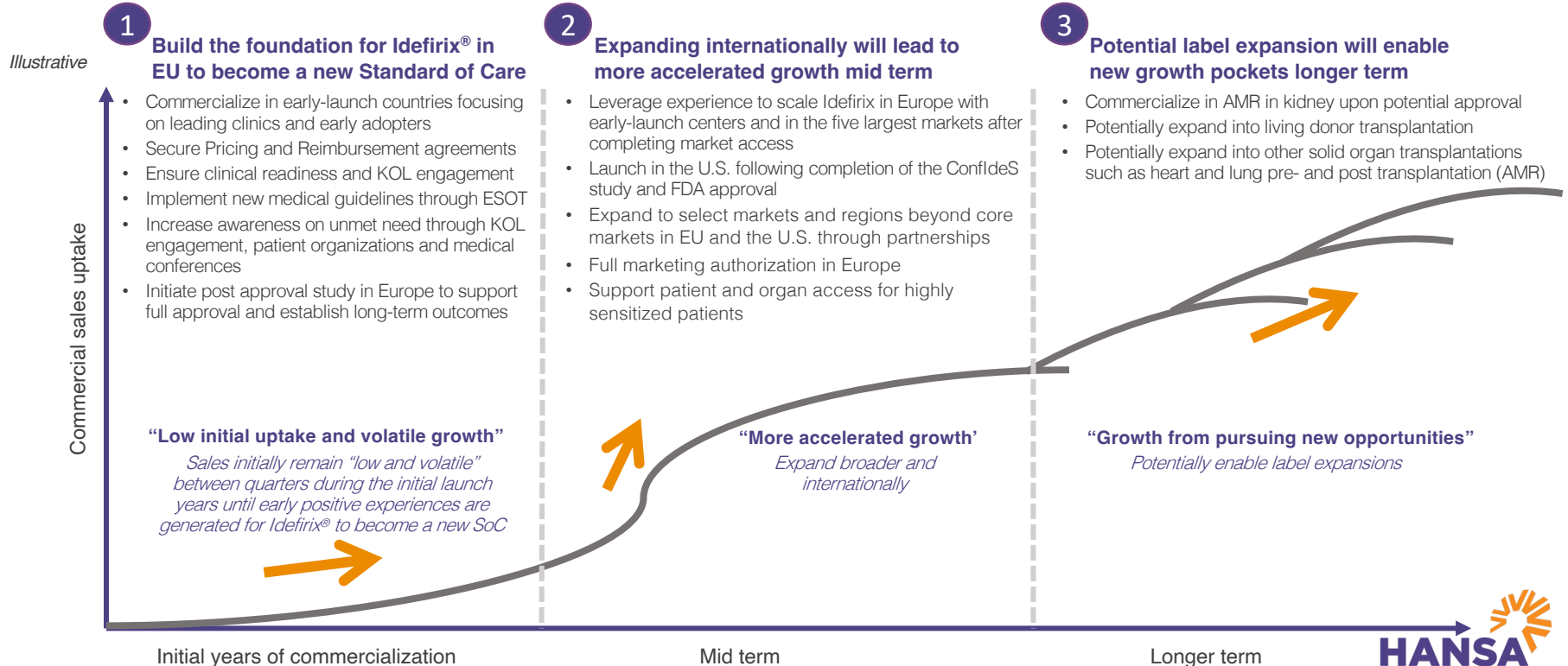
¹Annual kidney transplantations 2019 (pre-Corona)

²Transplantation data is from Global Observatory on Donation and Transplantation, 2019

³Pricing & reimbursement obtained in France on an early access basis

Our center focused and sequenced launch process will help build the foundation for Idefirix® to become a new Standard of Care in transplantation

Idefirix® is the first and only approved treatment in Europe for desensitization treatment of highly sensitized kidney transplant patients. The long-term market uptake is highly dependent on successful early experiences in key early adopter centers



Approximately 10-15% of patients on wait list are highly sensitized

Highly sensitized patients are difficult to match with an available kidney

Causes of sensitization include



Pregnancy



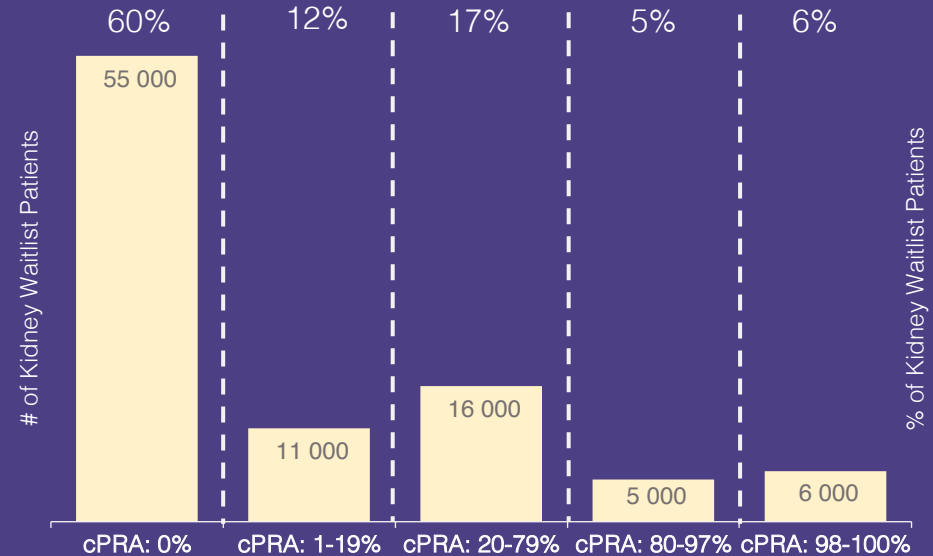
Blood transfusion



Previous transplantations

- Calculated Panel Reactive Antibodies (cPRA) is a measure for HLA-sensitization
- Inability to match or effectively desensitize patients remains a barrier for transplantation in highly sensitized patients
- Allocation Systems such as KAS and Eurotransplant rely on cPRA score to characterize patients for transplant

US Kidney Waitlist Patients by cPRA



Source: Organ Procurement and Transplant Network

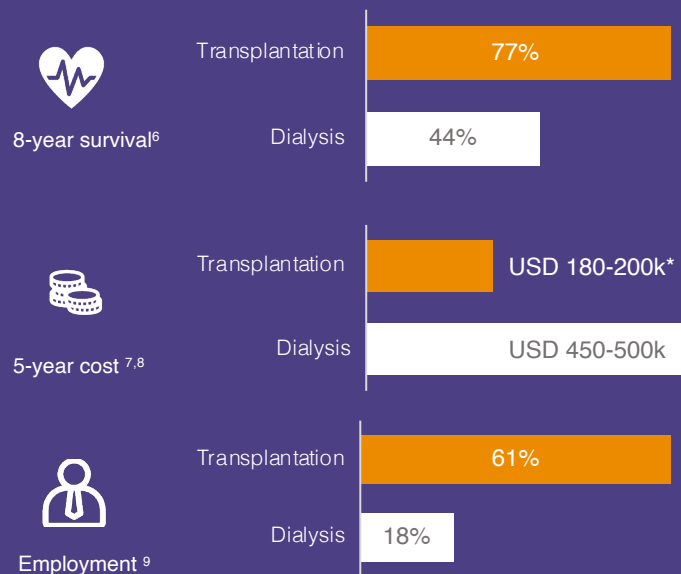
Transplantation leads to better outcomes

Saves lives, reduce costs and increase quality of life, incl. gains for the society

Several complications and risks with dialysis

- Undergoing dialysis treatment is associated with many complications and side effects incl. cardiovascular diseases¹. In the long term, patients may also eventually lose access to dialysis as a result of failed ports, bad veins, and other factors²
- In general, patients on the kidney transplant waiting list and who are on dialysis have a lower quality of life than non-dialysed patients or patients who have been transplanted³
- First study in Europe on labor market outcomes demonstrates societal gains of enabling transplantation with three times as many transplant patients employed compared to dialysis patients.
- Lastly, extended dialysis is also a high-risk factor for removal from the transplant wait list⁶

Better outcomes for transplantation patients



¹ Cozzolino et al., 2018

² Sinnakirouchenan and Holley, 2011 Shenoy, 2017

³ Wyld et al., 2012

⁴ Jarl et al. Transplantation, 2018, 102:1375-1381

⁵ NHS blood and transplant, 2018.

⁶ Orandi et al. N Engl J Med 2016;374:940-50

⁷ www.usrds.org

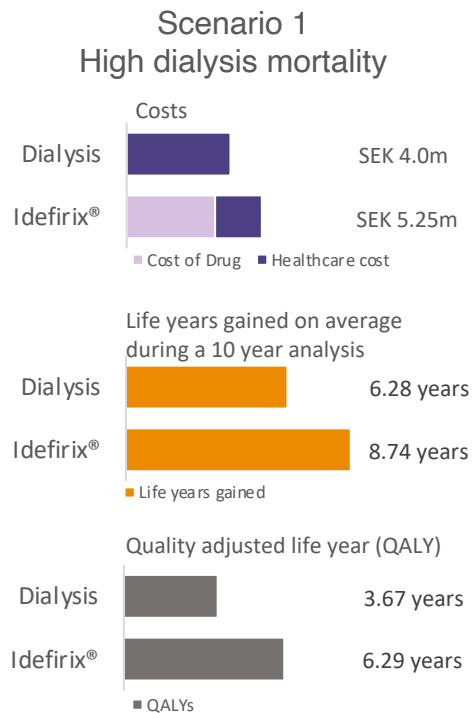
⁸ Shehata et al, Transfus Med Rev 201, 24 Suppl 1: S7-S27

⁹ Jarl et al. Transplantation, 2018, 102:1375-1381

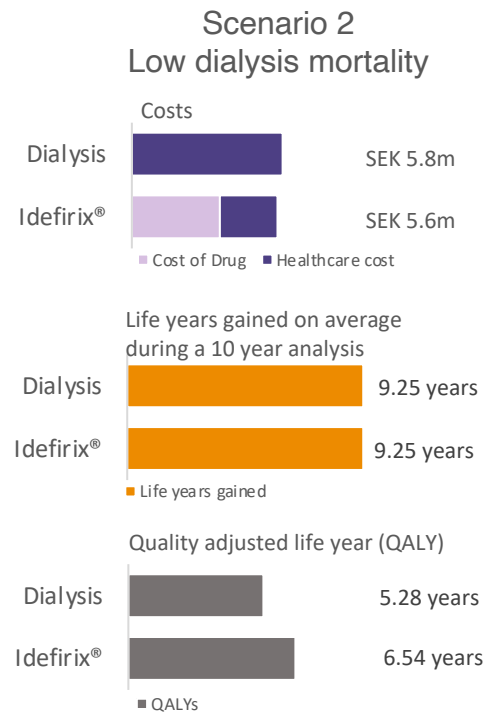
*Cost of kidney transplantation and 5 years of immuno-suppression treatment^{6,7}

First HTA report (TLV) published in Sweden favourable to the use of Idefirix® in highly sensitized patients incompatible to a deceased donor

Two cost-effectiveness scenarios presented – both within the accepted threshold for costs related to new drugs
 One scenario even concluding Idefirix treatment would lead to an overall cost saving – rare for orphan drugs



Costs per quality adjusted life year (QALY)
 SEK 460k (EUR 45k)



Costs per quality adjusted life year (QALY)
 SEK -170k (EUR -17k)

Scenario 2 supports Idefirix® as a cost saving drug

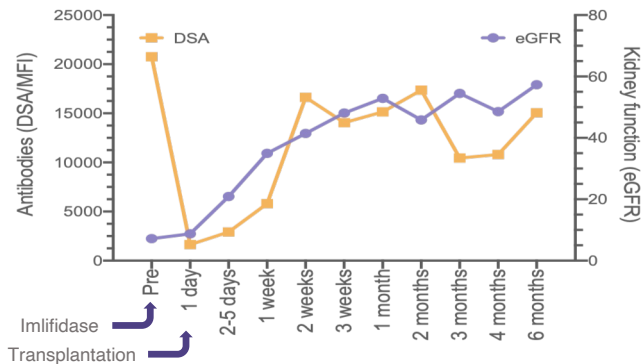
Completed and ongoing studies in kidney transplantation



Imlifidase enabled kidney transplantation in 46 highly sensitized patients during clinical trials

Pooled analysis from four Phase 2 trials

- Analysis included 46 patients
 - 50% had a cPRA of 100% (Average 99%)
 - 85% were crossmatch positive
 - 70% were retransplanted
- Donor Specific Antibody (DSA) levels rapidly decreased and all crossmatches were converted to negative, thus enabling transplantation in all patients
- At study completion, all patients alive and graft survival at 94% six months post transplantation



Study design of our four Phase 2 trials

- | | | |
|---------------------|-----------------------|---|
| Study 02
Phase 2 | Subjects | 8 patients |
| | Design | Single-center, single-arm, open-label |
| | Main objective | Efficacy defined as Imlifidase dosing scheme resulting in HLA antibody levels acceptable for transplantation, within 24 hours |
| Study 03
Phase 2 | Subjects | 10 patients |
| | Design | Single-center, single-arm, open-label, no prior desensitization |
| | Main objective | Safety in the transplantation setting and efficacy defined as HLA antibody levels acceptable for transplantation |
| Study 04
Phase 2 | Subjects | 17 patients |
| | Design | Investigator initiated, single-center, single-arm, open-label. All patients had prior desensitization with IVIG and/or PLEX |
| | Main objective | Safety in combination with Cedars Sinai's "standard protocol" for desensitization of highly sensitized patient |
| Study 06
Phase 2 | Subjects | 18 patients |
| | Design | Multicenter, multinational, single-arm, open-label |
| | Main objective | Efficacy in creating a negative crossmatch test |

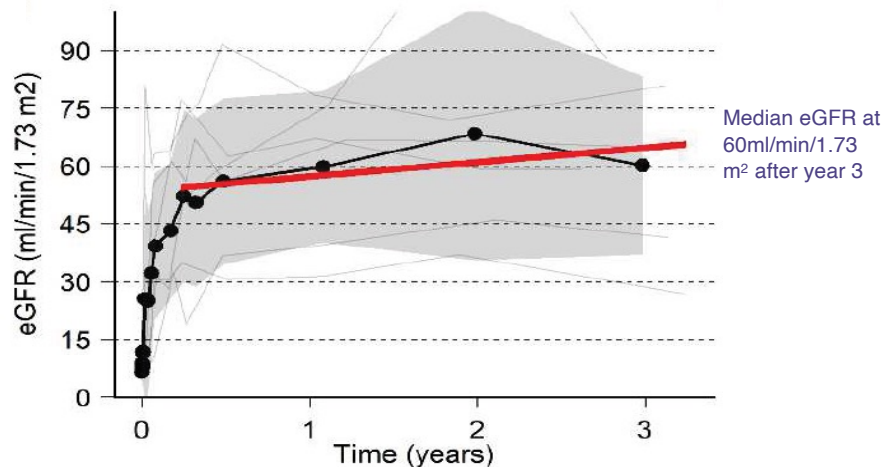
3-year follow up demonstrate graft survival of 84% after imlifidase treatment and transplantation

Data accepted for publication in the American Journal of Transplantation¹ [Link AJT article](#)
30 patients participating in follow-up study at year three

AMR frequency in line with other desensitization protocols

- Three-year follow-up data shows graft survival of 84% after imlifidase treatment and transplantation and a mean eGFR of 55 mL/min/1.73 m² (61 ml/min/m² for those without AMR)
- For a subgroup of patients (n=13) with cPRA of $\geq 99.9\%$ graft survival was 92% and improved kidney function for patients with a mean eGFR at 60ml/min/1.73 m² after year three
- 38% of the patients experienced active antibody mediated rejection episodes (AMR) within the first six months, which compares with 25-60% of patients in the literature for highly sensitized patients²
- Only two AMR episodes were reported beyond the first 6 months. All AMRs were treated with standard therapies and no graft losses were attributed to AMR
- Patient survival 90% (three deaths unrelated to imlifidase)
- Long-term safety profile indicates no increase in the rates of infection or malignancy
- Next milestone expected in 2023 on the 5-year follow-up data

Improved kidney function for patients with cPRA $\geq 99.9\%$



¹ American Journal of Transplantation - Outcomes at 3 years post-transplant in imlifidase-desensitized kidney transplant patients (AJT16754)

Link to AJT article <https://onlinelibrary.wiley.com/doi/epdf/10.1111/ajt.16754>

² Vo et al. 2013; Colvin 2007; Gloor et al. 2008; Haas et al. 2014; Jordan et al. 2010; Lefaucheur et al. 2010; Solez et al. 2007; Riella et al. 2014)

U.S. ConfideS study: First patient enrolled Dec'21; BLA submission expected H1 2024

U.S. trial design

64 highly sensitized kidney patients with the highest unmet medical need

- Patients with a cPRA score of $\geq 99.9\%$ will be enrolled
- First patients enrolled at Columbia University, NYC
- 22 patients enrolled across ten sites end of Q2 2022

1:1 Randomization

- When a donor organ becomes available and a positive crossmatch with the intended recipient indicates that the organ is not compatible, the patient will be randomized to either imlifidase or to a control arm, where patients either remain waitlisted for a match or receive experimental desensitization treatment*

Primary endpoint

- Mean estimated glomerular filtration rate (eGFR) "kidney function" at 12 months.
- For randomized patients who do not undergo transplantation, lose their graft or die before 12 months, eGFR will be set to zero, consistent with kidney failure

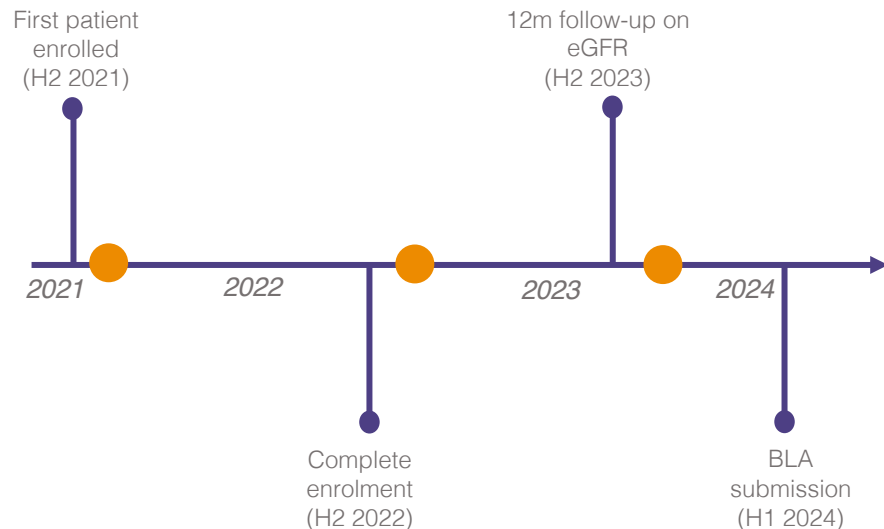
Secondary endpoint

- Patient survival at 12 months

Up to 15 leading transplantation centers in the U.S. will be engaged in the study

- Robert A. Montgomery, M.D. Professor of Surgery and Director, NYU Langone Transplant Institute, NYC is appointed to be the principal investigator

Timeline



*Experimental desensitization treatment can include any combination of plasma exchange (PLEX), intravenous IVIg, anti-CD20 antibody, and eculizumab. Link to the full protocol at [ClinicalTrials.gov](https://clinicaltrials.gov)

U.S. kidney transplantation landscape

Our ConfideS study is currently enrolling patients across ten leading transplantation centers across seven states covering ~10% of annual kidney transplants in the U.S.; Aim to have up to 15 centers recruiting patients

>23,000¹ annual kidney transplantations

~71%¹ deceased donor

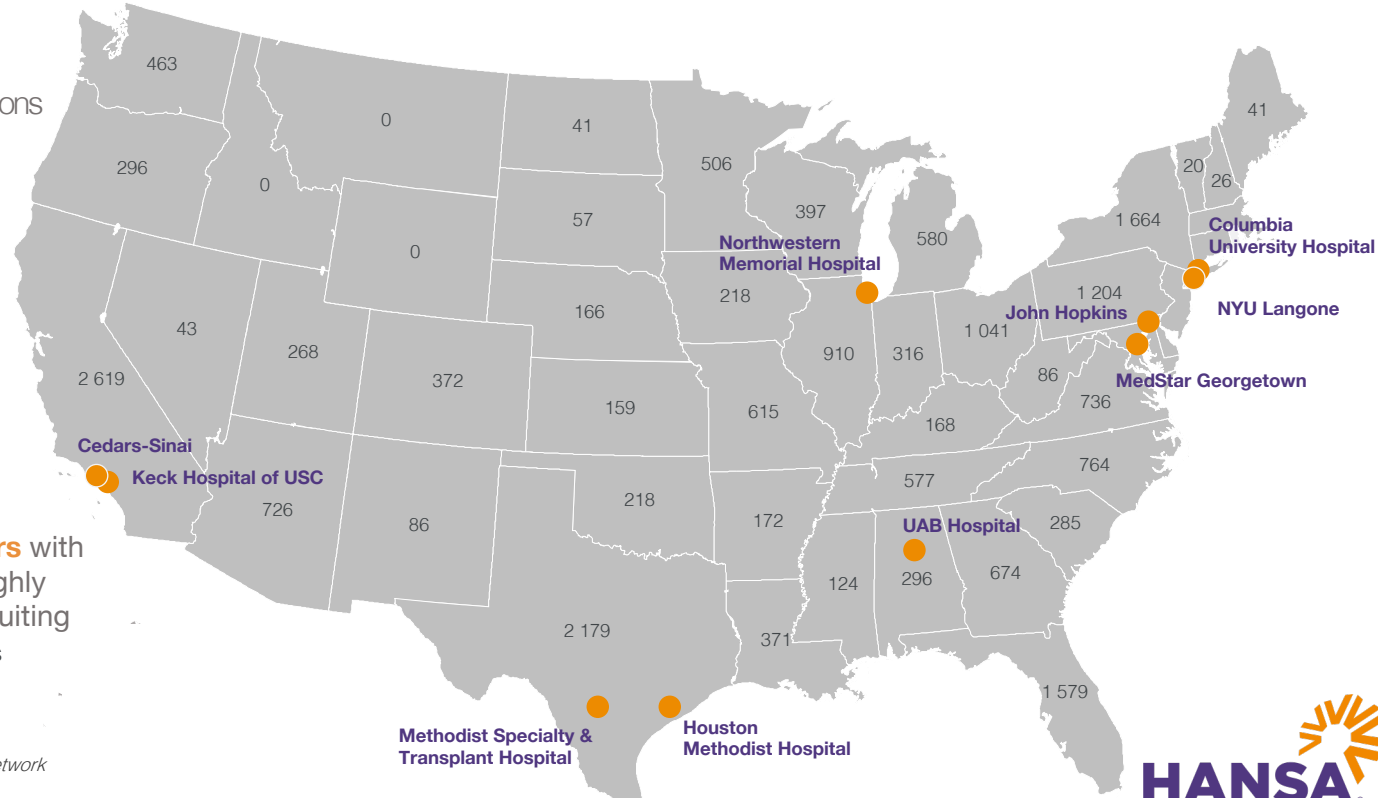
~90,000² waiting for a kidney transplant

10-15%³ of waitlisted patients are highly sensitized

Ten leading transplantation centers with experience in desensitization and highly sensitized patients are currently recruiting

2,542¹ combined annual kidney transplants

334¹ highly sensitized (>80% cPRA)



¹2019 data from Organ Procurement & Transplantation Network

²United Network for Organ Sharing

³EDQM. (2020). International figures on donation and Transplantation 2019 and SRTT Database and individual assessments of allocation systems

First patient treated in post-authorization efficacy study (PAES) of Idefirix® (imlifidase) in highly sensitized kidney transplant patients

The study will provide further important insights regarding Idefirix® desensitization treatment of highly sensitized kidney transplant patients

An open-label Phase 3 study in 50 patients

- First patient was treated by Dr. Oriol Bestard, Chair of Nephrology and Kidney Transplantation at Vall d'Hebron University Hospital in Barcelona
- Study will enroll 50 highly sensitized patients with positive crossmatch against an available deceased donor across multiple countries and centers in Europe
- The study is an obligation under the conditional marketing authorization for Idefirix® granted by EMA in August 2020, in order to complete a full marketing authorization in the EU
- The aim will be to confirm the long-term efficacy and safety of Idefirix® with the primary objective to determine the one-year graft failure-free survival of the Idefirix® treated and transplanted patients.
- In addition, a total of 50-100 patients undergoing compatible kidney transplantation at the participating centers will be included and serve as a non-comparative concurrent reference cohort, with no formal comparison, to contextualize the one-year graft failure-free survival of the Idefirix® treated patients



Study to assess imlifidase in combination to optimize patient outcome

in highly sensitized patients with donor specific antibodies (DSA) rebound and antibody mediated kidney transplant rejection

Trial design (ClinicalTrials.gov ID: NCT05049850)

The study is designed to assess if imlifidase in combination with bortezomib¹, belatacept², rituximab³ and IVIg⁴ can suppress donor specific antibodies (DSA) and the occurrence of antibody-mediated rejection (AMR) in transplant patients with a positive crossmatch towards their living donor.

Open label, single arm study

- Imlifidase is administered within the 24-hour prior to a living donor transplantation

Primary endpoint

- Proportion of patients with DSA rebound (up to 3 months after transplantation)
- Rebound of DSA may cause AMR and is thus a risk for graft loss

Secondary endpoint

- Proportion of patients with AMR (up to 6 months after transplantation)

The study will be run at the NYU Langone Transplant Institute and is expected to commence in 2022

¹ bortezomib, a proteasome inhibitor which has activity against mature plasma cells, the source of DSA

² belatacept, a fusion protein which is crucial in blocking T-cell co-stimulation and which is effective in reducing de novo DSA generation in humans

³ rituximab, an anti-CD20 monoclonal antibody that targets B-cells and which is an immunomodulatory agent

⁴ intravenous immunoglobulin (IVIg) which is commonly used in desensitization regimens and for the treatment of AMR

Link to the full protocol at [ClinicalTrials.gov](https://clinicaltrials.gov)



Study 01 Phase 1

The 01 study results

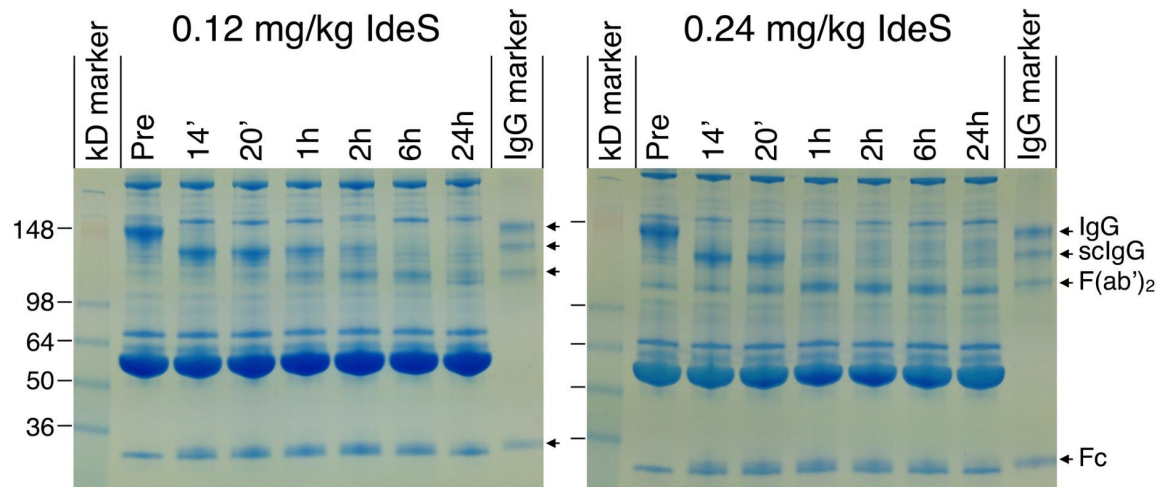
Data showed complete removal of IgG and a good tolerability profile

Efficacy

- ✓ Rapid degradation of IgG in serum in subjects dosed with 0.12 and 0.24 mg/kg imlifidase. Imlifidase had full effect within 6 hours. The entire IgG pool was converted into F(ab')₂ and Fc-fragments. Maximal effect was accomplished 2-6 hours after dosing.

Safety

- ✓ Newly synthesized intact IgG was clearly detectable in all subjects after 1-2 weeks after dosing. After 3 weeks the level of intact IgG constituted the main IgG fraction in serum



CLINICALTRIALS.GOV ID

NCT01802697 (2013/2014)

SUBJECTS

29 (20 active plus 9 placebo) healthy subjects (Sweden)

DOSES/FOLLOW UP TIME

The starting dose was 0.01 mg/kg BW and the highest dose group received 0.24 mg/kg BW

MAIN OBJECTIVES

- The objectives were to assess safety, efficacy in IgG cleavage, pharmacokinetics and immunogenicity of imlifidase following intravenous administration

STUDY DESIGN

- Randomized placebo-controlled dose-escalation study with 29 (20 active plus 9 placebo) healthy subjects

STATUS

Completed

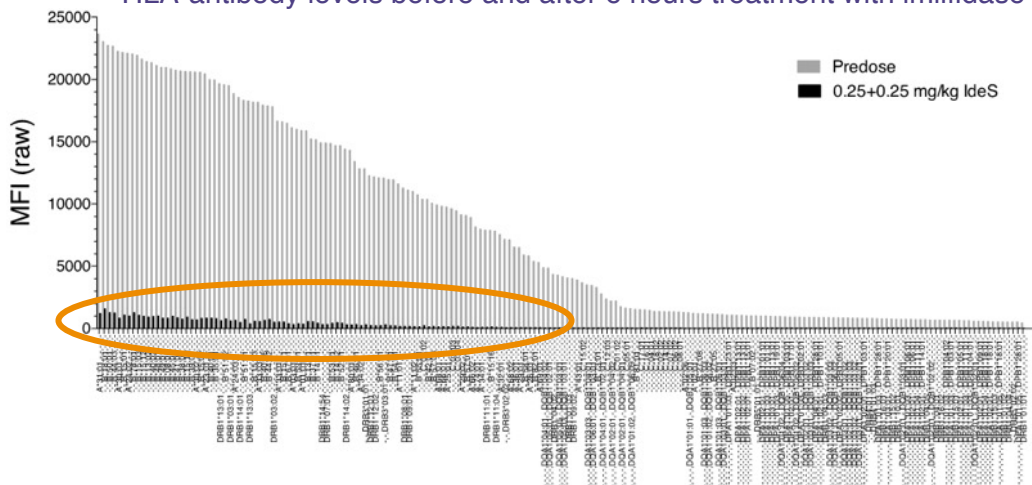
- The 01 study showed that Imlifidase was considered safe to use
48

The 02 study results

Data showed that 1-2 doses of imlifidase at 0.25 mg/kg BW resulted in HLA antibody levels acceptable for transplantation¹

- ✓ Imlifidase is well tolerated in patients with chronic kidney disease
- ✓ Efficacy results strongly support further development in the patient population
- ✓ The first HLA-incompatible transplantation ever after desensitization with imlifidase was performed in one of these patients (2014)

HLA-antibody levels before and after 6 hours treatment with imlifidase



¹ Lorant et al (2018) American Journal of Transplantation (2018)

- Efficacy defined as Imlifidase dosing scheme resulting in HLA antibody levels acceptable for transplantation, within 24 hours from dosing
- Safety

- Single-center, Single arm with ascending doses, open-label
- Transplantation not part of protocol

- Primary efficacy endpoint reached
- Safe and well tolerated

Study 03 Phase 2

The 03 study proved safety and efficacy

HLA antibodies at acceptable levels; enabling transplantation in all patients

CLINICALTRIALS.GOV ID

NCT02475551

SUBJECTS

10 Patients (Sweden)

DOSES/FOLLOW UP TIME

0.25 and 0.50 mg/kg during 180 days

MAIN OBJECTIVES

- Safety in the transplantation setting
- Efficacy defined as HLA antibody levels acceptable for transplantation

STUDY DESIGN

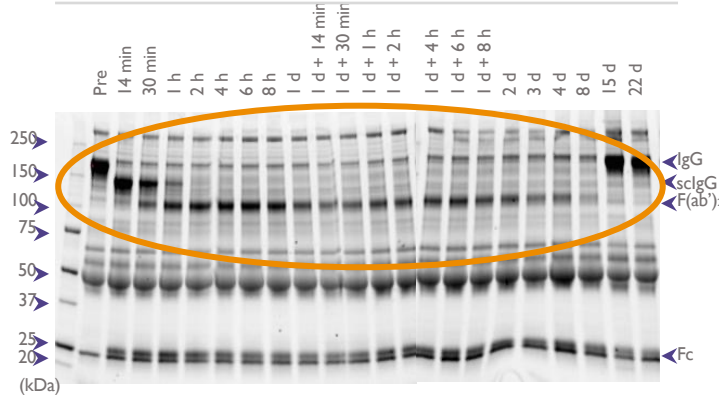
- Single-center, single-arm, open-label, no prior desensitization
- Similar design as 13-HMedIdeS-02 but transplantation part of protocol
- In deceased and living donors

STATUS

Completed

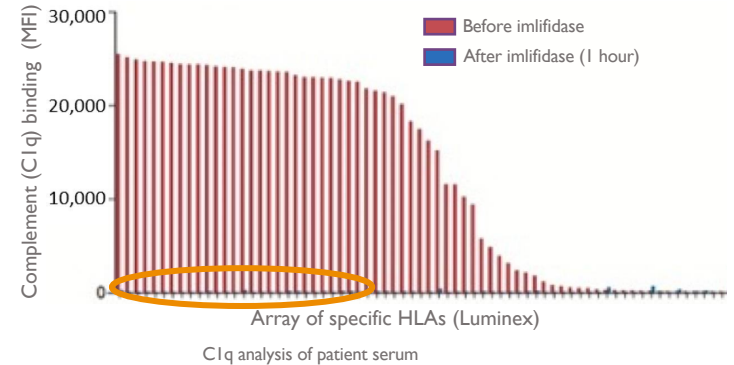
- Proved safety and efficacy with HLA antibodies at acceptable levels; enabling transplantation in all patients

Analysis of IgG in patient serum before and after imlifidase treatment



SDS-PAGE analysis of patient serum

Analysis of complement binding HLA antibodies before and after imlifidase



CIq analysis of patient serum

Protocol

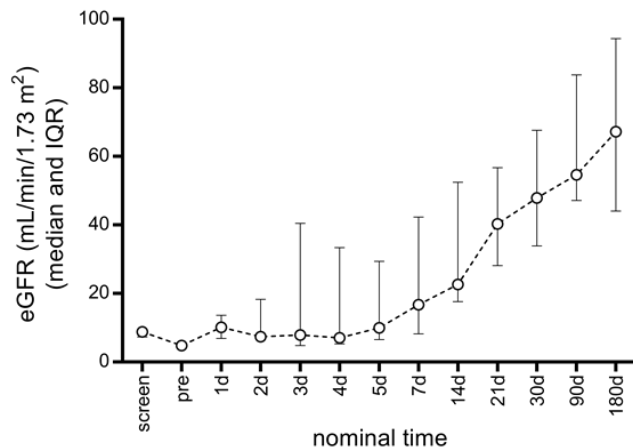


Study 04 Phase 2

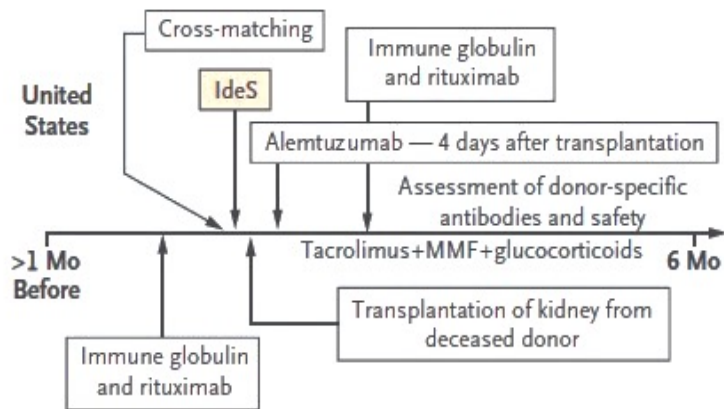
The 04 study results

Study proved safety and efficacy with Cedar Sinai's standard protocol (rituximab and IVIg)

Graft function (eGFR) post six months



Cedar's desensitization protocol in combination with imlifidase



CLINICALTRIALS.GOV ID

NCT024226684

SUBJECTS

17 Patients (US)

DOSES/FOLLOW UP TIME

0.24 mg/kg 180 days

MAIN OBJECTIVES

- Safety in combination with Cedars Sinai's "standard protocol" for desensitization of highly sensitized patients
- Efficacy in preventing AMR

STUDY DESIGN

- Investigator initiated study
- Investigator sponsored IND
- Imlifidase to desensitize patients previously treated with rituximab and IVIg
- Deceased donors only

STATUS

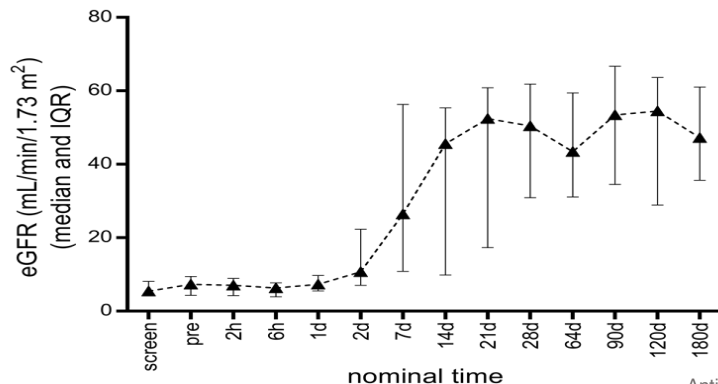
Completed

Study 06 Phase 2

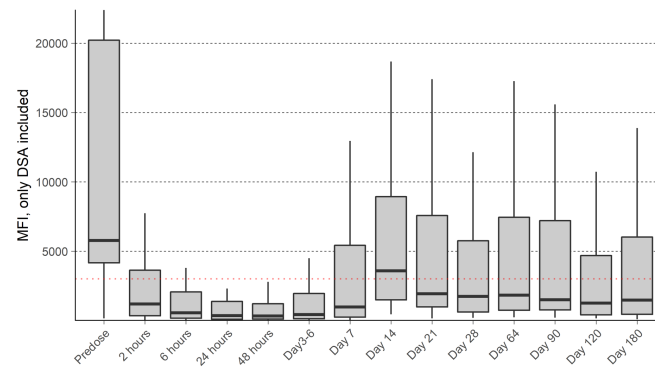
The 06 study results

Study showed proved safety and efficacy in making highly sensitized patients eligible for kidney transplantation

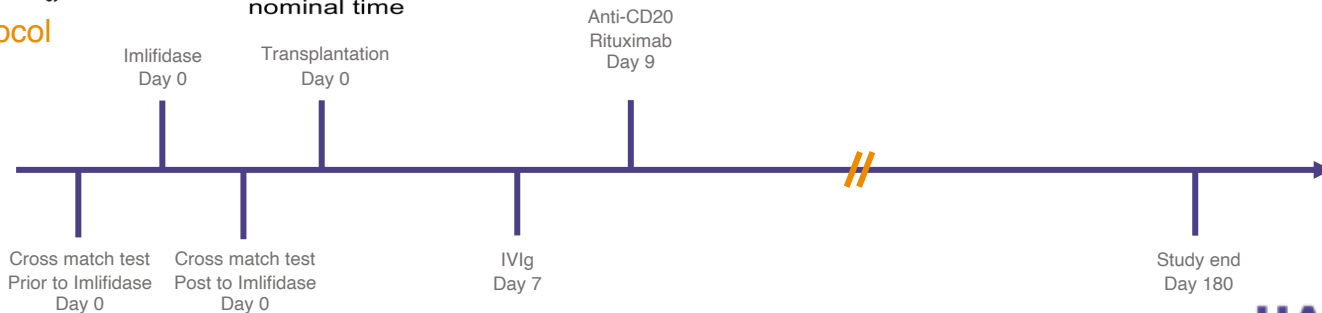
Graft function (eGFR) post imlifidase



DSA level pre-dose and post imlifidase



Protocol



CLINICALTRIALS.GOV ID

NCT02790437

SUBJECTS

18 Patients (US+Sweden+France)
19 safety set, 18 efficacy set

DOSES/FOLLOW UP TIME

0.25 mg/kg 180 days

MAIN OBJECTIVES

- Efficacy in creating a negative crossmatch test

STUDY DESIGN

- Multicenter, multinational, single-arm, open-label Included patients who may have had prior unsuccessful desensitization or patients in whom it was unlikely to be effective







STATUS

Completed

Jordan SC, et al. (2019).

Results from the international phase II study on the safety and efficacy of imlifidase in highly-sensitized kidney transplant patients. Abstract presented at ATC.

Completed studies with imlifidase in transplantation

STUDY	SUBJECTS/ COUNTRY	STUDY DESIGN	PRIMARY ENDPOINT	SECONDARY ENDPOINTS	STATUS/ PUBLICATION
Study 01 Phase 1	29 subjects 	<ul style="list-style-type: none"> Randomized placebo-controlled dose-escalation study with 29 (20 active plus 9 placebo) healthy subjects 	<ul style="list-style-type: none"> Safety and tolerability 	<ul style="list-style-type: none"> Efficacy in IgG cleavage, the pharmacokinetics (PK) and immunogenicity of imlifidase 	Complete PLOS ONE (2015) ¹
Study 02 Phase 2	8 subjects 	<ul style="list-style-type: none"> Single-center, single-arm, open-label 	<ul style="list-style-type: none"> Dosing resulting in HLA-antibody reduction (MFI<1100) 	<ul style="list-style-type: none"> Efficacy: HLA antibody reduction acceptable for transplantation (MFI <1100 as measured in SAB assay) 	Complete Lorant et al (2018) American Journal of Transplantation ²
Study 03 Phase 2	10 subjects 	<ul style="list-style-type: none"> Single-center, single-arm, open-label No prior desensitization 	<ul style="list-style-type: none"> Safety: AEs, clinical laboratory tests, vital signs, ECGs 	<ul style="list-style-type: none"> Efficacy: HLA antibody reduction acceptable for transplantation (MFI <1100 as measured in SAB assay) 	Complete The New England Journal of Medicine (2017) ³
Study 04 Phase 2	17 subjects 	<ul style="list-style-type: none"> Investigator initiated study, Single-center, single-arm, open-label All patients had prior desensitization with IVIG and/or plasmapheresis 	<ul style="list-style-type: none"> Assessment of efficacy in eliminating DSAs in DSA and flow cytometry positive, highly sensitized patients Assessment of safety Assessment of efficacy/kidney function 	<ul style="list-style-type: none"> Serum creatinine (0-6 months) Proteinuria (0-6 months) DSA at multiple timepoints posttransplant (day 0, D30, D90, D180) 	Complete The New England Journal of Medicine (2017) ³
Study 06 "Highdes" Phase 2	18 subjects 	<ul style="list-style-type: none"> Multicenter, multinational, single-arm, open-label Included pts who may have had prior unsuccessful desensitization or pts in whom it was unlikely to be effective 	<ul style="list-style-type: none"> Crossmatch conversion in DSA+ patients who have a positive XM test to their available LD or DD 	<ul style="list-style-type: none"> DSA reduction at multiple timepoints (2, 6, 24, 48 h after imlifidase) Time to create negative CDC XM test and/or flow cytometry (FACS) XM test Safety 	Complete Annals of Surgery (Lonze et al, only New York patients) Montgomery et al ATC abstract (2019) ⁴
Long-term follow-up study	Up to 46 subjects 	<ul style="list-style-type: none"> A prospective, observational long-term follow-up study of patients treated with imlifidase prior to kidney transplantation 	<ul style="list-style-type: none"> Long-term graft survival in patients who have undergone kidney transplantation after imlifidase administration 	<ul style="list-style-type: none"> Patient survival, kidney function, comorbidity, treatments and QoL Safety DSA Immunogenicity 	Ongoing

¹ Winstedt et al., "Complete Removal of Extracellular IgG Antibodies in a Randomized Dose Escalation Phase I Study with the Bacterial Enzyme IdeS – A Novel Therapeutic Opportunity", PLOS ONE 2015, 10(7)

² Lorant et al., "Safety, immunogenicity, pharmacokinetics and efficacy of degradation of anti-HLA antibodies by IdeS (imlifidase) in chronic kidney disease patients" Am J Transplant. 2018 Nov;18(11):2752-2762

³ Jordan et al., "IgG Endopeptidase in Highly Sensitized Patients Undergoing Transplantation", N Engl J Med 2017;377:442-53.

⁴ Montgomery et al., "Safety And Efficacy Of Imlifidase In Highly-sensitized Kidney Transplant Patients: Results From A Phase 2 Study" ATC Abstract, 2019

Medical Advisory Board in kidney transplantation



Professor Stanley Jordan

(Chairman) M.D., Ph.D., Director of Kidney Transplantation and Transplant Immunology, Kidney and Pancreas Transplant Center and Director of Division of Pediatric and Adult Nephrology, Cedars-Sinai Medical Center, Los Angeles, California



Professor Robert Montgomery

M.D., Ph.D., FACS, Director at NYU Langone Transplant Institute, New York, NY, USA



Professor Christophe Legendre

M.D., Ph.D. Professor at Paris Descartes University and Head of the Adult Nephrology and Transplantation unit at Necker Hospital in Paris.

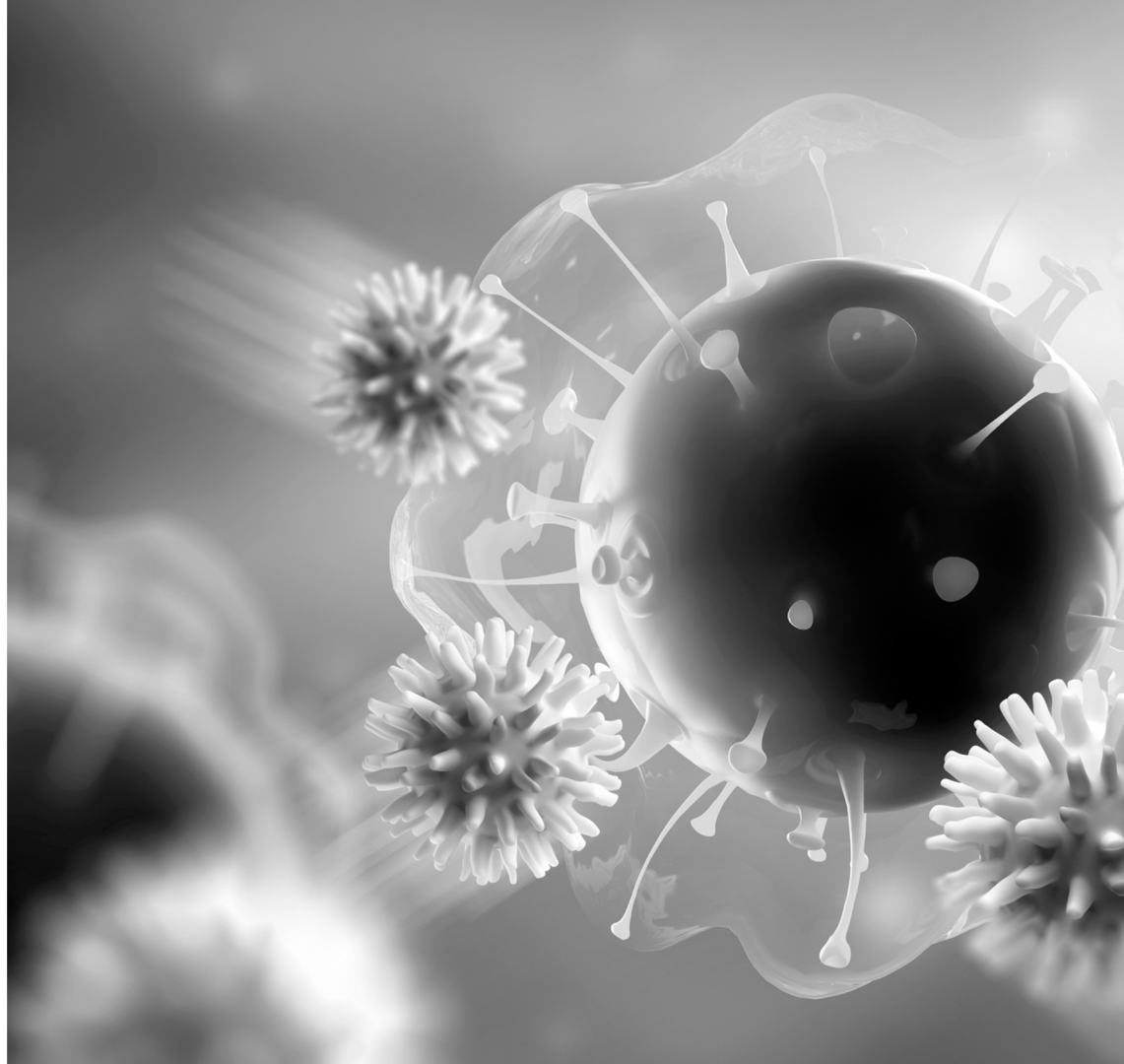


Professor Kathryn Wood

Ph.D. Fellow of the Academy of Medical Sciences, Professor of Immunology in the Nuffield Department of Surgical Sciences, University of Oxford, England, runs the Transplantation Research Immunology Group



Our antibody cleaving enzyme technology



Broad clinical pipeline in transplantation and auto-immune diseases

Candidate/ Program	Indication	Research/ Preclinical	Phase 1	Phase 2	Phase 3	Marketing Authorization	Marketed	Next Anticipated Milestone
Imlifidase	EU: Kidney transplantation in highly sensitized patients ^{1,2}	Completed	Completed	Completed	Planned	Completed	Marketed	EU: Additional agreements around reimbursement from H2'21
	US: Kidney transplantation in highly sensitized patients ^{1,2}	Completed	Completed	Completed	Ongoing			Completion of enrollment (64 patients) H2'22
	Anti-GBM antibody disease ³	Completed	Completed	Completed	Planned			Pivotal Phase 3 study expected to commence in 2022 (50 patients)
	Antibody mediated kidney transplant rejection (AMR)	Completed	Completed	Ongoing				First data readout H2'22
	Guillain-Barré syndrome (GBS)	Completed	Completed	Ongoing				Completion of enrollment (30 patients) H2 2022
	Pre-treatment ahead of gene therapy in Limb-Girdle (Partnered with Sarepta)	Ongoing						Preclinical phase
	Pre-treatment ahead of gene therapy in Duchenne (Partnered with Sarepta)	Ongoing						Preclinical phase
	Pre-treatment ahead of gene therapy in Pompe disease (Partnered with AskBio)	Ongoing						Preclinical phase
NiceR	Recurring treatment in autoimmune disease, transplantation and oncology	Ongoing						Completion of GLP toxicology studies in 2022
EnzE	Cancer immunotherapy	Ongoing						Research phase

¹ Results from the Phase 1 study have been published, Winstedt et al. (2015) PLOS ONE 10(7)

² Lorant et al American Journal of Transplantation and 03+04 studies (Jordan et al New England Journal of Medicine)

³ Investigator-initiated study by Mårten Segelmark, Professor at the universities in Linköping and Lund

Completed

Ongoing

Planned

Post approval study running in parallel with commercial launch

Continuous progress in our ongoing clinical Programs

Enrollment status
July 19, 2022



Antibody Mediated Rejection Phase 2 study

- 30/30 patients enrolled in the AMR phase 2 study
- Completion of enrollment expected H1 2022*
- First data read out expected in H2 2022*



- Patients enrolled
- Patients remaining

Guillain-Barré Syndrome Phase 2 study

- 18/30 patients enrolled in the GBS program
- Ten centers are active and open for recruitment
- Significant initiatives were implemented during H1 2022 to support the completion of enrollment in H2 2022*
- First data read out expected in H1 2023

Enrollment status
July 19, 2022



- Patients enrolled
- Patients remaining

Anti-GBM Phase 3 study

- FDA has accepted Hansa's Investigational New Drug (IND) application to proceed with a Phase 3 study
- The planned study will commence this year targeting 50 patients across the U.S. and Europe*



- Patients enrolled
- Patients remaining

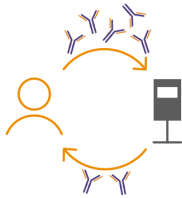
U.S. ConfIdeS Phase 3 study

- Randomized, controlled trial in highly sensitized kidney transplant patients across up to 15 centers
- 22/64 patients enrolled for randomization
 - Ten centers are active and open for recruitment
 - Completion of enrollment expected H2 2022*

Development of IgG-modulating technologies

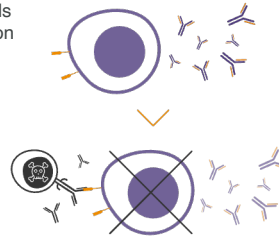
Mechanisms can be both complementary and competing

PLEX, plasmapheresis, immunoadsorption
Mechanically removes antibodies from circulation



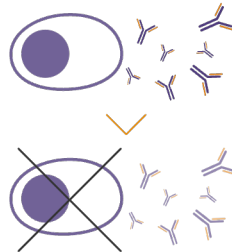
1950s

B-cell depleting mAbs
Lowering antibody levels through B-cell elimination



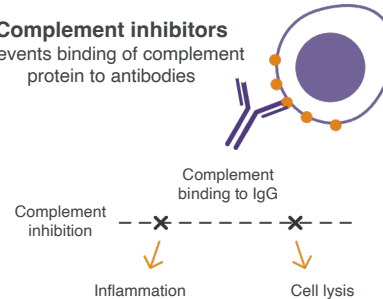
1990s

Proteasome inhibitors
Depletes antibody producing long-lived plasma cells and lowers overall immunoglobulin levels



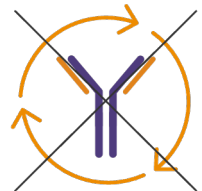
2000s

Complement inhibitors
Prevents binding of complement protein to antibodies



2010s

FcRn-inhibitors
Lowering IgG through blocking of antibody recycling

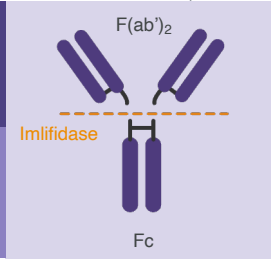


2020s

Imlifidase – IgG-cleaving enzyme

Deactivates IgG within 2-6 hours through enzymatic cleavage. IgG-free window for approximately one week

Unique mechanism-of-action is the basis for competitive advantage vs other IgG-modulating therapies

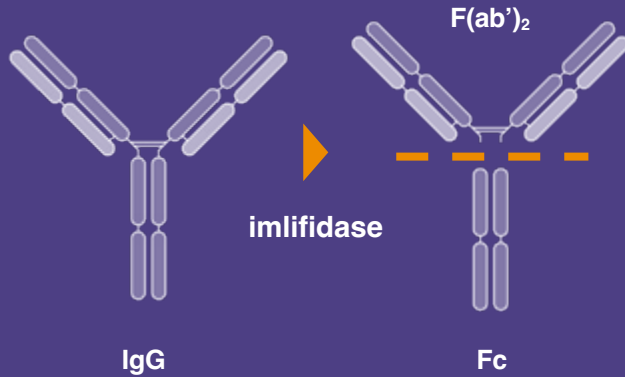


Imlifidase mode of action

Novel approach to effectively eliminate pathogenic IgG

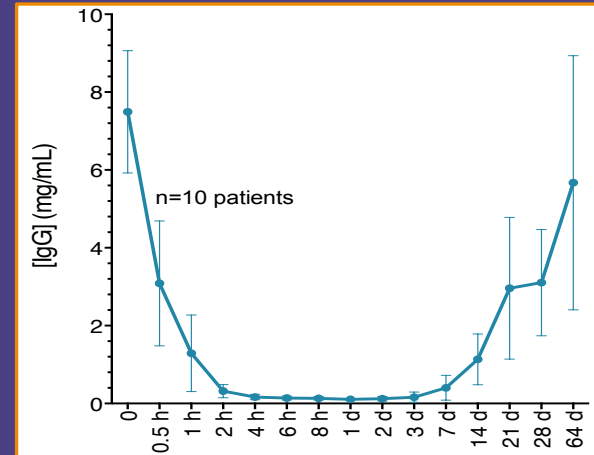
Proven mechanism of action

- Interacts with Fc-part of IgG with extremely high specificity
- Cleaves IgG at the hinge region, generating one F(ab')₂ fragment and one homo-dimeric Fc-fragment



Inactivation of IgG in human serum

- Rapid onset of action that takes down IgG below detectable level in 2-6 hours post 15 min infusion
- IgG antibody-free window for approximately one week



Our unique antibody cleaving enzyme technology may have relevance across a range of indications

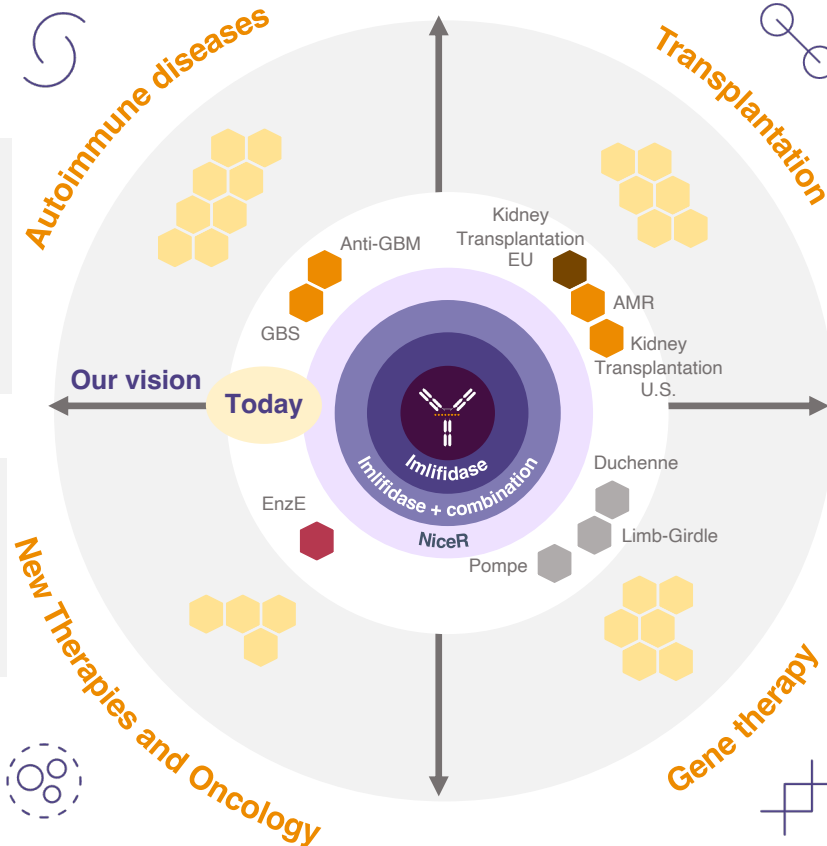
Targeting rare IgG mediated diseases

Anti-GBM paves the way for development in other autoimmune diseases

- Rapidly progressive glomerulonephritis
- Neurological disorders
- Skin and blood disorders

IgG-cleaving enzymes to enable or even potentiate cancer therapy

- Allogenic stem cell (bone marrow) transplantation (HSCT)
- Enzyme-based antibody Enhancement (EnzE)



Expanding our commercial franchises

- Regulatory approval (conditional)
- Clinical development
- Partnership (preclinical development)
- Preclinical development
- Potential indications (currently not pursued)

Shaping a new standard for desensitization will help enable new indications in transplantations

- Antibody mediated rejection (AMR) in kidney transplantation
- Other transplantation types

Exploring opportunities in gene therapy

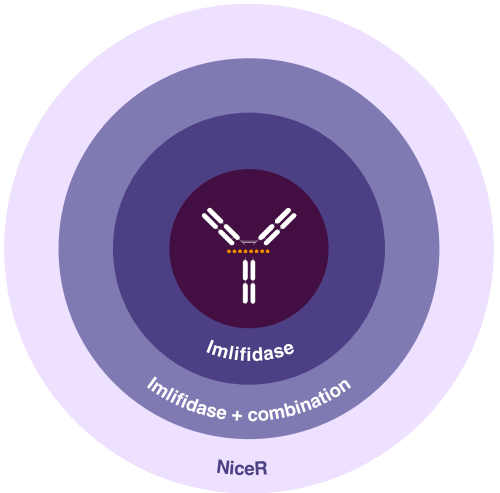
- Encouraging preclinical data published in Nature
- Validation through collaborations with Sarepta and AskBio
- Wide indication landscape beyond

The technology platform is the primary basis for achieving our vision

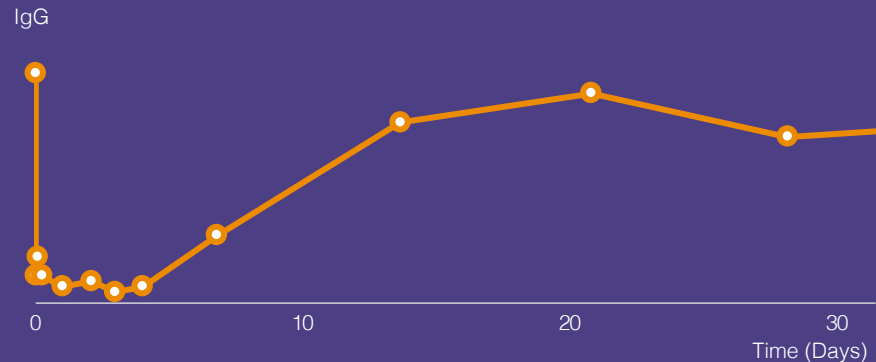
Targeting rare IgG mediated diseases and conditions

Key opportunities:

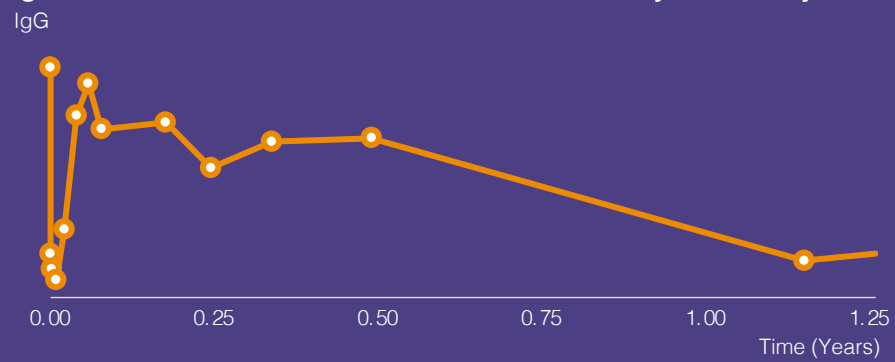
- Expanding into **new indications**
- Reduce immune response to IgG-cleaving enzyme, i.e. allow **repeated treatment**
- **Combination therapy**, i.e. induction and maintenance therapy



IgG levels after imlifidase treatment in humans – First 30 days



IgG levels after imlifidase treatment in humans – 1 year and beyond

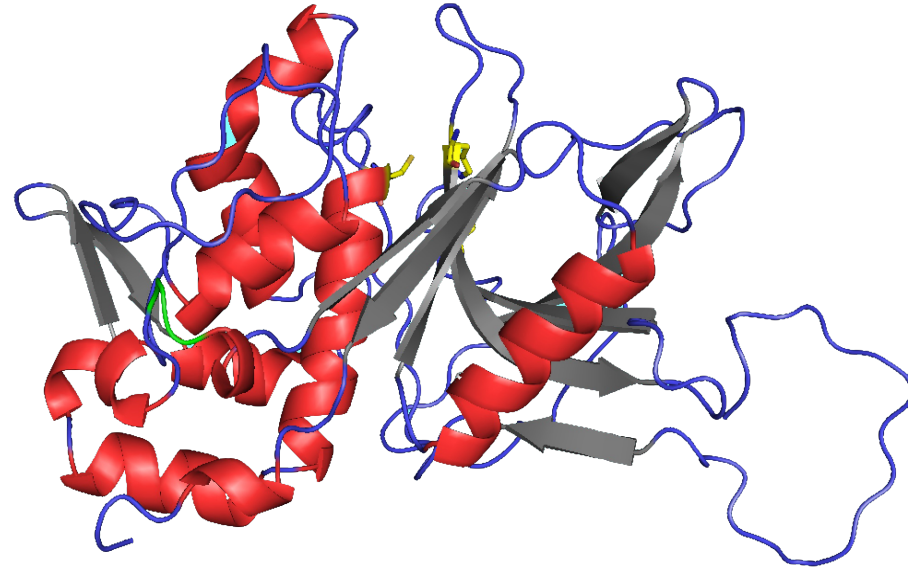


Our IgG antibody-cleaving enzyme, imlifidase

Origins from *Streptococcus pyogenes*

- Cysteine protease derived from an Immunoglobulin G (IgG)-degrading enzyme of *Streptococcus pyogenes*
- Contains only one cysteine - no disulfide bridges
- Monomeric protein with a molecular mass of 35 Kilo Dalton
- Isoelectric point of 6.1
- The coding gene for imlifidase is cloned and expressed in *Escherichia coli*

Imlifidase consists of 311 amino acids



Imlifidase is a lyophilized product formulation

Shelf life of 18 months at 2-8° Celsius storage

Imlifidase will be infused in 15 minutes

- The product for commercial supply will be a lyophilized (cold chain) powder concentrate (11 mg solution) for infusion currently with a claimed shelf life of 18 months at 2-8°C storage. Ongoing stability studies indicate a shelf life of at least 24 months.
- Each vial is filled with 1.2 mL of a 10 mg/mL solution before freeze drying (=12 mg). Extractable volume after reconstitution with 1.2 mL sterile water is 1.1 mL of 10 mg/mL solution - resulting in 11 mg product
- The protein concentration, 10 mg/mL, has desirable characteristics with respect to not form aggregates
- Continuous stability programs ongoing to study changes in protein characteristics and performance.
- Imlifidase dose is clinically set to 0.25 mg/kg bodyweight (11 mg / 0.25 mg/kg = 44 kg (BW) / vial content) 2R vial size is suitable for the content



Supply Chain

Imlifidase in kidney transplantation



Drug Development



Drug substance
Manufacturer (API)



Logistics of bulk product
- handling of drug substance product



Final product
(packaging and labelling)



Distribution



Clinics and hospitals



Patients



Drug product manufacturer
(upscaling)

Manufacturing process

Hansa has close collaborations with highly experienced European based third party CMOs

Drug substance production process (API)

Northway Biotech



Fermentation/ harvesting

- Working Cell Bank
- Pre-Cultivation
- Main Cultivation
- Cell Harvest

Protein purification

- Cell Disruption
- Protein Release

Protein purification cont.

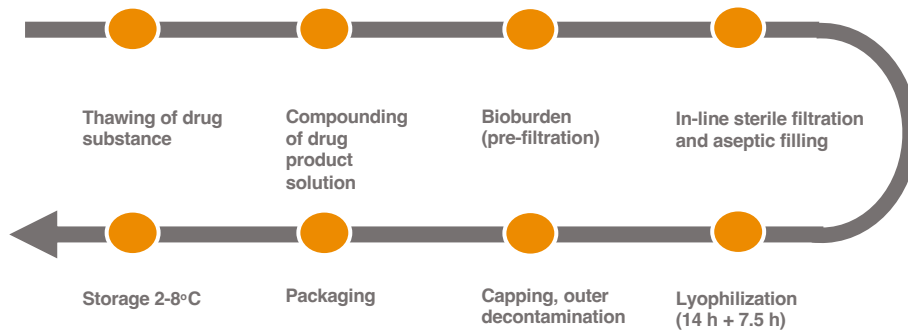
- Ion Exchange Chromatography
- Ceramic Hydroxy Apatite Chromatography
- Hydrophobic Interaction Chromatography
- Ultrafiltration/ Diafiltration

Filling

- Formulation, filtration, filling and storage (-80°C)

Drug product production process (upscaling)

Baxter



Thawing of drug substance

Compounding of drug product solution

Bioburden (pre-filtration)

In-line sterile filtration and aseptic filling

Storage 2-8°C

Packaging

Capping, outer decontamination

Lyophilization (14 h + 7.5 h)



Facts

- Based in Vilnius, Lithuania
- Start-up Year: 2004
- Capacity: 300 L fermentor (1000 L fermentor in 2020)
- Certifications: GMP compliance, Manufacturing authorization license
- Inspections: National regulatory agency (EU), EU/US customer inspections, FDA mock inspection



Facts

- Based in Halle/Westfalen Germany
- Start-up Year: 2001 (contract manufacturing)
- Capacity: 6-35 L drug product solution per batch (5,000-30,000 vials)
- Certifications: GMP compliance, Manufacturing authorization license
- Inspections: National regulatory agency (EU), FDA, EU/US customer inspections



Clinical development programs



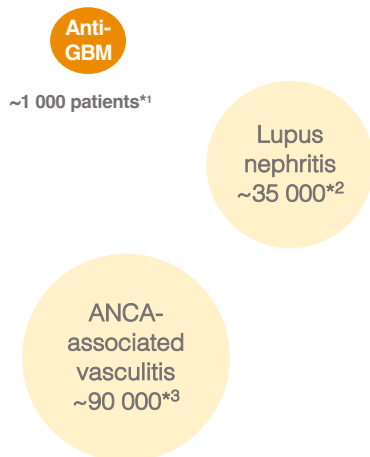
Hansa's antibody cleaving enzyme technology

may have relevance in several autoimmune diseases where IgG plays an important role in the pathogenesis

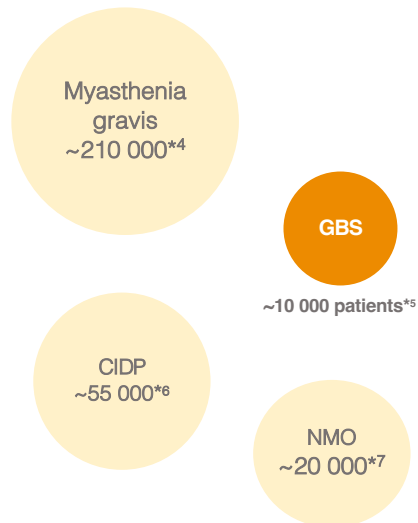
- Clinical programs
- Potential autoimmune indications (currently not pursued)

*Total disease populations in EU & US, based on prevalence and population data

Rapidly progressive glomerulonephritis



Neurological disorders



Skin disorders



Blood disorders



CIDP: Chronic inflammatory demyelinating polyradiculoneuropathy
NMO: Neuromyelitis optica
EBA: Epidermolysis bullosa acquisita
ITP: Immune thrombocytopenia
WAHA: Warm antibody hemolytic anemia
APS: Antiphospholipid syndrome

¹DeVrieze, B.W. and Hurley, J.A. *Goodpasture Syndrome*. StatPearls Publishing, Jan 2021.

²<https://www.ncbi.nlm.nih.gov/books/NBK459291/> [accessed 2021-03-29]

³Patel, M et al. *The Prevalence and Incidence of Biopsy-Proven Lupus Nephritis in the UK*. Arthritis & Rheumatism, 2006.

⁴Berti A, Cornec D, Crowson CS, Specks U, Matteson EL. *The Epidemiology of ANCA Associated Vasculitis in the U.S.: A 20 Year*

Population Based Study. Arthritis Rheumatol. 2017;69.

⁵*Myasthenia Gravis*. National Organization for Rare Disorders. <https://rarediseases.org/rare-diseases/myasthenia-gravis/> [accessed 2021-03-29]

⁶*Guillain-Barré syndrome*. Orpha.net. https://www.orpha.net/consor/cgi-bin/OC_Exp.php?lng=GB&Expert=2103 [accessed 2021-03-29]

⁷*Chronic Inflammatory Demyelinating Polyneuropathy: Considerations for Diagnosis, Management, and Population Health*. The American Journal of Managed Care. <https://www.ajmc.com/view/chronic-inflammatory-demyelinating-polyneuropathy-considerations-for-diagnosis-management-and-population-health> [accessed 2021-03-29]

⁸Marrie, R.A. *The Incidence and Prevalence of Neuromyelitis Optica*. International Journal of MS Care, 2013 Fall: 113-118

⁹Mehren, C.R. and Gniadecki, R. *Epidermolysis bullosa acquisita: current diagnosis and therapy*. Dermatol Reports, 2011-10-05

¹⁰Vertentil, S. et al. *Prevalence Estimates for Pemphigus in the United States*. JAMA Dermatol, May 2019: 627-629.

¹¹*Immune Thrombocytopenia*. National Organization for Rare Disorders. <https://rarediseases.org/rare-diseases/immune-thrombocytopenia/> [accessed 2021-03-29]

¹²*Warm Autoimmune Hemolytic Anemia*. National Organization for Rare Disorders. <https://rarediseases.org/rare-diseases/warm-autoimmune-hemolytic-anemia/> [accessed 2021-03-29]

¹³Litvinova, E. et al. *Prevalence and Significance of Non-conventional Antiphospholipid Antibodies in Patients With Clinical APS Criteria*. Frontiers in Immunology, 2018-12-14.

Anti-GBM, a rare acute autoimmune disease

Positive read-out from phase 2 study with 2/3 of anti-GBM patients achieving dialysis independence six months after treatment

Anti-GBM

- Indication: Antibodies are directed against an antigen intrinsic to the glomerular basement membrane (GBM) causing acute injury of kidney and/or lung
- Anti-GBM affects 1.6 in a million people annually with majority of patients losing their kidney function^{1,2}, requiring chronic dialysis and kidney transplantation
- Phase 2 study concluded that imlifidase leads to rapid clearance of anti-GBM antibodies, with two-thirds of patients achieving dialysis independence six months after treatment
- U.S. FDA has accepted Hansa's Investigational New Drug (IND) application to proceed with a Phase 3 study of imlifidase in 50 anti-GBM patients across U.S. and EU.
- First patient expected to be enrolled in 2022
- Our Anti-GBM program obtained Orphan Drug designation from both FDA and European Commission in 2018



Anti-GBM Phase 2

Imlifidase in Anti-GBM

The idea is that imlifidase in anti-GBM patients may cleave IgG bound to the GBM within a few hours and prevent further renal damage

Today only a fraction of the total IgG antibodies are removed with plasma exchange and IgG in the interstitial tissue and bound to the GBM remains

CLINICALTRIALS.GOV ID

NCT03157037 (Since March 2017)

SUBJECTS

15 patients targeted. Patients will be monitored for six months
Recruitment at 15 clinics

DOSES/FOLLOW UP TIME

Dosage 0.25mg/kg 180 days follow up

MAIN OBJECTIVES

- Primary objective is to evaluate the safety and tolerability of imlifidase on background of standard of care, and assess efficacy based on renal function at six months after treatment

STUDY DESIGN

- Open label, multicenter, single arm Phase 2 study with adverse renal prognosis
- Investigator initiated study

STATUS

Plans to initiate a Phase 3 study of imlifidase to treat 50 anti-GBM patients (FPI 2022)

Potential of using imlifidase vs. PLEX in anti-GBM



Results from Phase 2 study of imlifidase in anti-GBM disease published in Journal of American Society of Nephrology (JASN)¹

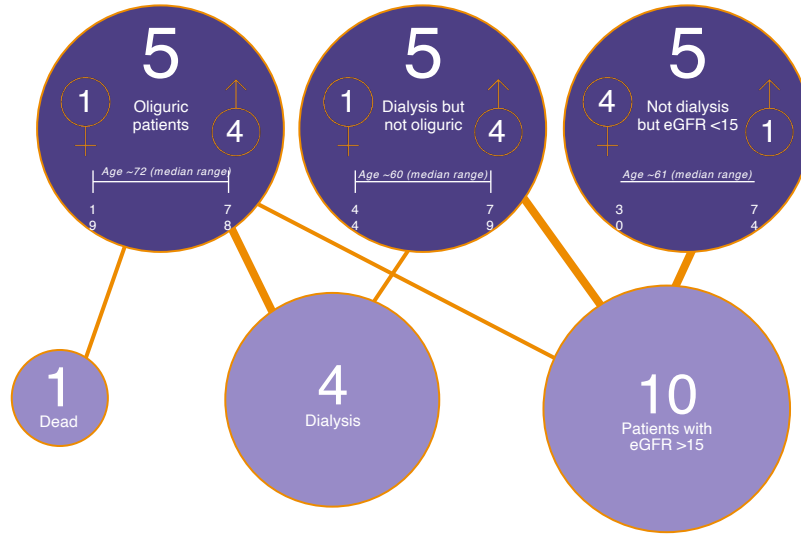
U.S. FDA has accepted Hansa's Investigational New Drug (IND) application to proceed with a Phase 3 across U.S. and EU with the first patient is expected to be enrolled in 2022

JASN recognises the potential in deactivation of autoantibodies in autoimmune diseases

10 out of 15 patients were dialysis independent after six months vs. the historical cohort, where only 18% had functioning kidney

New Anti-GBM Phase 3 study of imlifidase in 50 patients

- Global protocol in place and approved by FDA. Selection of investigators and site set up is now ongoing
- New Phase 3 trial will be an open-label, controlled, randomized, multi-centre trial comparing imlifidase and SoC with SoC alone
- EMA submission preparation in progress
- Plans to expand the trial to include Japan



Segelmark et al. JASN (2022)

¹ Journal of the American Society of Nephrology <https://pubmed.ncbi.nlm.nih.gov/35260419/>

² McAdoo et al.: Patients double-seropositive for ANCA and anti-GBM antibodies have varied renal survival, frequency of relapse, and outcomes compared to single-seropositive patients. *Kidney Int* 92: 693–702, 2017





Phase 2 study to evaluate safety, tolerability and efficacy of imlifidase

in patients with Guillain-Barré syndrome (GBS)

Design of the GBS trial

- Open-label, single-arm trial in combination with SoC treatment given within 10 days of onset of GBS
- Infusion of 0.25mg/kg imlifidase at Day 1, followed by IVIg (400 mg/kg) at Days 3-7, and follow-up of PK/PD for 14 days, safety and efficacy parameters at 6 months and 12 months
- 30 patients targeted and matched to controls based on geographical location, age, presence of diarrhea, severity of condition
- Outcome compared to matched controls (up to 4 controls per patients) from the IGOS¹ database

Main objective

- To evaluate safety, tolerability, PK/PD, and efficacy of imlifidase in GBS patients in combination with SoC intravenous immunoglobulin

Status

- 18/30 patients enrolled end of Q2 2022
- 10/10 sites are recruiting patients
- Recruitment will be done across France, UK and The Netherlands
- Enrollment is expected to be completed in H2 2022 (temporary halted during 2019 due to Covid-19)
- First data readout H1 2023

In 2018, the FDA granted Orphan Drug Designation to imlifidase for the treatment of GBS

Guillain-Barré syndrome

GBS is an acute autoimmune attack on the peripheral nervous system

GBS can affect anyone at any age

- GBS is an acute autoimmune attack on the peripheral nervous system, which rapidly and progressively weakens extremities.
- Only parts of the patients fully recover from GBS, thus a high unmet medical need for new treatments; 40% lose strength and have pain while mortality is 3-7%
- Addressable population of ~10,000¹ per year in 7MM²
- Current Standard of Care is treatment with IVIG or PLEX
- The Phase 2 study is an open-label, single arm, multi-center study evaluating the safety, tolerability and efficacy of imlifidase in GBS patients in combination with standard of care intravenous immunoglobulin (IVIg)
- 18/30 patients enrolled end of Q2 2022. Ongoing recruitment of patients at 10 centers across France, UK and the Netherlands.
- Initiatives implemented to support the completion of enrollment incl. simplification of the protocol and increased capacity
- Completion of enrollment expected in H2 2022* with a first data read-out in H1 2023
- In 2018, the FDA granted Orphan Drug Designation to imlifidase for the treatment of GBS

¹ McGrogan et al. Neuroepidemiology 2009;32(2):150-63.

² 7MM = Seven major markets – US, Germany, UK, France, Spain, Italy, and Japan



GBS Phase 2

New Phase 2 study initiated in GBS to evaluate safety, tolerability and efficacy of imlifidase in GBS patients

CLINICALTRIALS.GOV ID

NCT03943589 (2019)

SUBJECTS

30 patients targeted
Recruitment at ten clinics in Europe
(France, U.K. and the Netherlands)

DOSES/FOLLOW UP TIME

Dosage 0.25mg/kg follow up 180 days
and 12 months

MAIN OBJECTIVES

- safety and effectiveness of imlifidase in patients diagnosed with GBS

STUDY DESIGN

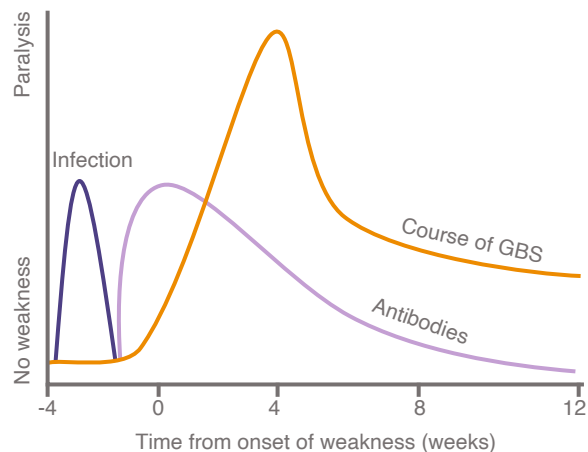
- Study is an open-label, single arm, multi-center trial evaluating safety, tolerability and efficacy of imlifidase, in combination with standard of care, IVIg, to treat GBS

STATUS

Ongoing recruitment

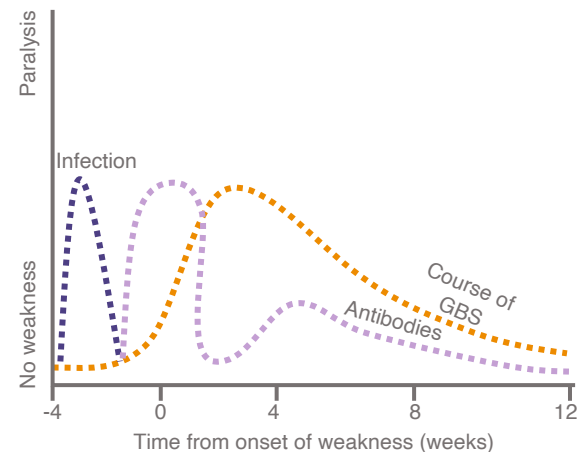
Today's Standard of Care IVIg or PLEX

Illustrative



Potential with imlifidase

Illustrative



Enrollment in Phase 2 program in Antibody Mediated Rejection (AMR) post kidney transplantation completed

Long term graft survival is challenged by AMR episodes post transplantation

Indication

- Acute antibody mediated rejection episodes post transplantation occurs in 5-7% of kidney transplants¹ annually and is a significant challenge to long term graft survival
- Today's standard of care include plasma exchange, steroids and IVIg.
- There is no approved treatment for AMR

¹ Puttarajappa et al., Journal of Transplantation, 2012, Article ID 193724.

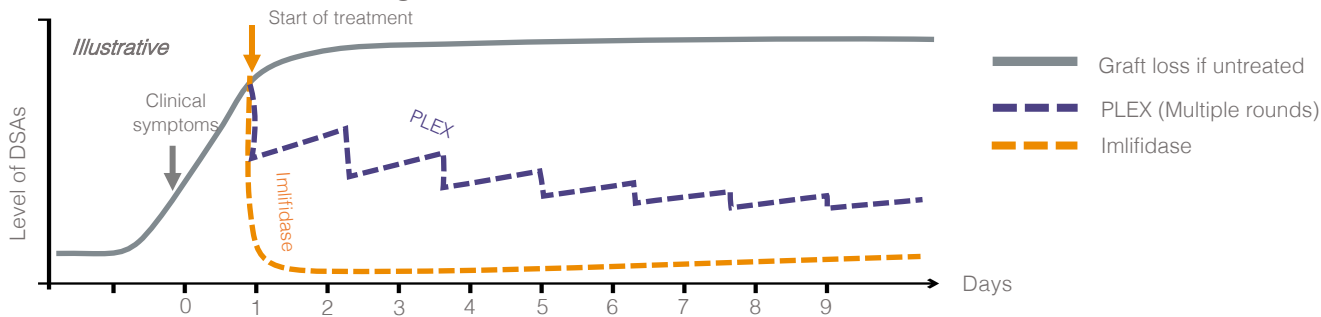
Phase 2 Study

- 30 patients with active or chronic active AMR episodes post kidney transplantation have been enrolled and randomized 2:1 to imlifidase vs. SoC
- The AMR phase 2 program is a randomized, open-label, multi-center and controlled study
- 20 individuals have been randomized to receive imlifidase treatment comprised of one intravenous dose of 0.25mg/kg, while 10 individuals in the active control arm received 5-10 sessions of plasma exchange (PLEX)
- Efficacy and safety is monitored over a six-month period post treatment.

Path forward

- ✓ **H1'22:** Completion of enrollment
- **H2'22:** Phase 2 data read-out
- ▼ **Decision on a regulatory path forward**

Potential of using imlifidase vs. PLEX in AMR



Antibody Mediated Rejection

Long term graft survival is challenged by AMR post transplantation

There is no approved treatment for AMR

- Active antibody mediated rejection after transplantation occurs in 5-7% of kidney transplants¹ annually⁴ and is a significant challenge to long term graft survival
- Today's standard of care include plasma exchange, and treatment with steroid and IVIg. AMR patients not treated successfully risk graft failure, dialysis and return to the waitlist
- The AMR Phase 2 study is a randomized, open-label, multi-center, active control study designed to evaluate the safety and efficacy of imlifidase in eliminating donor specific antibodies (DSAs) in the treatment of active episodes of acute AMR in kidney transplant patients
- Completion of enrollment in 30 patient at 14 centers across the US, Europe and Australia was done May 2022
- First data read out expected in H2 2022

¹ Puttarajappa et al., Journal of Transplantation, 2012, Article ID 193724.

² Seven major markets – US, Germany, UK, France, Spain, Italy, and Japan



AMR Phase 2

AMR Phase 2 study

Aim of the study is test imlifidase ability to reduce the amount of donor specific antibodies in AMR patients post transplantation

CLINICALTRIALS.GOV ID

NCT03897205 (2019)

SUBJECTS

30 patients targeted (20 patients will be treated with imlifidase and 10 with Plasma exchange). Recruitment from 11 sites in the U.S., EU and Australia.

DOSES/FOLLOW UP TIME

1 dose of imlifidase (0.25 mg/kg) or 5-10 sessions of plasma exchange

MAIN OBJECTIVES

- Imlifidase ability to reduce the amount of DSA in comparison with plasma exchange in patients who have an active AMR post transplantation
- Ensure safety for patients

STUDY DESIGN

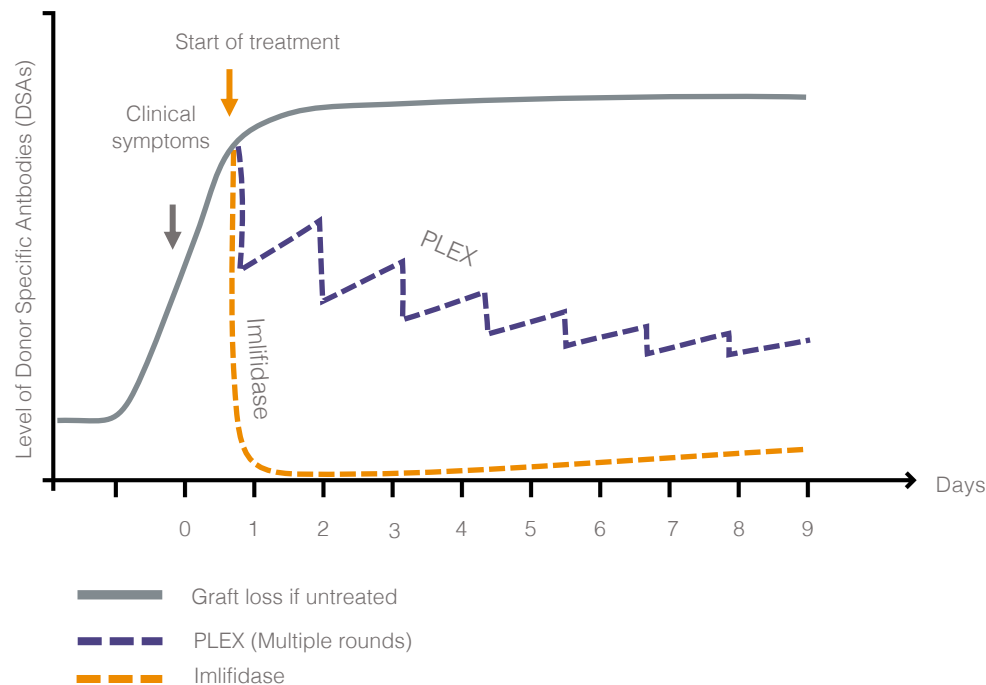
- Randomized, open-label multi-center, active control study, designed to evaluate the safety and efficacy of imlifidase in eliminating DSA in active AMR

STATUS

Completed enrollment awaiting first data readout H2 2022

Potential of using imlifidase vs. PLEX in AMR

Illustrative



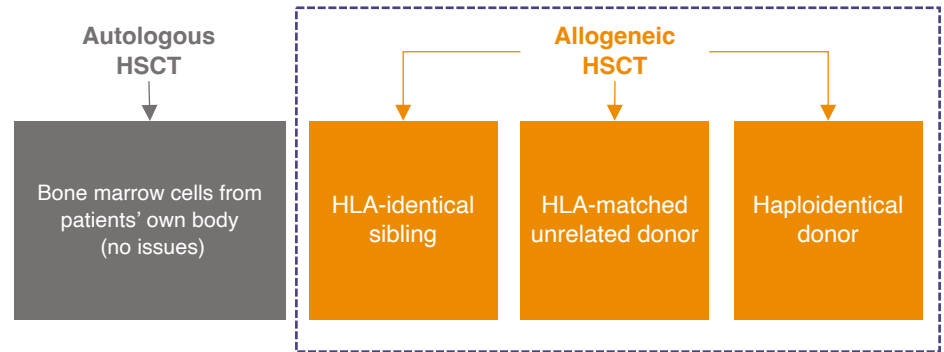
Exploring potential use of imlifidase in allogeneic hematopoietic stem cell transplantation (HSCT)

Desensitization treatment of patients with high levels of donor specific antibodies (DSA) prior to allogeneic HSCT transplant is a challenge; Imlifidase may have the potential to inactivate DSAs prior to transplantation

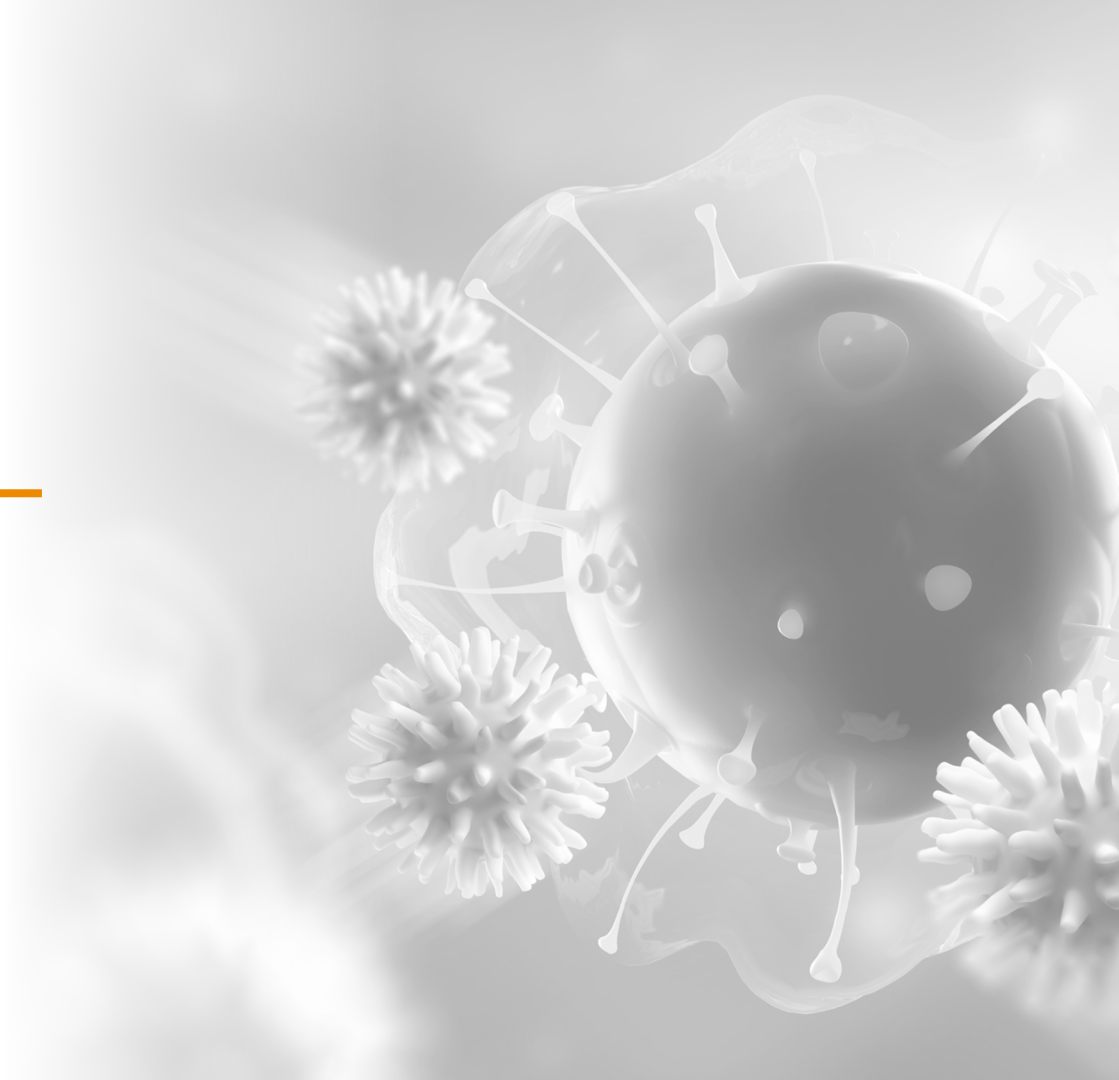
Transplantations are often acutely needed, which reduces the time available to find an adequately matched donor

- Haploidentical donors (e.g. parents, children) are often available and transplant outcome is good (e.g. engraftment, graft survival, survival)
- However, presence of donor specific antibodies (DSAs) have a negative impact on transplant outcome² (e.g. graft failure and survival) Prevalence of DSAs in allogeneic HSCT is typically between 10-21%¹.
- There are currently no approved drugs to manage patients with high levels of DSAs and current desensitization methods are inadequate, thus preventing patients from having a potentially life-saving HSCT
- Consensus recommendations published¹ by the EBMT³ on testing, monitoring and treatment of patients with donor specific antibodies recommend to desensitize all patients with DSAs
- Imlifidase may have the potential to transform the standard of care by enabling clinicians to inactivate DSAs prior to transplantation

Pre-existing DSAs may result in primary graft failure and poor survival after allogeneic hematopoietic stem cell transplantations



Pre-clinical programs





Preclinical collaboration with argenx BV

Collaboration to evaluate the potential combination of companies' IgG-modulating approaches

- A combination of Hansa's IgG antibody-cleaving enzyme, and efgartigimod, argenx's FcRn antagonist could potentially be used in both the acute and chronic setting of autoimmune diseases and transplantation to potentially unlock additional therapeutic value
- Under the agreement, both parties will contribute equally in terms of resource allocation and will share all IP and data developed through the collaboration
- Both parties will maintain exclusive rights to their respective technologies and products.

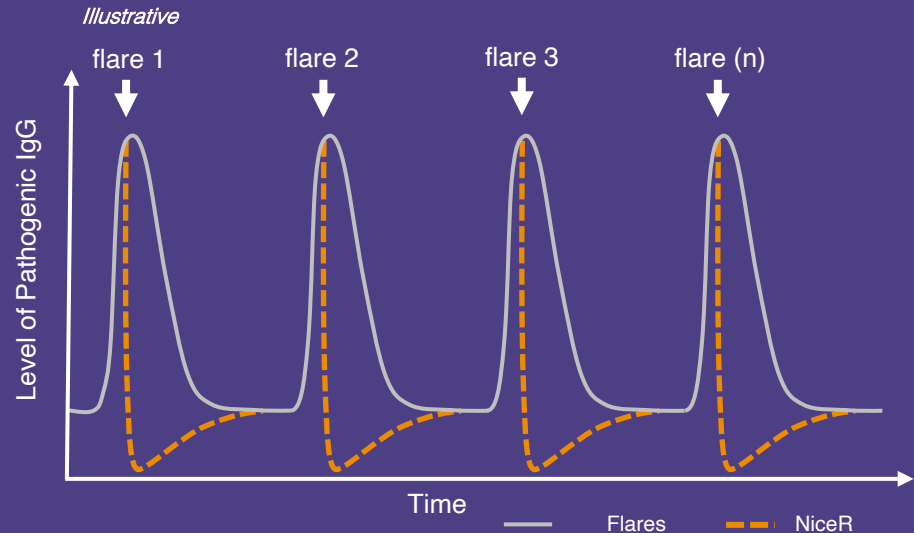
“NiceR” for repeat dosing

a new set of enzymes for repeat dosing for potential inactivation of flares in relapsing diseases

NiceR - Novel Immunoglobulin Cleaving Enzymes for Repeat dosing with lower immunogenicity

- Potential application for a broad array of indications, including reoccurring AMR, relapsing autoimmune diseases and oncology
- The first selected promising new drug candidate from the NiceR program is an IgG-cleaving enzyme (cysteine peptidase) with characteristics based on a homolog to imlifidase, but with lowered immunogenicity
- IND-enabling tox studies initiated in H1'21. Completion of GLP tox studies in 2022

NiceR can potentially inactivate flares



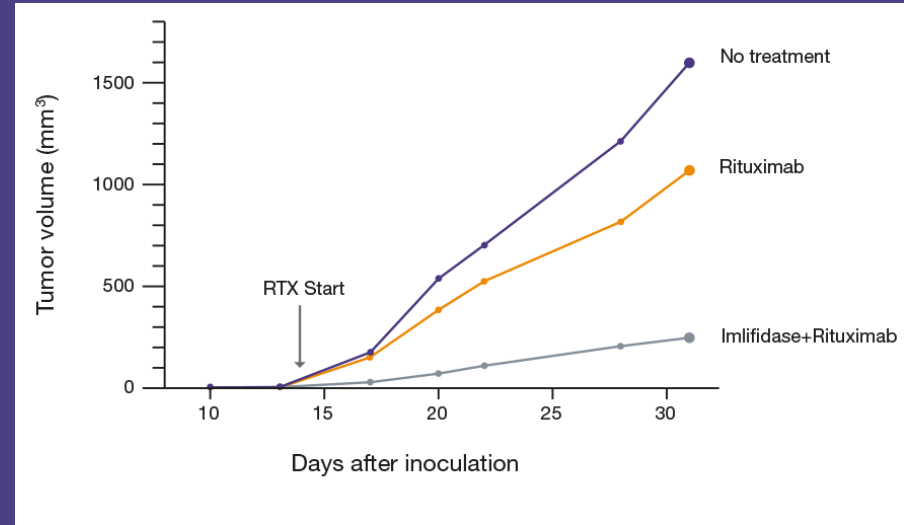
Our antibody cleaving enzymes

may potentially improve the therapeutic effect of immunotherapy in oncology (EnzE)

Proof of concept demonstrated in vivo for mice

- Enzyme based antibody enhancement through pre-treatment
- The abundance of normal IgG in blood interferes with therapeutic monoclonal antibodies
- Pre-treatment with imlifidase / NiceR has potential to significantly potentiate antibody-based cancer therapies
- Suppressive effect of IVIg on effector cell function abrogated by imlifidase
- Imlifidase can significantly improve the therapeutic effect of rituximab

Mice with human IgG (~9mg/mL)



¹ Järnum et al. Mol Cancer Ther 2017;16:1887-1897

Gene Therapy



Exploring opportunities in gene therapy

Exploring the opportunities in systemic administration of gene therapy for our unique antibody cleaving enzyme platform to potentially enable gene therapy treatment in NAb+ patients

A
revolutionary
approach

Significant
unmet need

Encouraging
pre-clinical
data

Partnership
strategy

Tropism and target tissue

AAV subtypes targets different tissues



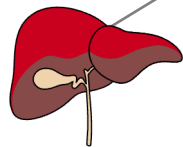
AAV 1, 2 & 5



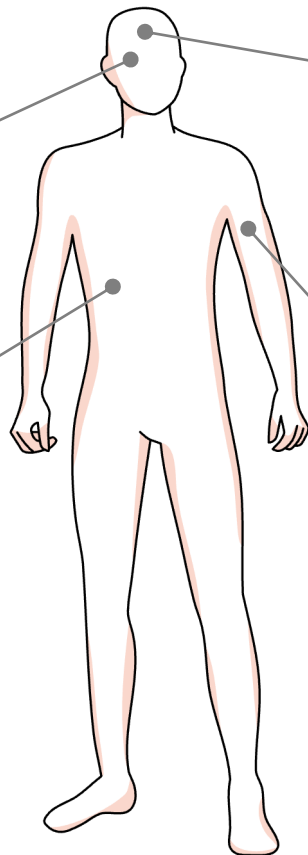
Eye (local target)
 $\sim 1 \times 10^{11}$ vg



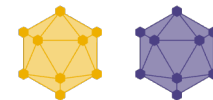
AAV 3, 7 & 8



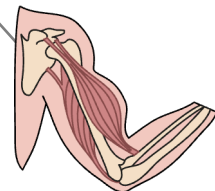
Liver (systemic)
 $\sim 1 \times 10^{14}$ vg



Brain (local target)
 $\sim 1 \times 10^{12}$ vg



AAV 4 & 8



Muscle (systemic)
 $\sim 1 \times 10^{15}$ vg



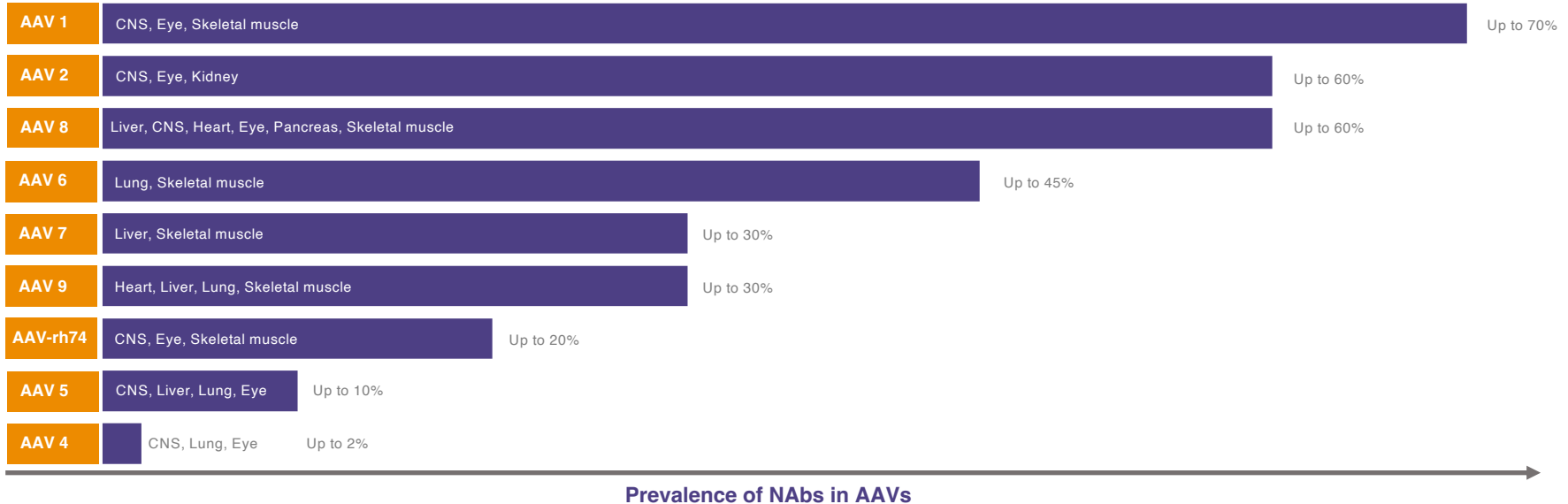
AAV 6, 7, rh74

Target tissues

Dose of gene therapy (vg)

Neutralizing antibodies are a barrier that precludes gene therapies

from working in a large group of patients. The prevalence of NABs varies significantly across the different vectors

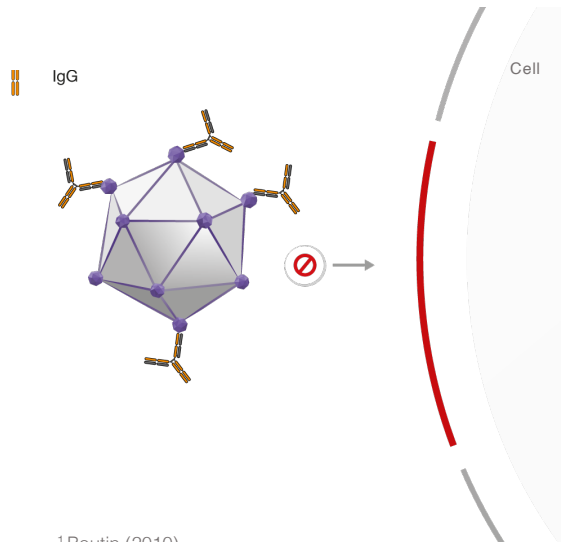


Source: Boutin et al (2010), Griffin et al (2019), Wang et al (2018), Calcedo & Wilson (2013), Falese et al (2017), Haiyan et al (2017), Ellsworth et al (2018), Greig et al (2017)

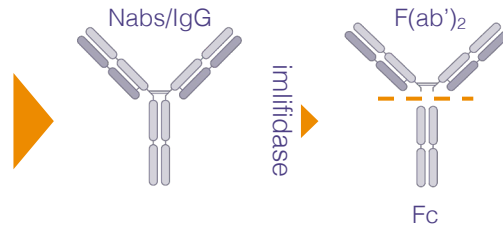
Neutralizing antibodies (Nabs) are immunological barriers in gene therapy; imlifidase may potentially eliminate Nabs

Between approximately 5% and 70%^{1,2} of patients considered for gene therapy treatment carry neutralizing anti-AAV antibodies forming a barrier for treatment eligibility

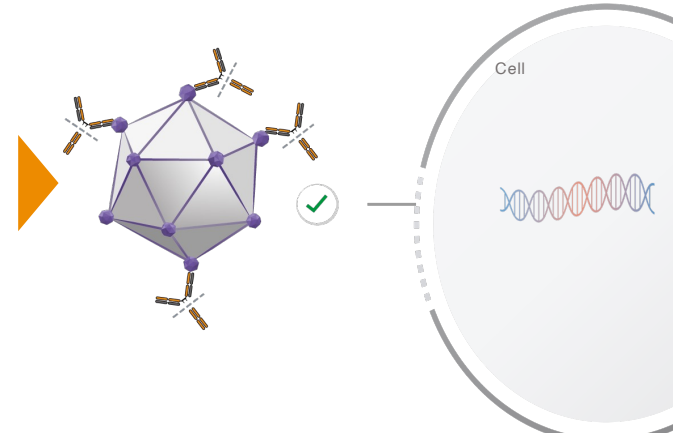
1 Antibodies prevent effective transfer of healthy gene sequence and can be a safety concern



2 Imlifidase is a unique IgG antibody-cleaving enzyme that cleaves IgG at the hinge region with extremely high specificity



3 The idea is to eliminate the neutralizing antibodies as a pre-treatment to enable gene therapy

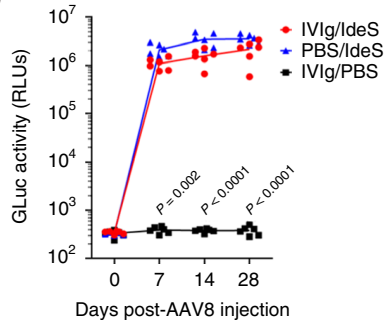


Imlifidase (IdeS) was highlighted in Nature Medicine¹

with encouraging outcome demonstrating imlifidase as a potential solution to overcome pre-existing antibodies to AAV-based gene therapy

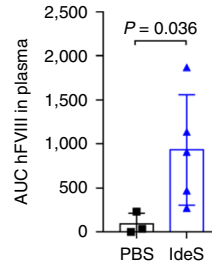
Imlifidase tested in a animal model

- Imlifidase decreased anti-AAV antibodies and enabled efficient gene transfer



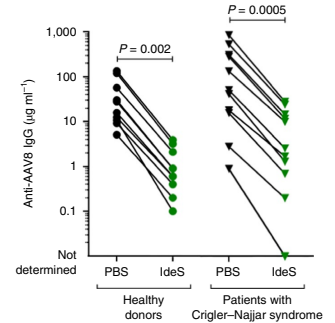
Imlifidase tested in NHP ahead of AAV vector infusion

- Pre-treatment with imlifidase in anti-AAV positive nonhuman primates (NHP) ahead of AAV vector infusion was safe and resulted in enhanced liver transduction and hFVIII plasma levels

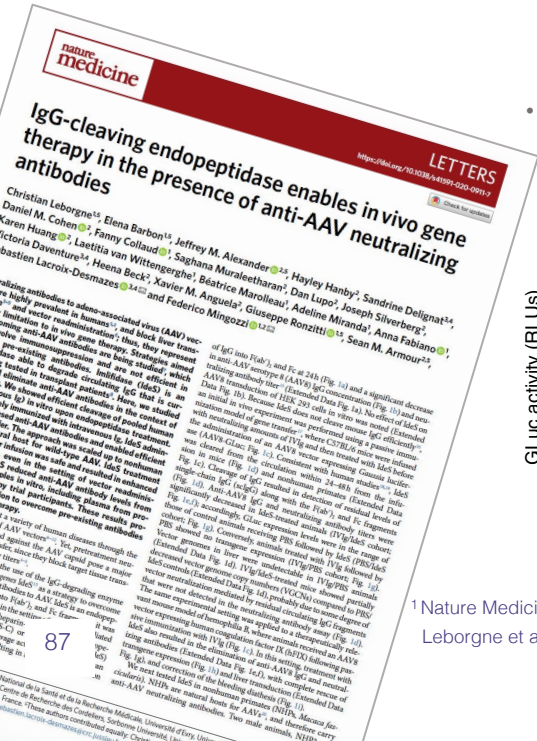


Imlifidase tested in human plasma samples (GT patients)

- Imlifidase reduced anti-AAV antibody levels from human plasma samples in vitro, incl. plasma from prospective gene therapy trial participants



¹ Nature Medicine <https://doi.org/10.1038/s41591-020-0911-7>
 Leborgne et al. Nat Med (2020)



Global and exclusive agreement with Sarepta Therapeutics

to develop and promote imlifidase as pre-treatment ahead of gene therapy in select indications



Indication exclusivity:

- Duchenne Muscular Dystrophy (DMD)
- Limb-Girdle Muscular Dystrophy (LGMD)

Hansa's key resources

- Imlifidase validated with positive clinical efficacy and safety data as well as European approval
- Positive preclinical data published in *Nature*
- Clear path to U.S. approval (kidney transplant)



Sarepta's key resources

- World leader within gene therapy targeted at muscular dystrophies
- Pre-clinical plan: PoC and IND-tox
- Clinical / Regulatory
- Promotion

Collaborative research, development and commercialization – working together at every stage



Collaboration with AskBio to evaluate imlifidase in gene therapy targeting Pompe disease

Feasibility program to evaluate imlifidase as pre-treatment ahead of gene therapy in Pompe disease for patients with pre-existing neutralizing antibodies (NAbs) to adeno-associated virus (AAV)



Hansa's key resources and deliverables

- Imlifidase validated with positive clinical efficacy and safety data as well as European approval
- Significant know-how around antibody cleaving enzymes
- Clear path to U.S. approval (kidney transplant)
- Hansa supplies material and provides additional support

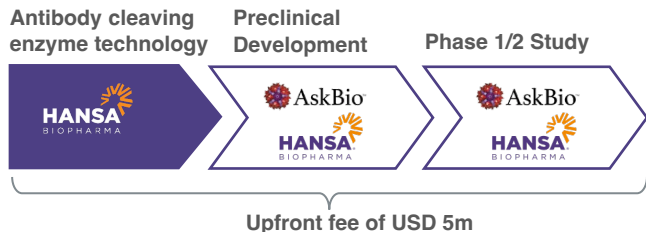


Fully owned subsidiary of Bayer AG

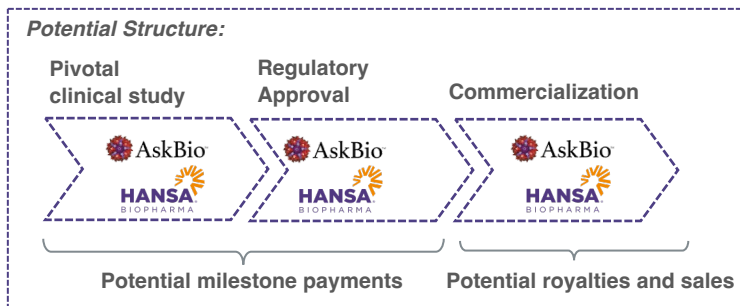
AskBio's key resources and deliverables

- Early innovator in the Gene Therapy space with AAV platform and ongoing clinical stage Pompe disease program
- Conducts pre-clinical and clinical trials according to agreed plan

Current agreement scoped around a feasibility program which covers preclinical work and a Phase 1/2 study



Exclusive option for AskBio to negotiate a potential full development and commercialization agreement



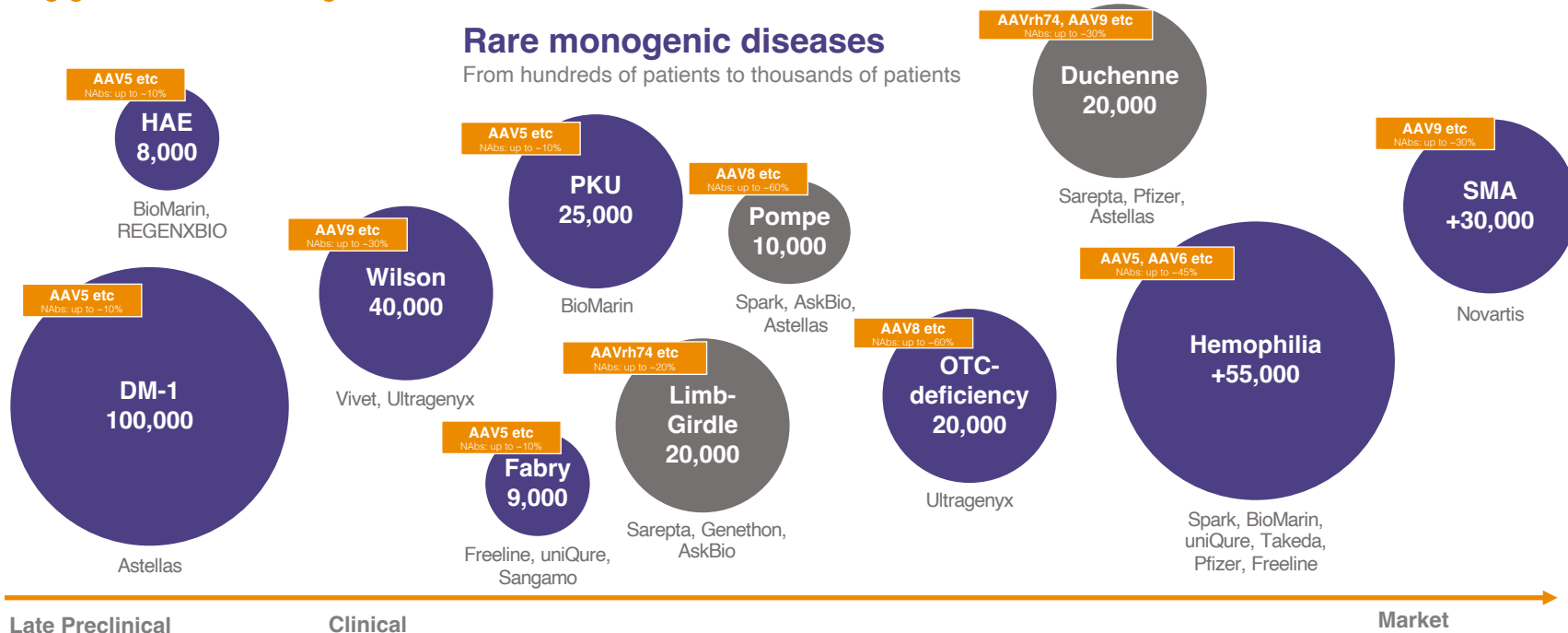
Systemic gene therapy is an emerging opportunity

with a focus on the potential to correct issues causing genes in rare monogenic diseases

- Preclinical programs with Sarepta and AskBio
- Potential gene therapy indications (currently not pursued)

Rare monogenic diseases

From hundreds of patients to thousands of patients



Late Preclinical

Clinical

Market

● Size of indication (US & EU)

Duchenne Muscular Dystrophy (DMD) SRP-9001

About Duchenne Muscular Dystrophy (DMD)¹

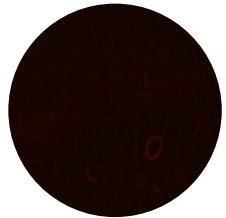
- Rare, fatal neuromuscular genetic disease
- Muscle weakness noticeable by age 3 to 5, and most patients use a wheelchair by the time they are 11
- Cardiac and respiratory muscle deterioration becomes life-threatening
- 1/3,500 to 5,000 male births (worldwide)
- Approximately 15% of patients have pre-existing IgG antibodies to rh74

SRP-9001 micro-dystrophin gene therapy for treatment of DMD

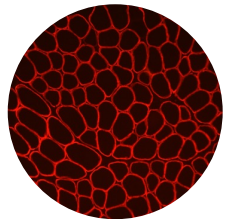
- AAVrh74 vector with micro-dystrophin transgene
- Broad patient experience (77 Duchenne trial participants dosed)
- 4 ongoing clinical trials – including recently initiated pivotal study
- Robust micro-dystrophin protein expression with commercially representative process material
- Functional benefits sustained up to 3 years after administration
- Observed safety profile is consistent

For further information regarding Sarepta's gene therapy programs, please refer to www.sarepta.com

Pre-treatment



Post-treatment



Source:

¹ Sarepta Therapeutics <https://investorrelations.sarepta.com/static-files/e9393c38-646f-45ee-9f56-955f3bfad71>

² National Institutes of Health. Genetics Home Reference. Duchenne and Becker muscular dystrophy. <https://ghr.nlm.nih.gov/condition/duchenne-and-becker-muscular-dystrophy>. Accessed Jan 2020.

³ Sarepta Therapeutics <https://investorrelations.sarepta.com/static-files/e9393c38-646f-45ee-9f56-955f3bfad7>

Limb-Girdle muscular dystrophy (LGMD) SRP-9003

About limb-girdle muscular dystrophy

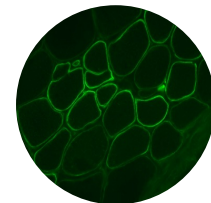
Limb-girdle muscular dystrophy is a group of diseases that cause weakness and wasting of the muscles

- Caused by defects in genes encoding for proteins residing within the sarcolemma, cytosol or nucleus of the muscle cell
- LGMD subtypes are often grouped according to which protein is affected
- Approximate global prevalence of 1.63 per 100,000 individuals; over 30 subtypes exist
- Approximately 15% of patients have pre-existing IgG antibodies to rh74

SRP-9003 β -sarcoglycan (SGCB) gene therapy for treatment of LGMD2E

- AAVrh74 vector with transgene β -sarcoglycan
- Open label study ongoing (N=6)
- Interim analysis disclosed in May 2021²:
 - Two dosing cohorts
 - Cohort 1 (n=3) - 1.85×10^{13} vg/kg; 2-year follow-up
 - Cohort 2 (n=3) - 7.41×10^{13} vg/kg; 1-year follow-up
 - No new safety signals, and treatment-related AEs occurred early and were transient and manageable
 - Robust, dose-dependent SGCB protein expression in all patients at Day 60, resulting in reconstitution of the sarcoglycan complex; SGCB expression sustained up to 2 years in cohort 1
 - Demonstrated functional improvements, including both NSAD and timed function tests, compared to baseline that were sustained for 2 years in cohort 1 and 1 year in cohort 2

β -sarcoglycan



For further information regarding Sarepta's gene therapy programs, please refer to www.sarepta.com

*Doses are based on titer method using supercoiled plasmid standard

Source:

- 1) National Institutes of Health. Genetics Home Reference. Duchenne and Becker muscular dystrophy. <https://ghr.nlm.nih.gov/condition/duchenne-and-becker-muscular-dystrophy>. Accessed Jan 2020
- 2) Rodino-Klapac et al. Presented at the annual meeting of the American Society of Cell and Gene Therapy May 11-14, 2021

Pompe Disease (PD) AAV2/8-LSPHGAA

About Pompe Disease

- Defect in a gene making an enzyme called acid alpha-glucosidase (GAA), which is used to break down glycogen
- Accumulation of glycogen result in severe impact on the normal organ and muscle function
- Current standard of care is enzyme replacement therapy (ERT)
- Approximate incidence is 1 per 40,000¹ births, or ~200 per year in the US + EU
- Prevalence is estimated to be around 10,000 in the US and Europe combined²
- Approximately 40-60%^{3,4} of patients have pre-existing IgG antibodies to AAV8

AskBio's AAV2/8-LSPHGAA gene therapy

- AAV2 vector genome cross-packaged as AAV8
- Liver-specific promoter to express GAA enzyme
- Open label Phase I/II study ongoing
- Study in 8 Late-Onset Pompe Disease patients
- ClinicalTrials.gov: [NCT03533673](https://clinicaltrials.gov/ct2/show/study/NCT03533673)

For further information regarding AskBio's gene therapy program, please refer to www.askbio.com

Sources:

¹Pompe Disease, <https://rarediseases.org/rare-diseases/nompe-disease/> [accessed 2022-02-08]

²Calculated by Hansa on the basis of incidence numbers from <https://rarediseases.org/rare-diseases/nompe-disease/> and life expectancy estimates from <https://nompediseaseaware.com/late-onset-nompe-disease/> as well as population statistics for the United States and European Union/Europe.

³ESGCT 27th Annual Congress Abstracts, Sensitivity of different AAV serotypes to pre-existing NAbs, https://www.esgct.eu/home/Barcelona%202019/NEW_Abs%20Barcelona%20Abstracts.pdf

⁴Boutin et al. Prevalence of serum IgG and neutralizing factors against adeno-associated virus (AAV) types 1, 2, 5, 6, 8, and 9 in the healthy population: implications for gene therapy using AAV vectors. Hum Gene Ther. 2010. <https://pubmed.ncbi.nlm.nih.gov/20095819/>

ESG Overview



Formalising our ESG approach

At Hansa, we have always strived to achieve sustainable business practices. We are now formalizing our approach to sustainability and ESG issues, starting with identifying our key material focus areas.



Our mission: We leverage our unique enzyme technology platform to develop innovative, lifesaving and life altering immunomodulating therapies, bring these to the patients with rare diseases who need them, and generate value to society at large.

Our key ESG material aspects

Environment

Climate & waste impacts of production and logistics

Hansa's environmental impact is small because our production is limited in volume. However, as we grow, we need to be transparent about, and make efforts to limit, our climate and waste impacts.



Social

Unmet needs and equity in health

Patients with rare conditions in general, and highly sensitized patients in particular, have many unmet needs which our therapies help address. These unmet needs can also be reinforced by ethnic or socio-economic status, particularly regarding access to organ transplants. Collaboration with patient groups can help us reach even more patients who can benefit from our treatments.



Putting patients first

In the biopharma industry, there is a risk that patient access to innovative treatment is delayed. Hansa can therefore provide bridge financing on a case-by-case basis to benefit patients who have limited treatment options.



Employee wellbeing, diversity and inclusion

Ensuring employee wellbeing, diversity and inclusion is a fundamental commitment at Hansa. It is also essential for attracting talent in a fast-growing organization and delivering on our strategy.



Third-party risks

We diligently select new business partners, as well as monitor our existing partners and require them to comply with all laws and regulations and our Code of Conduct.



Pricing

In Europe, value-based pricing and universal health coverage is common, but in other countries access is a pressing issue. We can expand access to unfunded patients through collaboration with patient groups.



Governance

Safety, efficacy and ethics

To build a successful company and achieve our mission to extend and enhance the lives of the patients we serve, we must hold ourselves to the very highest standards. Trust is at the core of everything we do.



Return to investors

Biopharma companies need to remain economically attractive as an investment, so as to continue to secure capital and develop new treatments.



UN Sustainable Development Goals

The Sustainable Development Goals (SDGs) were adopted by all UN Member States in 2015 as a universal call to action to end poverty, protect the planet and ensure that all people enjoy peace and prosperity by 2030. They have since become a gold standard for sustainability across businesses, and each of our recommended factors have been developed to align with relevant goals.



Capital Markets



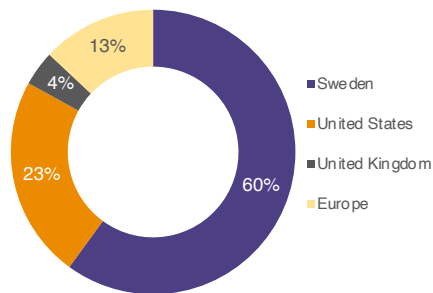
Ownership in Hansa Biopharma

Top 10 ownership as per March 31, 2021

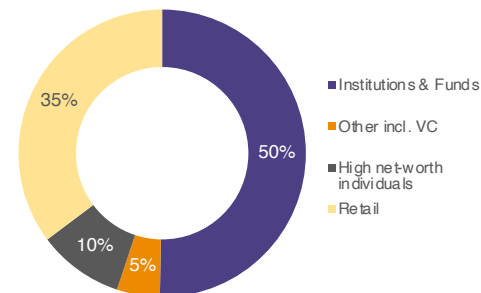
Name	No. of shares	Ownership
Redmile Group, LLC	5,380,863	12.1%
Fjärde AP-Fonden (AP 4)	2,207,397	4.9%
Nexttobe AB	2,155,379	4.8%
Invesco Advisers, Inc.	1,973,931	4.4%
Olausson, Thomas	1,820,500	4.1%
Försäkrings AB Avanza Pension	1,743,201	4.0%
Tredje AP-Fonden (AP 3)	1,389,650	3.1%
The Vanguard Group, Inc.	1,223,839	2.7%
Schroder Investment Management	888,132	2.0%
C WorldWide Asset Management	799,749	1.8%
Other	25,005,477	56.1%
Total	44,588,118	100.0%

Classification of ownership as per Dec 31, 2021

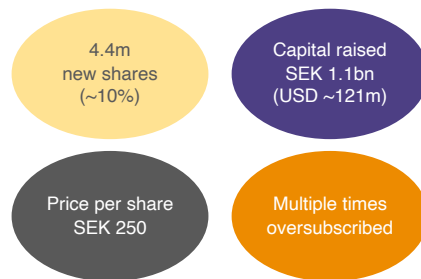
Ownership by country



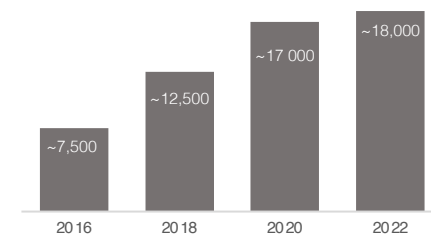
Ownership by type



Capital Raise July 2020



No. of shareholders



Company collected consensus

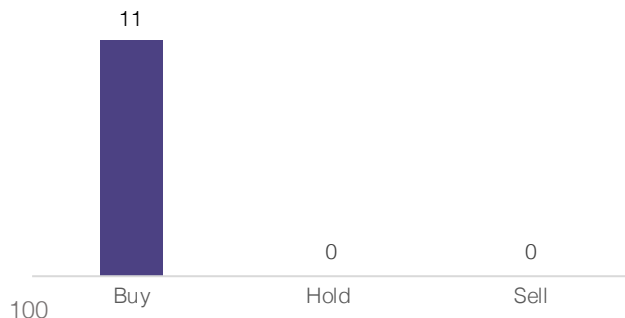


Consensus is based on a collection of analyst estimates pre our Q2 2022 report (July 19, 2022)

	Price Target, SEK	WACC	Patient uptake, EU			Revenue, SEKm		
			FY'22e	FY'23e	FY'24e	FY'22e	FY'23e	FY'24e
Average	236	11,0%	26	51	108	98	182	331
Median	230	11,0%	26	59	105	105	194	328
High	410	15,0%	35	66	195	128	280	543
Low	120	7,6%	15	22	30	53	72	96
<i>Number of contributions</i>	11	10	8	8	8	10	10	10

	EBIT, SEKm			Operating Cash Flow, SEKm			Cash position, SEKm		
	FY'22e	FY'23e	FY'24e	FY'22e	FY'23e	FY'24e	FY'22e	FY'23e	FY'24e
Average	-611	-669	-670	-583	-646	-674	408	413	-54
Median	-617	-645	-581	-577	-619	-596	347	429	195
High	-455	-489	-320	-454	-461	-296	1 122	1 318	535
Low	-755	-1 072	-1 424	-726	-1 105	-1 118	162	-285	-922
<i>Number of contributions</i>	10	10	10	10	10	10	10	7	7

Analyst recommendations

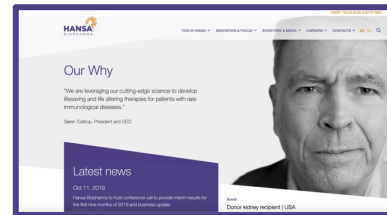


Bank/Research Institution	Analyst	Location	E-mail
SEB	Christopher Uhde, PhD	Stockholm	christopher.uhde@seb.se
ABG Sundal Collier	Adam Karlsson	Stockholm	adam.karlsson@abgsc.se
Carnegie	Erik Hultgård	Stockholm	erik.hultgard@carnegie.com
Redeye	Johan Unnerus	Stockholm	johan.unnerus@redeye.se
RBC	Zoe Karamanoli	London	zoe.karamanoli@rbccm.com
Kempen	Jacob Mekhael	Amsterdam	jacob.mekhael@kempen.com
Intron Health Research	Naresh Chouhan	London	naresh@intronhealthresearch.com
Ökonomiskt Ugebrev	Lars Hatholt	Copenhagen	hatholt@outlook.com
Danske Bank	Caroline Banér	Stockholm	caroline.baner@danskebank.se
Erik Penser Bank	Ludvig Svensson	Stockholm	ludvig.svensson@penser.se
H.C. Wainwright	Douglas Tsao	New York	dtsao@hcwresearch.com

Corporate Contacts

Investor Relations and
Corporate Communications

Visit our web site
www.hansabiopharma.com



Klaus Sindahl

Head of Investor Relations

Mobile: +46 (0) 709-298 269

Email: klaus.sindahl@hansabiopharma.com



Katja Margell

Head of Corporate Communications

Mobile: +46 (0) 768-198 326

Email: katja.margell@hansabiopharma.com

Calendar and events

Aug 9, 2022

BTIG Biotechnology Conference 2022, New York

Aug 10, 2022

Kempen US non-deal road show, New York

Aug 11, 2022

Canaccord Annual Growth Conference, Boston

Aug 18, 2022

Penserpodden "Focus on autoimmunity" (virtual)

Aug 24, 2022

Handelsbanken Life Science Innovation Day, Stockholm (virtual)

Aug 31, 2022

Redeye Late-stage Life Science conference, Stockholm

Sept 7, 2022

Pareto annual Healthcare Conference 2022, Stockholm

Sept 8, 2022

Citi's 17th Annual BioPharma Conference, Boston

Sept 12, 2022

H.C. Wainwright Global Investment Conference, New York

Sept 13-14, 2022

MorganStanley Global Healthcare Conference, New York

Sept 20, 2022

Redeye Afterwork presentation, Gothenburg

Sept 21, 2022

Redeye Lunch presentation, Stockholm

Sept 26, 2022

Aktiespararna Aktiedagen, Lund

Oct 20, 2022

Interim Report for January-September 2022

Nov 23, 2022

SEB Healthcare Seminar 2022, Stockholm

Nov 24, 2022

Redeye Life Science Day, Stockholm

Dec 1, 2022

Erik Pensers Banks Temadag - Health Care, Stockholm



HANSA

BIOPHARMA