



**CORPORATE PRESENTATION**  
**NOVEMBER 2020**

# SAFE HARBOR

This presentation contains forward-looking statements within the meaning of the Israeli securities law that involve risks and uncertainties. These forward-looking statements relating to future events and future performance of the Company and the portfolio companies, jointly or separately, such as statements regarding, but are not limited to, market opportunities, strategy, competition, the further development and potential safety and efficacy of the products, the projected revenue and expense levels and the adequacy of the available cash resources. Some of the information contained herein is based upon or derived from information provided by third-party consultants and other industry sources as well as by the portfolio companies. We have not independently verified and cannot assure the accuracy of any data obtained by or from these sources.

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The Company is continuously monitoring the impact of the worldwide spread of the corona virus (COVID-19) on its activities and the activities of its portfolio companies. At this time, there is a material uncertainty regarding the economic and other ramifications of the spread of COVID-19. This spread might have a negative effect on the activities of the Company and its portfolio companies, including, but not limited to, their market value, the ability to raise capital (governmental, private or public), the ability to materialize the Company's holdings, the possibility to advance strategic transactions, and the ability to carry out R&D and regulatory activities.

This presentation does not constitute or form part of, and should not be construed as constituting or forming part of, any offer or invitation to sell or issue, or any solicitation of any offer to purchase or subscribe for, any of the Company shares or its portfolio companies shares, nor shall any part of this presentation nor the fact of its distribution form part of or be relied on in connection with any contract or investment decision relating thereto, nor does it constitute a recommendation regarding our or our portfolio companies securities.

# WHAT TO LOOK FOR IN 2021



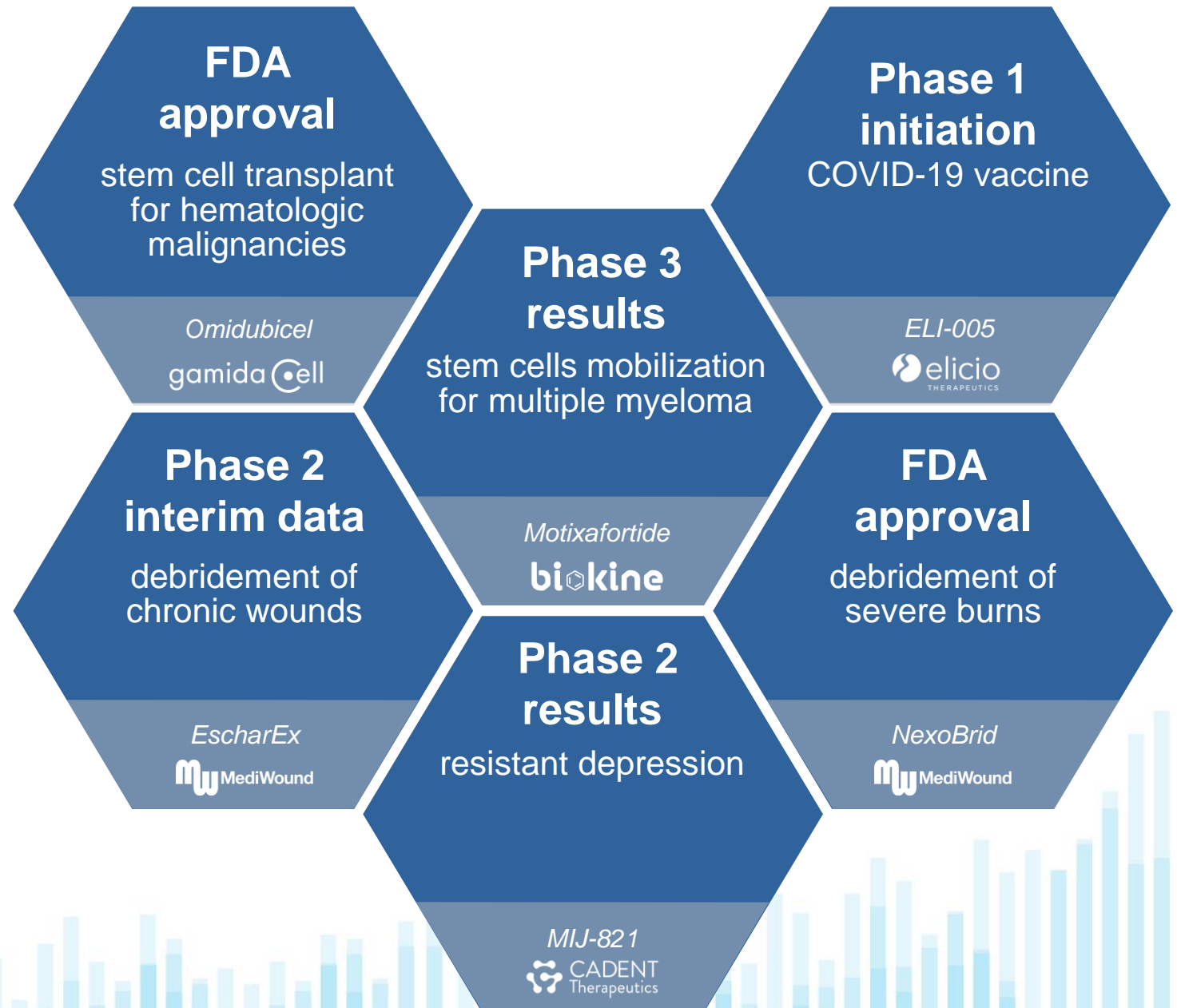
2 FDA marketing approvals



3 advanced clinical trial readouts



COVID-19 vaccine first-in-human trial



# CLAL BIOTECHNOLOGY INDUSTRIES

Leading life sciences investment company traded on the stock exchange (TASE: CBI)



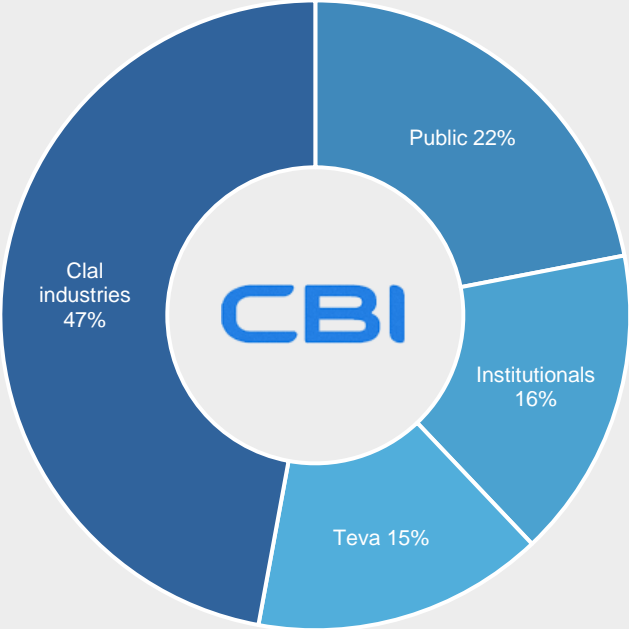
A member of Len Blavatnik's Access Industries group



Team based in Tel Aviv and Boston



## Shareholders



Collaborations with global healthcare companies and major investment funds




Portfolio of well-funded biotech/ med-tech companies; ~\$400M raised in 2019/2020





Advanced technologies from leading US/IL institutions addressing major unmet needs




# EXPERIENCED MANAGEMENT TEAM







**Avi Fischer - Chairman of the Board**

  **claltech**







**William Koster, PhD**

 **Neurogen Corporation**  **SciFluor**  **Bristol-Myers Squibb**



**Robert Connelly  
Strategic Advisor**

 **domantis**  **pulmatrix**  
 **axcella**  **FLAGSHIP PIONEERING**  
SCIENTIFIC VENTURES FOR LIFE





**Ofer Gonen - CEO**

 **gamida Cell**  **MediWound** 



**Assaf Segal - CFO**

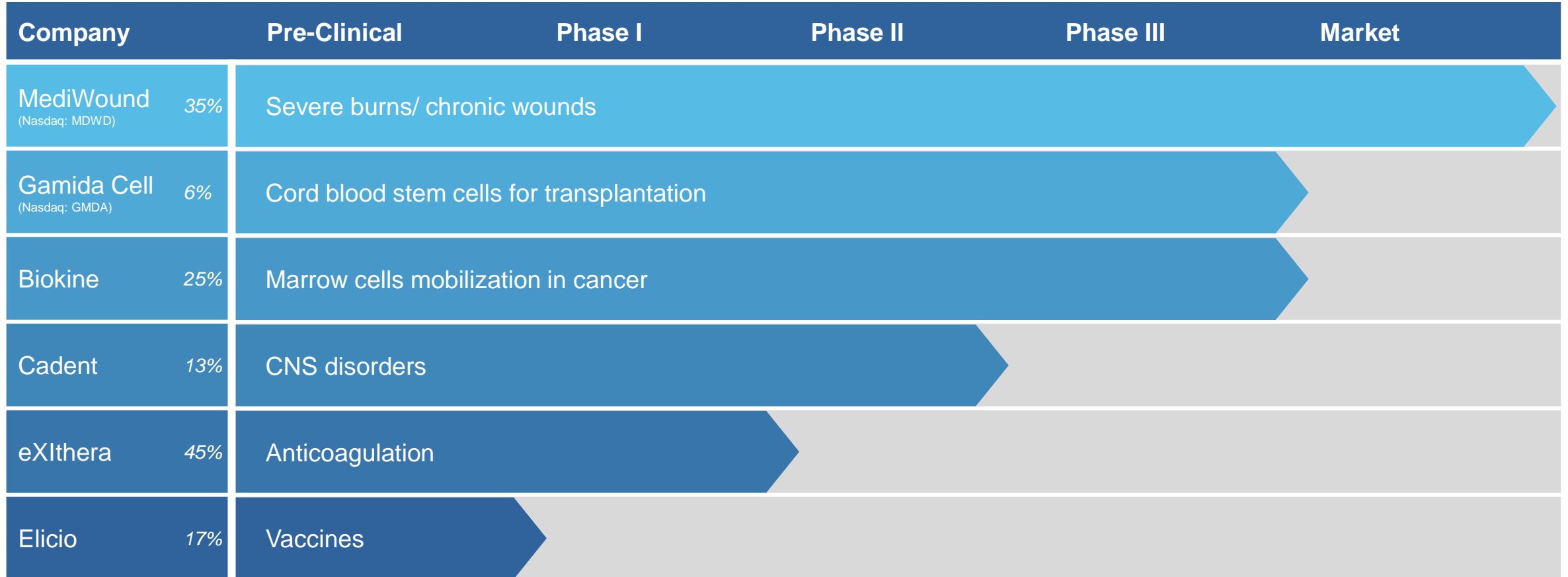
 **MediWound**  **pwc**  **amdocs**  
 **Variance**  
Financial Consulting



**Gilad Rosenberg, MD -  
Medical Director**

 **d PHARM**  
Pharmaceuticals  **MSD**

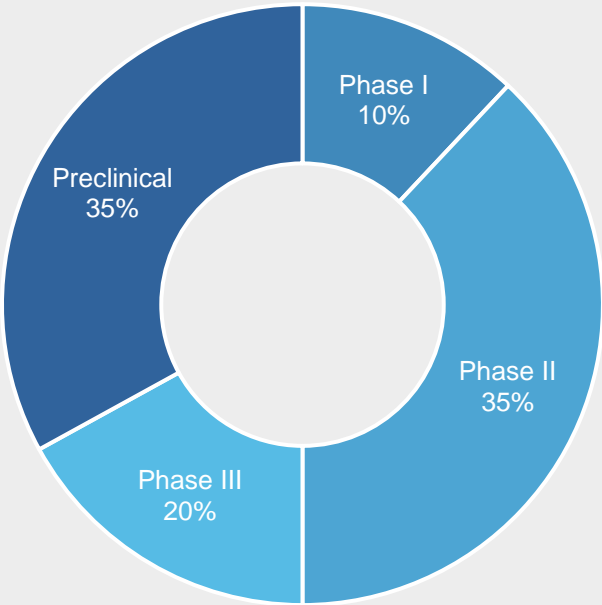
# DIVERSE CLINICAL DEVELOPMENT ACTIVITY



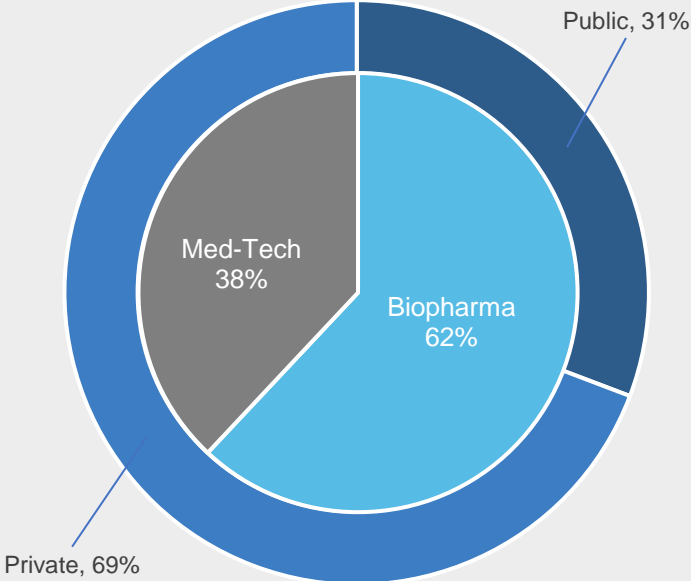


# BALANCED POTFOLIO

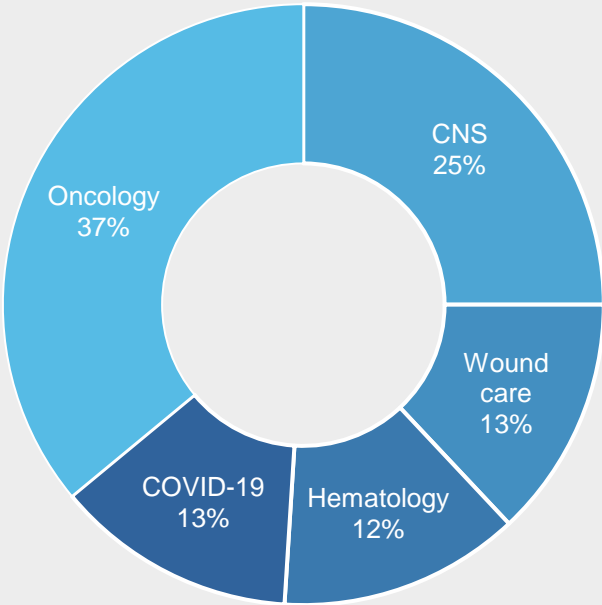
Stage of Development



Asset Distribution



Therapeutic Areas



The background of the slide is a dark blue color with a subtle, abstract pattern of interconnected circles and lines, resembling a molecular or network structure. The pattern is more prominent on the left side and fades towards the right.

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# **PORTFOLIO COMPANIES**



## Unique technology for the expansion of cord blood-derived stem cells

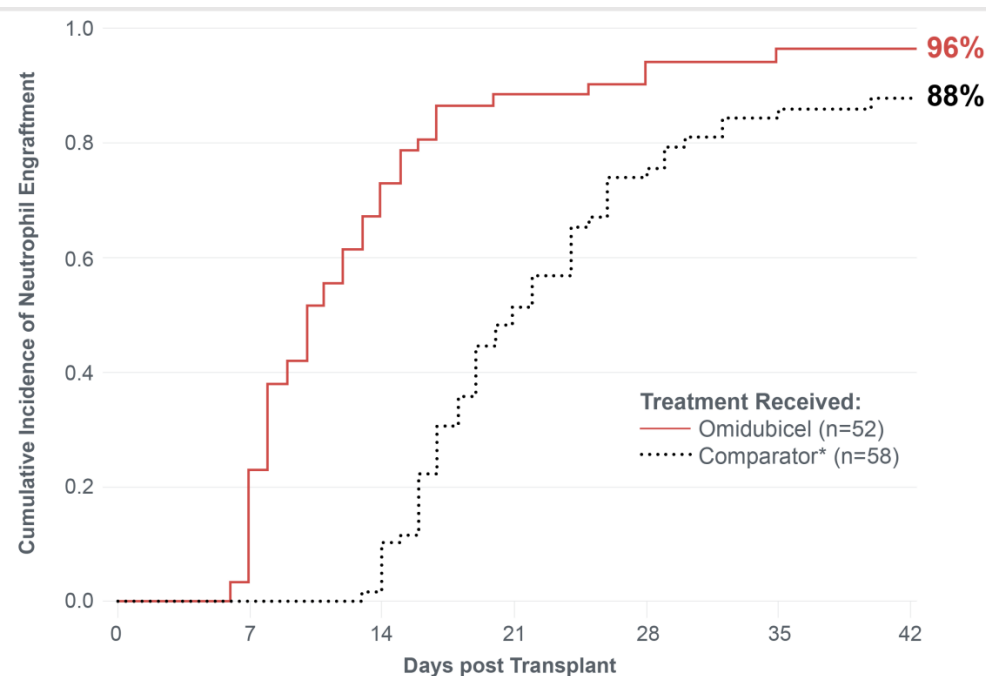
### Omidubicel – treatment for hematological malignancies

- Orphan drug status and 'Breakthrough Therapy' designation granted by FDA
- Phase III trial (n=125) completed
  - **Primary endpoint of neutrophil engraftment achieved**, with median time to engraftment of 12 days in the omidubicel group vs. 22 days in the comparator group ( $p < 0.001$ )
  - **Trial met all 3 secondary endpoints** (day 42 platelet engraftment, Grade 2/3 infections by day 100 post-transplant, days alive out of the hospital by day 100 post-transplant)
- Phase II in aplastic anemia is ongoing

**GDA-201** – phase I in hematological malignancies is ongoing; promising early evidence of clinical activity observed in advanced non-Hodgkin's lymphoma

(Nasdaq: GMDA)

## Phase III trial primary endpoint: cumulative incidence of neutrophil engraftment



Omidubicel is designed to enhance the life-saving benefits of cord blood stem cell transplant

## Develops, manufactures & commercializes drugs for burns and wound care

### NexoBrid - an enzymatic orphan drug for burn debridement

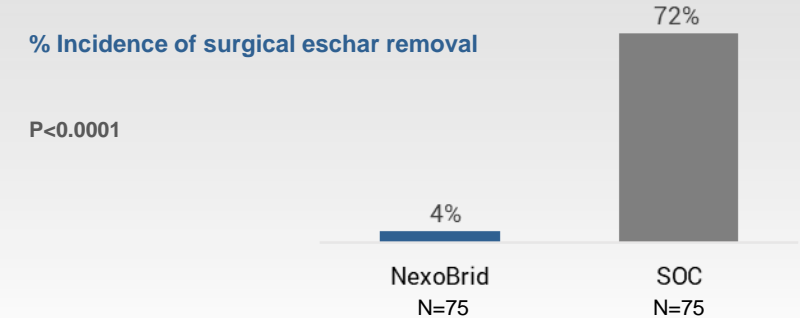
- Marketed in the EU and other territories; becoming standard of care
- Phase III trial for FDA approval has met the primary and all secondary endpoints; BLA submitted and accepted for review by FDA
- Strategic agreements of up to \$202 million with BARDA: funding all of NexoBrid's R&D activities and procuring NexoBrid for \$16.5 million
- License agreement with Vericel (Nasdaq: VCEL) for NexoBrid in North America

### EscharEx - an enzymatic drug for chronic wound debridement

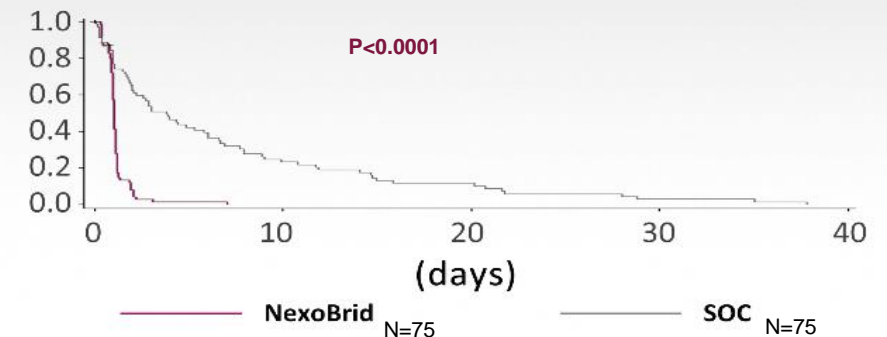
- Positive results in initial phase II trial
- A large phase II study comparing EscharEx to standard of care in venous leg ulcers initiated; interim data expected in H1/2021



(Nasdaq: MDWD)



**Kaplan-Meier estimates for time to complete eschar removal**



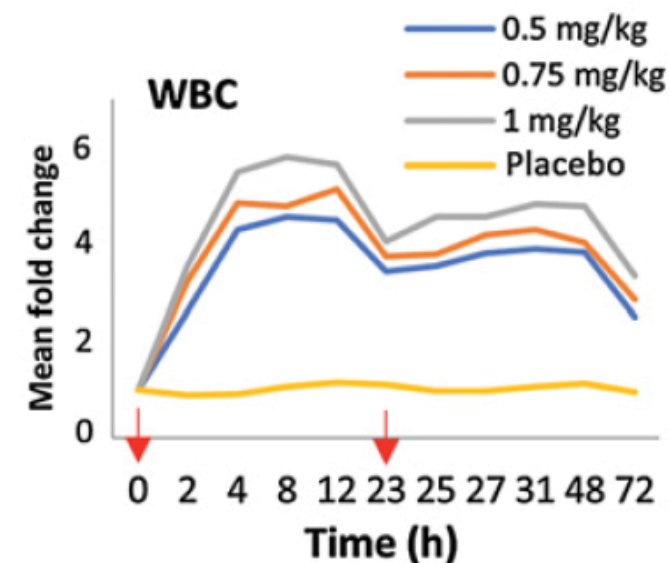
**Evidence-based breakthrough technology addressing large and growing markets**

## Small molecules and peptides disrupting cancer cell communications

**Motixafortide (BL-8040)** - a novel high-affinity antagonist of CXCR4, mobilizing hematopoietic stem cells; orphan drug status in US and EU

- Developed by BioLineRx (Nasdaq: BRLX) under a worldwide exclusive license
- Phase II has shown that a single injection of motixafortide mobilized sufficient cells for transplantation as 4-6 injections of G-CSF, the current standard of care
- **A phase III** trial in hematopoietic stem cell mobilization for autologous marrow transplantation in multiple myeloma has been **recently stopped for success** as the primary endpoint was met
- Motixafortide is also being developed as a therapy for solid tumors (phase I/II), acute myeloid leukemia (phase II) and for ARDS (phase Ib)

**BKT300** - a small molecule in pre-clinical research that interferes with a key cell cycle regulator over-expressed in tumor cells, aimed for solid and non-solid cancers



Motixafortide dose-dependent mobilization of white blood cells (WBC) from the bone marrow into blood stream; red arrows indicate the time of the two injections of motixafortide (n=6/group)  
*Clin Cancer Res, 2017, 23; 6790–801*

Innovative arsenal for combating tumor expansion  
with multiple potential applications

# NOVEL THERAPIES FOR NEUROLOGICAL AND PSYCHIATRIC CONDITIONS

## Developing drugs for CNS disorders

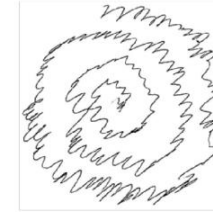
### Positive or negative modulation of brain NMDA receptors:

- **MIJ821** - for treatment-resistant depression; licensing deal with Novartis provides for milestone payments of up to \$386 million, plus royalties; a phase II trial recently completed enrolment
- **CAD-9303** - for schizophrenia; currently in phase I

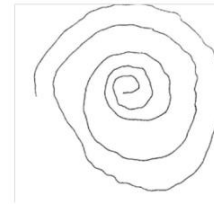
### Modulation of ion channels to treat movement disorders

- **CAD-1883** - drug for spinocerebellar ataxia (SCA) and essential tremor
  - A phase II trial in essential tremor has indicated proof of concept
  - Granted Orphan Drug status by FDA for SCA

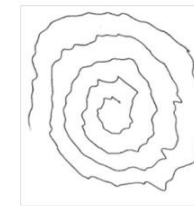
Pre-Dose



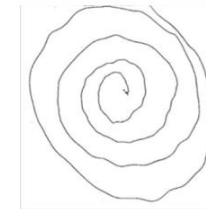
Day 14



Pre-Dose



Day 14



Pre-Dose



Day 14



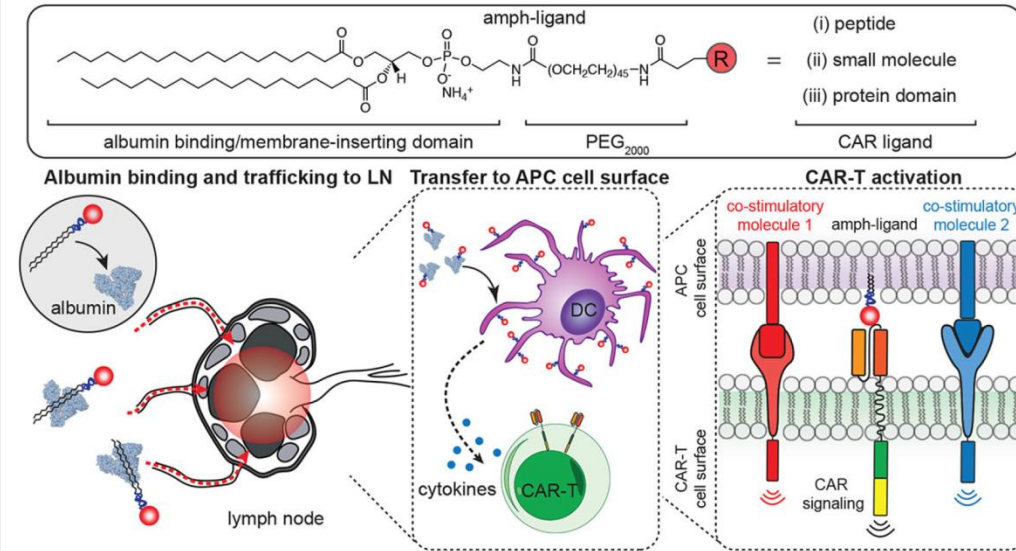
Robust improvements in spiral drawing in essential tremor patients who responded to CAD-1883 in a Phase 2a trial

*Presented at the 3<sup>rd</sup> Pan American Parkinson's Disease and Movement Disorders Congress, February 15, 2020*

**An attractive portfolio of products to treat CNS  
debilitating diseases; targeting multi billion \$  
markets**

## Developing potent next generation, lymph node-targeting, immuno-modulatory therapeutic and prophylactic vaccines for cancer and infectious diseases

- Lymph node targeting aims to achieve potent immune activation, robust T cell response, immune memory, and, in cancer, tumor eradication and durable cure
- Solid intellectual property; ongoing relationship with founding MIT laboratory
- Preclinical data from Elicio's COVID-19 vaccine indicate:
  - Up to 25-fold more T cell response over benchmark vaccines and >265-fold greater neutralizing antibody levels than recovering patients
  - Potent CD8 and CD4 T cell presence in lung tissue and respiratory fluid
- Expertise in cancer therapeutics; an executive team with a track record of success and exits with novel platform biotechnology companies
- ELI-002: anti-KRAS therapeutic vaccine for KRAS-positive pancreatic and colorectal cancers; phase I/II to start in Q1 2021
- Pipeline includes a combination with CAR-T for better efficacy in solid tumors



*Science. 2019, 12; 365(6449): 162–168*

**Lymph node targeting better unlocks the power of the immune response**

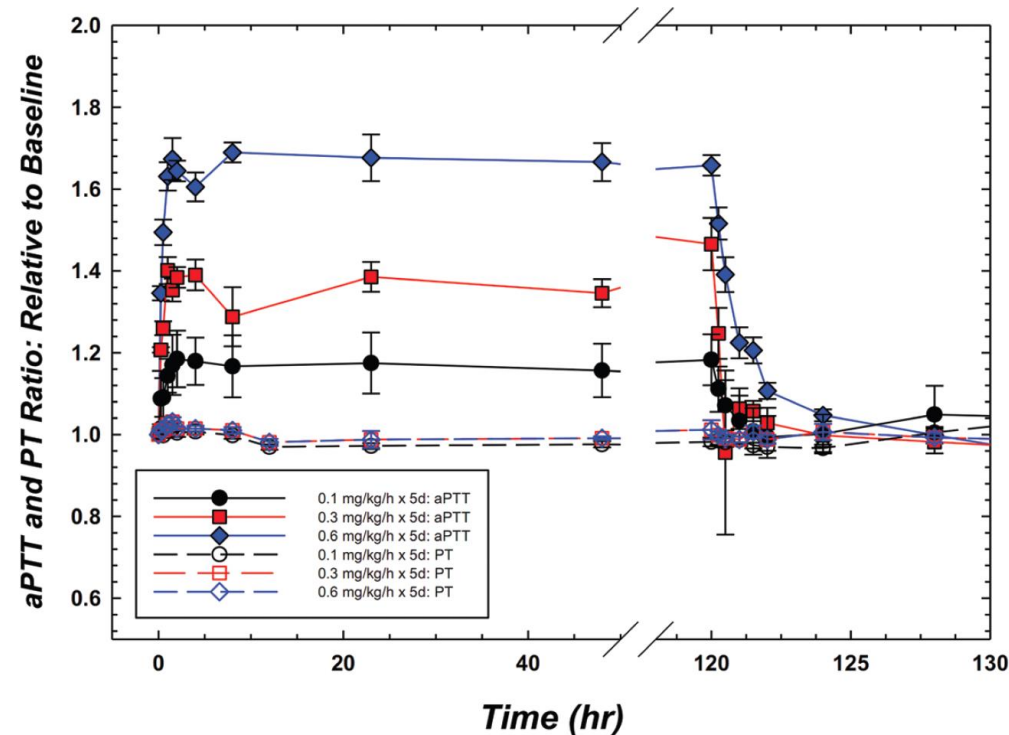
## World leader in developing novel, safer, first-in-class antithrombotic drugs

- Despite the benefits of available anticoagulants, all convey significant bleeding risk
- Factor XI antagonists inhibit the intrinsic clotting pathway, related to cardiovascular risk, but not the extrinsic pathway, critical to bleeding control in surgery and trauma
- Humans with genetically low levels of factor XI have decreased incidence of thrombosis, which is not associated with spontaneous bleeding

## EP-7041 - a novel, IV, selective small molecule Factor XIa inhibitor

- Phase I in healthy volunteers showed good tolerance, predictable dose-dependent increase in aPTT (efficacy marker) and rapid start and end of activity
- Strategic collaboration with Haisco Pharmaceutical Group (002653:CH)
- Phase II trial evaluating EP-7041 in COVID-19 patients in ICU, and a trial in the setting of extracorporeal membranous oxygenation (ECMO) are being planned

## Phase I trial coagulation data



**Factor XIa inhibition may finally dissociate anti-thrombotic effect from bleeding risk**



# MED-TECH PORTFOLIO



## Sight Diagnostics

Commercializing OLO™, a novel AI and optical, FDA approved, point-of-care system for complete blood count

Worldwide launch activities

Significant COVID-19 related opportunities

Recently completed \$71M financing



## Colospan

Developing CG-100, an anastomosis protection device for colorectal surgery to prevent anastomotic leaks, forgoing the need for a protective colostomy

CE Approved device

Clinical trial to support PMA is ongoing in US, EU and Israel



## Pi-Cardia

Developing Leaflex™, a novel catheter for treating calcified aortic stenosis without a need for valve replacement

Proof-of-concept clinical trial completed successfully; a trial to evaluate durability is ongoing

Recently completed \$27M financing; initiated a strategic collaboration with Venus (China)



## MinInvasive

Commercializing OmniCuff™ - a minimally invasive system for rotator cuff tear repair

Excellent clinical outcomes in clinical trials and in routine use

FDA Approved

Received NMPA (CFDA) approval; partnering with Microport in China



## FDNA

Deploying Face2Gene: AI-based platform to detect physiological patterns to reveal disease-causing genetic variants

Used by 70% of the world's geneticists (2,000 sites; over 130 countries)




















































Telehealth opportunity boosted by COVID-19

The background of the entire image is a dark blue field filled with a complex, abstract pattern of interconnected circles and lines, resembling a molecular structure or a network diagram. The circles vary in size and are connected by thin, light blue lines, creating a sense of depth and movement across the frame.

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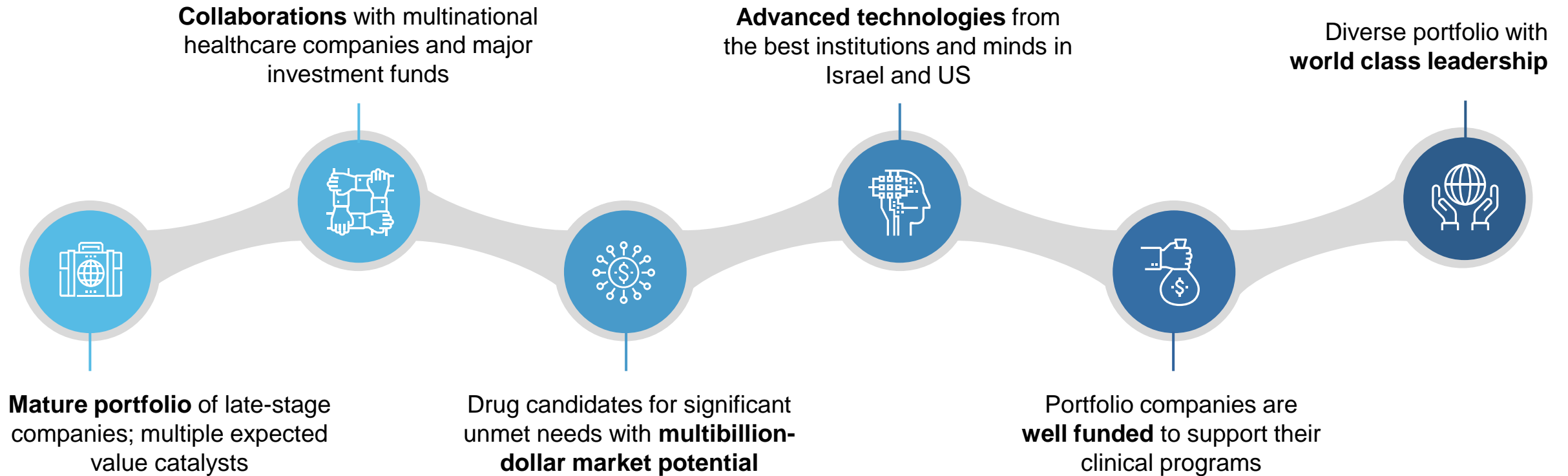
**LOOKING  
AHEAD**

# UPCOMING EVENTS - VALUE CREATING MILESTONES

	2020	2021
Gamida Cell	 <b>Omidubicel</b> phase 3 top line results (hematological malignancies)  <b>Financing</b> \$69M  <b>Omidubicel</b> phase 1/2 readout (aplastic anemia)  <b>GDA-201</b> phase 1 data (NHL)  <b>Omidubicel</b> phase 3 readout (hematological malignancies)  <b>Omidubicel</b> BLA submission (hematological malignancies)	 <b>Omidubicel</b> FDA approval (hematological malignancies)
MediWound	 <b>EscharEx</b> phase 2 (chronic wounds)  <b>Procurement</b> by <b>BARDA</b>  <b>NexoBrid</b> BLA submission (severe burns)  <b>NexoBrid</b> phase 3 enrollment completion (pediatric burns)  <b>NexoBrid</b> phase 3 results (severe burns)  <b>NexoBrid</b> phase 3 readout (pediatric burns)  <b>NexoBrid</b> FDA approval (severe burns)  <b>EscharEx</b> phase 2 interim data (chronic wounds)	
Biokine	 <b>Sale of additional</b> rights ( <b>BioLineRX</b> )  <b>Motixafortide + Keytruda</b> phase 2 readout (pancreatic cancer)  <b>Motixafortide</b> phase 3 interim analysis (marrow mobilization)  <b>Motixafortide</b> + <b>Libtayo</b> phase 2 (pancreatic cancer)  <b>Motixafortide</b> + <b>Keytruda</b> phase 2 readout (pancreatic cancer)  <b>Motixafortide</b> phase 2b interim data (AML)  <b>Motixafortide</b> phase 1b interim data (ARDS)  <b>Motixafortide</b> phase 3 readout (marrow mobilization)  <b>BKT-300</b> IND submission (oncology)	
Cadent	 <b>Milestone</b> payment ( <b>Novartis</b> )  <b>CAD-9303</b> phase 1/2 (schizophrenia cognitive deficits)  <b>Financing</b> \$40M  <b>MIJ-821</b> phase 2 results (resistant depression)  <b>IPO/</b> strategic collaboration  <b>CAD-9303</b> phase 1/2 readout (schizophrenia)  <b>CAD-1883</b> phase 1/2 (SCA)	
Elicio	 <b>Eli (002)</b> Pre IND meeting  <b>Financing</b> \$48M  <b>ELI-002</b> Pre-IND meeting  <b>ELI-005</b> pre-clinical data (COVID-19 vaccine)  <b>ELI-005</b> Strategic collaboration  <b>ELI-002</b> phase 1/2 (cancer vaccine)  <b>ELI-005</b> phase 1 (COVID-19 vaccine)  <b>ELI-002</b> phase 1/2 data (cancer vaccine)	
eXlthera	 <b>EP-7041</b> Haisco licensing deal (China)  <b>EP-7041</b> IND submission (COVID-19)  <b>EP-7041</b> licensing deal/ financing  <b>EP-7041</b> phase 2 (COVID-19)  <b>EP-7041</b> phase 2 (ECMO)	
Anatomy	 <b>Sight CBC</b> FDA approval  <b>Pi-Cardia</b> \$27M financing  <b>Sight</b> \$71M financing  <b>Pi-Cardia</b> strategic collaboration  <b>Pi-Cardia</b> Leaflex studies  <b>MinInvasive</b> OmniCuff CFDA  <b>Sight CBC</b> OLO launch  <b>FDNA</b> TeleHealth launch  <b>Colospan</b> financing  <b>MinInvasive</b> soft launch in China  <b>Colospan</b> US pivotal trial	

(\*) The above events may be delayed/ cancelled/ otherwise varied in relation to the spread of the COVID-19 pandemic

# READY TO MATERIALIZE





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