



Atossa Therapeutics, Inc.

Q4 inline. Positive breast cancer clinical trials (5 Phase 2) milestones and key data over the next year should be strong catalysts for stock. Raising P/T to \$7.25.

Q4 inline: Atossa recently (on March 25) reported its Q4 2024 (ending December) results. Net loss was \$6.3 million or EPS of \$(0.05), compared with our and consensus estimates of \$(0.06). There was no Q4 guidance. Atossa is a clinical stage drug development company so it generates no revenue.

Operating expenses: Operating expenses were \$7.2 million, up from \$6.4 million in Q3.

No guidance: Management did not provide forward guidance, but we believe ~\$7 million to be a reasonable near term quarterly cash burn rate.

Adjusting estimates: We are maintaining our 2025 EPS estimate of \$(0.25).

Endoxifen in development: Atossa has one main therapeutic drug candidate, Endoxifen for breast cancer (for 2 settings). The company recently (in March) announced that it will pursue Metastatic Breast Cancer Indication for (Z)-Endoxifen.

Breast cancer: Atossa's breast cancer drug under development is its proprietary form of Endoxifen which is being developed primarily in two settings: 1) to reduce tumor cell activity in breast cancer patients in the "window of opportunity" between diagnosis of breast cancer and surgery; and 2) for women with high mammographic breast density ("MBD") to reduce the density and/or to act as an adjunct to mammography.

5 clinical trial in progress: In December 2021, Atossa began to enroll patients in its clinical study (Karisma) of Endoxifen in Sweden. In February 2023, the first patient has been dosed in the Phase 2 EVANGELINE (Endoxifen Versus exemestane + goserelin) study. In March 2023, Atossa announced that endoxifen will be evaluated in a new study arm of the ongoing I-SPY 2 clinical trial. In conjunction with Quantum Leap Healthcare Collaborative, Atossa announced in August the first patient dosed in (Z)-endoxifen in combination with abemaciclib (VERZENIO) as part of I-SPY 2.

Positive Topline data from KARISMA-Endoxifen Phase 2 study: In November 2024, Atossa released positive topline data from the KARISMA-Endoxifen Phase 2 study of (Z)-endoxifen in premenopausal women with mammographic breast density (MBD). The study, which was conducted through the Karolinska Institute in Stockholm, Sweden, demonstrated that low doses of (Z)-endoxifen significantly reduced MBD and was well tolerated.

Positive initial data from Phase 2 I-SPY 2 Trial: In November 2024, Atossa released a preliminary analysis from its Phase 2 trial of (Z)-endoxifen in ER+/HER2- breast cancer, showing that (Z)-endoxifen meet the primary endpoint with 95 percent (19/20) of patients. The data showed a rapid reduction in key breast cancer biomarkers, including a 69 percent reduction in Ki-67 and a 30.4% reduction in functional tumor volume after three weeks.

Clinical data can be catalysts: Atossa anticipates finishing or making significant milestones in its various clinical trials over the next year. We believe achieving key clinical milestones and data will likely be strong catalysts for the stock.

Positive high risks versus high rewards: Overall, concerns outweighed by growth prospects and valuation. Atossa's drugs still have long development roads left and the high risks of clinical trials failures, but we believe the ~billion dollars market potential presents high rewards for the risks.

Current valuation attractive: We are maintaining our BUY rating, but raising our 12-month price target to \$7.25 from \$7.00 based on a NPV analysis. This represents significant upside from the current share price. We believe this valuation appropriately balances out the company's high risks with the company's high growth prospects and large upside opportunities.

Company Description

Based in Seattle, WA, Atossa Therapeutics is a clinical-stage biopharmaceutical company focused on COVID-19 and breast cancer drugs development.

United States
Healthcare

April 20, 2025

Edward Woo, CFA
(561) 327-9435
ewoo@ascendant.com

COMPANY UPDATE

Rating: **BUY**

Ticker: ATOS

Price: \$0.62

Target: \$7.25
(from \$7.00)

Stock Data

Exchange:	NasdaqCM
52-week Range:	\$0.55 – 1.81
Shares Outstanding (million):	129
Market cap (\$million):	\$80
EV (\$million):	\$9
Debt (\$million):	\$0
Cash (\$million):	\$71
Avg. Daily Trading Vol. (\$million):	\$1
Float (million shares):	129
Short Interest (million shares):	8
Dividend, annual (yield):	\$0 (NA%)

Revenues (US\$ million)

	2025E (Cur.)	2025E (Old)	2026E (Cur.)	2026E (Old)
Q1 Mar	OE		OE	
Q2 Jun	OE		OE	
Q3 Sep	OE		OE	
Q4 Dec	OE		OE	
Total	OE		OE	
EV/Revs	N/A		N/A	

Earnings per Share (pro forma)

	2025E (Cur.)	2025E (Old)	2026E (Cur.)	2026E (Old)
Q1 Mar	(0.06)E		(0.06)E	
Q2 Jun	(0.06)E		(0.06)E	
Q3 Sep	(0.06)E		(0.06)E	
Q4 Dec	(0.06)E		(0.06)E	
Total	(0.25)E		(0.25)E	
P/E	N/A		N/A	

Important Disclosures

Ascendant Capital Markets LLC seeks to do business with companies covered by its research team. Consequently, investors should be aware that the firm may have a conflict of interest that could affect the objectivity of this report. Investors should consider this report as only a single factor in making an investment decision.

For analyst certification and other important disclosures, refer to the Disclosure Section, located at the end of this report, beginning on page 14.

Exhibit 1: Atossa Therapeutics Investment Highlights

Investor Highlights



- Lead compound, (Z)-endoxifen being investigated in multiple ongoing and completed Phase 2 trials for breast cancer / breast conditions
- Deep intellectual property portfolio
- Large, unaddressed / underserved market opportunities in breast cancer prevention and treatment settings
- \$94M cash at 9/30/23, approximately three-year operating runway
- Experienced management team with extensive life sciences background
- World class R&D collaborators

Source: Company reports

Exhibit 2: Accomplishments and Upcoming Milestones (as of May 2024)

Near Term Catalysts / Capital Table



Near Term Catalysts	Key Metrics
<ul style="list-style-type: none"> • Mammographic Breast Density <ul style="list-style-type: none"> - Density reduction data available 2H '24 • I-Spy <ul style="list-style-type: none"> - 10mg neoadjuvant data available 2H '24 • EVANGELINE <ul style="list-style-type: none"> - 80mg PK run-in cohort completion 2Q '24 - Treatment arm initiated - 2H '24 • DCIS <ul style="list-style-type: none"> - Enrollment updates throughout '24 • Combinations <ul style="list-style-type: none"> - CDK 4/6 - enrollment updates throughout '24 - ADC - clinical start TBD 	<ul style="list-style-type: none"> • Cash (as of 12/31/23) <ul style="list-style-type: none"> - \$88.5M – represents approx. three years working capital - Zero debt • Nasdaq: ATOS (as of 4/26/24) <ul style="list-style-type: none"> - Market Cap - \$192M - Share Price - \$1.53 - 52 Week Range - \$0.59 - \$2.31 • Outstanding Warrants / Options (as of 12/31/23) <ul style="list-style-type: none"> - 11.0M warrants exercisable at \$1.00 or \$1.05/share - 10.5M warrants exercisable at \$2.88/share - 13.7M options exercisable at average \$2.04/share

Source: Company reports

Exhibit 3: Atossa Market Opportunities

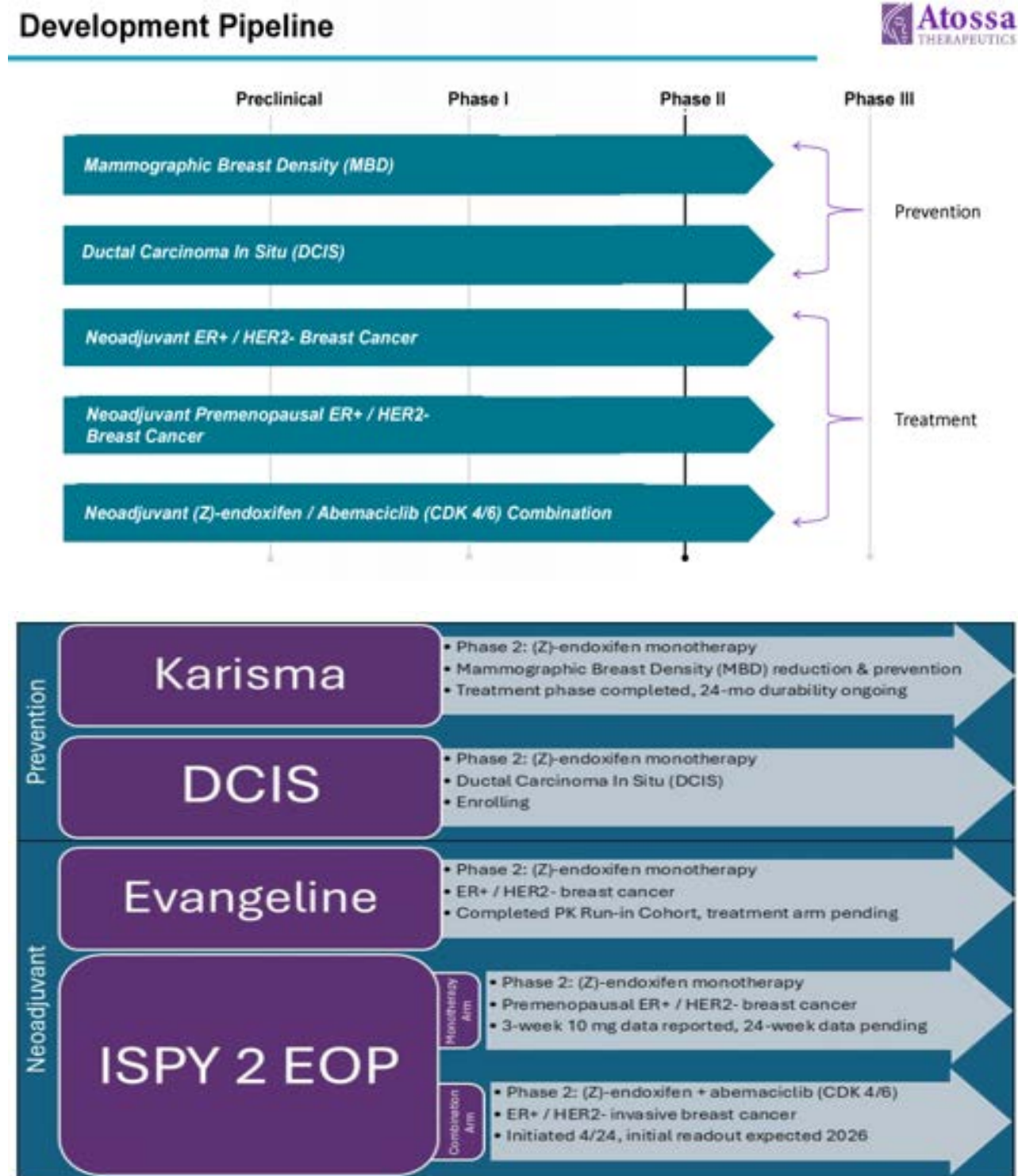
Large Market Opportunities



PROGRAM		OPPORTUNITY
Mammographic Breast Density	➔	27 Million Women in U.S. with dense breasts ¹
Ductal Carcinoma In Situ (DCIS)	➔	60,000 DCIS diagnoses annually in U.S. ²
Neoadjuvant	➔	250,000 ER+ Breast Cancers/Yr. in U.S. ²

Source: Company reports.

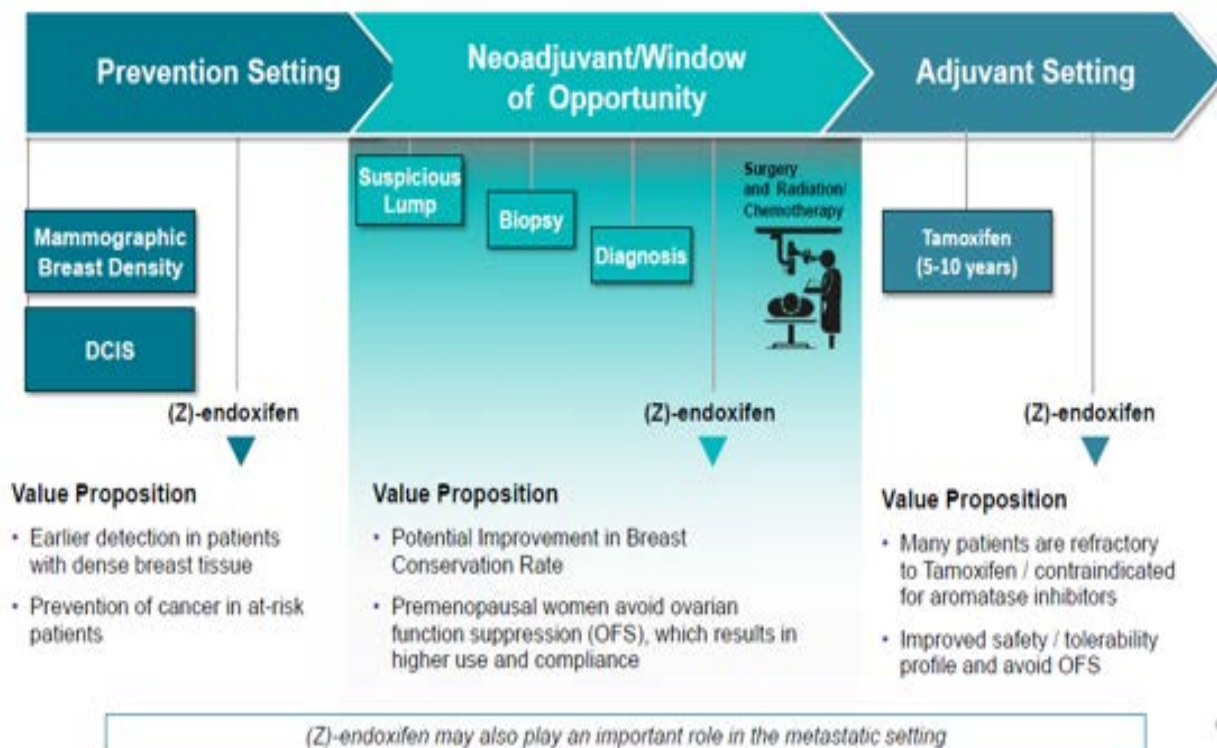
Exhibit 4: Atossa Drug Development Pipeline (as of March 2025)



Source: Company reports

Exhibit 5: ENDOXIFEN

Clinical Positioning In Breast Cancer



The Breast Cancer Problem



Source: Company reports

Exhibit 6: Endoxifen Clinical Trials

Karisma-Endoxifen Study



Issue

- 50% of the women in the world have dense breast tissue
 - Elevated density is a significant independent risk factor for developing breast cancer
 - Elevated density make mammograms less effective
 - Federal legislation goes into effect Sept. '24 requiring notification of density

Study

- Phase 2, randomized, double-blind, placebo-controlled, study of (Z)-endoxifen in premenopausal women with Measurable Breast Density (MBD)
 - (Z)-endoxifen 1 or 2 mg/day (or PBO) for 6 months
 - Endpoints – change from baseline in MBD at 3 and 6 months and durability of change at 24 months
 - Fully enrolled (n=240) Nov. '23
 - Six-month density reduction data 2H '24

KARISMA-Endoxifen Trial



- Significant decrease in mammographic breast density in the 1 mg and 2 mg (Z)-endoxifen arms
 - Decrease is similar to data with 20mg tamoxifen
- The discontinuation rates and adverse event profiles were similar between 1mg (Z)-endoxifen and placebo
 - Women in the 2mg (Z)-endoxifen arm reported significantly more hot flashes, night sweats, and vaginal discharge than the placebo group.
- (Z)-endoxifen adverse event profile appeared to be more favorable than KARISMA-tamoxifen data.
- No issues related to skin rashes, itching (other than genital), dry mouth, fatigue, depression, or sexual interest were reported with (Z)-endoxifen.

No. of participants, percent density change, (Z)-endoxifen concentrations, no. of early terminators and change in side effects by (Z)-endoxifen and tamoxifen doses

Variables	(Z)-endoxifen dose			Tamoxifen dose	
	0 mg	1 mg	2 mg	5 mg	20 mg
No. of premenopausal women	80	80	80	72*	79*
Percent density change (from a regression model)	0.27	-17.3*	-23.5*	-19.6*	-18.5*
Mean (Z)-endoxifen concentration at end of study ng/ml	-	4.8	9.7	2.5*	12.6*
No. of early terminations (%) because of an adverse event related to the IMP*	4 (5.0)	5 (6.3)	11 (13.8)	5 (6.9)	7 (8.9)
Mean change in Likert score from baseline to end of treatment					
Hot Flashes	0.10	0.26	0.45**	0.23*	0.67**
Night Sweats	0.10	0.26	0.58**	0.60*	0.70**
Cold Sweats	0.04	0.12	0.10	0.12	0.33**
Vaginal discharge	-0.06	0.05	0.03**	-0.12	0.18
Genital itching	0.05	0.21	0.12	0.16*	0.22
Pain or cramps in legs and feet	0.14	0.20	0.17	0.22*	0.38*

* premenopausal women in the intention to treat population, * p<0.01, **p<0.05, * for normal CYP2D6 metabolites, IMP= Investigational Medicinal Product

Source: Company reports

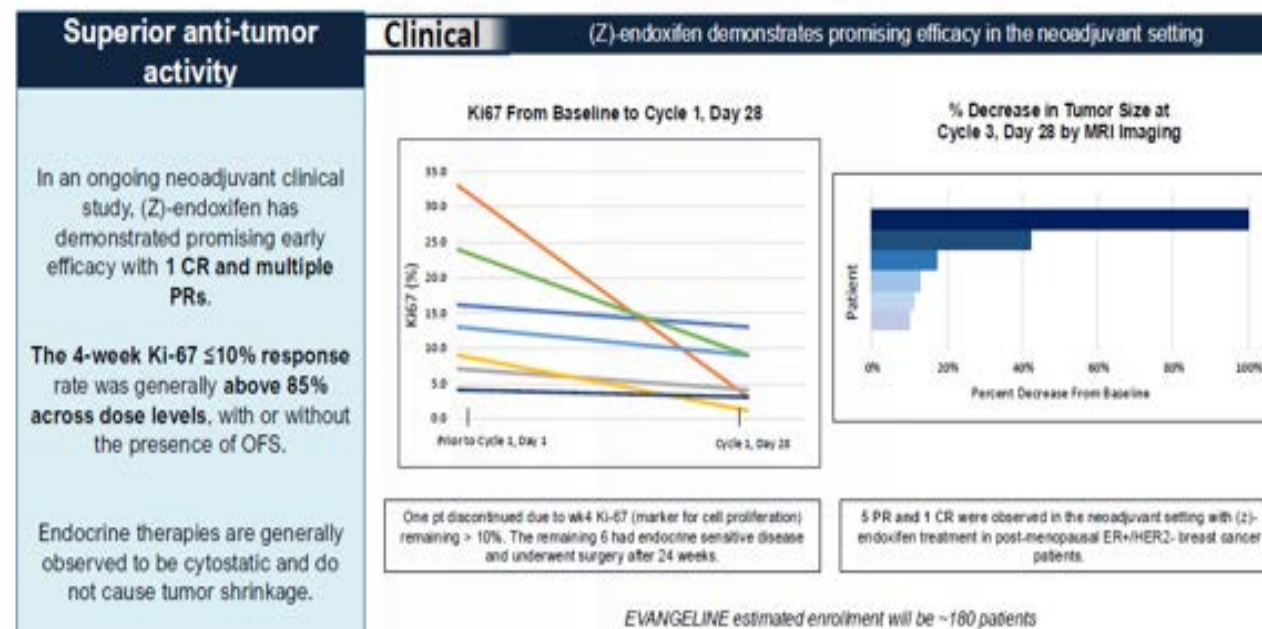
Exhibit 7: Endoxifen Clinical Trials

U.S. Phase 2 Study - EVANGELINE



- Open-label, randomized, Phase 2 study in premenopausal women with Grade 1 or 2 ER+/HER2- breast cancer – first patient was enrolled in February 2023
- Subjects are enrolled with the intent of surgical treatment in the involved breast(s) after completing neoadjuvant study treatment
- Expected to enroll approximately 175 patients at up to 25 sites across the United States
- Primary objective is to evaluate the endocrine sensitive disease rate, measured by Ki-67 compared to treatment with current standard of care
- Current SOC includes medication given to block the ovaries from making estrogen, which in premenopausal women is associated with significant morbidity and inadequate compliance, which compromises efficacy and increases the risk of mortality

EVANGELINE OVERVIEW



Source: Company reports

Exhibit 8: Endoxifen Clinical Trials

I-SPY 2 – Neoadjuvant Combination



Issue

- Women with high clinical stage but less proliferative tumors are particularly challenging to treat
 - High risk of late recurrence
 - Currently approved CDK 4/6 combination therapies have sub-optimal safety profile



Study

- Phase 2 neoadjuvant study of (Z)-endoxifen in combination with *abemaciclib* (VERZENIO) in women diagnosed with ER+ / HER2- invasive breast cancer
 - Co-sponsored by Atossa and Eli Lilly & Company
 - Part of the largest and most successful platform trial in history
 - Participants receive 40mg (Z)-endoxifen and 150mg *abemaciclib* daily for six months
 - Endpoints – Ki-67 reduction and objective response – assessed by MRI and pathology
 - Initiated April '24

U.S. Phase 2 Study – I-SPY 2



- Ground-breaking platform trial for neoadjuvant treatment of locally advanced breast cancer
- (Z)-endoxifen is being evaluated in the Endocrine Optimization Pilot Protocol targeting patients with newly diagnosed ER+ invasive breast cancer for whom chemotherapy is expected to provide little or no benefit
- These patients have substantial risk for recurrence
- Approximately 20 patients will be treated with (Z)-endoxifen for up to 24 weeks prior to surgery
- Enrolling patients at all 41 I-Spy sites across the United States

Source: Company reports

Exhibit 9: Q4 2024 Results and Recent Highlights (as of March 25, 2025)

Atossa Therapeutics Announces Full Year 2024 Financial Results and Provides Corporate Update

March 25, 2025 12:00 PM EDT

Ended 2024 with \$71.1 million of cash and cash equivalents and no debt

Conference Call and Webcast Scheduled for Tuesday, March 25, 2025, at 8:30 a.m. Eastern Time

SEATTLE, March 25, 2025 (GLOBE NEWSWIRE) -- Atossa Therapeutics, Inc. (Nasdaq: ATOS) ("Atossa" or the "Company"), a clinical-stage biopharmaceutical company developing innovative medicines for breast cancer, today announced its financial results for the full year ended December 31, 2024 and provided an update on recent company developments.

Fourth Quarter and Early 2025 Highlights

- **Strategic Decision to Pursue Metastatic Breast Cancer Indication:** Atossa plans to advance its lead program, (Z)-endoxifen, to target metastatic breast cancer. The Company believes this approach may offer a more streamlined regulatory pathway to deliver (Z)-endoxifen to patients with urgent unmet medical needs, as the current treatment options for metastatic breast cancer often provides limited durability of response and substantial side effects. (Z)-endoxifen—a potent and well-tolerated selective estrogen receptor modulator (SERM)—has shown encouraging signs in previous clinical trials, which Atossa believes supports its potential to fill this critical gap in treatment. Atossa also intends to continue engagement with the U.S. Food and Drug Administration (FDA) to advance additional indications, including breast cancer prevention and neoadjuvant therapy, which typically require larger and longer clinical trials.
- **Tolerability and Pharmacokinetic Data from Phase 2 Evangeline Trial:** Atossa presented three posters at the San Antonio Breast Cancer Symposium (SABCS) highlighting pharmacokinetic (PK) and tolerability data from the Phase 2 EVANGELINE trial. This randomized, non-inferiority study evaluates (Z)-endoxifen in premenopausal women with primary ER+/HER2- breast cancer as a neoadjuvant treatment. Substantial tumor suppression was observed across all dosing levels, with or without ovarian function suppression (OFS or goserelin). The 4-week Ki-67 \leq 10 percent response rate was generally above 85 percent across dose levels, with or without the presence of OFS. Overall, (Z)-endoxifen was well tolerated and target tissue steady state concentration (C_{ss}) levels were achieved without significant Grade 3 or 4 toxicities. Given the gynecologic events that were previously reported in the 80 mg groups, the study is expected to continue under an amended protocol as a randomized trial that compares (Z)-endoxifen 40 mg per day with OFS to exemestane plus OFS, using the 4-week Ki-67 reduction as the primary endpoint. Additionally, the trial is expected to include a single arm cohort with a 40 mg (Z)-endoxifen monotherapy neoadjuvant treatment, using a 24-week Ki-67 endpoint.
- **Full Results from Phase 2 KARISMA-Endoxifen Study:** Additional SABCS data showcased the potential of low-dose (Z)-endoxifen to significantly reduce mammographic breast density (MBD). In the trial, a 1 mg dose reduced MBD by 17.3 percentage points ($p < 0.01$), while a 2 mg dose achieved a 23.5 percentage-point reduction ($p < 0.01$), compared to just 0.27 percentage points in the placebo arm, which we believe highlights the effectiveness of the lower dose in achieving significant reductions. Importantly, no significant differences in adverse events of special interest were observed between the 1 mg dose and the placebo. The 2 mg dose was associated with higher rates of hot flashes, night sweats and vaginal discharge compared to the placebo group. We believe these findings further support the therapy's favorable safety and efficacy profile.

"Metastatic breast cancer remains an area of critical unmet need, where improved therapies are urgently required," said Steven Quay, M.D., Ph.D., President and Chief Executive Officer of Atossa. "(Z)-endoxifen has demonstrated promising anti-estrogenic and anti-tumor effects alongside a favorable tolerability profile, which we believe positions it as a potential next-generation therapy. We believe that pursuing an initial approval in metastatic breast cancer could offer a more efficient regulatory pathway, potentially enabling us to make (Z)-endoxifen available sooner to the patients who need it most. We also believe this strategy further supports our ability to expand the broader potential of (Z)-endoxifen to address multiple stages of breast cancer, from reducing tumor growth to preventing recurrence after successful treatment."

Source: Company reports

Exhibit 10: Atossa Therapeutics, Inc. Stock Price (5-years)



Source: <https://bigcharts.marketwatch.com/>

Exhibit 11: Consensus Expectations (as of March 25, 2025)

	Revenue (mils)			EPS	
	2024E	2025E		2024E	2025E
Q1 Mar	\$0A	\$0E	Q1 Mar	\$(0.05)A	\$(0.07)E
Q2 Jun	\$0A		Q2 Jun	\$(0.05)A	
Q3 Sep	\$0A		Q3 Sep	\$(0.06)A	
Q4 Dec	\$0E		Q4 Dec	\$(0.06)E	
Total	\$0E	\$0E	Total	\$(0.22)E	\$(0.27)E

*Quarterly estimates may not add to annual estimates due to variations in contributing estimates and rounding

Source: Company report, LSEG, and Ascendant Capital Markets estimates

FINANCIAL MODEL

Atossa Therapeutics, Inc.

Income Statement (\$ mils)	Mar-23	Jun-23	Sep-23	Dec-23	2023	Mar-24	Jun-24	Sep-24	Dec-24	2024	Mar-25	Jun-25	Sep-25	Dec-25	2025	Mar-26	Jun-26	Sep-26	Dec-26	2026
Fiscal Year End: December 31	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4A	FY-A	Q1E	Q2E	Q3E	Q4E	FY-E	Q1E	Q2E	Q3E	Q4E	FY-E
Total Revenue	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
<u>Cost of Revenues</u>	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Gross Profit	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Research and development	3.5	3.7	4.5	5.7	17.3	3.7	3.6	3.4	3.4	14.1	5.0	5.0	5.0	5.0	20.0	5.0	5.0	5.0	5.0	20.0
General and administrative	3.6	4.1	3.0	3.4	14.0	3.2	3.6	3.0	3.7	13.5	3.0	3.0	3.0	3.0	12.0	3.0	3.0	3.0	3.0	12.0
<u>Restructuring and other</u>					0.0					0.0					0.0					0.0
Total operating expenses	7.1	7.8	7.5	9.0	31.4	7.0	7.1	6.4	7.2	27.6	8.0	8.0	8.0	8.0	32.0	8.0	8.0	8.0	8.0	32.0
Operating income (loss)	(7.1)	(7.8)	(7.5)	(9.0)	(31.4)	(7.0)	(7.1)	(6.4)	(7.2)	(27.6)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)
Interest income (expense)	0.9	1.0	1.3	1.2	4.3	1.1	1.1	1.0	0.9	4.1	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
<u>Other income (expense)</u>	(0.0)	(3.0)	(0.0)	0.0	(3.1)			(1.8)	(0.1)	(1.9)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Income before income taxes	(6.3)	(9.8)	(6.2)	(7.8)	(30.1)	(5.9)	(6.0)	(7.2)	(6.3)	(25.5)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)
<u>Income taxes</u>					0.0					0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net income (loss)	(6.3)	(9.8)	(6.2)	(7.8)	(30.1)	(5.9)	(6.0)	(7.2)	(6.3)	(25.5)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)
<u>Nonrecurring/noncash adjustments</u>					0.0					0.0					0.0					0.0
Net income (pro forma)	(6.3)	(9.8)	(6.2)	(7.8)	(30.1)	(5.9)	(6.0)	(7.2)	(6.3)	(25.5)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)
EBITDA																				
Shares, Basic	126.6	126.6	125.8	125.5	126.1	125.3	125.7	125.8	126.7	125.9	129.2	129.3	129.4	129.5	129.4	129.7	129.8	129.9	130.0	129.9
Shares, Diluted	126.6	126.6	125.8	125.5	126.1	125.3	125.7	125.8	126.7	125.9	129.2	129.3	129.4	129.5	129.4	129.7	129.8	129.9	130.0	129.9
EPS Basic (pro forma)	(\$0.05)	(\$0.08)	(\$0.05)	(\$0.06)	(\$0.24)	(\$0.05)	(\$0.05)	(\$0.06)	(\$0.05)	(\$0.20)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.25)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.25)
EPS Diluted (pro forma)	(\$0.05)	(\$0.08)	(\$0.05)	(\$0.06)	(\$0.24)	(\$0.05)	(\$0.05)	(\$0.06)	(\$0.05)	(\$0.20)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.25)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.06)	(\$0.25)
Margins																				
Gross margin																				
Research and development																				
General and administrative																				
Operating margin	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
Tax rate, GAAP	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%	0%
Net margin	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM	NM
YY % change																				
Total Revenue																				
Gross margin																				
Research and development	134%	8%	-13%	13%	15%	7%	-4%	-24%	-40%	-19%	33%	41%	47%	47%	42%	0%	0%	0%	0%	0%
General and administrative	11%	29%	-1%	7%	11%	-10%	-13%	-1%	11%	-4%	-7%	-16%	1%	-20%	-11%	0%	0%	0%	0%	0%
Operating income (loss)	50%	18%	-9%	11%	13%	-2%	-9%	-15%	-21%	-12%	15%	13%	25%	12%	16%	0%	0%	0%	0%	0%
Net income (loss)	31%	47%	-22%	4%	12%	-6%	-38%	16%	-18%	-15%	36%	32%	11%	26%	25%	0%	0%	0%	0%	0%
EPS Diluted (pro forma)	31%	47%	-22%	4%	12%	-5%	-38%	16%	-19%	-15%	32%	29%	8%	23%	22%	0%	0%	0%	0%	0%

Source: Company reports and Ascendant Capital Markets estimates.

Atossa Therapeutics, Inc.

Balance Sheet (\$ mils)	Mar-23	Jun-23	Sep-23	Dec-23	Mar-24	Jun-24	Sep-24	Dec-24	Mar-25	Jun-25	Sep-25	Dec-25	Mar-26	Jun-26	Sep-26	Dec-26
Fiscal Year End: December 31	Q1A	Q2A	Q3A	Q4A	Q1A	Q2A	Q3A	Q4A	Q1E	Q2E	Q3E	Q4E	Q1E	Q2E	Q3E	Q4E
Assets																
Cash and cash equivalents	103.9	99.4	94.0	88.5	84.0	79.5	74.8	71.1	64.0	56.7	49.5	42.3	35.0	27.8	20.5	13.3
Short term investments									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Restricted cash	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Research and development tax rebat	0.7	0.7	0.0						0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Deferred income taxes									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Prepaid expenses and other	6.2	5.8	3.5	3.6	3.0	2.1	2.2	3.3	3.3	3.3	3.3	3.3	3.3	3.3	3.3	3.3
Total current assets	110.9	106.0	97.6	92.2	87.1	81.7	77.1	74.5	67.3	60.1	52.9	45.6	38.4	31.1	23.9	16.6
Property and equipment, net									(0.0)	0.0	0.0	0.0	0.0	0.1	0.1	0.1
Intangibles, net									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Deferred income tax									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other	5.3	2.4	4.0	4.0	4.0	4.1	2.4	2.0	2.0	2.0	2.0	2.0	2.0	2.0	2.0	2.0
Total assets	116.2	108.4	101.7	96.3	91.1	85.9	79.5	76.4	69.3	62.1	54.9	47.7	40.4	33.2	26.0	18.7
Liabilities and stockholders' equity																
Accounts payable	1.4	1.2	0.7	0.8	1.2	1.1	1.6	0.7	0.7	0.7	0.7	0.7	0.7	0.7	0.7	0.7
Accrued expenses	0.6	2.3	2.7	2.6	1.6	1.9	1.7	2.8	2.8	2.8	2.8	2.8	2.8	2.8	2.8	2.8
Deferred income tax									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Warrant liabilities									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other	0.9	0.0	0.0	1.8	2.5	2.7	2.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Short term debt									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total current liabilities	2.9	3.5	3.4	5.2	5.3	5.7	5.8	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
Deferred income taxes									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Warrant liabilities									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other long term liabilities									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Long term debt									0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Total other liabilities	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Preferred stock	0.6	0.6	0.6	0.0					0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Common stock	22.8	22.8	22.8	22.8	22.829	22.9	22.9	23.5	24.3	25.0	25.8	26.6	27.4	28.1	28.9	29.7
Additional paid-in capital	252.4	253.8	253.4	254.5	255.096	255.5	256.2	259.8	259.8	259.8	259.8	259.8	259.8	259.8	259.8	259.8
Retained earnings	(162.5)	(172.3)	(178.5)	(186.3)	(192.166)	(198.2)	(205.4)	(211.8)	(219.8)	(227.8)	(235.8)	(243.8)	(251.8)	(259.8)	(267.8)	(275.8)
Accumulated other comprehensive income									0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Total stockholders' equity	113.3	104.9	98.3	91.0	85.759	80.2	73.7	71.5	64.4	57.1	49.9	42.7	35.5	28.2	21.0	13.8
Total stockholders' equity and liabil	116.2	108.4	101.7	96.3	91.087	85.9	79.5	76.4	69.3	62.1	54.9	47.6	40.4	33.2	26.0	18.7

Balance Sheet Drivers

	Mar-23	Jun-23	Sep-23	Dec-23	Mar-24	Jun-24	Sep-24	Dec-24	Mar-25	Jun-25	Sep-25	Dec-25	Mar-26	Jun-26	Sep-26	Dec-26
	Q1A	Q2A	Q3A	Q4A	Q1A	Q2A	Q3A	Q4A	Q1E	Q2E	Q3E	Q4E	Q1E	Q2E	Q3E	Q4E
Book & Cash Value (per share)																
Book Value per Share (diluted)	\$0.89	\$0.83	\$0.78	\$0.73	\$0.68	\$0.64	\$0.59	\$0.56	\$0.50	\$0.44	\$0.39	\$0.33	\$0.27	\$0.22	\$0.16	\$0.11
Cash per Share (diluted)	\$0.82	\$0.78	\$0.75	\$0.70	\$0.67	\$0.63	\$0.59	\$0.56	\$0.50	\$0.44	\$0.38	\$0.33	\$0.27	\$0.21	\$0.16	\$0.10
Net cash per Share (diluted)	\$0.82	\$0.78	\$0.75	\$0.70	\$0.67	\$0.63	\$0.59	\$0.56	\$0.50	\$0.44	\$0.38	\$0.33	\$0.27	\$0.21	\$0.16	\$0.10

Source: Company reports and Ascendant Capital Markets estimates

Atossa Therapeutics, Inc.

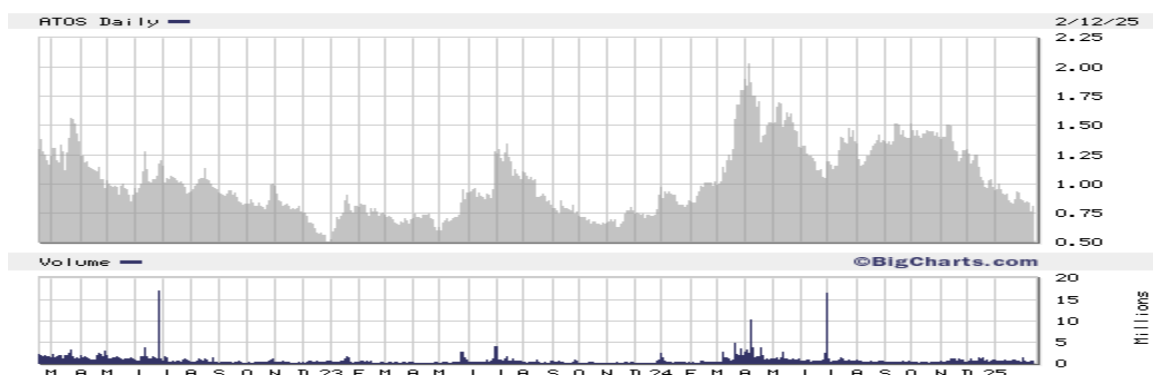
Cash Flow Statement (\$ mils)	Mar-23	Jun-23	Sep-23	Dec-23	2023	Mar-24	Jun-24	Sep-24	Dec-24	2024	Mar-25	Jun-25	Sep-25	Dec-25	2025	Mar-26	Jun-26	Sep-26	Dec-26	2026
Fiscal Year End: December 31	Q1A	Q2A	Q3A	Q4A	FY-A	Q1A	Q2A	Q3A	Q4A	FY-E	Q1E	Q2E	Q3E	Q4E	FY-E	Q1E	Q2E	Q3E	Q4E	FY-E
Cash flow from operating activities																				
Net income	(6.3)	(9.8)	(6.2)	(7.8)	(30.1)	(5.9)	(6.0)	(7.2)	(6.3)	(25.5)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)	(8.0)	(8.0)	(8.0)	(8.0)	(32.0)
Depreciation	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Amortization					0.0					0.0					0.0					0.0
Debt related amortization expense					0.0					0.0					0.0					0.0
Stock comp	1.6	1.6	0.9	0.5	4.6	0.4	0.4	0.7	0.8	2.3	0.8	0.8	0.8	0.8	3.1	0.8	0.8	0.8	0.8	3.1
Deferred rent					0.0					0.0					0.0					0.0
A/R reserves					0.0					0.0					0.0					0.0
Deferred income taxes					0.0					0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Change in fair value of warrant liability					0.0					0.0					0.0					0.0
Writedowns and impairments		3.0			3.0			1.7		1.7					0.0					0.0
Other gains/losses					0.0	0.0	0.0	(0.0)	0.0	0.0					0.0					0.0
Other					0.0					0.0					0.0					0.0
Changes in operating assets and liabilities:																				
Prepaid expenses & other current	(1.3)	(0.5)	2.3	2.2	2.8	0.7	0.9	(0.2)	(1.0)	0.4	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Research and development tax	0.0	0.0	0.7	0.0	0.7					0.0					0.0					0.0
Income tax					0.0					0.0					0.0					0.0
Other assets	1.6	0.9	(1.7)	(2.4)	(1.7)		(0.1)	0.1	0.4	0.3	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Accounts payable	(1.5)	(0.4)	(0.3)	0.1	(2.2)	0.4	(0.2)	0.5	(0.9)	(0.1)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Accrued expenses	(0.4)	0.2	0.4	(0.2)	(0.1)	(0.4)	1.3	(0.2)	(0.8)	(0.1)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Other liabilities	(0.7)	0.6	0.0	1.9	1.9	0.0	(0.7)	(0.2)	0.8	(0.1)	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Net cash (used in) provided by	(7.0)	(4.5)	(3.9)	(5.6)	(20.9)	(4.7)	(4.5)	(4.8)	(7.1)	(21.0)	(7.2)	(7.2)	(7.2)	(7.2)	(28.9)	(7.2)	(7.2)	(7.2)	(7.2)	(28.9)
Cash flow from investing activities																				
Purchases of property and equipment		(0.0)	(0.0)		(0.0)	(0.0)	(0.0)	(0.0)		(0.0)	0.0	(0.0)	0.0	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.1)
Purchases of short-term investments					0.0					0.0					0.0					0.0
Acquisitions					0.0					0.0					0.0					0.0
Other					0.0					0.0					0.0					0.0
Net cash used in investing activities	0.0	(0.0)	(0.0)	0.0	(0.0)	(0.0)	(0.0)	(0.0)	0.0	(0.0)	0.0	(0.0)	0.0	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.0)	(0.1)
Cash flow from financing activities																				
Issuance of debt					0.0					0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Repayment of debt					0.0					0.0					0.0					0.0
Issuance of stock			(1.5)		(1.5)					0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Proceeds from stock option exercises					0.0	0.2	0.1		3.4	3.7					0.0					0.0
Other					0.0					0.0					0.0					0.0
Dividends and distributions					0.0					0.0					0.0					0.0
Cash provided by (used in) financing	0.0	0.0	(1.5)	0.0	(1.5)	0.2	0.1	0.0	3.4	3.7	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Effect of exchange rate on cash					0.0					0.0					0.0					0.0
Net increase (decrease) in cash	(7.0)	(4.5)	(5.4)	(5.6)	(22.4)	(4.5)	(4.4)	(4.8)	(3.7)	(17.4)	(7.2)	(7.2)	(7.2)	(7.2)	(28.9)	(7.2)	(7.3)	(7.2)	(7.3)	(29.0)
Beginning cash and equivalents	111.0	104.0	99.5	94.1	111.0	88.6	84.1	79.6	74.9	88.6	71.2	64.0	56.7	49.5	71.2	42.3	35.0	27.8	20.5	42.3
Ending cash and equivalents	104.0	99.5	94.1	88.6	88.6	84.1	79.6	74.9	71.2	71.2	64.0	56.7	49.5	42.3	42.3	35.0	27.8	20.5	13.3	13.3

Source: Company reports and Ascendant Capital Markets estimates

ANALYST CERTIFICATION

Each analyst hereby certifies that the views expressed in this report reflect the analyst's personal views about the subject securities or issuers. Each analyst also certifies that no part of the analyst's compensation was, is, or will be, directly or indirectly, related to the specific recommendations or views expressed in this report. The analyst who prepared this report is compensated based upon the overall profitability of Ascendant Capital Markets, LLC, which may, from time to time, include the provision of investment banking, financial advisory and consulting services. Compensation for research is based on effectiveness in generating new ideas for clients, performance of recommendations, accuracy of earnings estimates, and service to clients.

Atossa Therapeutics, Inc.



Source: <https://bigcharts.marketwatch.com/>

	Report Date		Price
Report	Date	Rating	Target
1	9/17/2020	Buy	7.00
2	11/15/2020	Buy	7.50
3	4/6/2021	Buy	7.75
4	5/31/2021	Buy	8.00
5	9/7/2021	Buy	8.50
6	11/20/2021	Buy	8.75
7	3/3/2022	Buy	8.00
8	5/29/2022	Buy	7.50
9	8/17/2022	Buy	7.00
10	11/12/2022	Buy	6.00
11	3/27/2023	Buy	5.50
12	5/23/2023	Buy	5.25
13	9/6/2023	Buy	5.50
14	12/22/2023	Buy	5.75
15	5/1/2024	Buy	6.00
16	6/5/2024	Buy	6.25
17	9/10/2024	Buy	6.50
18	12/7/2024	Buy	7.00

- Ascendant Capital Markets, LLC has received compensation for advisory or investment banking services from the company in the past 12 months.

IMPORTANT DISCLOSURES

This report has been distributed by Ascendant Capital Markets, LLC and is for the sole use of our clients. This report is based on current public information that we consider reliable, but we do not represent it is accurate or complete, and it should not be relied on as such. This report contains information from various sources, including United States government publications, The Wall Street Journal and other periodicals, Yahoo! Finance and other sources, and is for informational purposes only and is not a recommendation to trade in the securities of the companies mentioned within the report. We seek to update our research and recommendations as

appropriate, but the large majority of reports are published at irregular intervals as we consider appropriate and, in some cases, as constrained by industry regulations.

We may have a business relationship with companies covered in this report. Ascendant Capital Markets, LLC may make a market in the securities of the subject company. We and our affiliates, officers, directors, and employees will from time to time have long or short positions in, act as principal in, and buy or sell, the securities or derivatives (including options and warrants) thereof of covered companies referred to in this report. This report is not an offer to sell or the solicitation of an offer to buy any security in any jurisdiction where such an offer or solicitation would be illegal. It does not constitute a personal recommendation or take into account the particular investment objectives, financial situations, or needs of individual clients. Clients should consider whether any information in this report is suitable for their particular circumstances and, if appropriate, seek professional advice, including tax advice. The price and value of the investments referred to in this report may fluctuate.

Following are some general risks that can adversely impact future operational and financial performance and share price valuation: (1) industry fundamentals with respect to legislation, mandates, incentives, customer demand, or product pricing; (2) issues relating to competing companies or products; (3) unforeseen developments with respect to management, financial condition or accounting policies or practices; or (4) external factors that affect the interest rates, currency, the economy or major segments of the economy. Past performance is not a guide to future performance, future returns are not guaranteed, and loss of original capital may occur. Certain transactions, including those involving futures, options, and other derivatives, give rise to substantial risk and are not suitable for all investors. Our report is disseminated primarily electronically, and, in some cases, in printed form. The information contained in this report is not incorporated into the contents of our website and should be read independently thereof. Copyright Ascendant Capital Markets, LLC. No part of this material may be copied, photocopied or duplicated by any means or redistributed without the prior written consent of Ascendant Capital Markets, LLC.

Risks & Considerations

Risks to attainment of our share price target include balance sheet/liquidity risks, failure of product candidates to demonstrate safety and efficacy in clinical trials, failure to gain regulatory approvals, ability to commercialize product, failure to obtain suitable reimbursement, competition, changing macroeconomic factors, investor sentiment for investing in biotech stocks, and changes in consumer or government priorities for healthcare.

Ascendant Capital Markets, LLC Rating System

BUY: We expect the stock to provide a total return of 15% or more within a 12-month period.

HOLD: We expect the stock to provide a total return of negative 15% to positive 15% within a 12-month period.

SELL: We expect the stock to have a negative total return of more than 15% within a 12-month period.

Total return is defined as price appreciation plus dividend yield.

Ascendant Capital Markets, LLC Distribution of Investment Ratings (as of April 11, 2025)

Rating	Count	Percent	Investment Banking Services Past 12 months	
			Count	Percent
Buy	52	98%	21	40%
Hold	0	0%	0	0%
Sell	1	2%	0	0%
Total	53	100%	21	40%

Other Important Disclosures

Our analysts use various valuation methodologies including discounted cash flow, price/earnings (P/E), enterprise value/EBITDAS, and P/E to growth rate, among others. Risks to our price targets include failure to achieve financial results, product risk, regulatory risk, general market conditions, and the risk of a change in economic conditions.

Dissemination of Research

Ascendant Capital Markets, LLC research is distributed electronically via the Thomson Reuters platforms, Bloomberg, Capital IQ and FactSet. Please contact your investment advisor or institutional salesperson for more information.

General Disclaimer

The information and opinions in this report were prepared by Ascendant Capital Markets, LLC. This information is not intended to be used as the primary basis of investment decisions and because of individual client objectives it should not be construed as advice designed to meet the particular investment needs of any investor. This material is for information purposes only and is not an offer or solicitation with respect to the purchase or sale of any security. The reader should assume that Ascendant Capital Markets, LLC may have a conflict of interest and should not rely solely on this report in evaluating whether or not to buy or sell securities of issuers discussed herein. The opinions, estimates, and projections contained in this report are those of Ascendant Capital Markets, LLC as of the date of this report and are subject to change without notice. Ascendant Capital Markets, LLC endeavors to ensure that the contents have been compiled or derived from sources that we believe are reliable and contain information and opinions that are accurate and complete. However, Ascendant Capital Markets, LLC makes no representation or warranty, express or implied, in respect thereof, takes no responsibility for any errors and omissions contained herein, and accepts no liability whatsoever for any loss arising from any use of, or reliance on, this report or its contents. Information may be available to Ascendant Capital Markets, LLC, or its affiliates that is not reflected in this report. This report is not to be construed as an offer or solicitation to buy or sell any security.

Additional Disclosures

Ascendant Capital Markets, LLC is a broker-dealer registered with the United States Securities and Exchange Commission (SEC) and a member of the FINRA and SIPC. Ascendant Capital Markets, LLC is not a Registered Investment Advisor nor is it an investment advisor registered with the Securities and Exchange Commission or with the securities regulators of any state, and at the present time is not eligible to file for federal registration.