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BriaCell

INVESTOR PRESENTATION

Spring 2026

Developing Novel Therapeutics to Destroy Cancer

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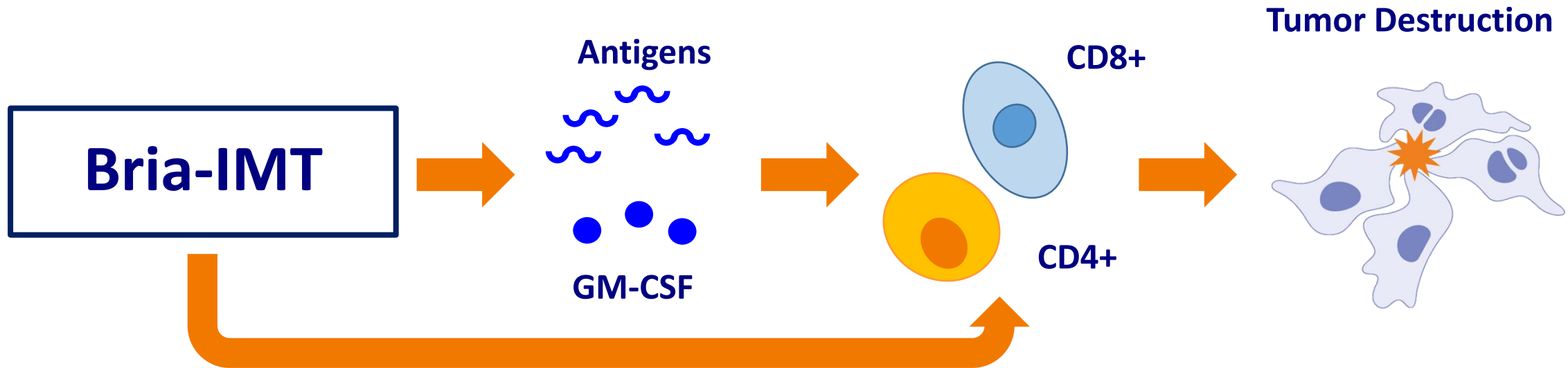
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- **Clinical stage immuno-oncology company developing an entirely new class of targeted immunotherapies to transform cancer care**
- **Lead drug candidate Bria-IMT™**
 - **Pivotal Phase 3 study underway in metastatic breast cancer (over 40K US deaths/year)**
 - Phase 2 study demonstrated 2-fold increase in survival vs comparable patients in the literature
 - Unprecedented clinical benefit in checkpoint inhibitor (CPI) and antibody-drug conjugate (ADC) resistant patients
 - Remarkable clinical efficacy in patients with central nervous system (CNS) metastases
 - Awarded Fast Track designation by FDA
 - Single agent and combination check point inhibitor (+ CPI) activity
- **Bria-OTS™ & Bria-OTS+™: Next generation, cell-based cancer immunotherapy platform**
 - Ongoing Phase 1/2a study in breast cancer
 - Initial positive results with single agent therapy
 - **Bria-OTS+ more potent version of Bria-OTS**
 - Next generation prostate and breast cancer candidates scheduled to enter clinic 2026
 - National Cancer Institute SBIR awards

Bria-IMT: Lead Candidate

- Bria-IMT - a cell based, patented, targeted immunotherapy
- Derived from a well characterized breast cancer cell line
- Expresses tumor antigens and GM-CSF to activate cancer fighting CD4+ and CD8+ T cells
- Stimulates the immune system to enhance targeted killing of cancer cells
- Off-the-shelf approach easy to distribute and administer
 - Cell are grown (cGMP), harvested, irradiated and cryopreserved for shipment to clinical sites where they are thawed and injected intradermally (upper back and thighs)



Evaluable Patients	HLA Match	Disease Control (CR, PR, and SD)	Disease Control in Immune Responders (DTH)
N=5	≥ 2	80% (4/5)	100% (4/4)
N=18	Any	50% (9/18)	60% (9/15)

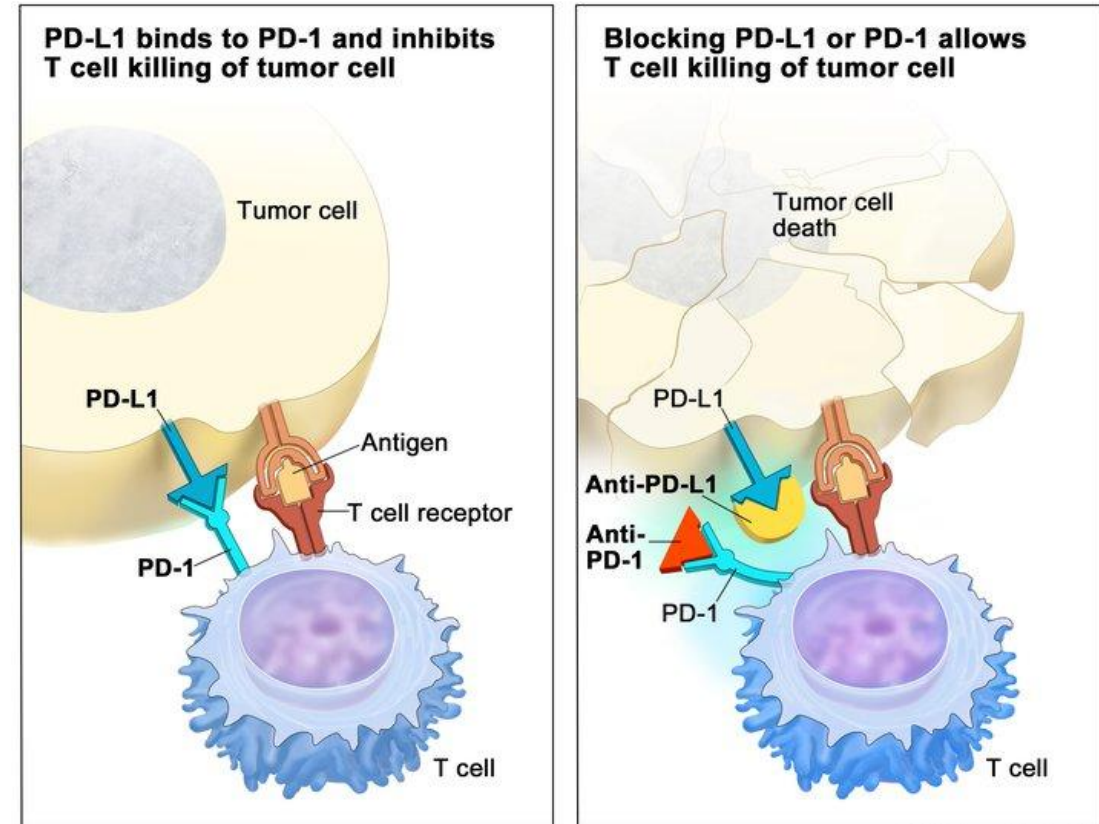
- 27 total heavily pre-treated (median 5 prior regimens) metastatic breast cancer patients treated with Bria-IMT monotherapy regimen, 18 evaluable
- Presence of HLA-type matching correlates with response to Bria-IMT
- Immune response measured by delayed-type hypersensitivity (DTH) to Bria-IMT correlates with disease control
- Tolerability excellent with no dose-limiting toxicities
- Clinical benefit demonstrated: 1 PR and 8 SD in 15 evaluable immune responders

How Do CPIs Work?

- PD-L1 expression protects cancer cells from tumor antigen driven T-cell attack
- PD-1 and PD-L1 inhibitors, also known as CPIs, neutralize this immune suppression

Why combine Bria-IMT with CPIs?

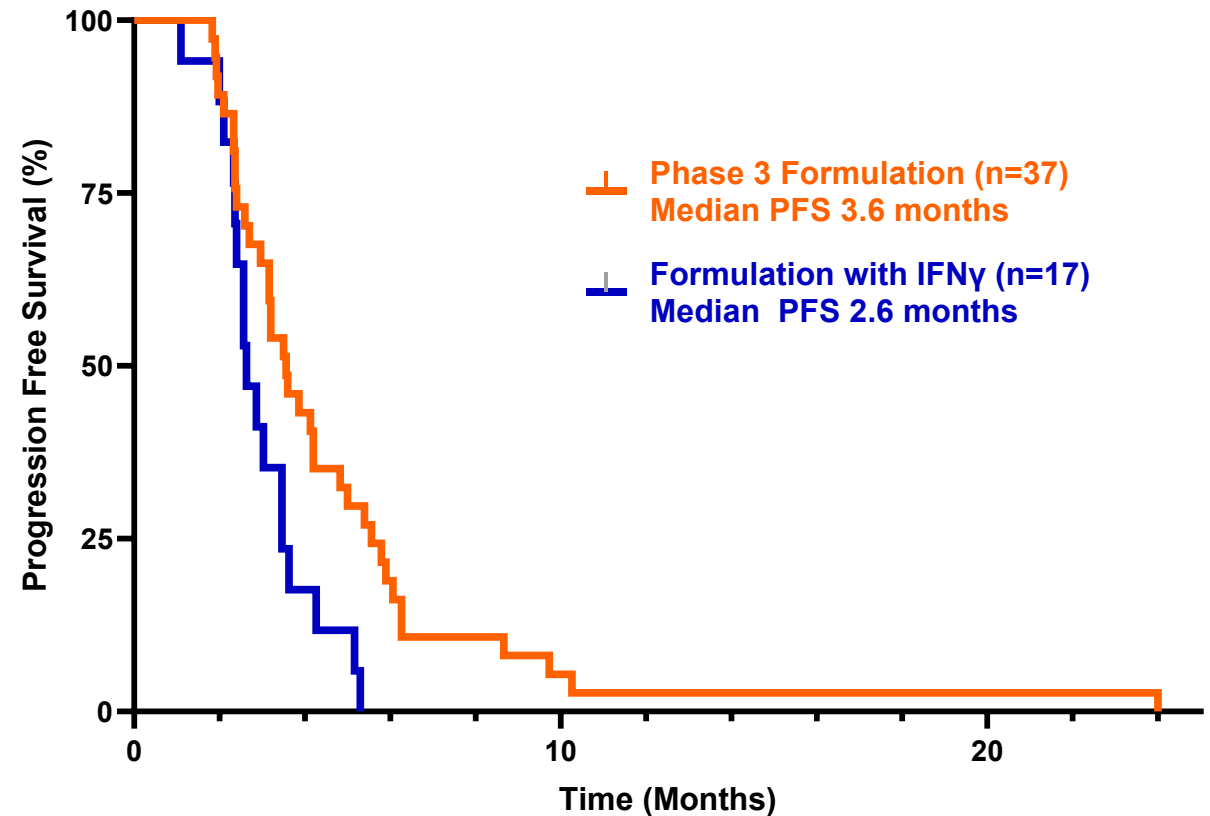
- >90% of patients express PD-L1 in our studies
- Potential synergy between Bria-IMT activated immune system and CPI's unblocking of immune system
- BriaCell's hypothesis: Combining CPIs with Bria-IMT providing powerful synergistic anti-tumor activity



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Bria-IMT + CPI Phase 2 Study

- Total of 54 patients enrolled
 - 11 treated in combination with pembrolizumab
 - 44 treated in combination with retifanlimab
 - one cross-over
- Median 6 prior regimens
- Evaluated 2 formulations:
 - Bria-IMT treated with IFN γ , n = 17
 - Bria-IMT not treated with IFN γ , n = 37
- Progression-free survival (PFS) 3.6 months vs 2.6 months favored no IFN γ (p<0.05)
- PFS of similar patients in the literature is 1.6-2.5 months¹
- Bria-IMT without IFN γ selected as formulation for Phase 3

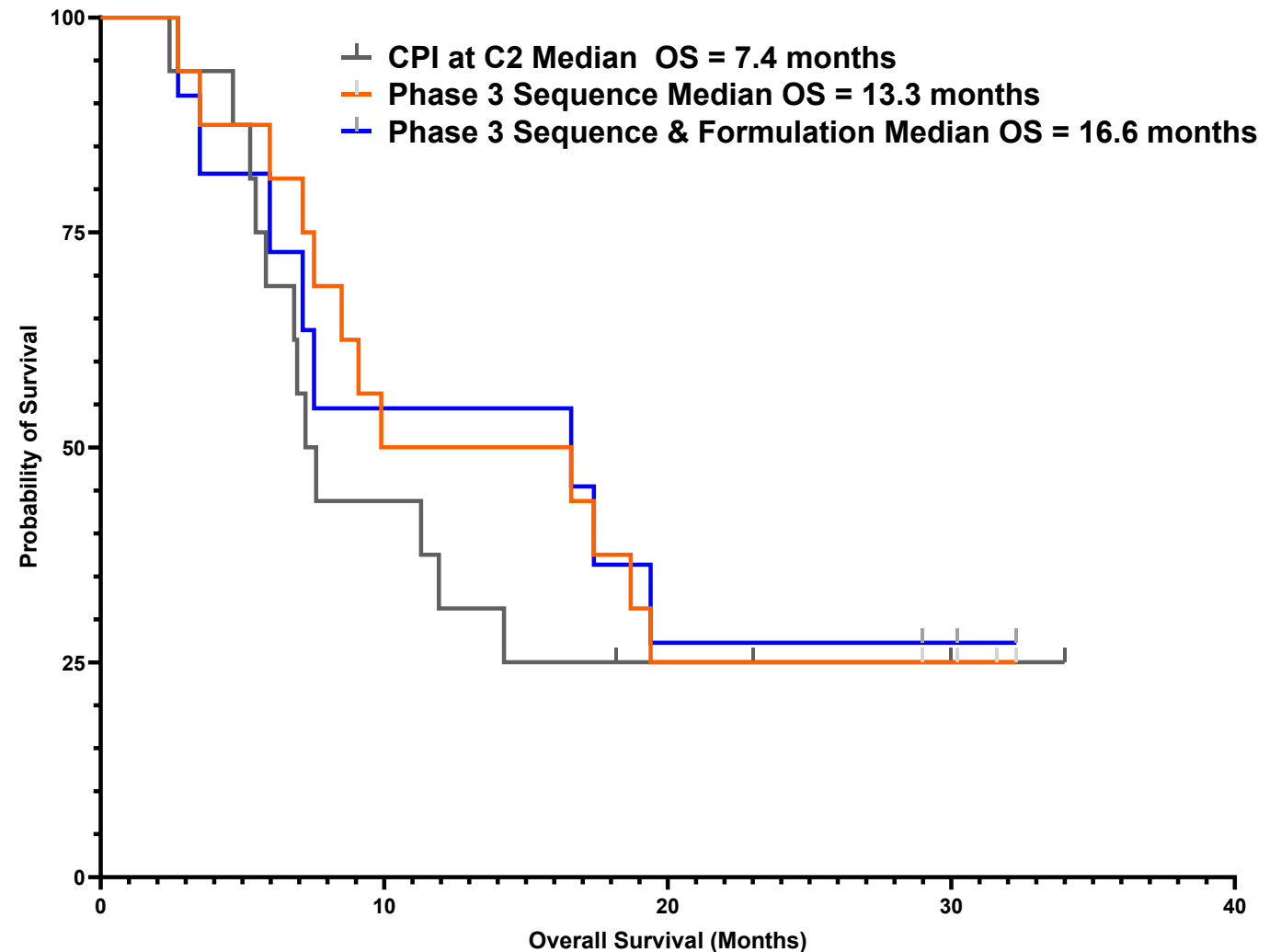


Data presented at ASCO 2026
<https://www.asco.org/abstracts-presentations/261300>

¹Cortes J, et al. Annals of Oncology 2018; Kazmi S, et al. Breast Cancer Res Treat. 2020 Aug 17; O'Shaughnessy J et al. Breast Cancer Res Treat. 2022; Tripathy D, et al. JAMA Oncol. 2022; Bardia A, et al. J Clin Oncol. 2024 May 20;42(15):1738-1744

- 32 patients treated with 2 sequences
 - 16 patients CPI in Cycle 1 (Phase 3 Sequence)
 - A subset also received the Phase 3 formulation
 - 16 patients with CPI in Cycle 2 (CPI in C2)
- Median 6 prior lines of therapy (range 2-13)
- Overall survival (OS) 13.3 months for the Phase 3 sequence
- Overall survival (OS) 16.6 months for the Phase 3 formulation and sequence
- OS compares favorably to 5.9-9.8 months¹ reported in comparable metastatic breast cancer patients
- No dose limiting toxicities to date

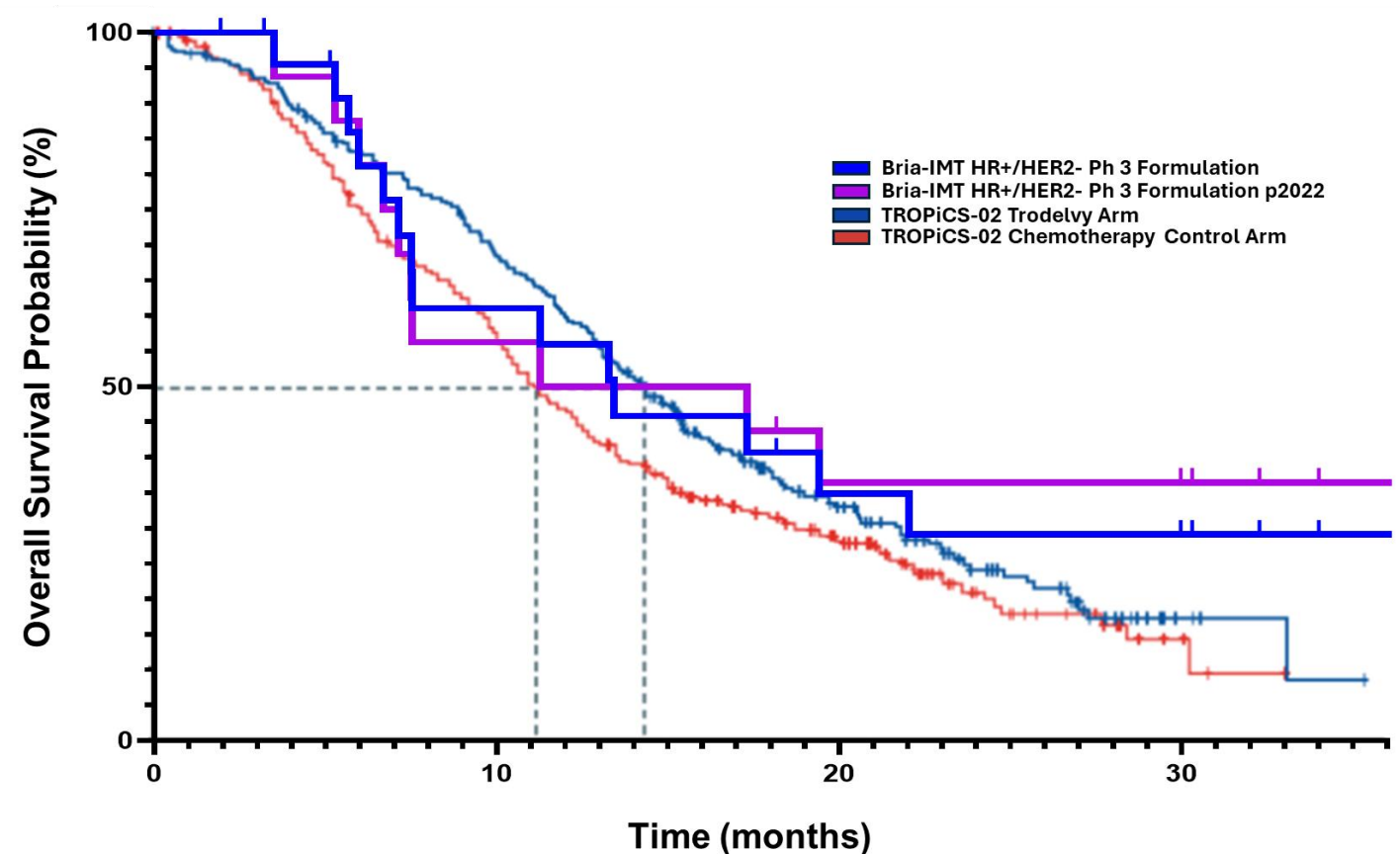
Overall Survival by Sequence and Formulation



¹Cortes J, et al. Annals of Oncology 2018; Kazmi S, et al. Breast Cancer Res Treat. 2020 Aug 17; O'Shaughnessy J et al. Breast Cancer Res Treat. 2022; Tripathy D, et al. JAMA Oncol. 2022; Bardia A, et al. J Clin Oncol. 2024 May 20;42(15):1738-1744

- 37 patients treated with Phase 3 formulation
 - Median 6 prior lines of therapy
 - 24 patients hormone receptor (HR)+
 - Overall survival (OS) 13.4 months for HR+ Patients
 - OS 14.3 months for HR+ Patients treated since 2022
- **Compares well to Trodelvy pivotal registration study in HR+ Patients**
 - Median 4 prior lines of systemic therapy
 - OS 14.4 months for Trodelvy
 - OS 11.2 months for single agent chemotherapy

Overall survival of Bria-IMT vs TROPiCS-02²



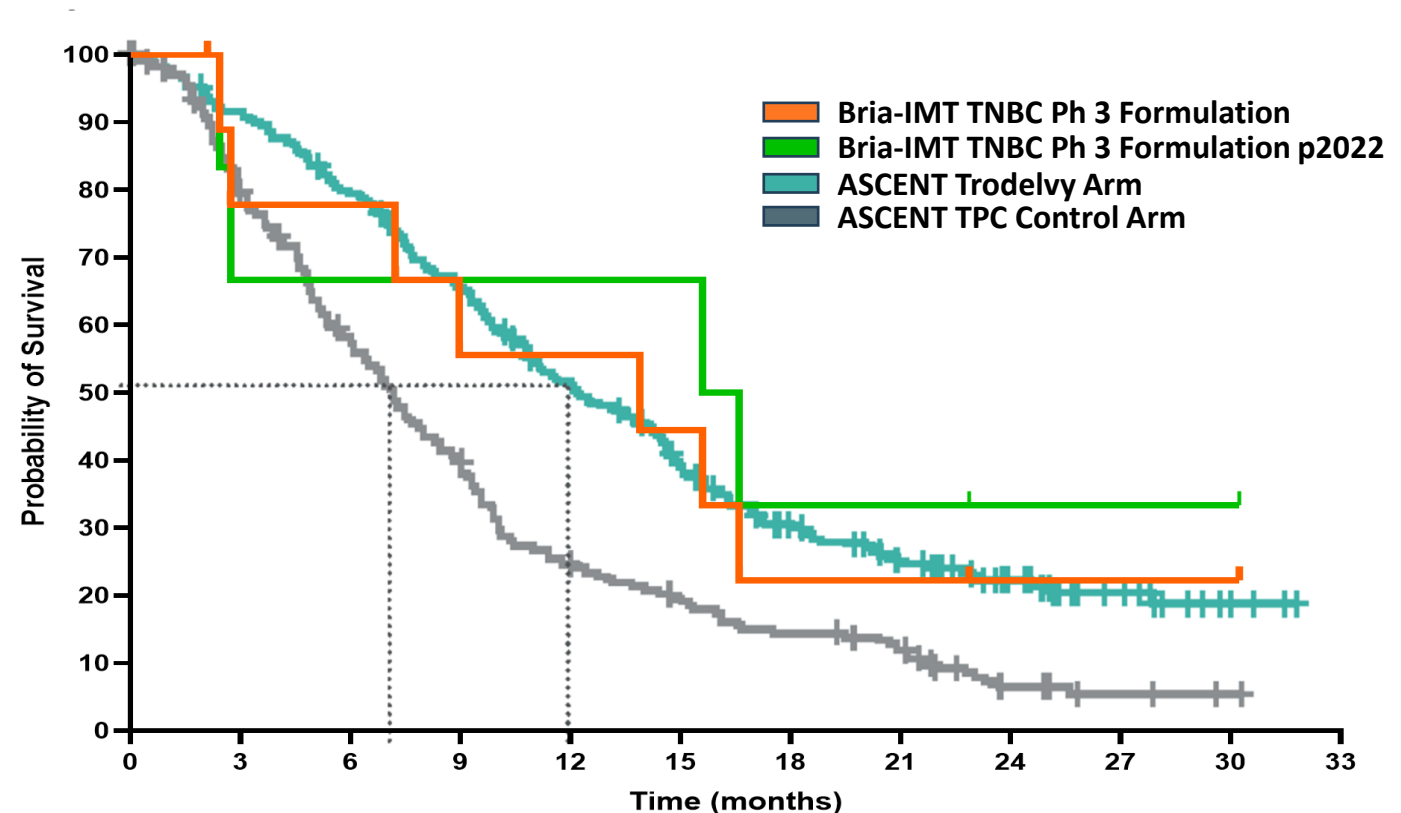
²Rugo, H. S., et al. The Lancet, 402(10411), 1423–1433.

ClinicalTrials.gov ID [NCT03328026](https://clinicaltrials.gov/ct2/show/study/NCT03328026)

Bria-IMT + CPI Phase 2 Overall Survival vs Trodelvy

- 37 patients treated with Phase 3 formulation
 - Median 6 prior lines of therapy
 - 10 patients with triple-negative breast cancer (TNBC)
 - Overall survival (OS) 13.9 months for TNBC
 - OS median 16.1 months for TNBC patients treated since 2022
- **Compares favorably to Trodelvy pivotal registration study in TNBC**
 - Median 4 prior regimens
 - OS 11.8 months for Trodelvy
 - OS 6.9 months for single agent chemotherapy

Overall survival of Bria-IMT vs ASCENT¹



¹Bardia, A., et al Journal of Clinical Oncology, 42(15), 1738–1744

Comparison to Other Similar Patients

- ~4 months median progression-free survival (PFS) twice that seen in comparable patients treated with best available therapy, including antibody-drug conjugate (ADC) resistant patients^{1,2,3}
- Clinical benefit seen in **53%** of evaluable patients in **all subtypes** of breast cancer who had failed an ADC
 - Compared with **7-10%** in comparable patients treated with best available therapy (the comparator for Phase 3)

Study	Progression-Free Survival (months)	Objective Response Rate (%)	Clinical Benefit Rate (%)
BriaCell's ADC Resistant Phase 2 patients who received pivotal Phase 3 study formulation (Bria-IMT regimen)	4.1	12**	53**
Bardia, A. et. al. ¹	1.7	4	8
Tripathy D. et. al. ²	1.9	3	10
O'Shaughnessy J. et. al. non-TNBC ³	2.3	4	7
O'Shaughnessy J. et. al. TNBC ³	1.6	5	10

*Data is for evaluable patients, n=42 with 12 not evaluable.

** Data is for evaluable patients, n = 17 with 6 not evaluable.

References: Data is shown for the intent to treat population for the control group treated with treatment of physician's choice, which is the comparator in the BriaCell phase 3 study

1. Bardia A, et al. Final Results From the Randomized Phase III ASCENT Clinical Trial in Metastatic Triple-Negative Breast Cancer and Association of Outcomes by Human Epidermal Growth Factor Receptor 2 and Trophoblast Cell Surface Antigen 2 Expression. J Clin Oncol. 2024 May 20;42(15):1738-1744.

2. Tripathy D, Tolaney SM, Tagliaferri M. Etrirnotecan Pegol Treatment for Patients With Metastatic Breast Cancer and Brain Metastases-Reply. JAMA Oncol. 2022 Nov 1;8(11):1700-1701. jamaoncol.2022.4346. PMID: 36136348. This paper describes patients with brain metastases.

3. O'Shaughnessy J, et al. Analysis of patients without and with an initial triple-negative breast cancer diagnosis in the 3 randomized ASCENT study of sacituzumab govitecan in metastatic triple-negative breast cancer. Breast Cancer Res Treat. 2022 Sep;195(2):127-139.

Robust Market Potential



Drug	Trial	Prior Lines	ORR (%)	PFS (months)	WW 2024 Sales* (Bil \$)
Bria-IMT™	Ph 2	5.5	10	4.1	
KISQALI® (ribociclib)	Ph 1/2a (MBC previously Rx trastuzumab, pertuzumab, and trastuzumab emtansine) ¹	5	0	1.3	\$ 3.0
KEYTRUDA® ** (pembrolizumab)	Ph 2 (MBC HER2 negative) ²	1	0	1.9	\$ 29.5
LYNPARZA® (Olaparib) + cediranib	Ph 1 (MBC triple negative) ³	3	0	3.7	\$ 3.7
IBRANCE® (palbociclib)	Ph 2a (MBC retinoblastoma+) ⁴	2	0	3.7	\$ 4.4
	Ph 2 (MBC HR+/HER2-) ⁵	2		6	
PERJETA® (pertuzumab)	Ph 1 (MBC) ⁶	2	0		\$ 4.0
PERJETA® (pertuzumab) + docetaxel	Ph 1b (MBC) ⁷	2	0		
PERJETA® (pertuzumab) + Herceptin	Ph 2 (MBC HER2+ progressed on trastuzamab) ⁸	1-3	0	5	
TRODELVY® (sacituzumab govitecan-hziy)	Ph 1/2 MBC (TNBC; HR+ HER2-) ⁹	3	0	5.5	\$ 1.3
Verzenio® (abemaciclib)	Ph 1/2 (HR+/HER2- MBC progressed on endocrine Rx and prior chemo) ¹⁰	4	0	6	\$ 5.3
Verzenio® (abemaciclib)	Ph 1/2 (MBC HR+ Intracranial mets) ¹¹	1-12	5.2/0	4.4/2.7	
KADCYLA® (ado-trastuzumab emtansine)	Ph 1 (MBC HER2+ failed trastuzamab) ¹²	4	21	~5.8	\$ 2.2
	Ph 1 (MBC HER2+ failed HER2 directed Rx) ¹³	5	26	4.6	
HERCEPTIN HYLECTA® (trastuzumab & hyaluronidase-oysk)	Ph 3 (HER2+ MBC) ¹⁴		26	4.6	\$ 1.5
ENHERTU® (HER2low) (fam-trastuzumab deruxtecan-nxki)	Ph 2 ¹⁵	1	52		\$ 3.8
ENHERTU® (HER2high) (fam-trastuzumab deruxtecan-nxki)	Ph 2 ¹⁵	5	60		

\$2-5 billion Bria-IMT opportunity#

*Worldwide sales figure is based on SEC filings

**Approved for multiple cancer indications

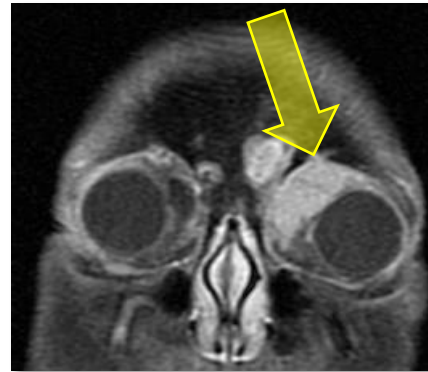
Independent market analysis

References:

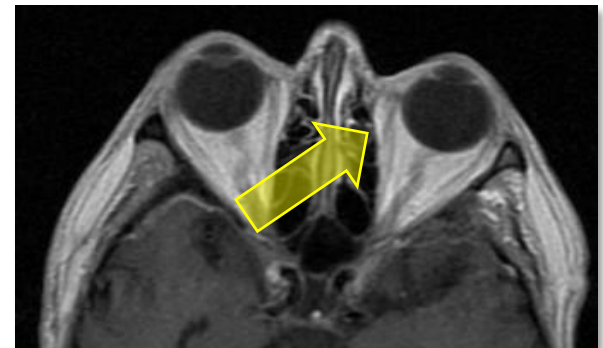
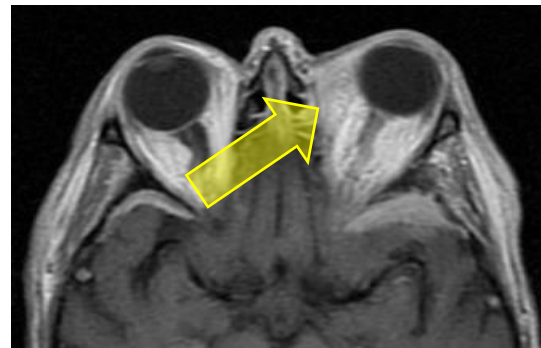
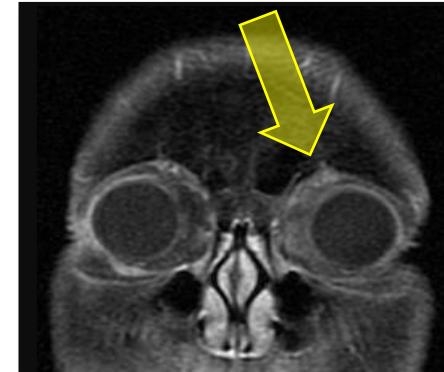
- [1. https://pubmed.ncbi.nlm.nih.gov/31235441/](https://pubmed.ncbi.nlm.nih.gov/31235441/)
 - [2. https://ascopubs.org/doi/10.1200/JCO.2023.41.16_suppl.1095](https://ascopubs.org/doi/10.1200/JCO.2023.41.16_suppl.1095)
 - [3. https://pubmed.ncbi.nlm.nih.gov/23810467/](https://pubmed.ncbi.nlm.nih.gov/23810467/)
 - [4. https://pubmed.ncbi.nlm.nih.gov/25501126/](https://pubmed.ncbi.nlm.nih.gov/25501126/)
 - [5. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8676999/](https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8676999/)
 - [6. https://pubmed.ncbi.nlm.nih.gov/15699478/](https://pubmed.ncbi.nlm.nih.gov/15699478/)
 - [7. https://pubmed.ncbi.nlm.nih.gov/18000498/](https://pubmed.ncbi.nlm.nih.gov/18000498/)
 - [8. https://pubmed.ncbi.nlm.nih.gov/20124182/](https://pubmed.ncbi.nlm.nih.gov/20124182/)
 - [9. https://www.nejm.org/doi/10.1056/NEJMoa1814213?url_ver=Z39.88-2003&rfr_id=ori:rid:crossref.org&rfr_dat=cr_pub%20%20pubmed](https://www.nejm.org/doi/10.1056/NEJMoa1814213?url_ver=Z39.88-2003&rfr_id=ori:rid:crossref.org&rfr_dat=cr_pub%20%20pubmed)
 - [10. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5581697/](https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5581697/)
 - [11. https://aacrjournals.org/clincancerres/article/26/20/5310/82934/](https://aacrjournals.org/clincancerres/article/26/20/5310/82934/)
 - [12. https://pubmed.ncbi.nlm.nih.gov/20421541/](https://pubmed.ncbi.nlm.nih.gov/20421541/)
 - [13. https://pubmed.ncbi.nlm.nih.gov/21172893/](https://pubmed.ncbi.nlm.nih.gov/21172893/)
 - [14. https://www.thelancet.com/journals/lanonc/article/PIIS1470-2045\(12\)70329-7/abstract](https://www.thelancet.com/journals/lanonc/article/PIIS1470-2045(12)70329-7/abstract)
 - [15. https://pubmed.ncbi.nlm.nih.gov/36780610/;](https://pubmed.ncbi.nlm.nih.gov/36780610/)
- <https://medicalinformation.astrazeneca-us.com/home/prescribing-information/enhertu.html>

- Patient on 13 prior failed regimens
- Baseline breast cancer metastases
 - Behind the left eye (orbit)
 - Outside lining of the brain (dura mater)
 - Adrenal gland
- 6 months of treatment
 - Orbital tumor completely resolved
 - The patient judged an overall partial responder

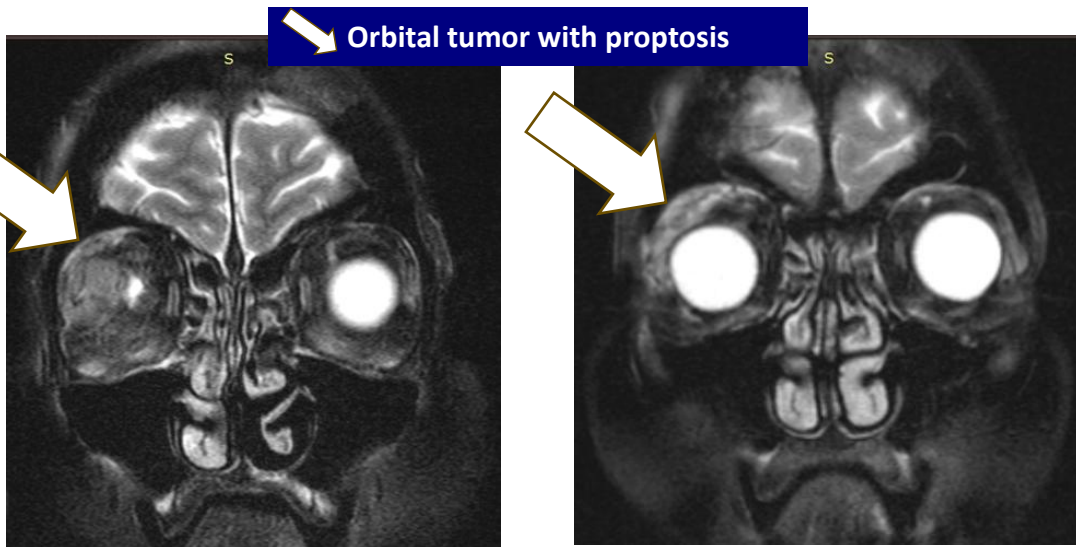
Baseline Scans



On Treatment Scans



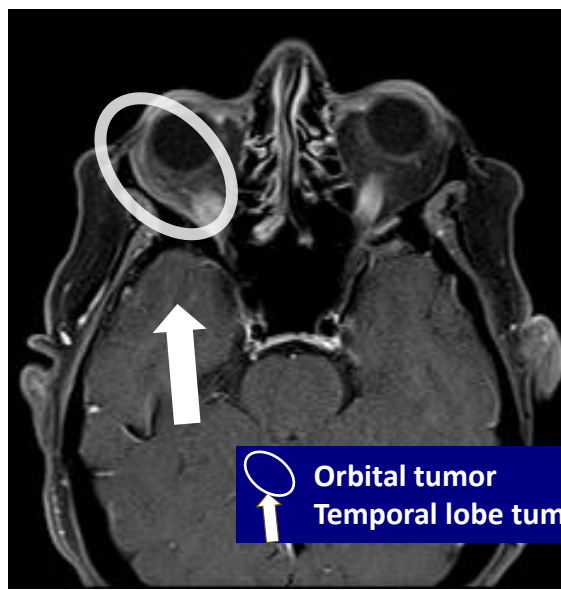
Bria-IMT + CPI Remarkable CNS Responder Case #2



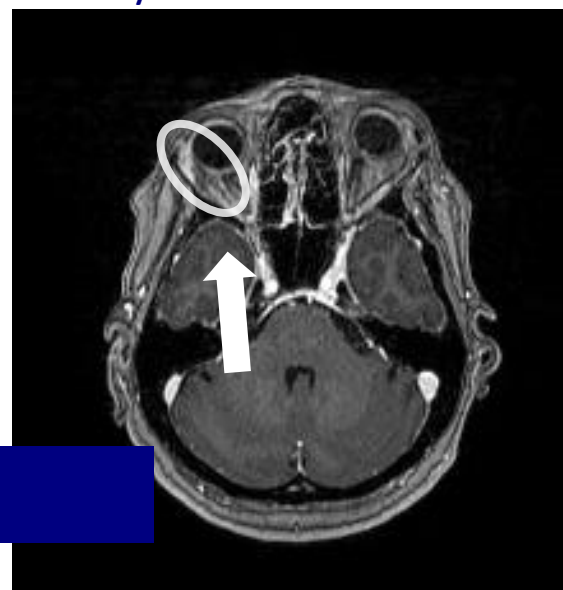
- Patient on 8 prior failed regimens including ADC Enhertu®
- Baseline tumor status
 - Extensive proptosis (eye bulging) & brain (temporal lobe) metastasis
- Marked tumor reduction at 11 months
 - Near complete proptosis resolution at 3 months with orbit (eye socket) tumor: 42 mm → 28 mm
 - Brain temporal lobe tumor: 22 mm → 0
 - Overall, 56% reduction (PR)
 - Improvement in eye pain and reduction in tumor markers
- On study for 24 months



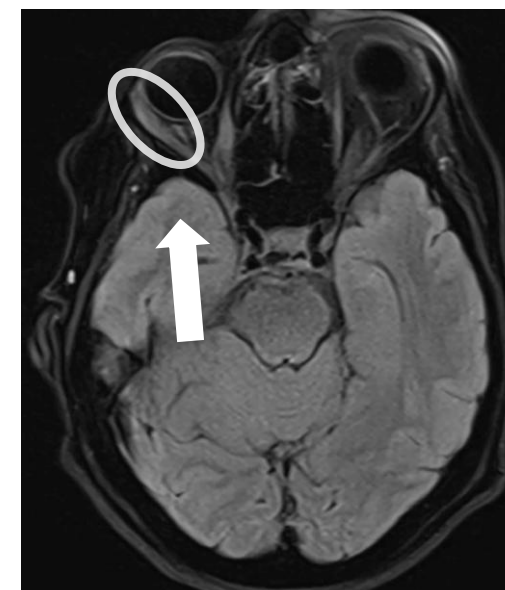
Pre-treatment



6 Months



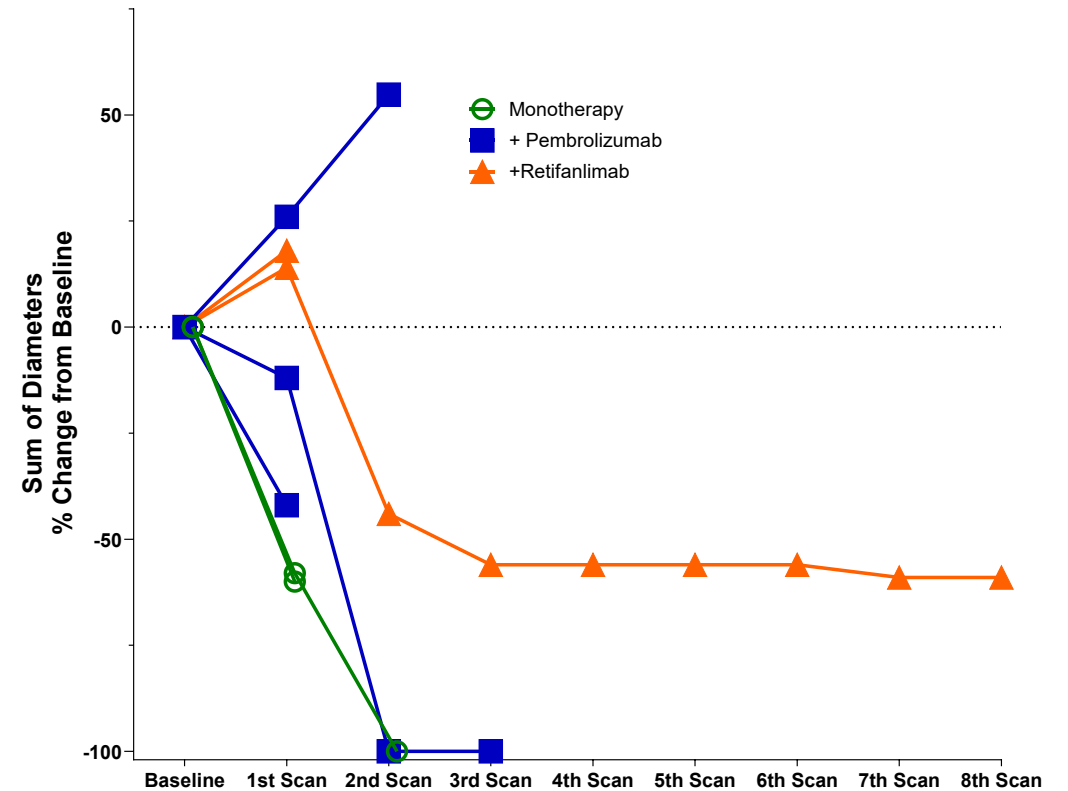
12 Months



20 Months

- **71% (5/7) intracranial objective response rate (iORR)**
 - Positive results in both Bria-IMT monotherapy and CPI combination
- **iORR in comparable patients typically <20%**^{1,2}
- Response across all subtypes of breast cancer
- Heavily pre-treated population includes 1 ADC resistant patient
- Planned CNS disease subgroup analysis in pivotal phase 3
- **Data highlight strong Bria-IMT potential in CNS metastases**

Intracranial Tumor Responses*



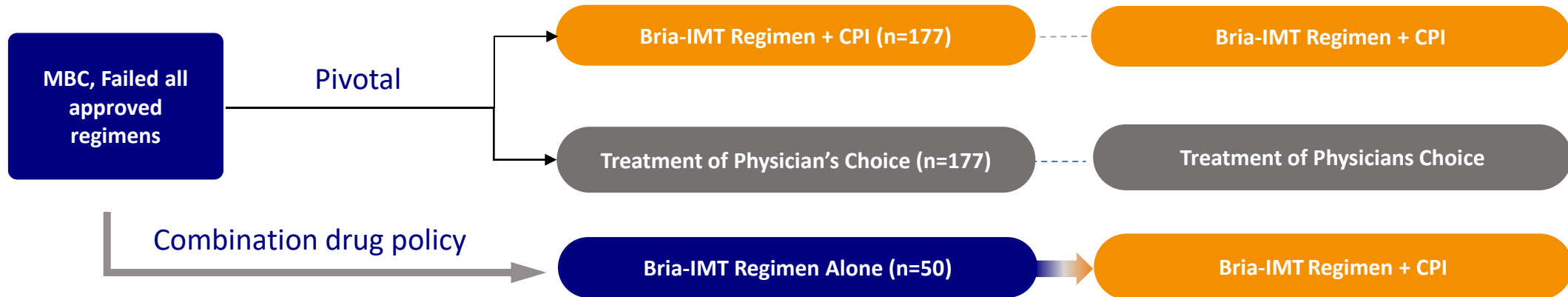
¹Niwinska A, Pogoda K, Jagiello-Gruszfeld A, Duchnowska R. Intracranial Response Rate in Patients with Breast Cancer Brain Metastases after Systemic Therapy. *Cancers (Basel)*. 2022 Feb 15;14(4):965. doi: 10.3390/cancers14040965. PMID: 35205723; PMCID: PMC8869862.

²Tripathy D, Tolaney SM, Seidman AD, Anders CK, Ibrahim N, Rugo HS, Twelves C, Diéras V, Müller V, Du Y, Currie SL, Hoch U, Tagliaferri M, Hannah AL, Cortés J; ATTAIN Investigators. Treatment With Etirinotecan Pegol for Patients With Metastatic Breast Cancer and Brain Metastases: Final Results From the Phase III ATTAIN Randomized Clinical Trial. *JAMA Oncol*. 2022 Jul 1;8(7):1047-1052. doi: 10.1001/jamaoncol.2022.0514. PMID: 35552364; PMCID: PMC9100460.

*Data in evaluable patients

Bria-IMT + CPI Pivotal Phase 3 Study

- Ongoing pivotal Phase 3 study in Advanced Breast Cancer (Bria-ABC)
- **Primary endpoint of overall survival**
 - Interim efficacy at 144 events
 - 97.5% powered to detect a 40% reduction in mortality
- **Positive interim results could support FULL approval of Bria-IMT + CPI**



Analyze at 144 events. If hazard ratio (HR) is ≤ 0.6 , submit BLA. If > 0.6 , continue to completion with HR target of 0.7

Eleven clinical trials that will shape medicine in 2026

Nature Medicine asks leading researchers to name their top clinical trial for 2026, from long-awaited vaccines for infectious diseases to new treatments for advanced cancers and long COVID.

Table 1 | Clinical trials to watch in 2026

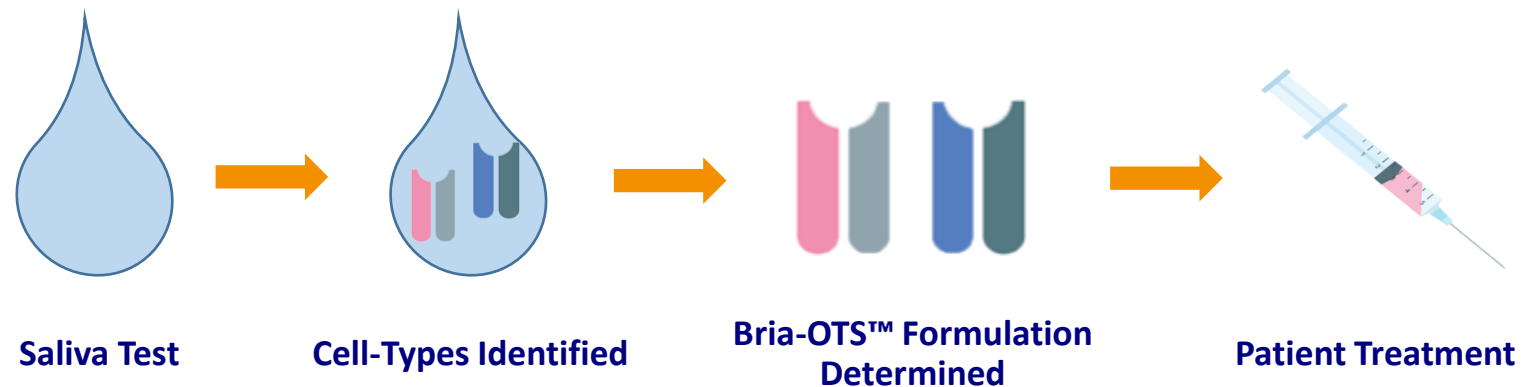
Treatment	Organization	Description	Phase	Indication
Bria-IMT	BriaCell Therapeutics Corporation	Cell-based immunotherapy plus an immune checkpoint inhibitor	Phase 3	Metastatic breast cancer

Nature Medicine | Volume 31 | December 2025 | 3943–3947 | **3943**

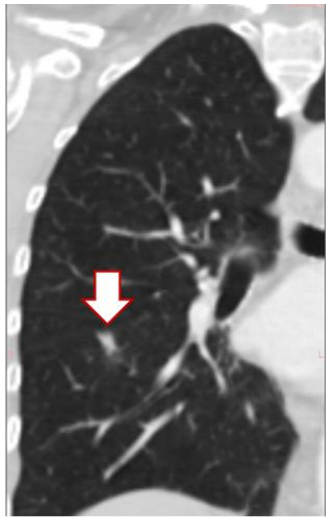
- Currently over 70 sites open in the US for the Phase 3 study including Mayo Clinic, U of Arizona, Cedar-Sinai, UCLA, UC San Diego, Yale, U of Miami, Northwestern, Dartmouth, Cleveland Clinic,
- Collaborations with MD Anderson, Memorial Sloan Kettering, Yale, Mayo Clinic, University of Pennsylvania, National Cancer Institute, and other top-tier institutions

Listed in Nature Medicine as one of the Clinical Trials that will SHAPE MEDICINE IN 2026

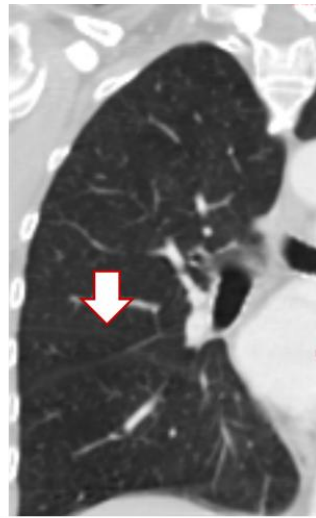
- BriaCell immunotherapy is most effective in human leukocyte antigen (HLA) type matched patients
- HLA typing identifies patient specific T cell presenting antigens necessary for immune response
- Bria-OTS expresses 15 HLA types in 4 cell lines, providing matched treatment to >99% of patients
- Simple saliva test delivers personalized Bria-OTS immunotherapy
- **Breast cancer Phase 1/2a study ongoing**



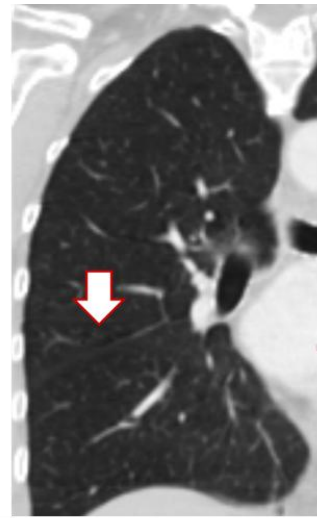
Bria-OTS Metastatic Breast Cancer Phase 1/2a



Pre-Treatment



2 months



4 months



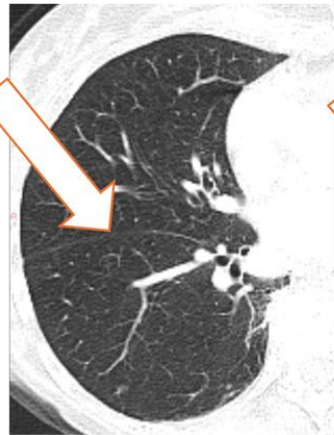
6 months



11 months



Pre-Treatment



2 months



4 months



6 months



11 months

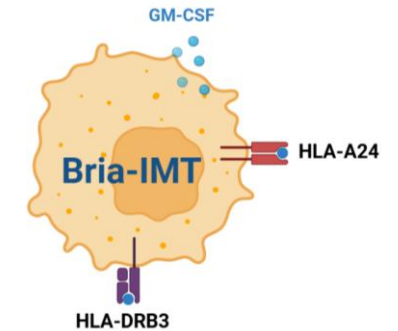
- Initial monotherapy dose escalation then dose expansion in combination with CPI
 - Up to 18 patients including 9 patient dose expansion
- Results of first patient treated:
 - Complete resolution of lung lesion at two months sustained for 11 months
 - Stable disease elsewhere
 - Multiple prior therapy failures

From Bria-IMT to Bria-OTS to Bria-OTS+: Engineering Stronger, Broader Immune Activation



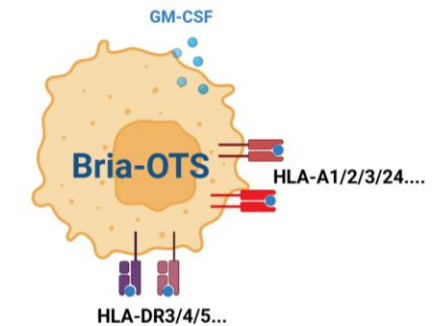
Bria-IMT™ (1st Generation)

- ❑ Breast cancer cell line engineered to express GM-CSF
- ❑ Demonstrated clinical activity and survival benefit in Phase 1/2a
- ❑ Validates whole-cell immunotherapy and provides the foundation for OTS



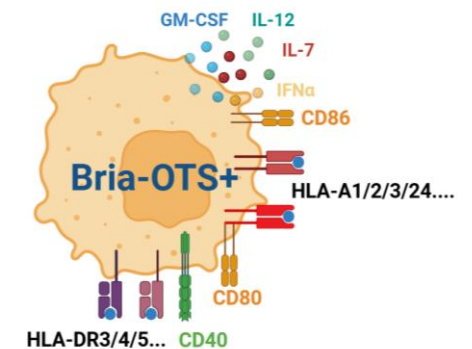
Bria-OTS™ (2nd Generation: HLA-Matched Platform)

- ❑ Evolution of Bria-IMT™ engineered to express 15 HLA alleles
- ❑ Four semi-allogeneic, off-the-shelf cell lines providing >99% population coverage
- ❑ Enables scalable, rapid, and personalized matching without autologous manufacturing



Bria-OTS+™ (3rd Generation: Engineered for Potency)

- ❑ Adds co-stimulatory molecules (CD80, CD86, CD40) and immune-modulatory cytokines (GM-CSF, IL-12, IL-7, IFN- α)
- ❑ Broad activation of CD4⁺, CD8⁺, NK, NKT, and dendritic cells and “serial killer” activity
- ❑ Supported by a National Cancer Institute Small Business Innovative Research award
- ❑ OTS+ variants now in development for multiple indications (prostate, lung, melanoma)
- ❑ *Bria-BRES+ and Bria-PROS+ scheduled to enter the clinic 2026*
- ❑ *IND approved and open for Bria-BRES+*



Expected Development Timeline and Catalysts

Bria-IMT + CPI

- Phase 2 safety & efficacy data
- Presentations at scientific conferences,
- Quarterly DSMB updates on pivotal Phase 3 study



Bria-OTS/OTS+

- OTS Phase 1/2a dose escalation in breast cancer
- Ongoing data readouts from breast cancer study
 - Initiation of Bria-BRES+
 - Bria-PROS+ IND Submission
 - Initiation of Clinical Study in prostate cancer
- Identify lead indications and initiate Bria-OTS+ pivotal registration studies
 - First Indication
 - Second Indication
 - Third Indication
- Continuous study readouts with additional indications (lung, melanoma and others)



William V. Williams, MD, FACP

President & CEO, Director

- Incyte, GlaxoSmithKline
- University of Pennsylvania



Giuseppe Del Priore, MD, MPH

Chief Medical Officer

- Cancer Treatment Centers of America
- NYU School of Medicine, New York Presbyterian



Gadi Levin, CA, MBA

CFO & Corporate Secretary

- Arthur Andersen
- University of Cape Town, Bar Ilan University



Miguel A. Lopez-Lago, PhD

Chief Scientific Officer

- Memorial Sloan-Kettering Cancer Center
- Stony Brook University, New York



Clinical Strategy Team involved in 20 previous drug or device approvals



Jamieson Bondarenko, CFA, CMT
Chairman of the Board

- Eight Capital, Dundee Securities, Wellington West Capital Markets and HSBC Securities



William V. Williams, MD, FACP
President & CEO, Director

- Incyte, GlaxoSmithKline



Vaughn Embro-Pantalony, MBA, FCPA, FCMA, CDir, ACC
Director

- Teva Novopharm Limited, Bayer Healthcare, Zeneca Pharma Inc.



Jane Gross, PhD
Director

- aTyr Pharma Inc., ZymoGenetics Inc. (acq. by Bristol Myers Squibb)



Martin Schmieg, CPA
Director

- Clear Intradermal Technologies, Inc., Sirna Therapeutics, Inc., Advanced Bionics Corporation



Rebecca A. Taub, MD
Director

- Madrigal Pharmaceuticals, Hoffmann-La Roche Company, Bristol-Myers Squibb;

Capitalization Structure*

	Nasdaq: BCTX, BCTXL TSX: BCT
Share Price:	US\$3.09
Shares Outstanding:	8.7M
Market Cap:	US\$27M
Options (US\$33.06 WAEP):	337k
Warrants (US\$16.22 WAEP):	7.2M

(*) as of June 8, 2026

- **Bria-IMT™ + CPI:** Pivotal phase 3 study underway in metastatic breast cancer.
 - Up to 2-fold increase in survival compared to comparable patients in the literature.
 - Phase 3 interim analysis expected in 2026.
- **Bria-OTS/OTS+™:** Ongoing phase 1/2a study for Bria-BRES™. Bucket trial with other cancer indications to be added.
 - Bria-PROS+ IND submission and initiation of clinical study in prostate cancer expected in 1H 2026.
- **Proven Management Team:** Clinical strategy team involved in 20 previous drug or device approvals.



Developing Novel Therapeutics to Destroy Cancer

Thank-you!