

**EACH  
LIFE IS  
UNIQUE**



**KAMADA**

# **KAMADA INVESTOR PRESENTATION**

NASDAQ & TASE: KMDA

October 2018

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# FORWARD LOOKING STATEMENT



This presentation is not intended to provide investment or medical advice. It should be noted that some products under development described herein have not been found safe or effective by any regulatory agency and are not approved for any use outside of clinical trials.

This presentation contains forward-looking statements, which express the current beliefs and expectations of Kamada's management. Such statements involve a number of known and unknown risks and uncertainties that could cause Kamada's future results, performance or achievements to differ significantly from the prospected results, performances or achievements expressed or implied by such forward-looking statements. Important factors that could cause or contribute to such differences include, but are not limited to, risks relating to Kamada's ability to successfully develop and commercialize its pharmaceutical products, the progress and results of any clinical trials, the introduction of competing products, the impact of any changes in regulation and legislation that could affect the pharmaceutical industry, the difficulty of predicting, obtaining or maintaining U.S. Food and Drug Administration, European Medicines Agency and other regulatory authority approvals, the regulatory environment, restraints related to third parties' IP rights and changes in the health policies and structures of various countries, environmental risks, changes in the worldwide pharmaceutical industry and other factors that are discussed under the heading "Risk Factors" of Kamada's 2017 Annual Report on Form 20-F as well as in Kamada's recent Forms 6-K filed with the U.S. Securities and Exchange Commission.

This presentation includes certain non-IFRS financial information, which is not intended to be considered in isolation or as a substitute for, or superior to, the financial information prepared and presented in accordance with IFRS. The non-IFRS financial measures may be calculated differently from, and therefore may not be comparable to, similarly titled measures used by other companies. In accordance with the requirement of the SEC regulations a reconciliation of these non-IFRS financial measures to the comparable IFRS measures is included in an appendix to this presentation. Management uses these non-IFRS financial measures for financial and operational decision-making and as a means to evaluate period-to-period comparisons. Management believes that these non-IFRS financial measures provide meaningful supplemental information regarding Kamada's performance and liquidity.

Forward-looking statements speak only as of the date they are made, and Kamada undertakes no obligation to update any forward-looking statement to reflect the impact of circumstances or events that arise after the date the forward-looking statement was made, except as required by applicable securities laws. You should not place undue reliance on any forward-looking statement and should consider the uncertainties and risks noted above, as well as the risks and uncertainties more fully discussed under the heading "Risk Factors" of Kamada's 2017 Annual Report on Form 20-F as well as in Kamada's recent Forms 6-K filed with the U.S. Securities and Exchange Commission.

# INVESTMENT HIGHLIGHTS



## COMMERCIAL STAGE BIOPHARMA

- Leader in plasma-derived protein therapeutics, focused on Alpha-1 Antitrypsin (AAT) and specific hyper-immune IgGs
- 2 FDA approved products
  - *Glassia® for AAT Deficiency (AATD); first FDA-approved liquid, ready-to-use IV AAT. Commercialized in the U.S. through Shire plc. Estimated revenues: \$177-\$228 MM (2018-2020); followed by 20 years of royalties.*
  - *KedRAB® for anti-rabies prophylaxis treatment. Commercialized in the U.S. through Kedrion Biopharma. Launched in Q1/2018.*

## BROAD PIPELINE/ IP

- Focused on global leadership in AATD
- Inhaled AAT for AATD
  - *Completed Ph2 (U.S.) and Ph2/3 (EU), MAA withdrawn June 2017;*
  - *EMA accepted new pivotal Phase 3 design;*
  - *FDA discussions ongoing re development path forward;*
  - *New pivotal Phase 3 pending IND/CTA approval*
- AAT IV for other indications developed through strategic collaborations
- Fully integrated propriety manufacturing technology for protein purification from human plasma
- Distributed biopharmaceutical products segment in Israel

## COMPELLING FINANCIAL PROFILE

- 2017 Revenue: \$102.8 MM (+33% YoY)
- 2018 Revenue Guidance: \$102-\$108MM<sup>1</sup>; Profitable; Cash flow positive
- Cash: \$44.6 MM (June 30, 2018); No Debt
- Strong balance sheet allows execution on pipeline and business development initiatives
- Listed on TASE (2005) & Nasdaq (2013)

1. The previously provided 2018 revenue guidance of \$116-\$120 million has been revised; As a result of a recently settled labor strike, part of 2018 planned product shipments are likely to be delayed and supplied in early 2019

# HIGH VALUE PRODUCT PORTFOLIO AND PIPELINE



Product	Indication	Phase I	Phase II	Phase III	Market
<b>Glassia® (IV AAT) *</b>	AAT Deficiency	FDA approved (2010)	----->		U.S. distribution through 
<b>KamRab®/KedRab® (IM Anti-Rabies)</b>	Prophylaxis for Rabies	FDA approved (2017)	----->		U.S. distribution through 

## Clinical Development

<b>Inhaled AAT</b>	AAT Deficiency <sup>1</sup>	EU Phase 2/3 (completed) MAA withdrawn (June 2017) EMA accept new Ph3 design US Phase 2 (completed) FDA review of path forward	----->		May seek partner upon IND/CTA approval
<b>G1-AAT (IV)</b>	Graft vs Host Disease (GvHD) <sup>1</sup>	Phase 1/2 (completed) Phase 2 (ongoing)	----->		Ph2 in collaboration with MAGIC <sup>3</sup>
<b>L1-AAT (IV)</b>	Lung Transplant	Phase 2 (ongoing)	----->		In collaboration with Shire plc
<b>D1-AAT (IV)</b>	Type 1 Diabetes <sup>2</sup>	Phase 2 (completed)	----->		Seeking partner for further development

## Early Stage Development

<b>Recombinant AAT</b>	AAT Deficiency	Early development			
<b>AAT (liquid)</b>	Organ preservation	Ex-Vivo Study			Massachusetts General Hospital

1. Orphan drug designation (US & EU); 2. Orphan drug designation (US only);  
3. Mount Sinai Acute GVHD International Consortium

# EXPERIENCED MANAGEMENT TEAM



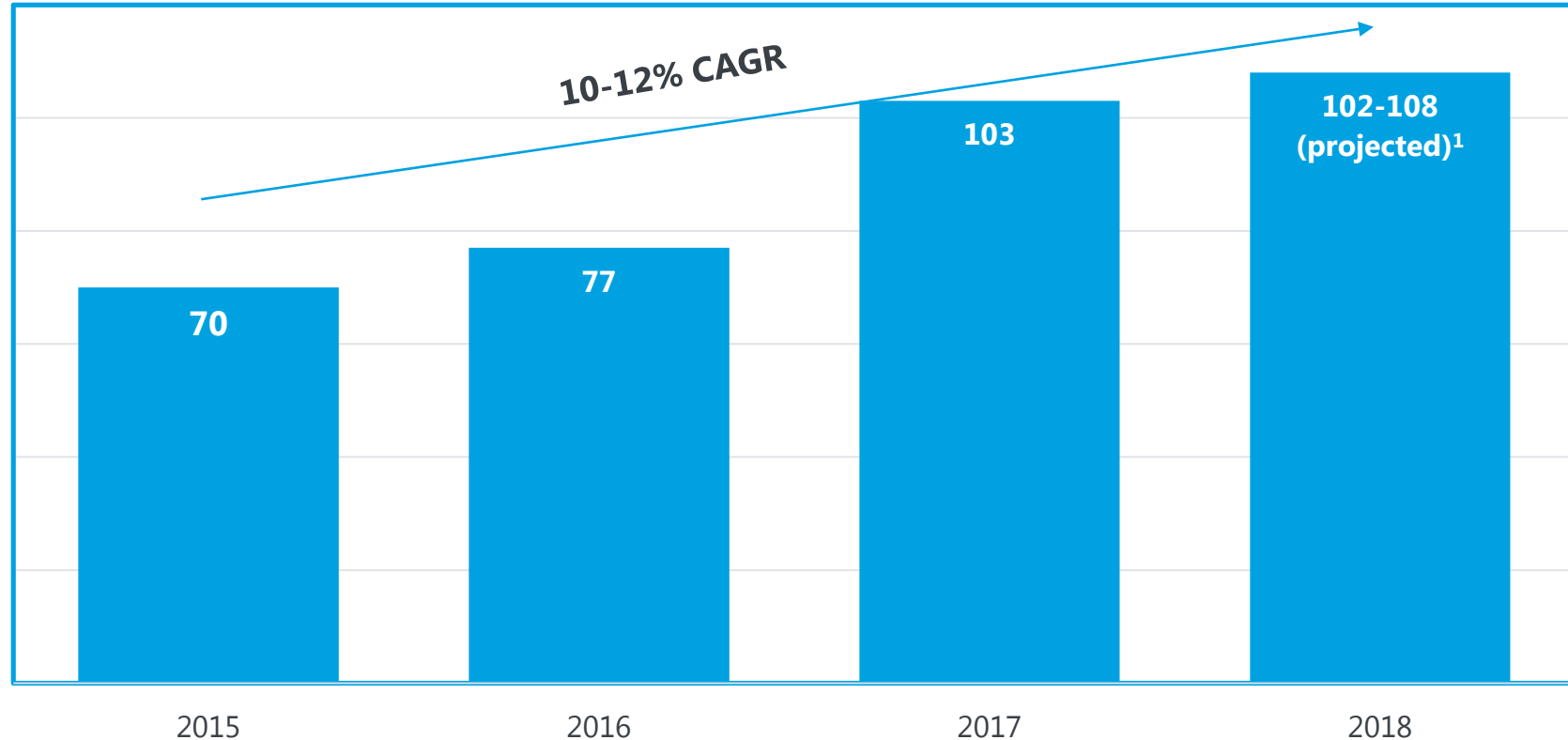
<b>Amir London</b>	CEO
<b>Chaime Orlev</b>	CFO
<b>Eran Nir</b>	VP Operations
<b>Yael Brenner</b>	VP Quality
<b>Liliana Bar, PhD</b>	VP Research and Development
<b>Orit Pinchuk</b>	VP Regulatory Affairs & PVG
<b>Naveh Tov, MD, PhD</b>	VP Clinical Development & Medical Director (Pulmonary)
<b>Michal Stein, MD</b>	VP Medical Director (Immunology)
<b>Ariella Raban</b>	VP Human Resources



# DEMONSTRATED STRONG SALES GROWTH DRIVEN BY GLASSIA



## Revenues US\$MM



1. The previously provided 2018 revenue guidance of \$116-\$120 million has been revised; As a result of a recently settled labor strike, part of 2018 planned product shipments are likely to be delayed and supplied in early 2019

# INCREASING REVENUE GROWTH AND PROFITABILITY



US \$ MM	FY 2015 Audited	FY 2016 Audited	% Change 2016/2015	FY 2017 Audited	% Change 2017/2016	1H 2017 Unaudited	1H 2018 Unaudited	% Change 2018/2017
Proprietary Products	43	56	30%	80	42%	34	38	14%
Distribution Products	27	21	-22%	23	8%	11	13	22%
<b>Total Revenues</b>	<b>70</b>	<b>77</b>	<b>10%</b>	<b>103</b>	<b>33%</b>	<b>44</b>	<b>51</b>	<b>16%</b>
<b>Gross Profit</b>	<b>15</b>	<b>21</b>	<b>39%</b>	<b>32</b>	<b>50%</b>	<b>14</b>	<b>18</b>	<b>26%</b>
<b>Gross Profit (%)</b>	<b>22%</b>	<b>28%</b>		<b>31%</b>		<b>32%</b>	<b>34%</b>	
R&D	(17)	(16)		(12)		(7)	(5)	
S&M and G&A	(10)	(11)		(13)		(6)	(6)	
<b>Operating Profit (Loss)</b>	<b>(11)</b>	<b>(5)</b>		<b>7</b>		<b>1</b>	<b>6</b>	
<b>Net Profit (Loss)</b>	<b>(11)</b>	<b>(7)</b>	<b>40%</b>	<b>7</b>	<b>204%</b>	<b>1</b>	<b>7</b>	<b>667%</b>
<b>Adjusted EBITDA<sup>1</sup></b>	<b>(6)</b>	<b>(1)</b>		<b>11</b>		<b>4</b>	<b>9</b>	

2017 Revenue Exceeded Guidance of \$100M

2018 Revenue Guidance of \$102-108M<sup>2</sup>;  
Continued Profitability and Positive Cash Flow

1. See Appendix A for Adjusted EBITDA reconciliation

2. The previously provided 2018 revenue guidance of \$116-\$120 million has been revised.

# Alpha-1 Antitrypsin Deficiency (AATD)





# AAT DEFICIENCY

Potentially Lethal and Often Undiagnosed



AAT  
Level



**Genetic/Hereditary  
condition causing  
decreased levels of  
AAT in blood and  
tissues**

U.S. ●



● EU



**Affects more than  
100,000 people in  
the U.S. and slightly  
lower number in  
Europe**

AAT  
Deficiency



**Predisposes to lung  
and liver diseases**

AAT deficiency-associated lung disease is characterized by airway obstruction and destructive changes in the lungs (Emphysema)

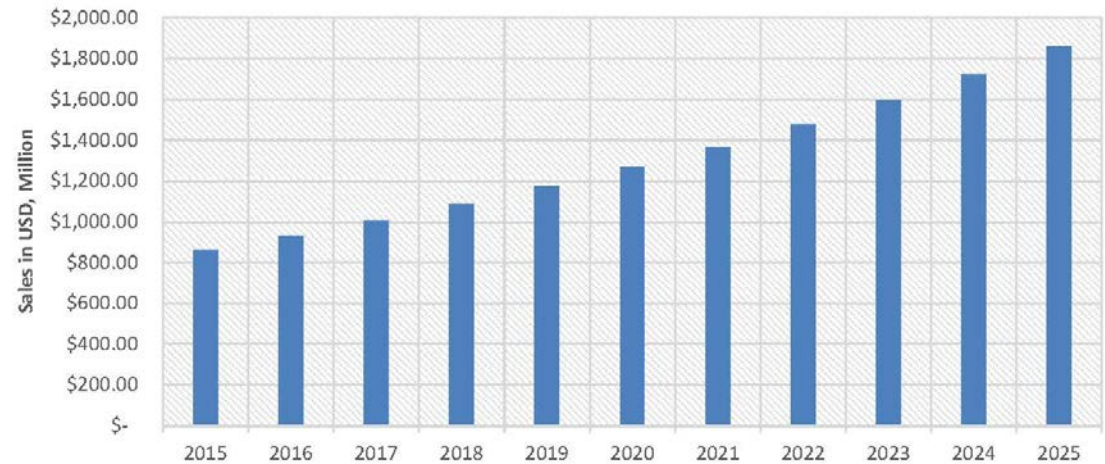
# AAT DEFICIENCY (AATD) MARKET

## Significant expansion opportunity



- Majority of patients suffering from AATD still remain undiagnosed & untreated
- Better diseases awareness and expanded diagnostics is contributing to increased demand
- Expected growth of U.S. market – approx. 6-8% annually<sup>1</sup>
- Chronic therapy creates sustainable product revenue opportunity
- Average annual reimbursement (U.S.) ~\$80-\$100K per patient
- Greater AAT use in Europe and other regions could further accelerate market growth

Global Market Size(2015-2025) <sup>1</sup>



Source: DelveInsight

- **AATD prevalence <sup>1</sup>: ~115,000 (U.S.); ~72,000 (EU5)**  
but only ~7,300 (U.S.) or ~1,800 (EU5) patients are treated <sup>1</sup>
- **Current market size is ~ \$1B WW**
- **Expected to reach \$1.8B by 2025**

**GLASSIA®:**  
**Liquid AAT**  
**for the**  
**Treatment**  
**of AAT**  
**Deficiency**



# GLASSIA® IS A DIFFERENTIATED PRODUCT



- Glassia® is the first liquid, FDA-approved ready-to-use, plasma-derived AAT product:
  - No reconstitution required
  - Reduces treatment time
  - Reduces risk of contamination and infection
- Kamada's highly purified liquid product is manufactured through a proprietary process
- Glassia® is sold in the U.S. by Shire plc
- Self-infusion approved by FDA in 2016

# GROWTH OF GLASSIA® DRIVEN BY STRATEGIC PARTNERSHIP WITH SHIRE

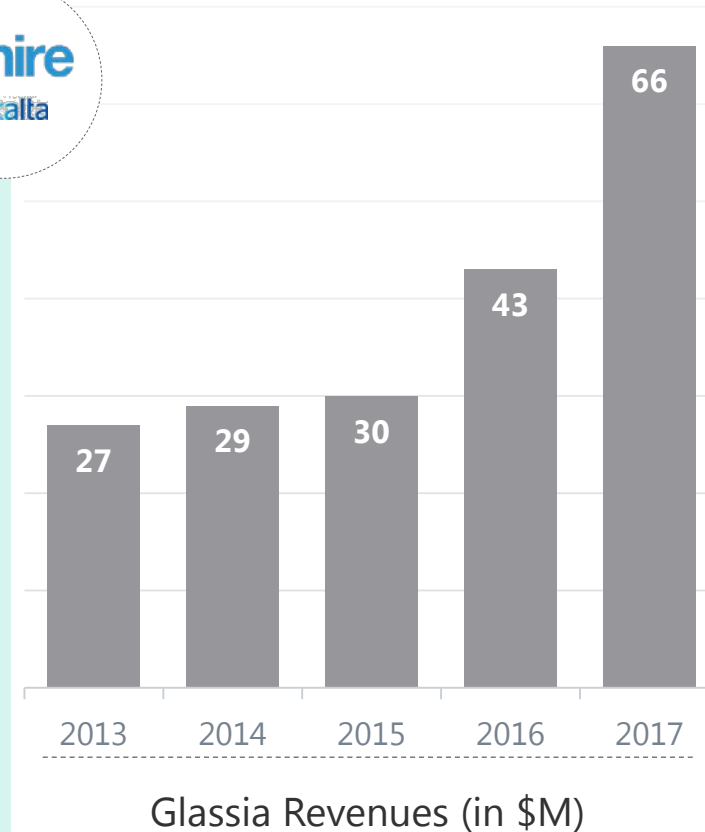


## Significant Revenues to Kamada through 2020, followed by 20 Years of Royalties

- Minimum/max revenues of \$177MM/\$228MM to Kamada expected for 2018-2020
- Kamada manufactures and supplies Glassia to Shire through 2020
- Commencing in 2021, Shire has option to manufacture Glassia and pay royalties to Kamada through 2040
- Territories – U.S., Canada, Australia, and New Zealand
- Agreement covers all future AAT-IV indications in the territories



Glassia® is sold in 5 countries, with majority of sales in the U.S.



**KamRAB/  
KedRAB:  
Human  
Rabies  
Immune  
Globulin**



# KamRAB/KedRAB

## Human Rabies Immune Globulin



### U.S. Opportunity:

Strategic agreement with Kedrion for the clinical development and marketing of KedRAB in U.S.

**KEDRION**  
BIOPHARMA

### Substantial WW Market (WHO estimates)

~10 million people worldwide require medical treatment against rabies each year after being exposed to an animal suspected of rabies infection

U.S.

### U.S. Market

- FDA Approval - August 2017
- Product launch: Q1/2018 in collaboration with Kedrion
- **~40,000 post-exposure prophylaxis treatments** administered each year, representing **~\$100 million market opportunity<sup>1</sup>**
- **Limited competition in US**

### Worldwide

- More than 1.5MM vials sold by Kamada to date = **~ 300,000 people treated WW**
- Major markets: India, Thailand, Israel, Russia
- Approved Supplier of the WHO
- November 2017: Signed new \$13 MM supply agreement with an international organization for 2018-2020



1. The plasma protein market in the United States, 2015, The Marketing Research Bureau Inc

**Inhaled  
AAT  
to Treat  
Alpha-1  
Antitrypsin  
Deficiency  
(AATD)**








**Alpha-1 Foundation survey<sup>1</sup> confirms high level of patients' interest in Inhaled-AAT**


**Inhaled AAT opportunity is estimated by Kamada at over \$1B world wide**



**Improved Quality Of Life (QOL)**



**ELF<sup>2</sup> levels 2-5 fold greater than IV**

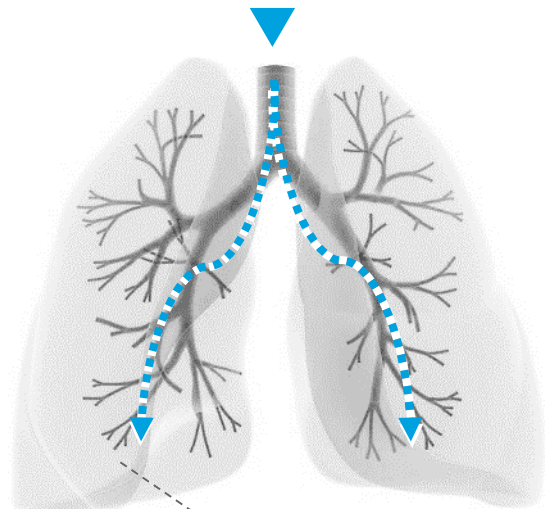


**Most effective mode of treatment for reaching primary site of injury<sup>3</sup>**

# INHALATION ENABLES DELIVERY OF AAT 5X HIGHER THAN INTRAVENOUS

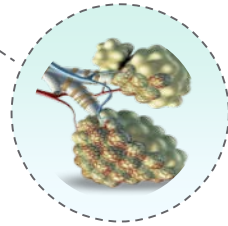


**Inhaled AAT ~ 50%** of the dosage reaches **airway tree** and alveoli

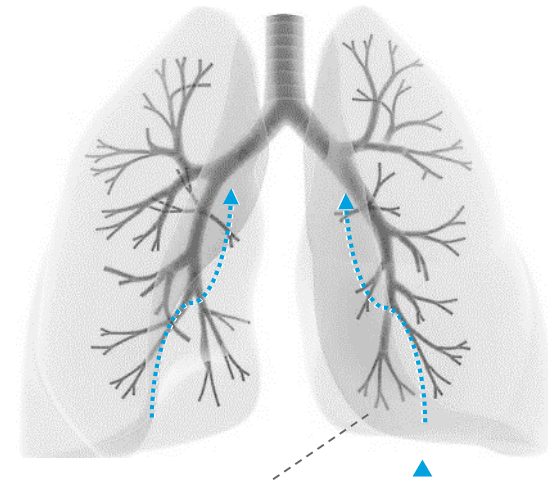


## Airway Obstruction

- FEV1/SVC<sup>1</sup>
- FEV1

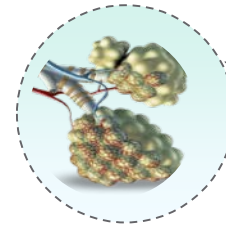


**Intravenous AAT ~ 2%** of the dosage reaches alveoli and **airway tree**



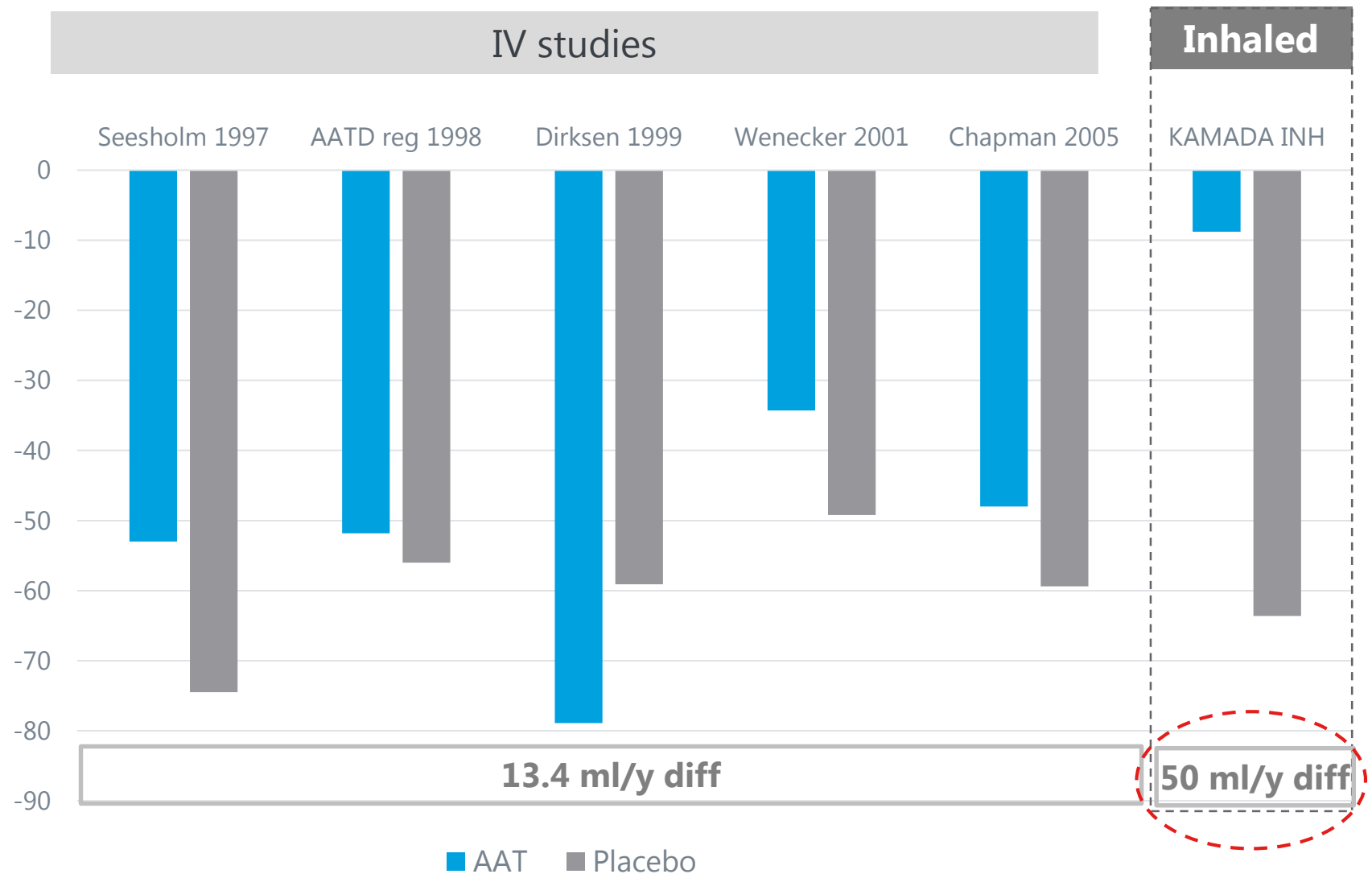
## Alveoli Emphysema

- DLCO
- CT densitometry



1. FEV = Forced Expiratory Volume. SVC = Slow Vital Capacity.

# INHALED AAT SLOWED FEV1<sup>1</sup> DETERIORATION BETTER THAN PREVIOUS AAT-IV TRIALS



1. FEV = Forced Expiratory Volume



## U.S.



- Ongoing discussions with the FDA addressing concerns and questions regarding the safety and efficacy of Inhaled AAT for the treatment of AATD
- Revised proposed Ph3 protocol and additional information provided to the FDA during Q3 2018
- Continued clinical development pending IND approval by FDA

## EU



EUROPEAN MEDICINES AGENCY  
SCIENCE MEDICINES HEALTH

- Phase 2/3 completed; Study endpoints were not met; statistical significant lung function improvement was observed
- MAA submitted based on data showing Lung Function Improvements; MAA withdrawn (June 2017) EMA viewed data as insufficient for approval
- Proposed new Ph3 protocol accepted by EMA in a Scientific Advice meeting held July 2018

Considering all strategic options for Inhaled-AAT, including seeking a partner

# IMMUNE-MODULATORY INDICATIONS



01

**Graft versus  
Host Disease**



02

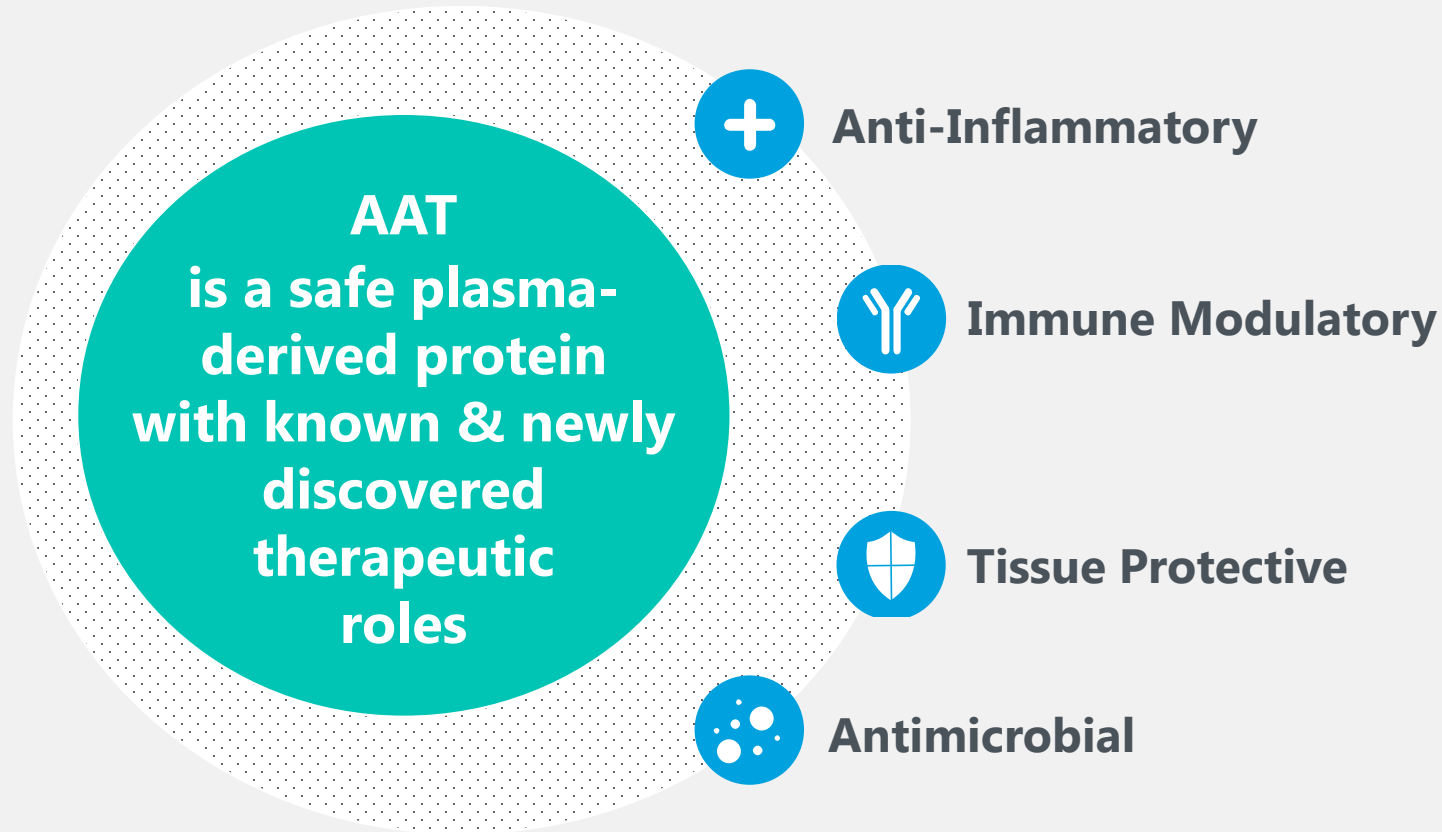
**Lung  
Transplantation**



03

**Type-1  
Diabetes**

# AAT REPRESENTS AN EXCITING POTENTIAL THERAPY FOR MULTIPLE INDICATIONS



Excellent safety profile, encouraging clinical and pre-clinical experience coupled with biochemical rationale may position AAT as a high-potential future treatment in various indications.



**01**

## **AAT to Treat Graft versus Host Disease**

# GRAFT VERSUS HOST DISEASE (GVHD):

A Major Complication in Hematopoietic Cell Transplantation



## DEADLY SIDE EFFECTS

**30-40%**

of bone marrow transplantations will develop acute GvHD

**40-50%**

of acute GvHD will not respond to steroid treatment (SR-aGvHD)

**~70%**

mortality rate of patients with SR-aGvHD



## SEARCHING FOR AN EFFECTIVE TREATMENT

Standard of prophylaxis care exhibits poor efficacy/severe AE's

No FDA-approved specific drug for GvHD indication – An Unmet Medical Need

Estimated Market Size<sup>1</sup>: ~ **\$500 MILLION**

1. Company estimates



# AN EARLY-BIOMARKER PREDICTS LETHAL GVHD



Study performed by Matthew J. Hartwell, et al., the Icahn School of Medicine at Mount Sinai <sup>1</sup> suggests that a biomarker algorithm can identify patients at high risk of lethal GvHD and non-relapse mortality in advance of symptoms onset

## Background

No laboratory test can predict the risk of non-relapse mortality (NRM) or severe GvHD after hematopoietic cellular transplantation (HCT) prior to the onset of GVHD symptoms.

## Method

Patient blood samples on day 7 after HCT were obtained from a multicenter set of 1,287 patients, and 620 samples were assigned to a training set. We measured the concentrations of 4 GVHD biomarkers (ST2, REG3 $\alpha$ , TNFR1, and IL-2R $\alpha$ )

## Results

A 2-biomarker model (ST2 & REG3 $\alpha$ ) concentrations identified patients with a cumulative incidence of 6-month NRM of 28% in the high-risk group and 7% in the low-risk group ( $P < 0.001$ ). GVHD-related mortality was greater in high-risk patients (18% vs. 4%,  $P < 0.001$ ), as was severe gastrointestinal GVHD (17% vs. 8%,  $P < 0.001$ ). The same algorithm can be successfully adapted to define 3 distinct risk groups at GVHD onset.

## Conclusion

A biomarker algorithm based on a blood sample taken 7 days after HCT can consistently identify a group of patients at high risk for lethal GVHD and NRM.

# COLLABORATION WITH MAGIC<sup>1</sup> TO EVALUATE AAT FOR PREEMPTION OF GVHD



## Proof-of-Concept Study:

- Open label single arm multicenter study to be conducted in 5 US centers which are members of Mount Sinai Acute GVHD International Consortium (MAGIC)<sup>1</sup>
- Study is co-funded by Mount Sinai and Kamada, and is sponsored by the Icahn School of Medicine at Mount Sinai (ISMMS) and Led by Prof James L.M. Ferrara, MD, and Prof. John Levine, MD, MS
- Top-line results expected to be available during the second half of 2019

Kamada has exclusive rights to develop and commercialize AAT for preemption of GvHD using the MAGIC Biomarkers

## Innovative approach

- Biomarker based algorithm to diagnose patients at risk to develop steroid-resistant GvHD (SR-GvHD) at day 7 after bone marrow transplantation (BMT).
- Early intervention could prevent patients from further disease deterioration

## Study objective

- To assess the safety and preliminary efficacy of IV AAT as preemptive therapy in patients at high risk for the development of SR-GvHD after BMT

## Design

- 30 patients treated with IV AAT for 2 months with a follow-up period of 1 year after BMT

## Endpoints

- Proportion of High Risk patients who develop SR-GvHD by day 100 post BMT, as well as safety, severity of GvHD, mortality, etc.

1. A consortium of 23 BMT centers in the USA, Europe and Asia that conducts clinical trials to prevent and treat acute GVHD (aGvHD).



**02**

## **AAT to Treat Lung Transplantation**

# ADVANCING THE LUNG TRANSPLANTATION OPPORTUNITY



## Lungs have the highest rate of rejection among transplanted solid organs

- ~33% will experience acute rejection within the first year
- ~50% will develop chronic rejection within the first 5 years

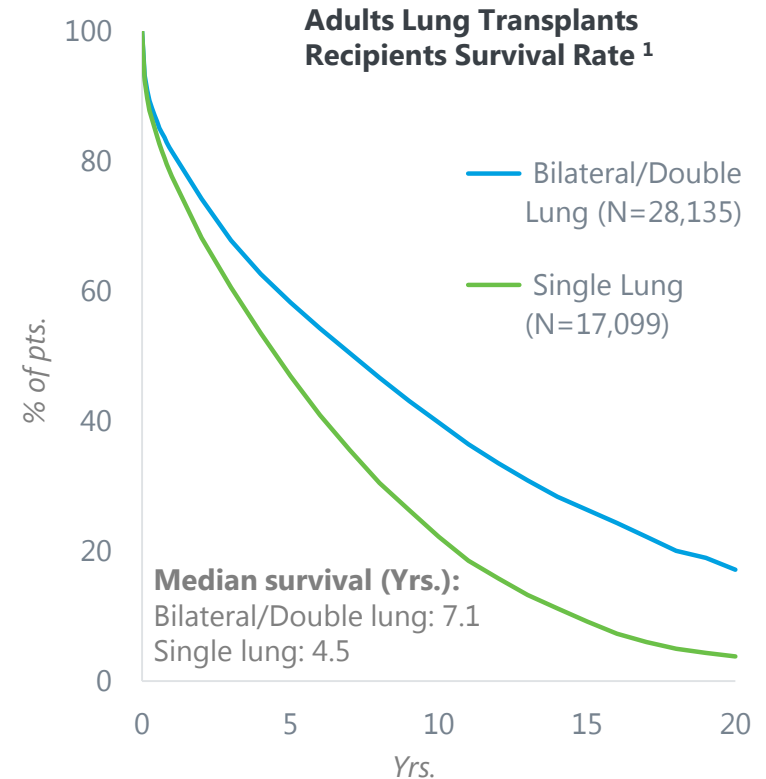


## No new treatment options have been made available for years

Physician feedback indicates strong need for improved post-transplant therapies over existing options (toxicity, immunosuppressive)



## Kamada initiated the first clinical trial designed specifically to prevent lung transplant rejection



**Potential Market Size<sup>2</sup> ~ \$400-500 MILLION**

# ON-GOING PHASE II STUDY WITH AAT IV

## For Prevention of Lung Transplant Rejection



### Phase II:

- Prospective, open label, standard-of-care (SOC) controlled, randomized, parallel group single center study
- In collaboration with Baxalta/Shire led by Prof. Mordechai Kramer, Rabin Medical Center, Israel
- Next interim report will include data from one-year of treatment for all patients, and is expected in H2 '18
- Top-line data from the Phase 2 trial anticipated in H2 '19.

### Study objective

- To assess the safety of AAT IV and the effect on rate and severity of acute and chronic lung rejection as well as pulmonary infections, in subjects undergoing first lung transplantation

### Design

- 30 lung transplant recipients randomized 2:1 to receive AAT IV on top of standard-of-care (SOC) or SOC alone, for 48 weeks plus 12 months of follow-up period

### Endpoints

- Safety: Related adverse events (AEs)
- Efficacy: Changes in FEV1 from baseline and overall effect, incidence and rate of acute lung rejection

### Interim results (16 Pts; 6 months)

- IV AAT demonstrated favorable safety and tolerability profile in 10 patients during first six months of treatment, consistent with previously observed results in other indications.



**03**

## **AAT to Treat Newly Diagnosed Type-1 Diabetes**

# AAT (IV) AS POTENTIAL TREATMENT FOR NEWLY DIAGNOSED TYPE-1 DIABETES (T1D) PATIENTS



MARKET OPPORTUNITY	AAT IMPACT	EXPECTED BENEFITS
<p><b>Type-1 Diabetes</b> Occurs when the immune system attacks and destroys beta cells in the pancreas</p>	<p><b>Studies have shown that AAT protects beta cell islets</b></p>	<p><b>Preservation of beta cells correlates with reduced risk of long-term complications</b></p>
<ul style="list-style-type: none"><li>• More than 10 million suffer from Type 1 diabetes globally</li><li>• 100,000 new patients/year diagnosed globally</li><li>• In the U.S. alone: 3 million patients, with 30,000 new patients diagnosed annually<sup>1</sup></li></ul>	<ul style="list-style-type: none"><li>• Delays the progression of autoimmune diabetes</li><li>• Inhibits insulinitis and beta-cell apoptosis</li><li>• Decreases beta-cell inflammation</li></ul>	<ul style="list-style-type: none"><li>• DCCT<sup>2</sup> indicated that patients with C-peptide on MMTT <math>\geq 0.2</math> pmol/mL were less likely to develop retinopathy and hypoglycemia complications<sup>3</sup></li><li>• Higher / sustained levels of C-peptide correlate with reduced incidences of the microvascular complications<sup>3</sup></li></ul>

1. JDRF publication; 2. The Diabetes Control and Complications Trial (DCCT)  
3. Greenbaum et al, 2012; 3. Steffes et al, 2013



## Phase II Completed: Double-Blind, Randomized, Placebo-Controlled, Multicenter Study



### Study objective

- To evaluate efficacy and safety of AAT) in treatment of newly diagnosed Type 1 Diabetes patients

### Design

- Two doses, placebo controlled, randomized with 70 pediatric and young adult patients. One year study

### Endpoints

- Beta cell preservation (C-peptide AUC), HbA1C, hypoglycemic events and insulin daily dose

### Results

- In overall study population, no significant treatment effect was observed.
- In the pre-determined subgroup of patients between the ages of 12-18 years old, a trend toward better efficacy was demonstrated in the high dose arm of AAT (120mg/kg)
- Presented results in an oral session at 78th Scientific Sessions of the American Diabetes Association (ADA)

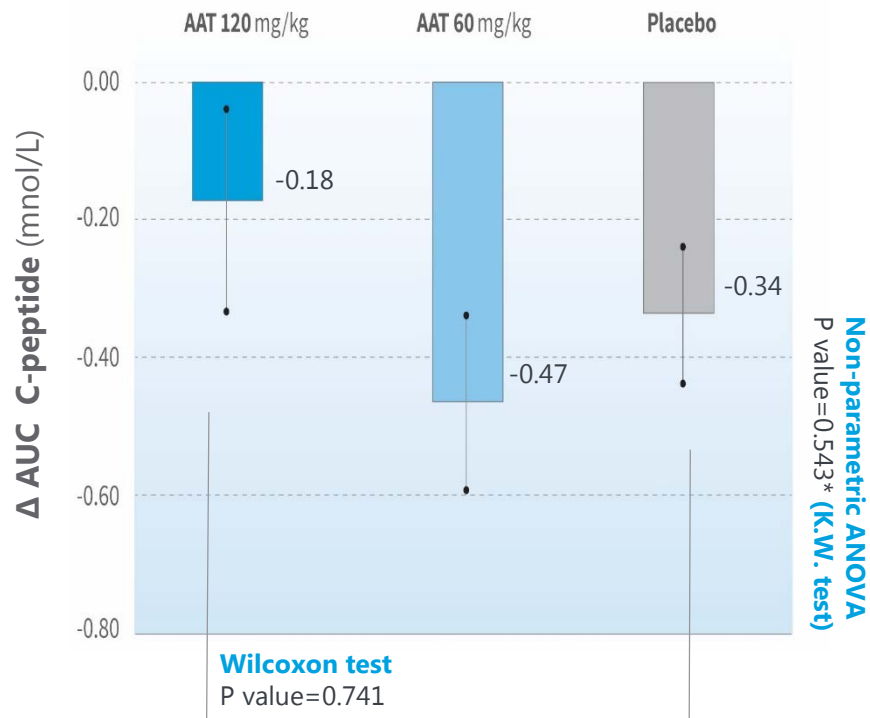


# BETA CELL FUNCTION AND INSULIN AT 1 YEAR

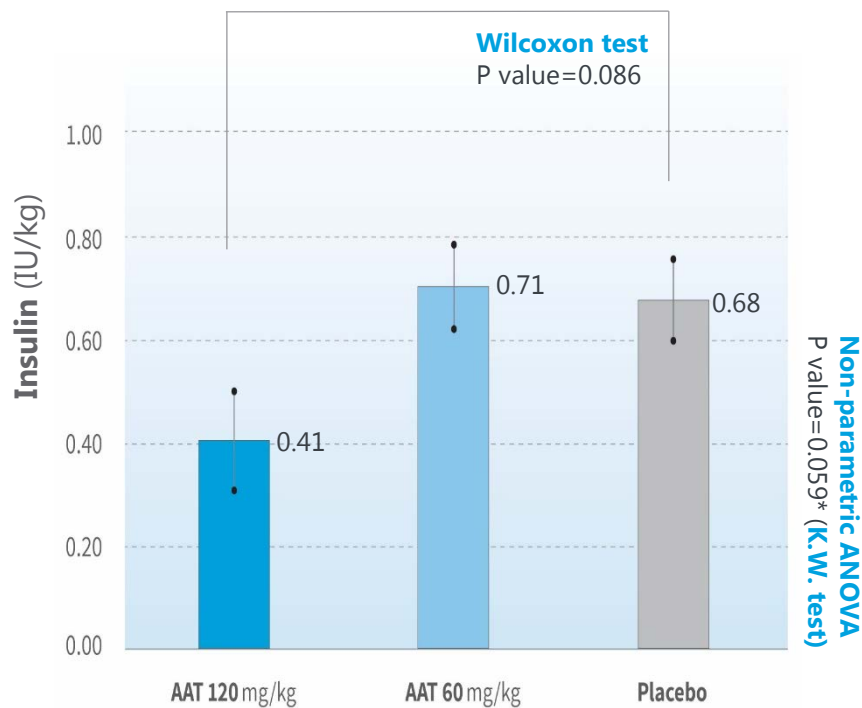
## Subgroup Analysis, Ages 12-18



### Beta-Cell Function by MMTT AUC C-peptide at 1 Year $\Delta$ Stimulated AUC



### Insulin Requirement at 1 Year

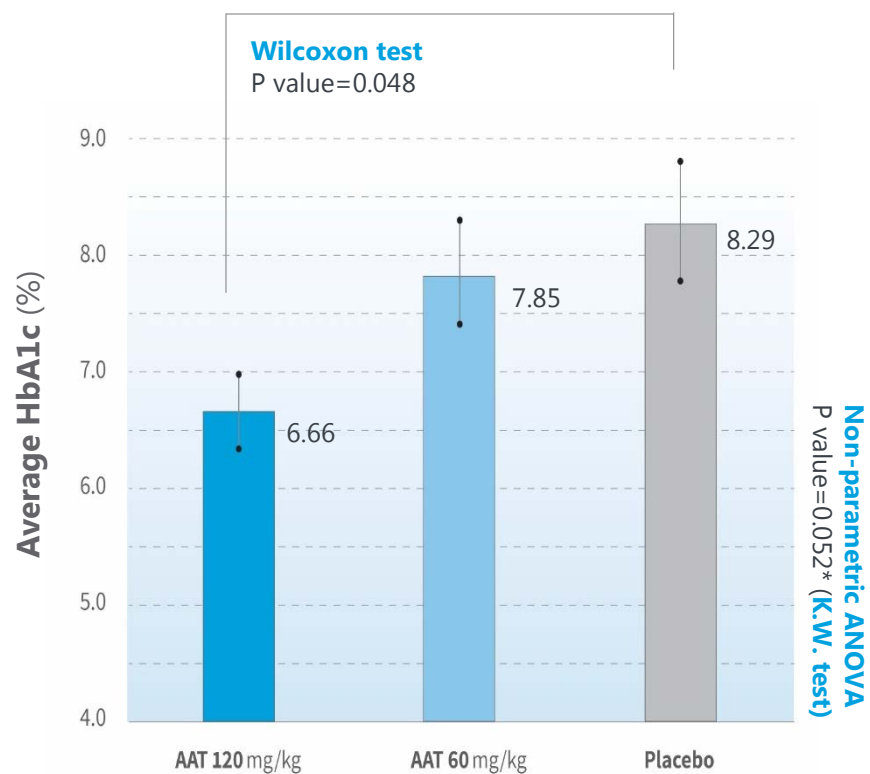


# GLYCEMIC CONTROL RESULTS AT 1 YEAR

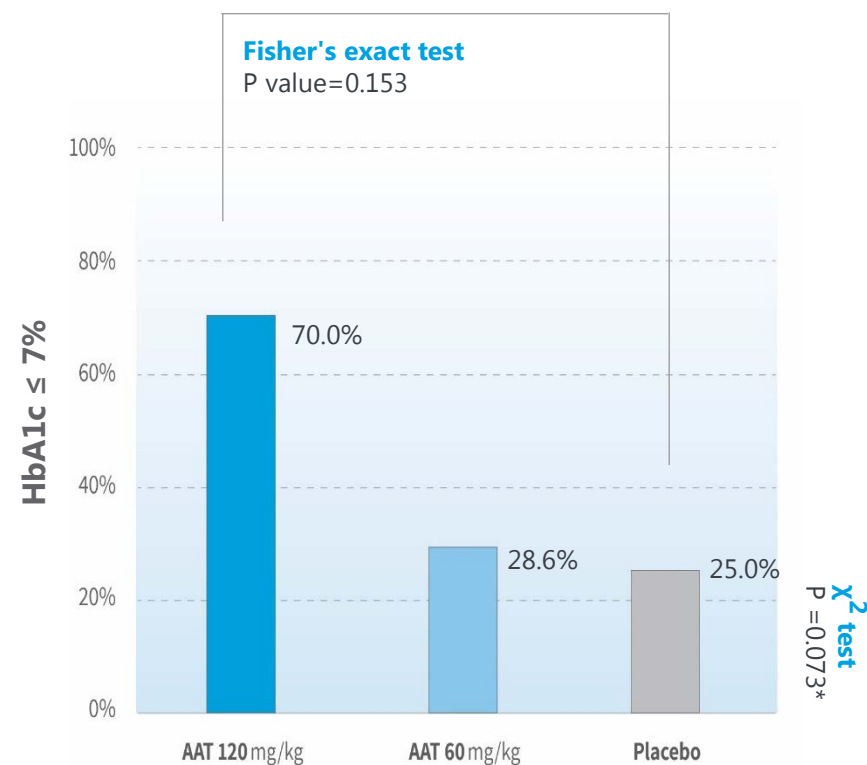
Subgroup Analysis, Ages 12-18



## % HbA1c



## Patients with HbA1c ≤ 7%





- Full data set analysis and publication
- Regulatory advice (US/EU)
- Seek strategic partner for collaboration in further product development



**Distribution  
Product  
Segment**



# DISTRIBUTION SEGMENT

Exclusive distributor in Israel of leading biopharmaceutical companies



Medical Field	Product/Brand Name	Description
Immunology	Intratect & Gammaplex <sup>1</sup>	Gamma-globulins 5% IV
Hospital & Critical Care	Vialabex/Zenalb/Albiomin <sup>1</sup>	Human serum Albumin
	Heparin sodium injection	Heparin sodium 5000 IU/ml
Infectious Disease	Ixiaro	Japanese encephalitis vaccine
	Varitect <sup>1</sup>	Varicella zoster IgG
	Megalotect <sup>1</sup>	CMV IgG
Hematology and Hemophilia	Optivate <sup>1</sup>	Coagulation Factor VIII (human)
	Replenine <sup>1</sup>	Coagulation Factor IX (human)
Respiratory	Foster	Beclometasone+ Formoterol inhaled
	Bramitob	Tobramycin, inhaled
	Provocholine	Methacholine, inhaled
Liver	Zutectra <sup>1</sup>	Hepatitis B IgG S.C
	Hepatect <sup>1</sup>	Hepatitis B IgG I.V

Additional products are under registration with the Israeli MOA



# EXPECTED 2018 MILESTONES



<b>Initiating next GvHD study in collaboration with Mt. Sinai Hospital and the MAGIC consortium</b>	<b>✓ Q1/2018</b>
<b>Rabies product launch in the U.S.</b>	<b>✓ Q1/2018</b>
<b>Inhaled AAT for AATD: Scientific Advice in EU</b>	<b>✓ Q3/2018</b>
<b>Inhaled AAT for AATD: Continued FDA discussion targeting approved IND</b>	<b>Q3/2018</b>
<b>Initiating inhaled AAT for AATD phase III study</b>	<b>Post IND/CTA approval</b>
<b>Interim report for Phase II for lung transplant trial (1 year treatment)</b>	<b>2H/2018</b>
<b>Seeking collaboration to advance type-1 diabetes program</b>	<b>2018</b>
<b>Achieve \$102-108 million in annual revenues, profitable, cash flow positive</b>	<b>2018</b>



# INVESTMENT HIGHLIGHTS



## COMMERCIAL STAGE BIOPHARMA

- Leader in plasma-derived protein therapeutics, focused on Alpha-1 Antitrypsin (AAT) and specific hyper-immune IgGs
- 2 FDA approved products
  - *Glassia® for AAT Deficiency (AATD); first FDA-approved liquid, ready-to-use IV AAT. Commercialized in the U.S. through Shire plc. Estimated revenues: \$177-\$228 MM (2018-2020); followed by 20 years of royalties.*
  - *KedRAB® for anti-rabies prophylaxis treatment. Commercialized in the U.S. through Kedrion Biopharma. Launched in Q1/2018.*

## BROAD PIPELINE/ IP

- Focused on global leadership in AATD
- Inhaled AAT for AATD
  - *Completed Ph2 (U.S.) and Ph2/3 (EU), MAA withdrawn June 2017;*
  - *EMA accepted new pivotal Phase 3 design;*
  - *FDA discussions ongoing re development path forward;*
  - *New pivotal Phase 3 pending IND/CTA approval*
- AAT IV for other indications developed through strategic collaborations
- Fully integrated propriety manufacturing technology for protein purification from human plasma
- Distributed biopharmaceutical products segment in Israel

## COMPELLING FINANCIAL PROFILE

- 2017 Revenue: \$102.8 MM (+33% YoY)
- 2018 Revenue Guidance: \$102-\$108MM<sup>1</sup>; Profitable; Cash flow positive
- Cash: \$44.6 MM (June 30, 2018); No Debt
- Strong balance sheet allows execution on pipeline and business development initiatives
- Listed on TASE (2005) & Nasdaq (2013)

1. The previously provided 2018 revenue guidance of \$116-\$120 million has been revised; As a result of a recently settled labor strike, part of 2018 planned product shipments are likely to be delayed and supplied in early 2019



**THANK YOU**  
[www.kamada.com](http://www.kamada.com)





## **Appendix A: Reconciliation of Non-IFRS Measures**

Adjusted EBITDA is defined as net income (loss), plus income tax expense, plus financial expense, net, plus depreciation and amortization expense, plus non-cash share-based compensation expenses, plus or minus income or expense in respect of exchange and translation differences and derivatives instruments not designated as hedging, and plus one-time management compensation payment.

We present adjusted EBITDA because we use this non-IFRS financial measure to assess our operational performance, for financial and operational decision-making, and as a means to evaluate period-to-period comparisons on a consistent basis. Management believes this non-IFRS financial measure is useful to investors because: (1) they allow for greater transparency with respect to key metrics used by management in its financial and operational decision-making; and (2) they exclude the impact of non-cash items that are not directly attributable to our core operating performance and that may obscure trends in the core operating performance of the business.

Non-IFRS financial measures have limitations as an analytical tool and should not be considered in isolation from, or as a substitute for, our IFRS results. We expect to continue reporting non-IFRS financial measures, adjusting for the items described below, and we expect to continue to incur expenses similar to certain of the non-cash, non-IFRS adjustments described below. Accordingly, unless otherwise stated, the exclusion of these and other similar items in the presentation of non-IFRS financial measures should not be construed as an inference that these items are unusual, infrequent or non-recurring. Adjusted EBITDA is not a recognized term under IFRS and does not purport to be an alternative to any other IFRS measure. Moreover, because not all companies use identical measures and calculations, the presentation of adjusted EBITDA may not be comparable to other similarly titled measures of other companies.

(US\$K, Unaudited)	YE2015	YE2016	YE2017
Net Income (Loss)	(11,270)	(6,733)	6,901
Taxes on income	0	1,722	269
Financial expenses (income), net	471	(343)	(338)
Depreciation and amortization expense	3,227	3,501	3,523
Share-based compensation charges	1,907	1,071	483
Expense (income) in respect of currency exchange and translation differences and derivatives instruments, net	(625)	(127)	612
Adjusted EBITDA	<u>(6,290)</u>	<u>(909)</u>	<u>11,450</u>

# INHALED AAT – IN THE WORDS OF THE KEY OPINION LEADERS



## EU Phase 2/3:

**“The study results demonstrated primarily that the overall treatment effect on lung functions, is of significant clinical value. This study is the first study ever that is indicative of inhaled AAT’s ability to potentially reduce lung inflammation as expressed by its preservation of lung function and the changes shown in symptoms.”**

*Prof. Jan Stolk, MD,*  
Department of Pulmonology,  
Leiden University Medical  
Center, Principal Investigator  
of the Phase 2/3 clinical trial  
and acting Chairman of the  
Alpha 1 International Registry  
(AIR)

**“The study analysis suggests exciting results that may lead to wider acceptance of the inhaled route of administration of alpha- 1 antitrypsin augmentation therapy, which could be a real breakthrough for AATD patients.”**

*Robert A. Sandhaus, Ph.D., M.D., FCCP,* Founder and  
Director of the Alpha1-Antitrypsin Deficiency  
Program at National Jewish Health in Denver,  
Colorado, and the Clinical Director of the Alpha-1  
Foundation

**“These new analyses confirm the clinically-meaningful lung function improvement seen with inhaled AAT patients in this study. These results are impressive and underscore the initial findings from this study. In my opinion, inhaled AAT has shown to be an efficacious treatment for this orphan disease.”**

*Prof. Kenneth Chapman, M.D.,* Director of the  
Canadian Registry for the Alpha-1 Antitrypsin  
Deficiency (Asthma and Airway Centre in Toronto  
Western Hospital, University of Toronto) and an  
investigator in the Phase 2/3 clinical trial.

## US Phase 2:

**“The results of this study are extremely compelling. Based on the results of this study, it is clear that inhaled AAT is the most effective mode of treatment for reaching the primary sites of potential lung injury, and restoring AAT inhibitory capacity. I look forward to the start of a pivotal study in the U.S. to confirm these results.”**

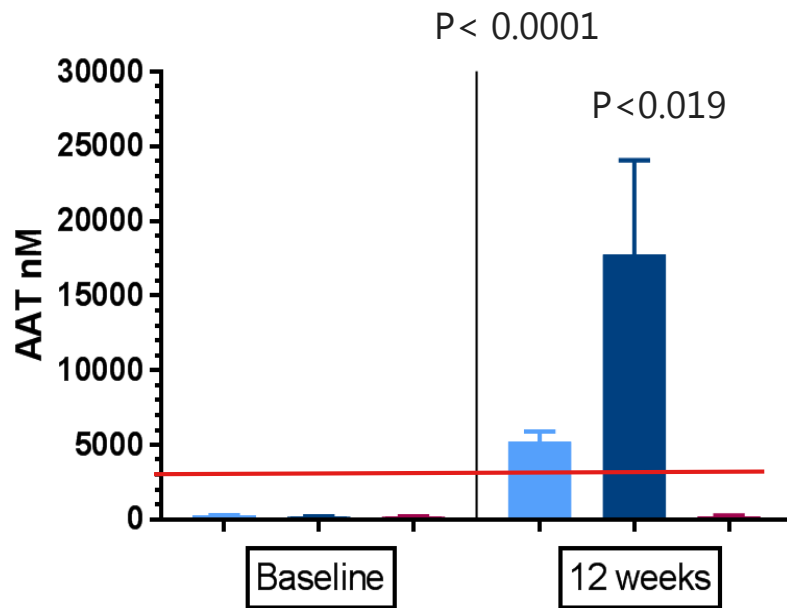
*Professor Mark Brantly, MD,* the  
Primary Investigator in this study  
who serves as a Vice Chair of  
Research, Department of Medicine,  
Chief Division of Pulmonary, Critical  
Care and Sleep Medicine, Professor  
of Medicine, Molecular Genetics  
and Microbiology at the University  
of Florida College of Medicine and  
Alpha One Foundation Research  
Professor.

# INHALED AAT PHASE II U.S.

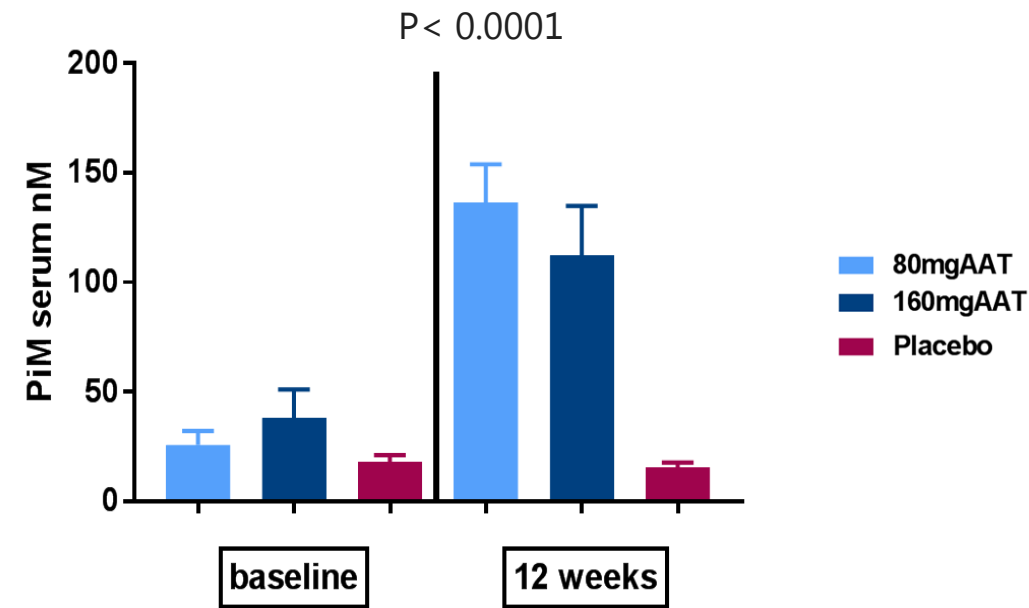
ELF AAT Antigenic Level & Inhibitory Capacity Increased Significantly



## ELF<sup>1</sup> AAT Antigenic Level



## PiM serum level



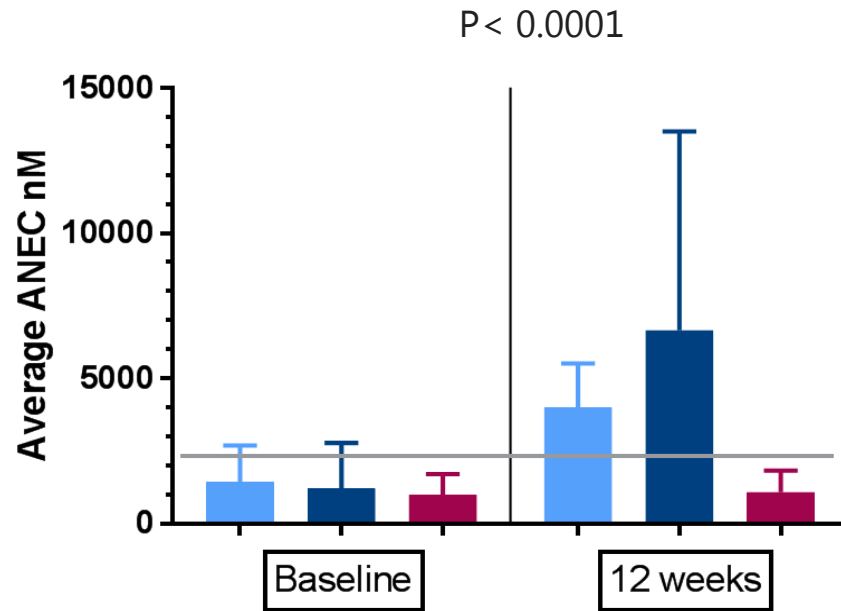
Higher AAT ELF level is reasonably likely to predict clinical benefit

# INHALED AAT PHASE II U.S.

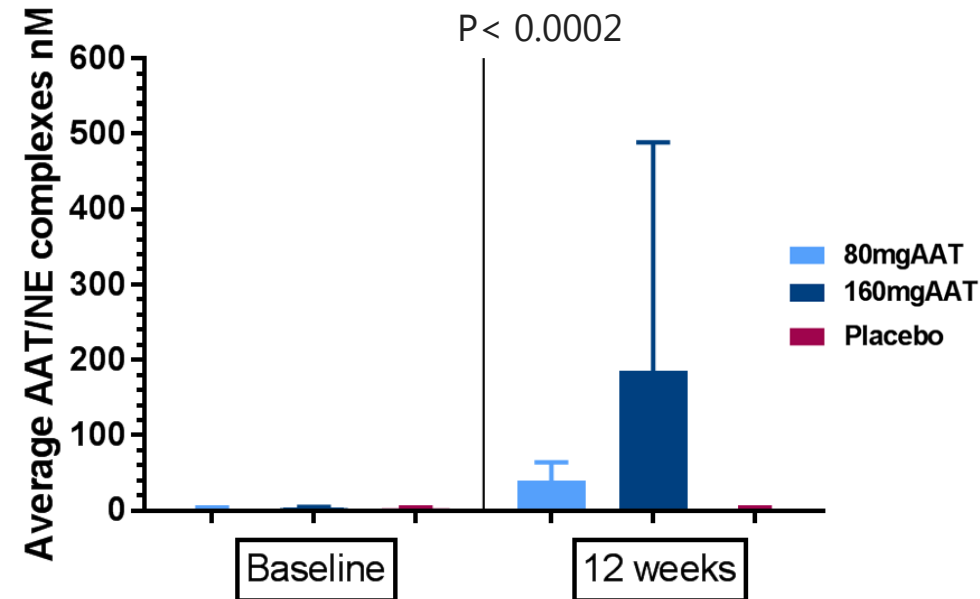
ELF AAT-NE  $\epsilon$ C Complexes & Inhibitory Capacity Increased Significantly



## ANEC<sup>1</sup>



## ELF AAT – NE Complexes



Inhaled AAT is the most effective means to restore AAT inhibitory capacity in the airways (ANEC<sup>1</sup> & AAT-NE Complexes)

# GVHD PROOF-OF-CONCEPT STUDY WITH AAT (IV)

For Graft-Versus-Host Disease (published 1/2016)



## Phase I/II study:

Open label of 24 patients with steroid-resistant GvHD bone- following allogeneic marrow stem cell transplant

### Study Design

4 dose groups - 15 day regimen.  
Doses given on days:  
1, 3, 5, 7, 9, 11, 13 and 15

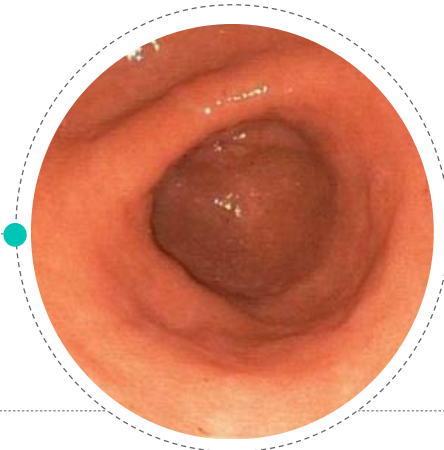
### Primary Endpoint

% of patients who experience no toxicity and in whom GVHD is stable or improved

### Results

- Encouraging preliminary clinical results;
- Stool AAT levels showed a decrease in intestinal AAT loss, suggesting healing of the bowel mucosa

**BEFORE** Duodenitis suspect severe upper and lower GVHD



**AFTER 8 DOSES OF AAT** Moderate mucosal denudement and edema noted throughout the duodenum



# “AAT COULD BE AN EFFECTIVE TREATMENT OPTION FOR NEWLY DIAGNOSED 12-18 YEARS OLD T1D PATIENTS”



**Peter Gottlieb, M.D.,**  
Professor of Pediatric and Medicine,  
Barbara Davis Center for Diabetes,  
University of Colorado School of  
Medicine and a leading member in  
TrialNet, an NIH-sponsored network of  
institutions and researchers dedicated  
to the prevention of type-1 diabetes.

“Given this study was not powered to show efficacy, the results are very encouraging.

These findings suggest that **administration of AAT could be an effective treatment option for newly diagnosed T1D patients who are 12-18 years old.** The results of this subgroup are intriguing and warrant further studies in a larger population.

Subgroup segmentation by age is common in this complicated disease, and the fact that we see the same positive trend in this age group for all three measures – C-peptide, daily insulin requirement, and HbA1C – suggests that the **results are consistent and could be promising.**”