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UPDATE REPORT

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BioCardia Inc. (BCDA-NASDAQ)

August 13, 2020

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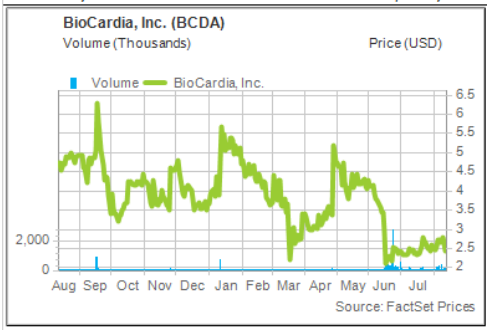
BUY: Reduced Burn Preserves Capital

BioCardia reported 2Q20 results. The burn rate was reduced from $4.6M in 1Q to $3.6M in 2Q. However, the net cash used in 2Q was even lower at just $2.3M. The Company ended the period with $11M in cash. Recall that BioCardia successfully closed a public offering of 4.7M shares of common stock (at $2.10 per share). The Company laid out their catalysts in the press release, which we present in our note.

Current Price	**$2.45**
Price Target	**$24.00**

Estimates	F2019A	F2020E	F2021E
Expenses ($000s)	$ 15,192	$ 16,254	$ 16,158
1Q March	$ 3,903	$ 4,647	$ 3,792
2Q June	$ 3,848	$ 3,601	$ 3,957
3Q September	$ 3,421	$ 3,924	$ 4,122
4Q December	$ 4,020	$ 4,081	$ 4,287
	F2019A	F2020E	F2021E
EPS (diluted)	$ (2.13)	$ (1.50)	$ (0.41)
1Q March	$ (0.08)	$ (0.67)	$ (0.12)
2Q June	$ (0.77)	$ (0.46)	$ (0.09)
3Q September	$ (0.63)	$ (0.18)	$ (0.10)
4Q December	$ (0.65)	$ (0.19)	$ (0.10)
EBITDA/Share	($0.96)	($1.11)	($0.41)
EV/EBITDA (x)	0.0	0.0	0.0

Stock Data	
52-Week Range	$2.01 - $7.25
Shares Outstanding (mil.)	12.4
Market Capitalization (mil.)	$30
Enterprise Value (mil.)	$30
Debt to Capital	0%
Book Value/Share	$2.62
Price/Book	0.9
Average Three Months Trading Volume (K)	14
Insider Ownership	35.6%
Institutional Ownership	1.7%
Short interest (mil.)	0.1%
Dividend / Yield	$0.00/0.0%



Investment Highlights

Catalysts Ahead (*from the press release*):

- Q3 2020: First patient treated in CardiAMP Chronic Myocardial Ischemia (CMI) Trial (BCDA-02)
- Q3 2020: FDA acceptance of Investigational New Drug application for CardiALLO Neurokinin-1 Receptor-Positive Mesenchymal Stem Cell Therapy (BCDA-03), the Company's allogeneic therapeutic platform, for the treatment of ischemic heart failure
- Q4 2020: Prespecified Data Safety Monitoring Board Review of all patients enrolled in the CardiAMP Heart Failure Trial, including futility analysis, based on sixty (60) patients that will have reached the primary one-year follow-up endpoint at the time of analysis (BCDA-01)
- Q4 2020: Prespecified Data Safety Monitoring Board Review of safety data from the roll-in cohort in CardiAMP CMI Trial (BCDA-02)
- Q4 2020: FDA acceptance of Investigational New Drug application for Neurokinin-1 Receptor-Positive Mesenchymal Stem Cell Therapy (BDCA-04) for the treatment of Acute Respiratory Distress Syndrome as a result of COVID-19
- Q4 2020: Targeted commercial availability of Avance Transseptal Sheath

Valuation: Our product models run out to the year 2030. For CardiAMP and CardiALLO and all the related cardiac indications, each of which represent blockbuster markets, we haircut the revenues by 70% (assume only a 30% probability of success). In addition, in our free cash flow (FCFF), discounted EPS (EPS), and sum-of-the-parts (SOP) models, we apply a risk rate (r) of 30% on top of the 90% risk cut in our models. Our share count is projected for 2030 and assumes multiple raises. Our models are then equal-weighted, averaged, and rounded to the nearest whole number to derive our 12 months price target of $24.00. **See the Valuation section in this report for some comparable valuations and recent transactions.**

Risks: Partnership risks, Commercialization risks, Financial risks, Clinical and Regulatory risks, and Legal and Intellectual Property risk.

Heart Disease – Treating the Underlying Cause. BioCardia strongly believes that stem cells have the power to change the course of cardiovascular disease. This may occur as a result of the trophic effects of these cells when administered directly into the local environment (heart muscle). The cells act like micro-drug factories secreting factors that help to reduce inflammation, reduce scarring, and promote micro-angiogenesis (formation of new blood supply to the tissue). In doing such, it is hoped in the case of an acute ischemic event that the cells can help limit the initial damage and, in the case of chronic disease, may help to arrest the damage, even reverse it partially.

Autologous vs. Allogeneic: BioCardia is developing treatment using both autologous and allogeneic cells. BioCardia developed a diagnostic assay to determine which treatment is best suited for each patient. Autologous stem cell therapy is the company's leading product, CardiAMP, which is currently in a pivotal Phase 3 trial. Through CardiAMP, the patient's own bone marrow is extracted and is directly administered to the patient's heart to allow the most efficacy. If a patients CD34 cell count does not reach the requirement for CardiAMP, BioCardia can offer CardiALLO. CardiALLO stands out as an allogeneic stem cell therapy that is based on donated marrow. CardiALLO is designed to be a fast follower behind CardiAMP.

What's New? As the company plans its next generation of cell therapy, it's building the science to develop a differentiated allogeneic product (marrow-based cells), in CardiALLO. The specific MSCs used in BioCardia's allogenic cell therapy are expanded from cells selected for the presence of the NK1 receptor, which is known to bind to substance P, an important neuropeptide associated with inflammation throughout the body and a primary mediator of inflammation in the airways. Preclinical data (N=26 animals), were treated with both low dose and high dose NK1R+ MSC, echocardiographic measures of cardiac ejection fraction, fractional shortening and cardiac outflow were meaningfully improved, with all three measures being statistically significant for both dosage levels over control animals.

Delivery Matters. One of the characteristics that differentiates BioCardia from competitors in the field is the way the cells are delivered. The cells are administered directly to the patient's heart through the use of their Helix Biotherapeutic Delivery System and the Morph steerable guide. The end result is that more cells stay where they are needed versus other systems where the majority of cells are washed away in the dynamic blood flow associated with the heart. **See Exhibit 5. Cell Quality Matters Too.** BioCardia is talking more and more about its potency assay and the ability to determine in advance of therapy that cells are consistent and viable with its requirements. **See Exhibit 8.**

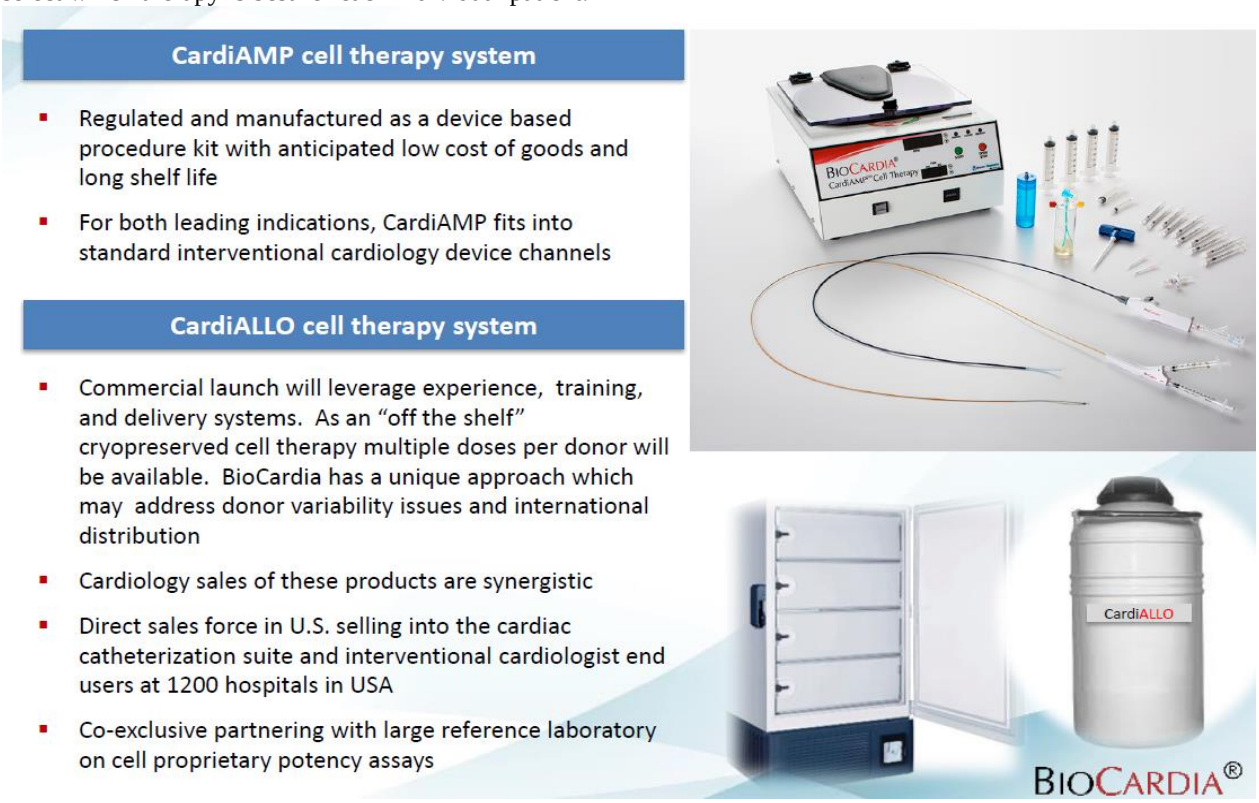
Roughly $20B is spent per year for hospitalization costs. Following an operation done on a patient's heart, on average, they are in the hospital for three to four nights spending at least $10,000 a night. CardiAMP can allow the patient to be released the same day with anticipated savings for the system of up to $40,000 of unwanted hospital costs. Cell Therapy represents the ability to intervene in heart failure outside of treatment paradigms that exist today. **See Exhibit 2.**

BioCardia is Assisted. The company is **c**onducting numerous clinical trials in heart disease and specifically heart ischemia. Focusing on heart failure consumes a significant amount of a company's resources. BioCardia gets an extra hand as the Centers for Medicare & Medicaid Services (CMS) has funded the ongoing Phase 3 trial of CardiAMP cell therapy system in ischemic heart failure.

Autologous vs. Allogeneic: BioCardia is developing treatment using both autologous and allogeneic cells. BioCardia developed a diagnostic assay to determine which treatment is best suited for each patient. Autologous stem cell therapy is the company's leading product, CardiAMP, which is currently in a pivotal Phase 3 trial. Through CardiAMP, the patient's own bone marrow is extracted and is directly administered to the patient's heart to allow the most efficacy. If a patients CD34 cell count does not reach the requirement for CardiAMP, BioCardia can offer CardiALLO. CardiALLO stands out as an allogeneic stem cell therapy that is based on donated marrow. CardiALLO is designed to be a fast follower behind CardiAMP.

Phase 3 Update: DSMB Say's No Safety Issues – Recommends that the trial should continue: BioCardia announced that the independent Data Safety Monitoring board (DSMB) has completed its prespecified data review for the Phase 3 pivotal CardiAMP Heart Failure Trial, which included safety follow-up results on 35 patients and all additional data available on the 77 patients randomized in the trial. The DSMB indicated there were no safety concerns with the CardiAMP study results and recommended that the trial continue, as planned. The trial is an ongoing multi-center, double-blinded, randomized (3:2), sham-controlled pivotal CardiAMP Heart Failure Trial that is expected to enroll 260 patients at up to 40 centers nationwide. The trial's primary efficacy endpoint is Six Minute Walk distance at 12 months' post-treatment, a measure of a patient's exercise capacity, and incorporates the impact of MACE and other clinically meaningful events. Secondary efficacy endpoints include quality of life as measured by the Minnesota Heart Failure Quality of Life self-assessment, and superiority relative to MACE and survival.

Exhibit 1. We view BioCardia as a unique company in the world of stem cells as the company has "The Best of Both Worlds", that is an Autologous and an Allogenic approach. A patient assay can determine if a patient's own cells are viable to treat disease and select which therapy is best for each individual patient.

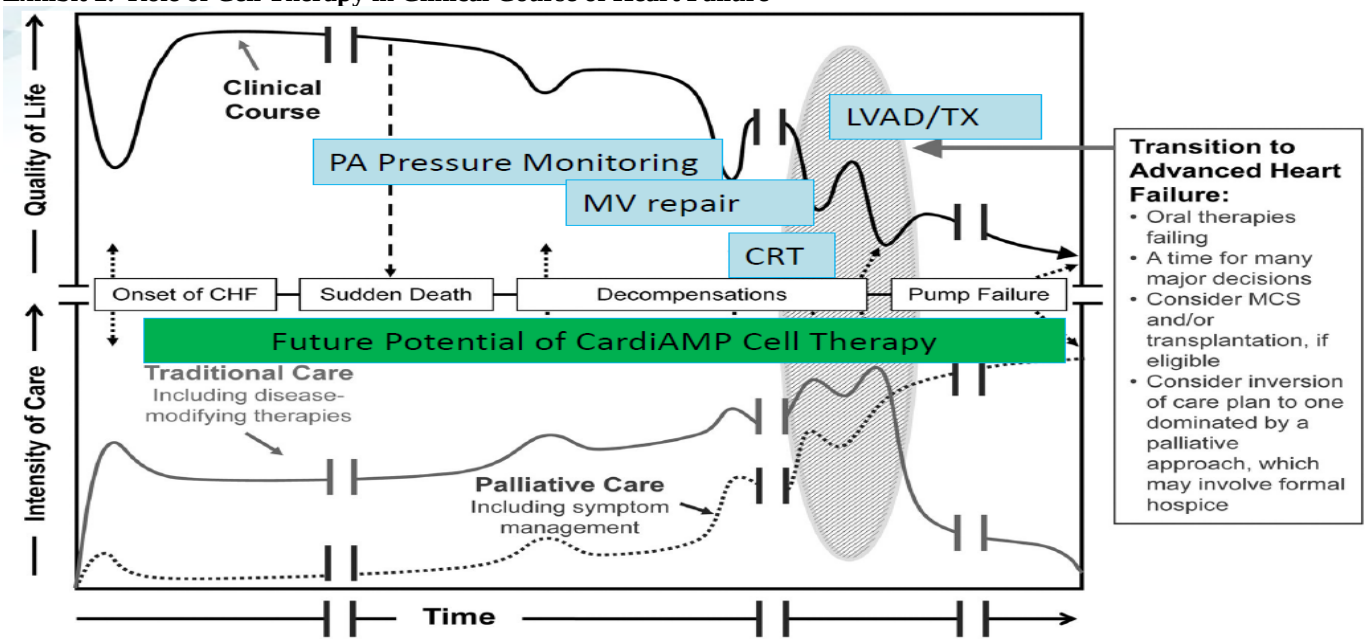


Source: BioCardia July 2019 Presentation.

Company Overview

BioCardia is a clinical stage regenerative medicine company developing therapies for cardiovascular diseases. It is well known that the cardiovascular market has large unmet medical needs and BioCardia believes CardiAMP and CardiALLO are part of the solutions for patients following a heart attack and or diagnosed with chronic myocardial ischemic and or ischemic related cardiac disease. The company's lead program, CardiAMP, incorporates a proprietary molecular model in order to determine if a patient's own bone marrow qualifies as a candidate for this therapy. The company hopes with this therapy to improve the quality of life for patients following treatment for heart disease. CardiAMP was designed to be delivered in a 60-90-minute procedure. CardiALLO is an option for those patients who are not able to produce enough viable cells to manufacture CardiAMP. Both heart failure trials, BCDA-01 and BCDA-02 have been supported by CMS (Medicare) which has offset the financial burden for expensive HF trials. The interest in cell-based therapy for heart failure is rising for multiple reasons including the lack of adverse events, high safety profile, and its ability to change the treatment paradigm in cardiovascular disease.

Exhibit 2. Role of Cell Therapy in Clinical Course of Heart Failure



Source: Modified from Decision Making in Advanced Heart Failure, A Scientific Statement From American Heart Association, Circ 2012

Exhibit 3. Upcoming Catalysts for BioCardia

Product	Indication	Event	Timeline	Impact
BCDA-01: CardiAMp Cell Therapy	Ischemic Heart Failure	Complete enrolling pivotal Phase 3 trial	2H20	++
BCDA-01: CardiAMp Cell Therapy	Ischemic Heart Failure	Complete Phase 3 and submit for US and EU approval	3Q21	+++
BCDA-01: CardiAMp Cell Therapy	Ischemic Heart Failure	Approval and Launch	2H22	+++
BCDA-02: CardiAMP Cell Therapy	Chronic Myocardial Ischemia	Complete enrollment for pivotal Phase 3	2H22	++
BCDA-02: CardiAMP Cell Therapy	Chronic Myocardial Ischemia	Complete Phase 3 and submit for US and EU approval	4Q23	+++
BCDA-02: CardiAMP Cell Therapy	Chronic Myocardial Ischemia	Approval and Launch	1Q24	+++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Seek Orphan Status	2021	+
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Phase 1/2 Investigational New Drug approved	4Q19	+
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Begin enrollment of Phase 1/2 Study	3Q20	+
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Complete enrollment for Phase 1/2 trial	3Q22	++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Complete study and file for pivotal Phase 3 trial	3Q23	++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Begin enrollment of pivotal Phase 3 trial	4Q23	++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Complete enrollment of Phase 3 trial	2H25	+++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Complete Phase 3 and submit for US and EU approval	3Q26	+++
BCDA-03 CardALLO Cell Therapy	Ischemic Heart Failure	Approval and Launch	1Q27	+++

Stock Significance Scale: + of moderate importance; ++ higher level; +++ highly

Source: Dawson James and BioCardia

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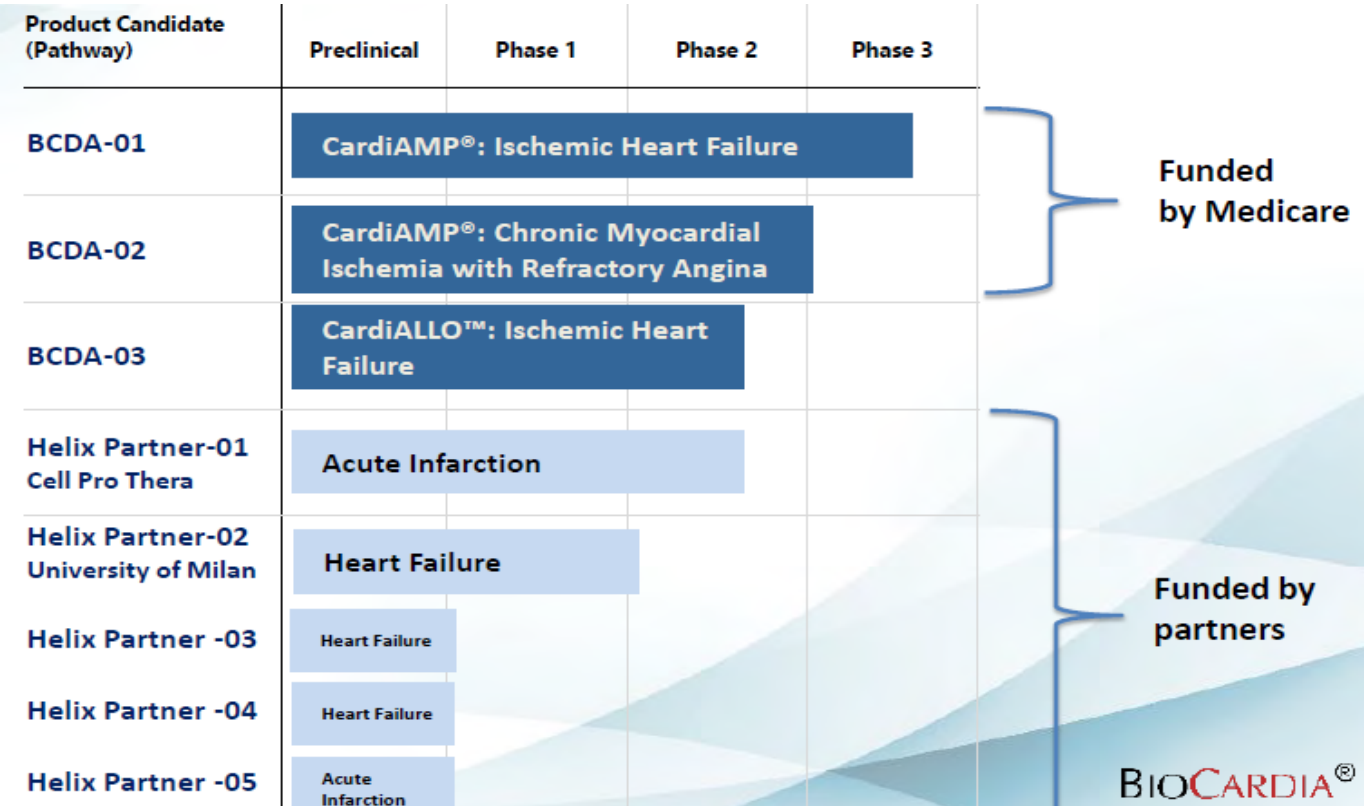
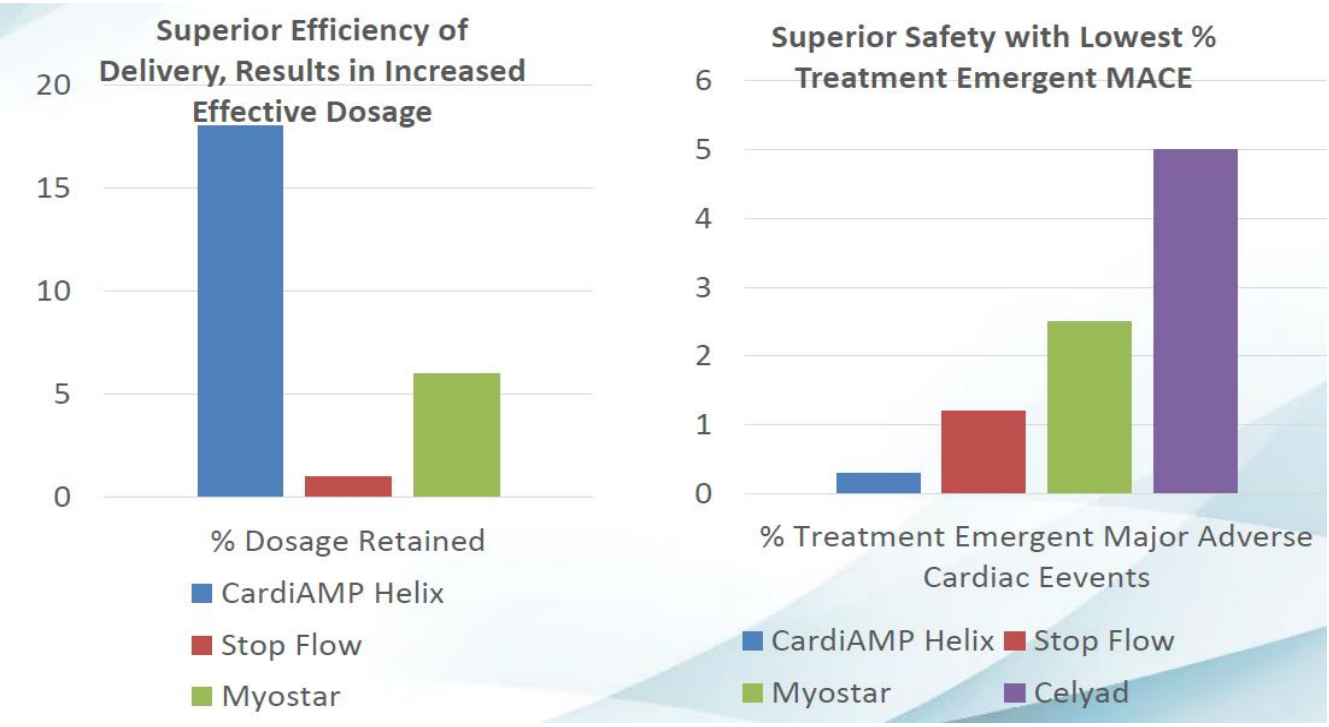
Exhibit 4. Advanced clinical pipeline



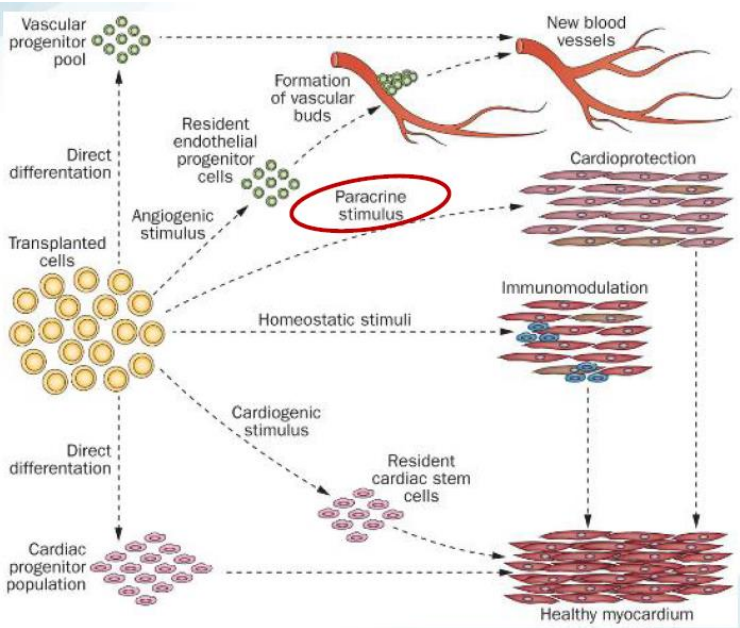
Source: BioCardia Presentation.

Exhibit 5. Performance of CardiAMP Cell Delivery with Proprietary Helix – The Ability to Deliver and Retain Dose on Target



Source: Mitsutake et al, Int.Heart J. (2017), Duckers H et al, Transcatheter Therapeutics 2018.

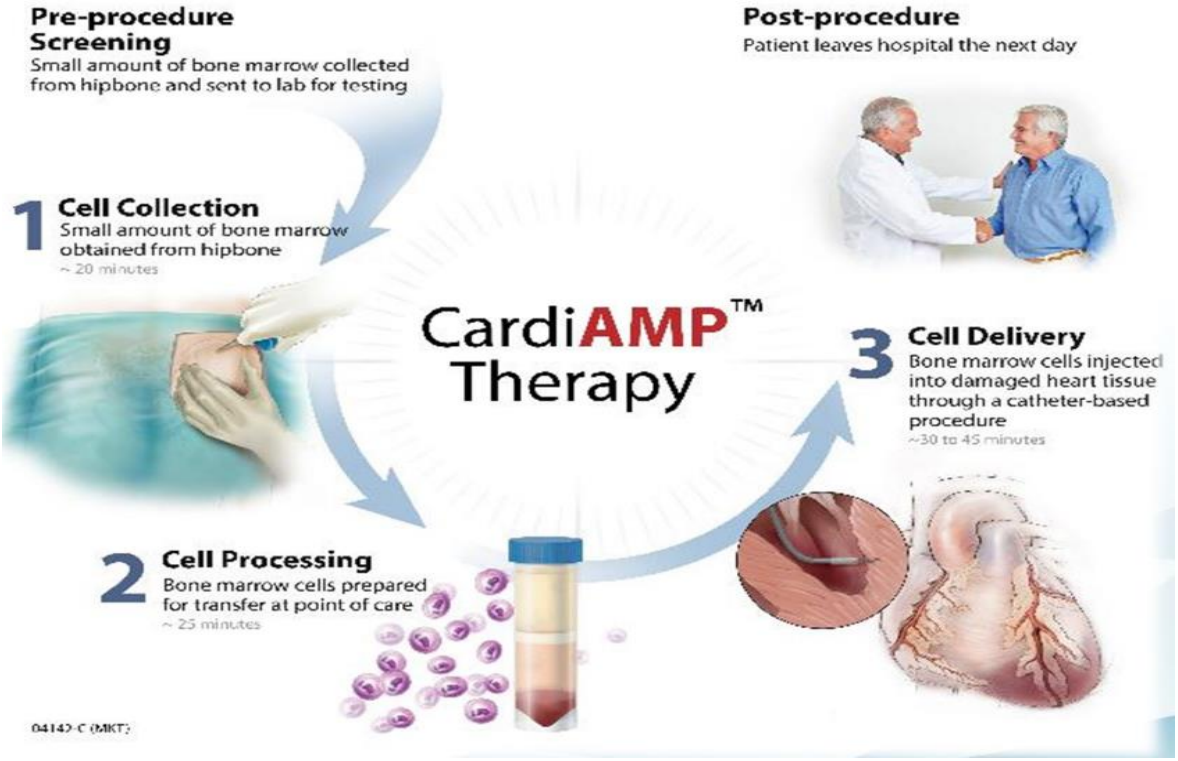
Exhibit 6. Mechanistic basis for regeneration. Transplanted cells are hypothesized to benefit the heart through direct and indirect pathways.



- Direct Regeneration: Transplanted cells actively home to injury sites and differentiate into new functional tissue to augment organ function.
- Indirect Regeneration: Transplanted cells secrete stimulatory cytokines to instigate an innate regenerative response from resident stem cells.
- Accelerating a natural process: Cells from bone marrow home to injury in the heart. Animal work supports reduced fibrotic scar and enhanced micro vessel density. Clinical work supports that there is benefit in this approach.

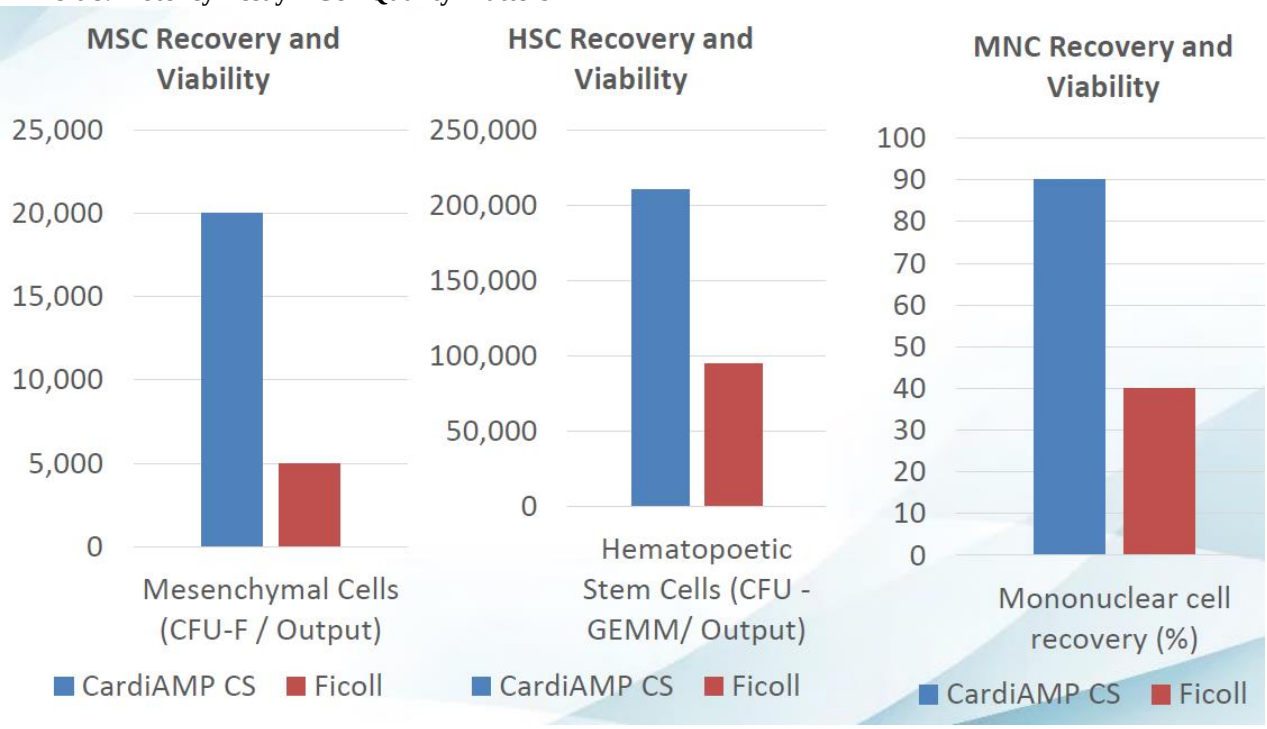
Source: BioCardia July 2019 Presentation.

Exhibit 7. Investigational CardiAMP (Autologous Model) cell therapy system, from collection and screening to delivery with a specialized catheter.



Source: BioCardia July 2019 Presentation.

Exhibit 8. Potency Assay – Cell Quality Matters

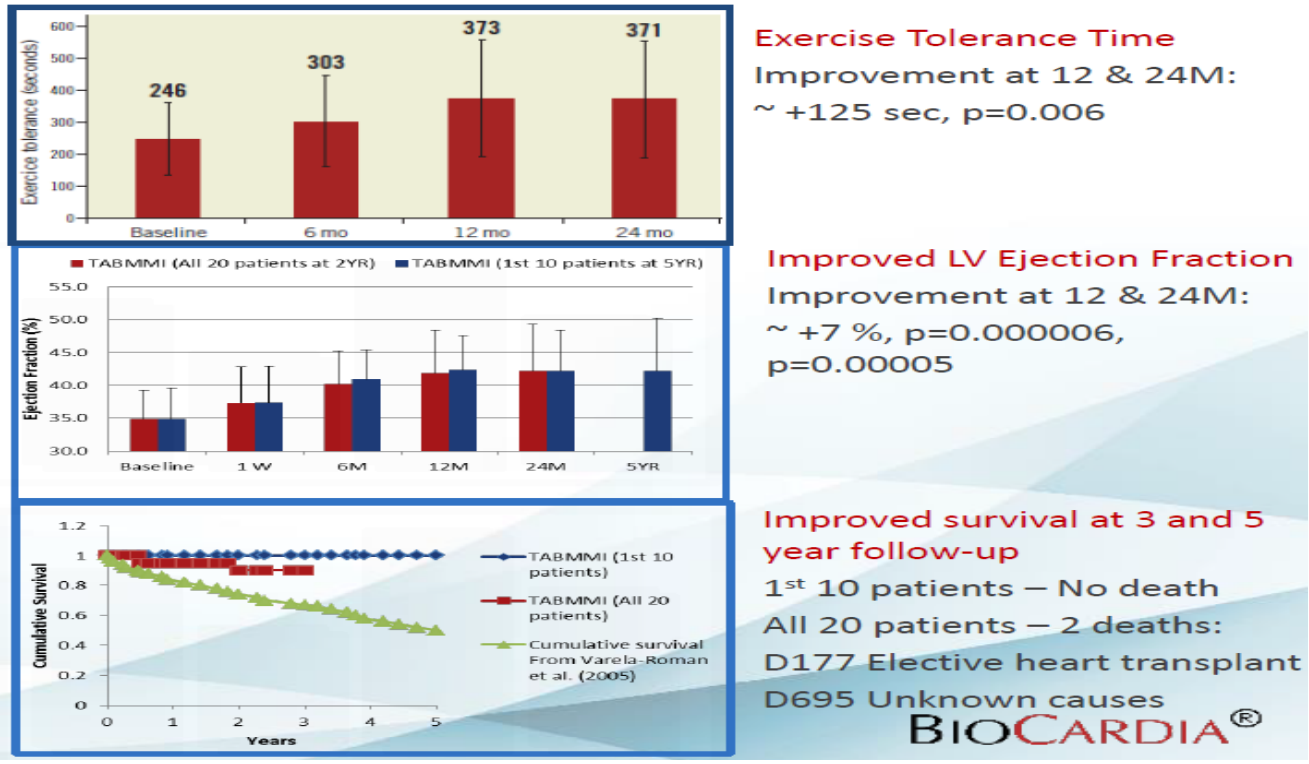


Source: BioResearch Open Access, Volume 4.1, 2015

Exhibit 9. Investigational CardiAMP cell therapy system pre-procedure screening selects patients likely to respond to CardiAMP.

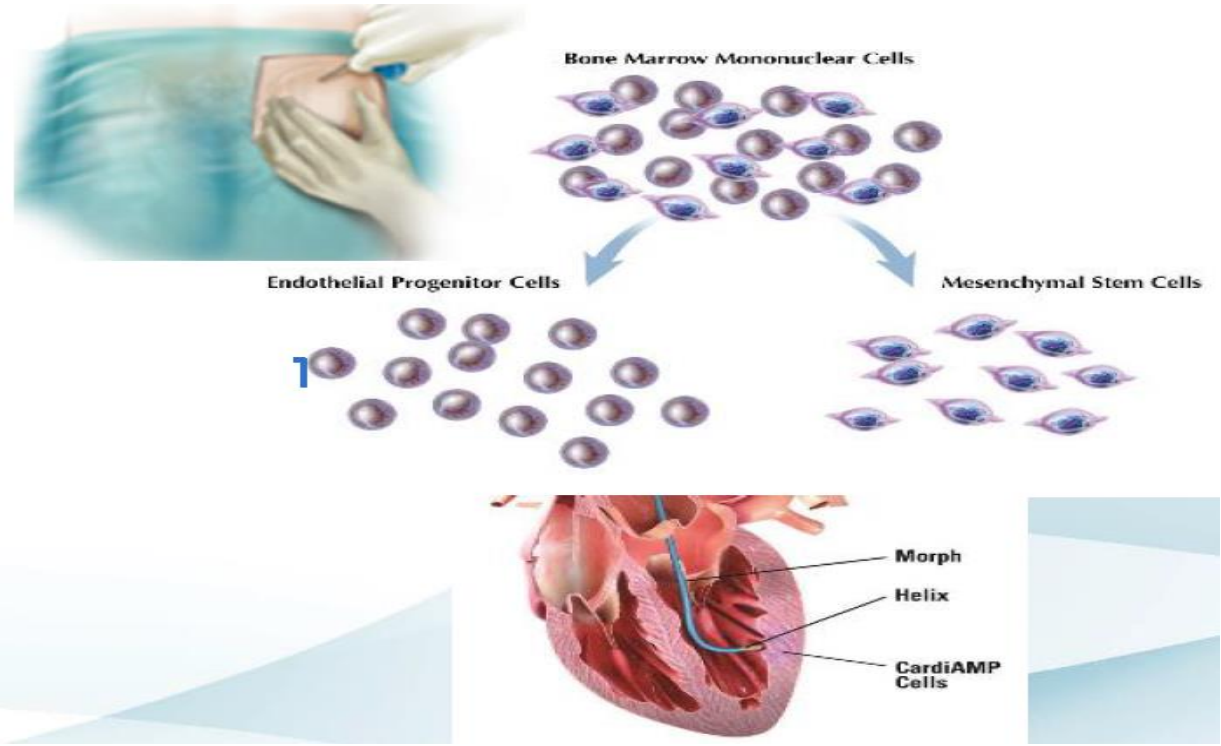
Source: BioCardia July 2019 Presentation.

Exhibit 10. On All Key Metrics (used for approval by regulators), CardiAMP has Shown Improvement.



Source: BioCardia July 2019 Presentation.

Exhibit 11. Transendocardial Autologous Cells in Heart Failure Trial (TAC-HFT)



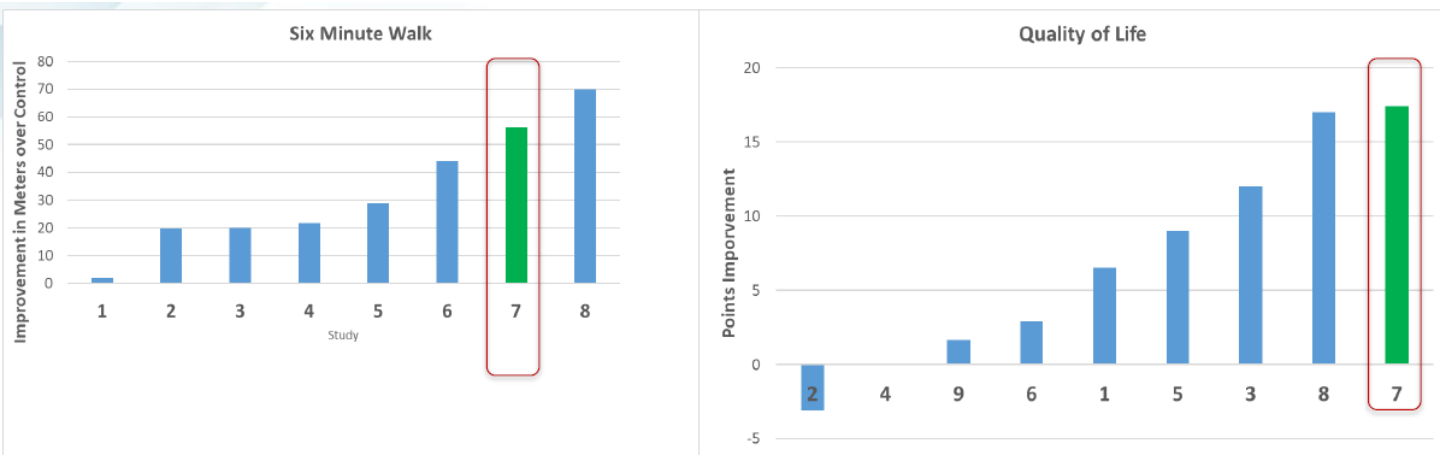
Source: BioCardia July 2019 Presentation.

Exhibit 12. Phase 2 Placebo Controlled Randomized Trial Results: Improvement Shown in Key Metrics.

Secondary Efficacy Endpoints	Active (Mean)	Placebo (Mean)	Treat. Difference	Favors CardiAMP Therapy	P-value
6 minute walk (meters) N=28, Mean ± St Dev	+14.3 ± 59.6	-42.0 ± 18.1	+56.3	✓	0.049
MLHF quality of life (pts) N= 29, Mean ± St Dev	-7.7 ± 17.8	+9.7 ± 24.8	-17.4	✓	0.038
Maximum Oxygen Use (mL/kg·min)	+0.16	-0.870	+1.03	✓	0.321 NS*
NY Heart Association Class	-0.42	-0.25	-0.17	✓	0.638 NS
LV End Systolic Volume (ml)	+3.2	+47.2	-44	✓	0.129 NS
LV End Diastolic Volume (ml)	+4.5	+51.2	-46.7	✓	0.149 NS
LV Ejection Fraction (%)	+0.97	-2.38	+3.35	✓	0.252 NS

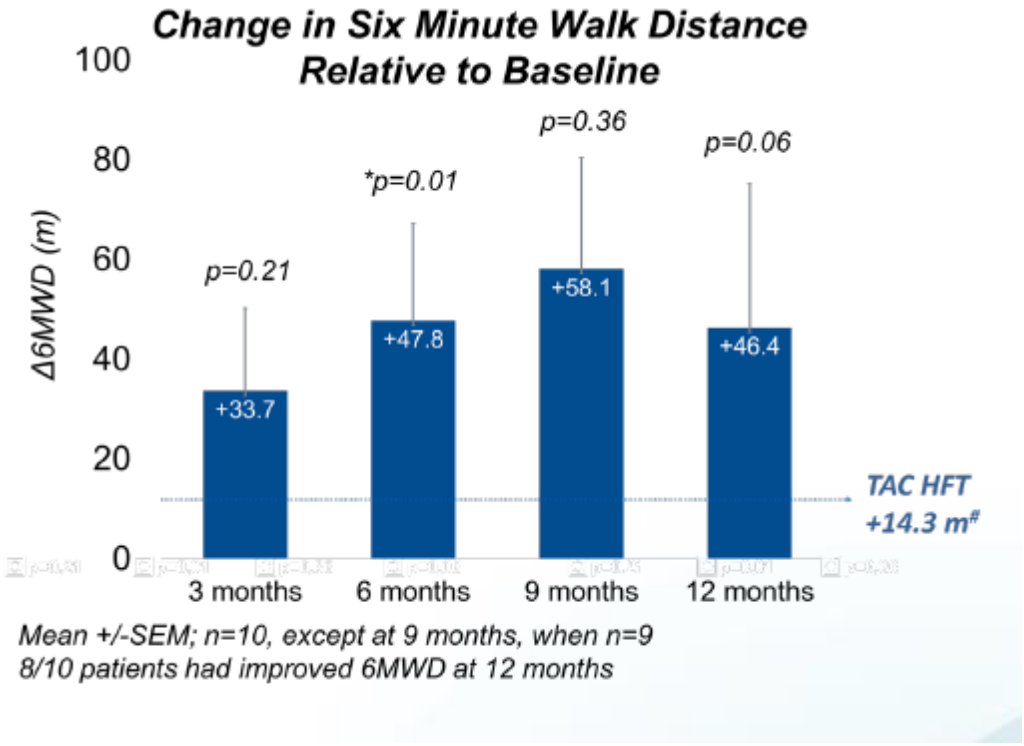
Source: BioCardia July 2019 Presentation.

Exhibit 13. Phase 2 Efficacy Results in Six Minute Walk Relative to Other (Entresto) CRT and Heart Failure Therapies.



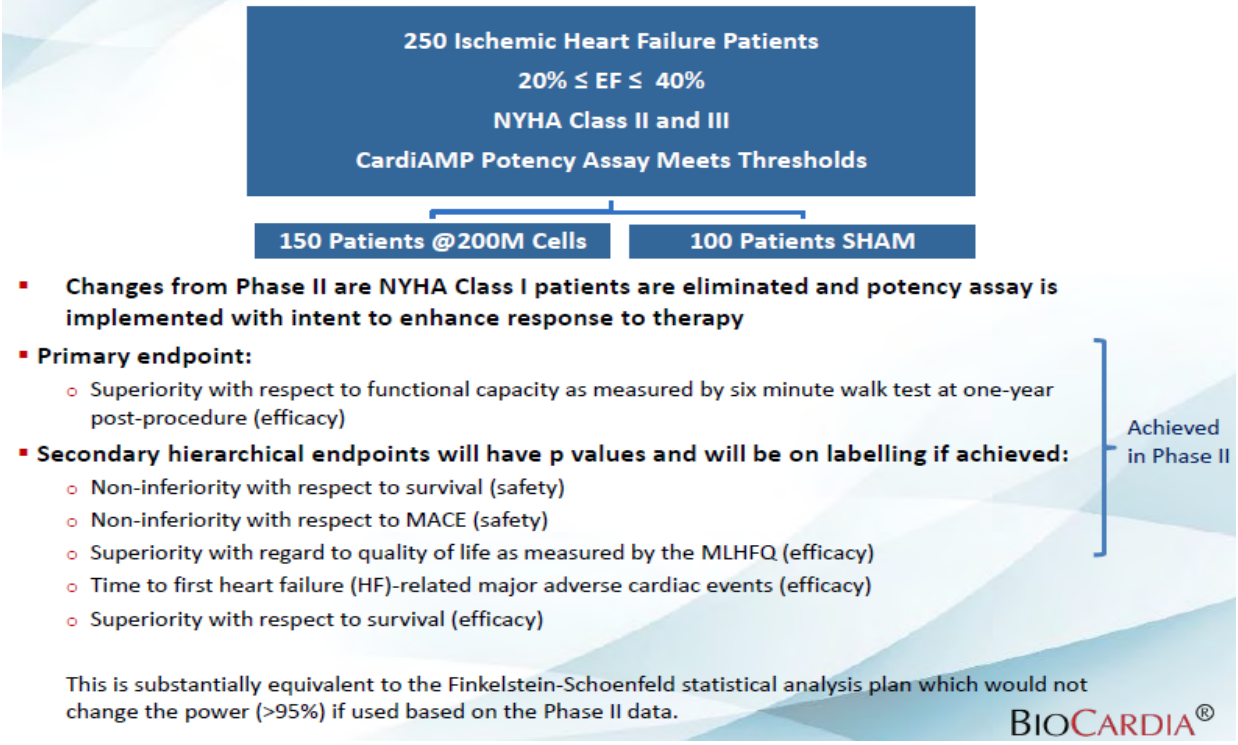
Source: BioCardia July 2019 Presentation.

Exhibit 14. Phase 3: Change in Six Minute Walk Distance (relative to base line). Data Safety Monitoring Board (DSMB) prespecified interim analysis of safety outcomes for the first 10 patients treated in the Phase 3 trial of its investigational CardiAMP cell therapy product no significant safety concerns with the CardiAMP study results and recommended that the trial continue, as planned.



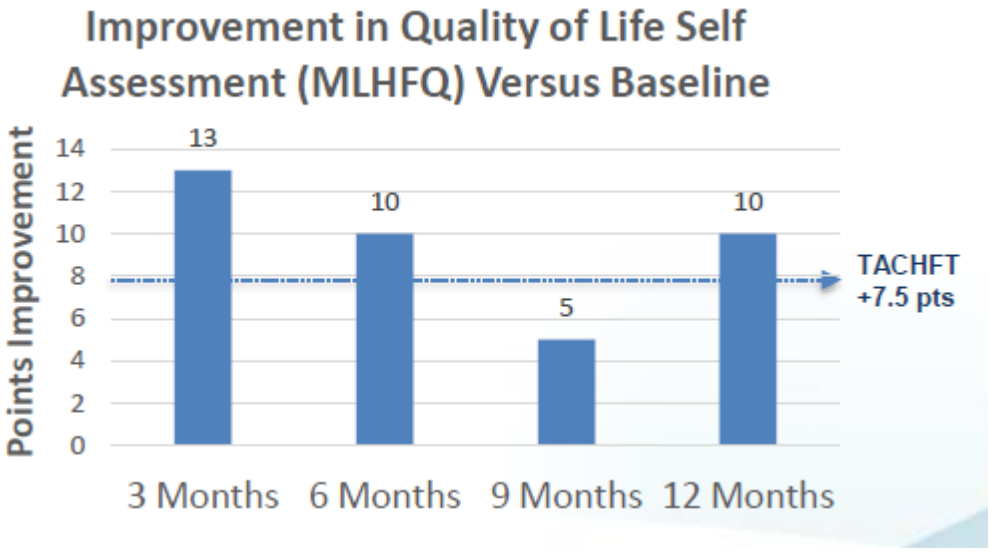
Source: BioCardia July 2019 Presentation.

Exhibit 15. Phase 3 Trial Design

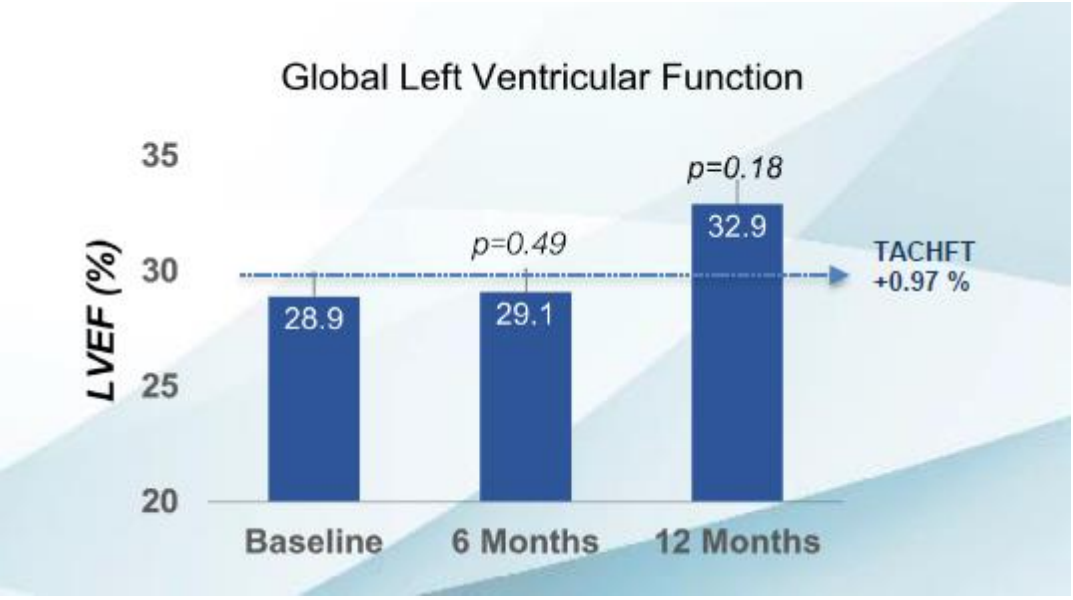


Source: BioCardia Presentation

Exhibit 16. Phase 3: QOL Measures. Results from CardiAMP-HF Study Roll-In Phase: 12-month follow-up results compared to baseline (shown as mean ±sem). N=10 for baseline, 3 months, and 6 months; N=9 for 9 months where one patient was hospitalized, N = 10 for 12 months. Results compared to improvements in Phase 2 TAC-HFT Study.



Source: BioCardia July 2019 Presentation.



Source: BioCardia July 2019 Presentation.

Product Modeling Assumptions

1. We base our market share estimates for BioCardia's therapies on multiple assumptions around the product attributes associated with product delivery and outcomes. Given the blockbuster size of the U.S. and EU HF markets we apply a 70% risk reduction to the net revenues, suggesting on success there is a lot of upside in our estimates.
2. We assume CardiAMP and CardiALLO will initially launch at $50,000 per operation. Our projected market share grows over the six-year launch cycle, with CardiAMP consuming up to 25% of the total market in the year 2028 and CardiALLO achieves a 15% share of the total market by the year 2030.
3. Based on BioCardia's estimation of completing enrollment of the pivotal Phase 3 trial of BCDA-01 in 2H20, we assume standard FDA review time of 10 months with the launch in 1Q22. We expect BCDA-02 to be a fast follower with approval and launch a year after BCDA-01, in 1Q23. To adjust for the risk of approval, we apply a 70% risk cut to our CardiAMP revenue model.
4. We expect that BCDA-03 (CardiALLO) could reach the market by 2027, however, for conservatism, we apply a therapeutic risk cut of 70% in our product model. This suggests that clinical progress could make our numbers too conservative.

Exhibit 17. Market Models:

BCDA-01 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
U.S. Prevalance CHF	5,000,000	5,005,000	5,010,005	5,015,015	5,020,030	5,025,050	5,030,075	5,035,105	5,040,140	5,045,180	5,050,226	5,055,276
Market Size Growth (Annual)	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%
Candidates (Class II & III) for Therapy	500,000	500,500	501,001	501,502	502,003	502,505	503,008	503,511	504,014	504,518	505,023	505,528
Market Share Penetration	0.0%	0.0%	0.0%		1.0%	3.0%	6.0%	7.0%	8.0%	10.0%	14.0%	15.0%
Number of Patients Procedures	0	0	0	0	5,020	15,075	30,180	35,246	40,321	50,452	70,703	75,829
Cost of Therapy	$ 20,000	$ 20,040	$ 20,080	$ 20,120	$ 20,160	$ 20,201	$ 20,241	$ 20,282	$ 20,322	$ 20,363	$ 20,404	$ 20,444
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ 30	$ 91	$ 183	$ 214	$ 246	$ 308	$ 433	$ 465

BCDA-01 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
E.U. Prevalance CHF	9,000,000	9,009,000	9,018,009	9,027,027	9,036,054	9,045,090	9,054,135	9,063,189	9,072,253	9,081,325	9,090,406	9,099,496
Market Size Growth (Annual)	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%
Candidates (Class II & III) for Therapy	900,000	900,900	901,801	902,703	903,605	904,509	905,414	906,319	907,225	908,132	909,041	909,950
Market Share Penetration	0.0%	0.0%	0.0%				1.0%	3.0%	5.0%	10.0%	12.0%	15.0%
Number of Patients Procedures	0	0	0	0	0	0	9,054	27,190	45,361	90,813	109,085	136,492
Cost of Therapy	$ 30,000	$ 30,060	$ 30,120	$ 30,180	$ 30,241	$ 30,301	$ 30,362	$ 30,423	$ 30,483	$ 30,544	$ 30,605	$ 30,667
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ -	$ -	$ 82	$ 248	$ 415	$ 832	$ 1,002	$ 1,256

BCDA-02 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
U.S. Prevalance CHF	5,000,000	5,075,000	5,151,125	5,228,392	5,306,818	5,386,420	5,467,216	5,549,225	5,632,463	5,716,950	5,802,704	5,889,745
Market Size Growth (Annual)	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%	1.5%
Candidates (Chronic Myocardial Ischemia) for Therapy	200,000	203,000	206,045	209,136	212,273	215,457	218,689	221,969	225,299	228,678	232,108	235,590
Market Share Penetration	0.0%	0.0%	0.0%	0.0%	2.0%	5.0%	10.0%	15.0%	20.0%	25.0%	25.0%	25.0%
Number of Patients Procedures	0	0	0	0	4,245	10,773	21,869	33,295	45,060	57,169	58,027	58,897
Cost of Therapy	$ 20,000	$ 20,040	$ 20,080	$ 20,120	$ 20,160	$ 20,201	$ 20,241	$ 20,282	$ 20,322	$ 20,363	$ 20,404	$ 20,444
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ 26	$ 65	$ 133	$ 203	$ 275	$ 349	$ 355	$ 361

BCDA-02 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
E.U. Prevalance CHF	9,000,000	9,072,000	9,144,576	9,217,733	9,291,474	9,365,806	9,440,733	9,516,259	9,592,389	9,669,128	9,746,481	9,824,453
Market Size Growth (Annual)	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%	0.8%
Candidates (Chronic Myocardial Ischemia) for Therapy	495,000	498,960	502,952	506,975	511,031	515,119	519,240	523,394	527,581	531,802	536,056	540,345
Market Share Penetration	0.0%	0.0%	0.0%	0.0%			2.0%	6.0%	10.0%	15.0%	20.0%	20.0%
Number of Patients Procedures	0	0	0	0	0	0	10,385	31,404	52,758	79,770	107,211	108,069
Cost of Therapy	$ 30,000	$ 30,060	$ 30,120	$ 30,180	$ 30,241	$ 30,301	$ 30,362	$ 30,423	$ 30,483	$ 30,544	$ 30,605	$ 30,667
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ -	$ -	$ 95	$ 287	$ 482	$ 731	$ 984	$ 994

BCDA-03 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
U.S. Prevalance CHF	5,000,000	5,005,000	5,010,005	5,015,015	5,020,030	5,025,050	5,030,075	5,035,105	5,040,140	5,045,180	5,050,226	5,055,276
Market Size Growth (Annual)	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%
Candidates (Class II & III) for Therapy	150,000	150,150	150,300	150,450	150,601	150,752	150,902	151,053	151,204	151,355	151,507	151,658
	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	2.0%	5.0%	10.0%	15.0%
Number of Patients Procedures	0	0	0	0	0	0	0	0	3,024	7,568	15,151	22,749
Cost of Therapy	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ 22,500	$ 22,545	$ 22,590	$ 22,635
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ 20	$ 51	$ 103	$ 154

BCDA-03 Heart Failure	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030
E.U. Prevalance CHF	9,000,000	9,009,000	9,018,009	9,027,027	9,036,054	9,045,090	9,054,135	9,063,189	9,072,253	9,081,325	9,090,406	9,099,496
Market Size Growth (Annual)	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%	0.1%
Candidates (Class II & III) for Therapy	270,000	270,270	270,540	270,811	271,082	271,353	271,624	271,896	272,168	272,440	272,712	272,985
Market Share Penetration	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	2.0%	5.0%	10.0%	15.0%
Number of Patients Procedures	0	0	0	0	0	0	0	0	5,443	13,622	27,271	40,948
Cost of Therapy			$ -	$ -	$ -	$ -	$ -	$ -	$ 29,500	$ 29,559	$ 29,618	$ 29,677
Price Growth	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%	0.2%
Probability of Success	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%	30%
U.S. Annual Sales (M)	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ -	$ 48	$ 121	$ 242	$ 365

Source: Dawson James Estimates

Exhibit 18. Comparables. The attached chart reviews some of the comparable regenerative medicine-based companies. We believe notable mentions for Blue Rock Therapeutics and Sana Biotechnology. Blue Rock was acquired in August 2019 by Bayer for $1 billion. What's important is that Blue Rock is developing a universal pluripotent stem cells into authentic, functional cells that can be used as allogeneic cellular therapies to treat a broad array of diseases. Bayer announced the acquisition of their remaining stake for approximately $240 million in cash to be paid upfront at closing and an additional $360 million payable upon achievement of pre-defined development milestones. With Bayer currently holding 40.8% stake, the investment corresponds to a total company value of Blue Rock Therapeutics of approximately $1 billion. Sana was founded by former executives of June and includes Steve Harr MD and a former biotechnology analyst and Hans Bishop and has been backed by ARCH Ventures and F-Prime Capital.

Company Name	Ticker	Share Price	Market Cap ($MM)	Cash ($MM)	Enterprise Value ($MM)	Last QTR's Burn Rate	QTR's of Cash Remaining	R&D ($MM)	SG&A ($MM)	Shares Outstanding (MM)
Athersys	ATHX	$2.70	$529	$33	$406	($16)		$12	$16	196
BioCardia	BCDA	$2.29	$28	$3	$14	($5)		$3	$5	12
Mesoblast	MESO	$13.00	$1,449	$104	$1,360	($8)		$13	$8	117
Pluristem	PSTI	$8.78	$219	$7	$143	($8)		$6	$7	25
Vericel	VCEL	$14.99	$674	$46	$619	($5)		$4	$22	45
Average (s)	¯X	$8.35	$580	$38	$508	($8)		7	11	79
BioCardia	BCDA	$2.29	$28.29	$3	$14	($5)		3	5	12

Source: Data is from FactSet as of 7/7/2020

Valuation. We value BioCardia based on the revenues in our product models, which are reduced by 70% rate cut or a 30% probability of success. This suggests we are leaving a lot of upside in our estimates on good data. We apply assumptions for FDA product approvals, launch dates, and product attributes to estimate revenues. These estimates feed into our income statement through the year 2030. Our price target is derived from an equal-weighted average of free cash flow (FCFF), discounted EPS (EPS), and sum-of-the-parts (SOP) models. For companies that are well established with mature products and revenues, we typically discount at a 10% rate, for companies in the early stages of product commercialization we typically use a higher rate, 15%. For BioCardia, we use a 30% risk rate as the company is not yet profitable. This risk rate is in addition to the therapeutic cut (70%) in our product models. The result of this methodology is a price target of $24.00 per share.

Exhibit 19. Discounted Free Cash Flow Model

	Average	$ 24

	Price Target	$ 23
	Year	2020

DCF Valuation Using FCF (mln):

units ('000)	2019E	2020E	2021E	2022E	2023E	2024E	2025E	2026E	2027E	2028E	2029E	2030E
EBIT	(14,709)	(15,544)	(15,189)	(27,353)	10,270	84,137	339,759	497,501	1,120,811	1,849,568	2,452,570	2,831,575
Tax Rate	0%	0%	0%	10%	18%	20%	24%	24%	24%	28%	30%	34%
EBIT(1-t)	(14,709)	(15,544)	(15,189)	(24,618)	8,422	67,309	258,217	378,101	851,816	1,331,689	1,716,799	1,868,839
CapEx												
Depreciation	78	22	-	-	-	-	-	-	-	-	-	-
Change in NWC												
FCF	(14,631)	(15,522)	(15,189)	(24,618)	8,422	67,309	258,217	378,101	851,816	1,331,689	1,716,799	1,868,839
PV of FCF	(6,660)	(5,435)	(4,091)	(5,100)	1,342	8,251	24,350	27,427	135,751	163,251	161,893	135,562

Discount Rate	30%	
Long Term Growth Rate	1%	
Terminal Cash Flow	6,508,717	
Terminal Value YE2030	472,130	
NPV	1,115,332	
NPV-Debt	1,016	
Shares out (thousands)	48,143	2030E
NPV Per Share	$ 23	

Source: Dawson James

Exhibit 20. EPS Model

Current Year		2020
Year of EPS		2030
Earnings Multiple		10
Discount Factor		30%
Selected Year EPS	$	38.82
NPV	$	28.00

Source: Dawson James estimates.

Discount Rate and Earnings Multiple Varies, Year is Constant							
		2030 EPS					
Earnings Multiple		5%	10%	15%	20%	25%	30%
	0	$0.00	$0.00	$0.00	$0.00	$0.00	$ -
	5	$119.00	$75.00	$48.00	$31.00	$21.00	$ 14.00
	10	$238.00	$150.00	$96.00	$63.00	$42.00	$ 28.00
	15	$357.00	$224.00	$144.00	$94.00	$63.00	$ 42.00
	20	$477.00	$299.00	$192.00	$125.00	$83.00	$ 56.00
	25	$596.00	$374.00	$240.00	$157.00	$104.00	$ 70.00
	30	$715.00	$449.00	$288.00	$188.00	$125.00	$ 84.00
	35	$834.00	$524.00	$336.00	$219.00	$146.00	$ 99.00

Source: Dawson James

Exhibit 21. Sum-of-the-Parts Model

BioCardia	LT Gr	Discount Rate	Yrs. to Mkt	% Success	Peak Sales MM's	Term Val
BCDA-01 CardiAMP cell therapy US	1%	30%	3	30%	$1,550	$5,346
NPV						$6.07
BCDA-01 CardiAMP cell therapy US	1%	30%	3	30%	$4,186	$14,434
NPV						$16.38
BCDA-02 CardiALLO cell therapy US	1%	30%	4	30%	$1,204	$4,152
NPV						$3.62
BCDA-02 CardiALLO cell therapy EU	1%	30%	4	30%	$3,314	$11,428
NPV						$9.97
BCDA-03 CardiALLO cell therapy US	1%	50%	5	30%	$515	$1,051
NPV						$0.34
BCDA-02 CardiALLO cell therapy EU	1%	50%	5	30%	$1,215	$2,480
NPV						$0.81
Net Margin						40%
MM Shrs OS (2030E)						48
Total						$21

Source: Dawson James

Risk Analysis

In addition to the typical risks associated with development stage specialty pharmaceutical companies, potential risks specific to BioCardia are as follows:

Partnership risk. The company is also expected to make agreements with partners for additional products, but there can be no assurances that the company will be able to secure favorable partnerships.

Commercial risk. There are no assurances that the company will be able to achieve significant sales, market share, or become profitable.

Clinical and regulatory risk. Lead products need to complete clinical trials. It is difficult to complete enrollment which could lead to a delay of the trial. Trials may not produce the results expected from previous research or be sufficient for regulatory approval.

Financial risk. The company may need to raise capital in the marketplace, and there can be no assurances that the company will be able to successfully raise capital and or do so, at favorable terms.

Legal and intellectual property risk. The company may have to defend its patents and technical know-how, and there can be no assurances that the patents will not be infringed or will be held as valid if challenged, and or that the company may infringe on third parties' patents.

Exhibit 22. Income Statement

Biocardia Inc: Income Statement ($000)																	
BCDA: YE December	**2018A**	**2019E**	**1Q20A**	**2Q20E**	**3Q20E**	**4Q20E**	**2020E**	**2021E**	**2022E**	**2023E**	**2024E**	**2025E**	**2026E**	**2027E**	**2028E**	**2029E**	**2030E**
Net product revenue	282	300	5	0	1	1	7	7									
Collaboration agreement revenue	343	382	33	27	25	25	110	116									
BCDA-01 CardiaAMP Cell Therapy revenues US		0					0	0	0	30,362	91,359	183,267	214,453	245,825	308,203	432,780	465,085
BCDA-01 CardiaAMP Cell Therapy revenues EU									0	0	0	82,470	248,153	414,829	832,149	1,001,577	1,255,729
BCDA-02 CardiaAMP Cell Therapy revenues US										25,677	65,286	132,796	202,586	274,714	349,241	355,188	361,237
BCDA-02 CardiaAMP Cell Therapy revenues EU										-	-	94,590	28,661	482,474	730,959	984,374	994,234
BCDA-03 CardALLO Cell Therapy revenues US													-	20,413	51,185	102,677	154,477
BCDA-03 CardALLO Cell Therapy revenues EU	-												-	48,174	120,796	242,317	364,566
Total Product Sales	**625**	**599**	**38**	**27**	**26**	**26**	**117**	**123**	**-**	**56,039**	**156,645**	**493,123**	**693,853**	**1,486,428**	**2,392,533**	**3,118,913**	**3,595,329**
Product Sales & Royalties & Milestones	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**56,039**	**156,645**	**316,062**	**417,039**	**589,125**	**829,425**	**1,132,962**	**1,345,366**
Expenses																	
Cost of goods sold	517	543	4	0	8	8	20	37	0	14,010	39,161	118,349	159,586	327,014	502,432	623,783	719,066
			30%	30%	30%	30%	17%	30%	#DIV/0!	25%	25%	24%	23%	22%	21%	20%	20%
Research and Development	8,453	8,876	2,786	2,224	2,330	2,423	9,319	9,785	10,275	10,788	11,328	11,894	12,489	13,113	13,769	14,457	15,180
Selling, general and administrative	5,757	6,045	1,857	1,377	1,587	1,650	6,347	6,664	18,000	22,000	23,100	24,255	25,468	26,741	28,078	29,482	30,956
Total expenses	**14,727**	**15,192**	**4,647**	**3,601**	**3,924**	**4,081**	**16,254**	**16,158**	**27,709**	**46,142**	**72,901**	**153,776**	**196,784**	**366,072**	**543,442**	**666,843**	**764,280**
Operating income (Loss)	(14,102)	(14,593)	(4,609)	(3,574)	(3,898)	(4,055)	(16,137)	(16,035)	(27,709)	9,897	83,745	339,347	497,069	1,120,357	1,849,091	2,452,069	2,831,049
Interest expense																	
Interest Income	118	23	16														
Other expense	(3)	(1)	(1)														
Total other income	**115**	**(1)**	**(1)**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**	**-**
Pretax Income	**(13,987)**	**(14,571)**	**(4,594)**	**(3,574)**	**(3,898)**	**(4,055)**	**(16,137)**	**(16,035)**	**(27,709)**	**9,897**	**83,745**	**339,347**	**497,069**	**1,120,357**	**1,849,091**	**2,452,069**	**2,831,049**
Income Tax Benefit (Provision)		-	-	-	-	-	-	-	(2,771)	1,781	16,749	81,443	119,297	268,886	517,745	735,621	962,557
Tax Rate	0%	0%	0%	0%	0%	0%	0%	0%	10%	18%	20%	24%	24%	24%	28%	30%	34%
GAAP Net Income (loss)	**(13,987)**	**(14,571)**	**(4,594)**	**(3,574)**	**(3,898)**	**(4,055)**	**(16,137)**	**(16,035)**	**(24,938)**	**8,115**	**66,996**	**257,904**	**377,772**	**851,471**	**1,331,346**	**1,716,449**	**1,868,492**
Deemed Dividend on Preferred Stock																	
GAAP-EPS	**(0.37)**	**(0.92)**	**(0.67)**	**(0.46)**	**(0.18)**	**(0.19)**	**(1.11)**	**(0.52)**	**(0.78)**	**0.25**	**2.07**	**7.94**	**11.59**	**26.02**	**40.52**	**52.04**	**56.42**
GAAP EPS (dil)	(0.37)	(0.96)	(0.67)	(0.46)	(0.18)	(0.19)	(1.11)	(0.41)	(0.59)	0.19	1.58	6.05	8.83	19.81	30.86	39.62	42.96
Weighted shares basic	38,285	15,761	6,832	7,711	21,861	21,883	14,572	30,699	32,076	32,204	32,333	32,463	32,593	32,723	32,854	32,986	33,118
Weighted shares dil	38,285	15,136	6,832	7,711	21,861	21,883	14,572	39,460	42,126	42,294	42,464	42,634	42,805	42,976	43,148	43,321	43,495

Source: Dawson James estimates.

Companies mentioned in this report:

Bayer (BAYRY): Not Covered
Blue Rock Therapeutics (acquired by Bayer)
June (acquired by Celgene – CELG, Not Covered).
Sana Biotech (Private)

Important Disclosures:

Price Chart:

Price target and ratings changes over the past three years:
Initiated – Buy – September 17, 2019 – Price Target $24.00
Update – Buy – November 20, 2019 – Price Target $24.00
Update – Buy – February 18, 2020 – Price Target $24.00
Update – Buy – May 5, 2020 – Price Target $24.00
Update – Buy – July 7, 2020 – Price Target $24.00
Update – Buy – August 13, 2020 – Price Target $24.00

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	Company Coverage		Investment Banking	
Ratings Distribution	# of Companies	% of Total	# of Companies	% of Totals
Market Outperform (Buy)	23	85%	4	17%
Market Perform (Neutral)	4	15%	1	25%
Market Underperform (Sell)	0	0%	0	0%
Total	27	100%	5	19%

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