

DISCLAIMER

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.

INVESTMENT OPPORTUNITIES IN HEALTHCARE

Large and Growing Industry

- Revenues > \$1.3 trillion
- High profit margins, strong cash-flows, significant multiples
- Resistant to economic cycles
- Multiple opportunities



Investment Approach

- Transatlantic deal sourcing
- Active lead/co-lead investor
- Long-term investment expertise
- Exit-driven investments

Compelling Exit Markets

- Attractive M&A environment
- Cash-rich corporate acquirers
- Significant premiums
- IPOs: tangible exit option

CLAL BIOTECHNOLOGY INDUSTRIES

Leading, publicly traded, life sciences investment company (TASE: CBI)







Team and portfolio based in Tel Aviv and Boston



Collaborations with global healthcare companies and major investment funds







♦ MicroPort

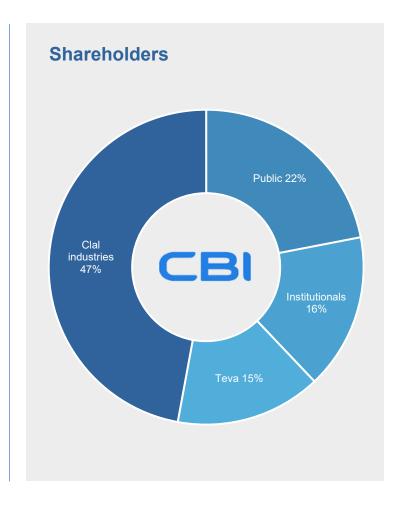


Portfolio of well-funded biotech/ med-tech companies; ~\$450M raised in 2019/2020



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







PERFORMANCE IN 2020

Cadent acquired by Novartis for **\$210-770M**



\$300M raised by portfolio companies

Positive phase 3 readouts



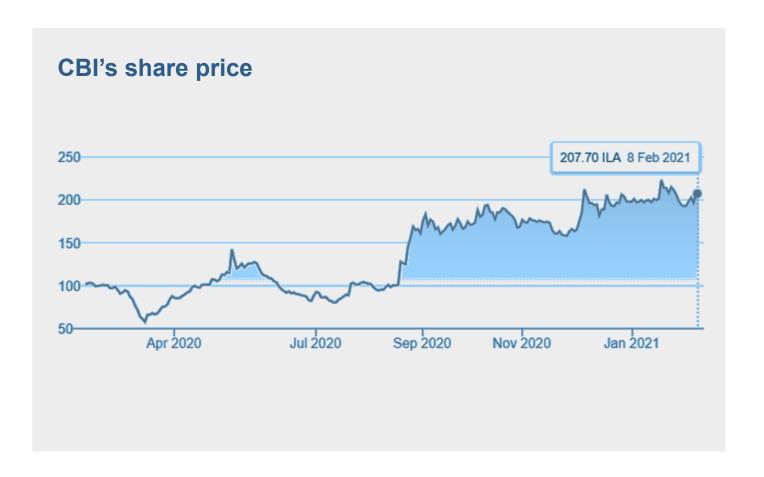
Neon acquired by BioNTech for **\$67M**



NIS **36M** dividend and up to NIS **10M** buyback underway

BLA submission





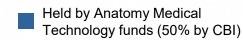


DIVERSE CLINICAL DEVELOPMENT ACTIVITY

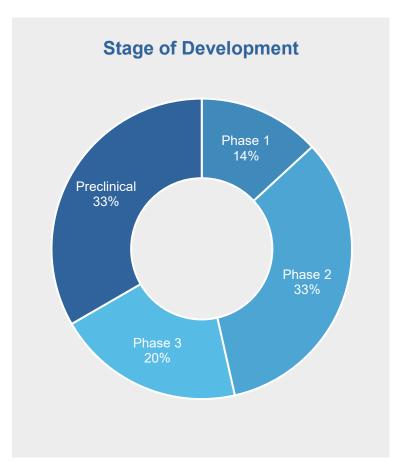
| Company | | Pre-Clinical | Phase 1 | Phase 2 | Phase 3 | Market | | | | |
|-------------------------------|-----|-------------------------------------|---------|---------|---------|--------|--|--|--|--|
| MediWound (Nasdaq: MDWD) | 35% | Severe burns/ chronic wounds | | | | | | | | |
| FDNA | 2% | Genetic diagnostics | | | | | | | | |
| MinInvasive | 20% | Orthopedics | | | | | | | | |
| Sight | 4% | Blood diagnostics | | | | | | | | |
| Gamida Cell (Nasdaq: GMDA) | 5% | Cell therapy | | | | | | | | |
| Biokine | 25% | Marrow cells mobilization in cancer | | | | | | | | |
| Colospan | 21% | GI surgery | | | | | | | | |
| Cadent | | CNS disorders | | | | | | | | |
| eXIthera | 45% | Anticoagulation | | | | | | | | |
| Anchiano (Nasdaq: ANCN) | 19% | Oncology | | | | | | | | |
| Pi Cardia | 8% | Cardiac valve repair | | | | | | | | |
| Elicio | 17% | Vaccines | | | | | | | | |

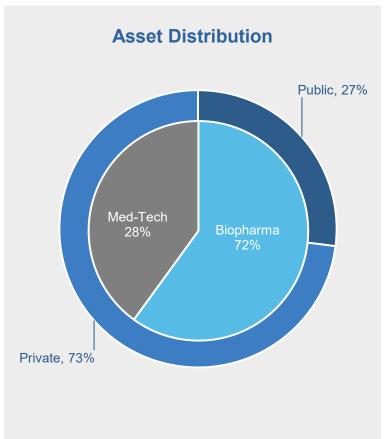


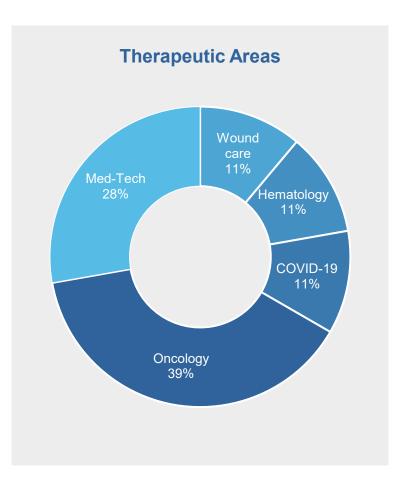




BALANCED AND DIVERSIFIED PORTFOLIO







VALUE CREATION OVER 3 YEARS

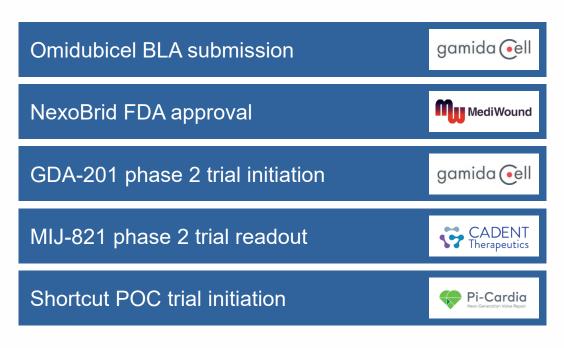
2018 2019 2020 PO 3 IPOs on Nasdaq Neon Anchiano Pi Cardia Gamida Cell MediWound Neon Pharma 5 Pharma / strategic Neon eXIthera Cadent deals Biokine Colospan **FDNA** Elicio Gamida Cell Pi Cardia M&A 2 M&As Sight Diagnostics Sight Diagnostics Cadent Financing eXIthera Gamida Cell ~\$700 million raised Elicio



MULTIPLE NEAR-TERM PIPELINE CATALYSTS

H1 2021 H2 2021







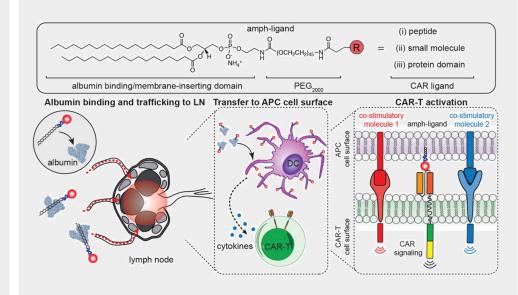




DRIVING THE IMMUNE SYSTEM TO ELIMINATE CANCER

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 durable eradication
- Solid intellectual property; ongoing relationship with founding MIT laboratory
- ELI-002: anti-mKRAS therapeutic vaccine for mKRAS-positive pancreatic and colorectal cancers; phase 1b to start in Q1 2021
- Pipeline includes a combination with CAR-T for better efficacy in solid tumors
- 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



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

Lymph node targeting better unlocks the power of the immune response





REVOLUTIONIZING WOUND CARE

Develops, manufactures & commercializes drugs for burns and wound care

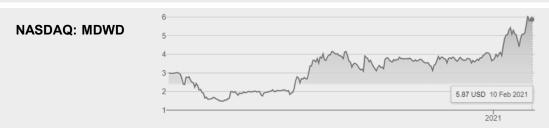
NexoBrid - an enzymatic orphan drug for burn debridement

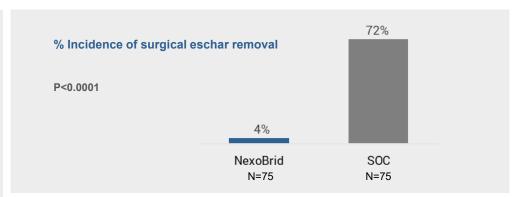
- Marketed in the EU and other territories; becoming standard of care
- Phase 3 trial for FDA approval has met the primary and all secondary endpoints;
 BLA submitted and accepted for review by FDA
- Strategic agreements of up to \$202 million with BARDA: funding all of NexoBrid's R&D activities and procuring NexoBrid for \$16.5 million
- License agreement with Vericel (Nasdaq: VCEL) for NexoBrid in North America

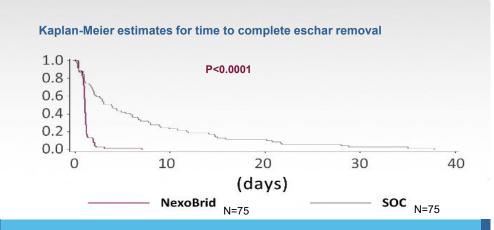
EscharEx - an enzymatic drug for chronic wound debridement

- Positive results in initial phase 2 trial
- A large phase 2 trial of EscharEx vs. standard of care in venous leg ulcers initiated; interim data expected in H1/2021









Evidence-based breakthrough technology addressing large and growing markets





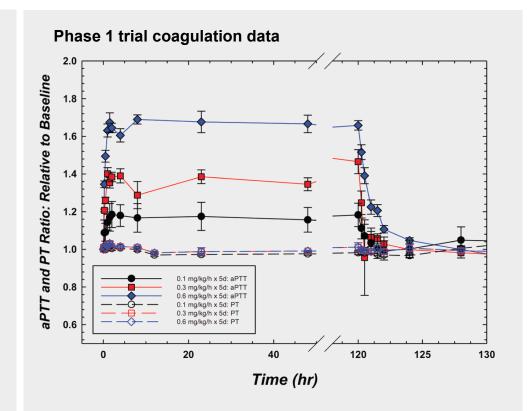
THROMBOSIS PREVENTION AVOIDING BLEEDING RISK

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

- Despite the benefits of available anticoagulants, all have significant bleeding risk
- Factor XI antagonists inhibit the intrinsic clotting pathway, related to cardiovascular risk, not the extrinsic pathway, critical to bleeding control in surgery and trauma
- Humans with genetically low levels of factor XI have deceased incidence of thrombosis, without associated spontaneous bleeding

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

- Phase 1 trial in healthy volunteers showed good tolerance, predictable dosedependent increase in aPTT (efficacy marker) and rapid start and end of activity
- Strategic collaboration with Haisco Pharmaceutical Group (002653:CH)
- Phase 2 trial evaluating EP-7041 in COVID-19 patients in ICU, and a trial in the setting of extracorporeal circulation (e.g., ECMO), are being planned



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



gamida **e**ll

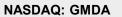
THE WORLD LEADER IN CORD BLOOD STEM CELL THERAPY

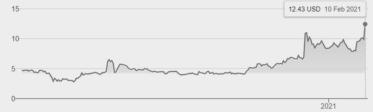
Unique technology for cellular expansion

Omidubicel - cord blood-derived stem cell therapy for hematological malignancies

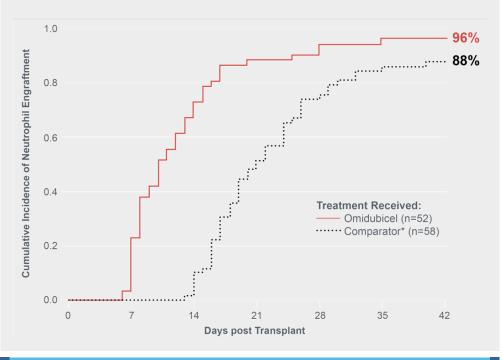
- Orphan drug status and 'Breakthrough Therapy' designation granted by FDA
- Phase 3 trial (n=125) completed
 - Primary endpoint of neutrophil engraftment achieved, with median time to engraftment of 12 days in the omidubicel group vs. 22 days in the comparator group (p<0.001)
 - Trial met all 3 secondary 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 2 in aplastic anemia is ongoing

GDA-201 – phase 1 trial in hematological malignancies is ongoing; promising early evidence of clinical activity observed in advanced non-Hodgkin's lymphoma





Phase 3 trial primary endpoint: cumulative incidence of neutrophil engraftment



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





UNIQUE NON-IMPLANT-BASED AORTIC VALVE REPAIR

Innovative catheter-based therapies for stenotic aortic valves

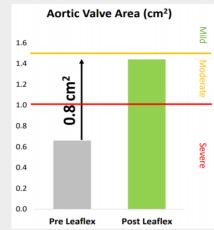
Leaflex™: a novel catheter-based approach for treating calcified stenotic aortic valves through leaflet scoring

- Successfully completed a proof-of-concept trial in humans
- A trial to evaluate Leaflex™'s long term effects is ongoing; trial completion expected in 2022
- Partnered with Sofinnova and Venus Medtech (China)

ShortCut™: a catheter-based device designed to avert coronary blockade during Transcatheter Aortic Valve Replacement (TAVR) by prior splitting of the aortic valve leaflets

First patients successfully treated

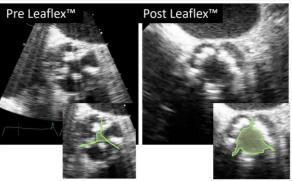




Results from firstin-human trial (n=11)

Presented at euroPCR 2019

(https://2d812c99-b1a6-4c6f-96d6acd1d700cf1d.filesusr.com/ugd/1f43fd_400d1de e54ea4fccab6cd286b78e4f6b.pdf?index=true)



Leaflex™: next generation valve repair, foregoing the need for valve replacement





BIRD'S-EYE VIEW

| Company | Field | Lead Product | Stage | Market Cap | % ⁽²⁾ | Next Catalyst | Timing |
|-------------|----------------------|---------------|-------------|-----------------------|------------------|-------------------------------|---------|
| MediWound | Wound care | NexoBrid | Market | \$151M ⁽¹⁾ | 35% | EscharEx interim readout | H1/2021 |
| Gamida Cell | Cell Therapy | Omidubicel | Phase 3 | \$699M ⁽¹⁾ | 5% | Omidubicel BLA submission | H2/2021 |
| Cadent | CNS disorders | MIJ-821 | Phase 2 | \$210-770M | 12% | MIJ-821 phase 2 results | H2/2021 |
| Elicio | Vaccines | ELI-002 | Preclinical | NA | 17% | ELI-002 phase 1/2 initiation | H1/2021 |
| Sight | Blood diagnostics | OLO | Market | NA | 2% | OLO launch | H1/2021 |
| Biokine | Marrow Transplant | Motixafortide | Phase 3 | NA | 25% | Motixafortide phase 3 readout | H1/2021 |
| Pi Cardia | Cardiac valve repair | Leaflex | Phase 2 | NA | 6% | Leaflex interim readout | H2/2022 |
| eXIthera | Anticoagulation | EP-7041 | Phase 2 | NA | 45% | EP-7041 phase 2 initiation | H1/2021 |
| Anchiano | Oncology | CM-101 | Phase 2 | \$32M ⁽¹⁾ | 19% | Anchiano/ Chemomb M&A | H1/2021 |
| FDNA | Genetic diagnostics | Face2Gene | Market | NA | 1% | Telehealth launch | H1/2021 |
| Colospan | GI surgery | CG-100 | Phase 3 | NA | 11% | CG-100 pivotal trial readout | H1/2024 |
| MinInvasive | Orthopedics | OmniCuff | Market | NA | 10% | Financing | H1/2021 |

⁽¹⁾ Based on Nasdaq quotes as of February 12, 2021(2) Represents ownership percentage by CBI only (direct and indirect ownership)



REASONS TO INVEST IN CBI

Collaborations with multinational healthcare companies and major investment funds

Mature portfolio of late-stage companies; multiple near-term

Diverse public and private the best institutions and minds in Israel and US

Drug candidates for significant unmet needs with multibillion
Portfolio of private and public companies; multiple near-term

Diverse public and private portfolio with world class leadership

Portfolio of private and public companies well funded to support

dollar market potential

their clinical programs



value catalysts

EXPERIENCED MANAGEMENT TEAM

Transatlantic well-connected team with sound scientific, medical, and commercial expertise



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