

Zacks Small-Cap Research

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Revelation Bios

(REVB-NASDAQ)

REVB: Company Making Great Progress Toward Commercialization

Revelation Biosciences is a life sciences company whose focus is the development of immunologic-based therapies. We value REVB at \$19.50 using the comparable method involving companies with similar characteristics.

OUTLOOK

Revelation Biosciences is a life sciences company whose development of immunologic-based therapies is based on the well-established biology of phosphorylated hexaacyl disaccharide (PHAD) and its effect on the innate immune system.

The company announced 2Q financial results that show a clinical stage company with good cash and a solid balance sheet. The company is making great progress in its Gemini program.

Current Price (08/06/26)

\$1.12

Valuation

\$19.50

SUMMARY DATA

52-Week High	\$11.72
52-Week Low	\$0.95
One-Year Return (%)	-89.02
Beta	-0.15
Average Daily Volume (sh)	42,163

Shares Outstanding (mil)	4
Market Capitalization (\$mil)	\$4
Short Interest Ratio (days)	N/A
Institutional Ownership (%)	13
Insider Ownership (%)	5

Annual Cash Dividend	\$0.00
Dividend Yield (%)	0.00

5-Yr. Historical Growth Rates

Sales (%)	N/A
Earnings Per Share (%)	N/A
Dividend (%)	N/A

Risk Level

High

Type of Stock

Small-Value

ZACKS ESTIMATES

Revenue

(in millions of \$)

	Q1 (Mar)	Q2 (Jun)	Q3 (Sep)	Q4 (Dec)	Year (Dec)
2023	0 A	0 A	0 A	0 A	0 A
2024	0 A	0 A	0 A	0 A	0 A
2025	0 A	0 A	0 A	0 A	0 A
2026	0 A	0 A	0 E	0 E	0 E

Earnings Per Share

	Q1 (Mar)	Q2 (Jun)	Q3 (Sep)	Q4 (Dec)	Year (Dec)
2023	1.96A	-0.25A	-0.26A	-8.33A	-10.80A
2024	-2.46A	-5.13A	-0.84A	-10.06A	-18.49A
2025	-2.11A	-7.01A	-1.77A	-1.65A	-23.95A
2026	-2.71A	-0.87A	-0.82E	-0.77E	-5.17E

*numbers don't add due to reverse stock split

Update

Revelation Biosciences (REVB) has continued to strengthen its position as a clinical-stage biotechnology company focused on developing therapies that address immune dysfunction associated with acute and chronic diseases. While the company remains in the development stage without commercial revenue, the past quarter has been marked by meaningful regulatory, clinical, and organizational milestones that have the potential to greatly increase shareholder value by reducing development risk and advancing its lead product candidate, Gemini, toward pivotal testing.

The most significant development during the quarter was the company's continued progress with Gemini, its immunomodulatory therapy being developed for acute kidney injury (AKI). Management announced that the U.S. Food and Drug Administration has agreed to a single adaptive Phase 2/3 clinical study design for Gemini in AKI patients. This represents an important regulatory achievement because it provides a potentially streamlined path toward registration while allowing the company to collect efficacy and safety data in a more efficient manner than separate Phase 2 and Phase 3 trials. For a small biotechnology company, regulatory alignment of this nature can substantially reduce execution risk and accelerate the timeline to potential commercialization.

Supporting that progress, Revelation also presented additional data from its PRIME clinical study demonstrating that Gemini continued to normalize inflammatory biomarkers and restore immune responsiveness in patients with chronic kidney disease. These findings reinforce the company's scientific hypothesis that correcting immune dysfunction may improve outcomes in patients at high risk for acute kidney injury and other inflammatory conditions. Importantly, the data demonstrated durable biological activity following a single administration, providing additional evidence supporting Gemini's mechanism of action.

Another encouraging development was the expansion of Revelation's Acute Kidney Injury Advisory Board with internationally recognized experts in nephrology and critical care medicine. The addition of experienced clinical leaders should help optimize the design and execution of the upcoming adaptive Phase 2/3 study while increasing confidence that the program is being guided by physicians with extensive expertise in treating critically ill patients.

The company also adopted a stockholder rights plan during the quarter. While such plans can be viewed differently by investors depending on circumstances, management indicated that the objective is to protect shareholders from coercive takeover tactics and help ensure that all investors have the opportunity to realize the full value of the company should a strategic transaction ever be considered.

Revelation's recently released quarterly financial results reflected a company continuing to invest aggressively in advancing its lead clinical asset. Research and development expenses increased as Gemini moved closer to late-stage testing, contributing to a quarterly net loss of approximately \$3.3 million. While higher spending is expected during clinical development, investors should take comfort in the company's strengthened balance sheet. Cash and cash equivalents stood at approximately \$11.5 million following the successful January warrant inducement financing, and management stated that existing cash resources are expected to fund operations into the first quarter of 2027. This provides financial flexibility as the company prepares for its next major clinical milestones.

Although Revelation remains an early-stage biotechnology company and continues to face the normal risks associated with drug development, the recent quarter demonstrated meaningful execution across several critical areas. The company strengthened its cash position, secured an important FDA agreement regarding

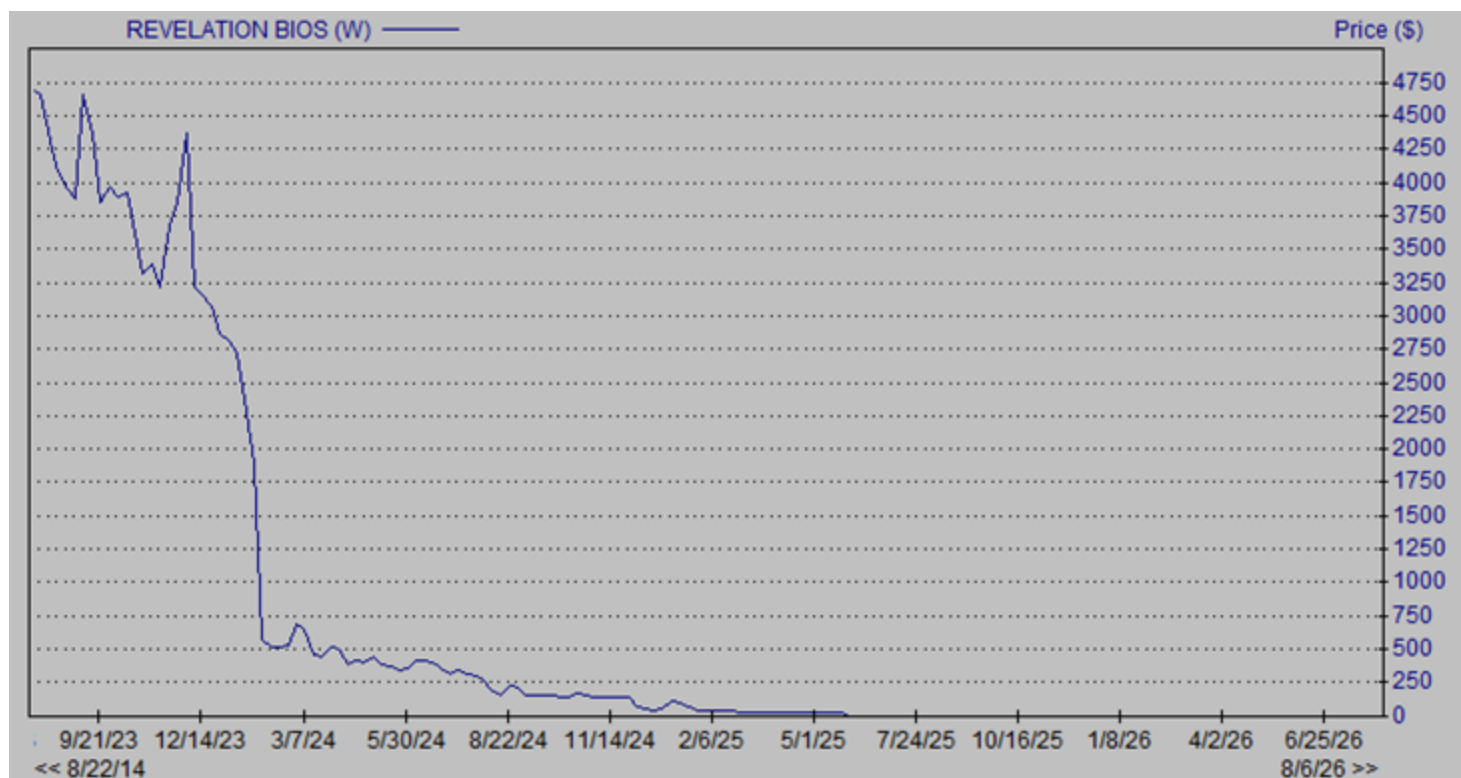
the clinical pathway for Gemini, generated additional supportive clinical data, expanded its scientific advisory capabilities, and continues preparing for what could become the most important value-creating stage of development.

Looking ahead, we are focusing on the initiation of the adaptive Phase 2/3 AKI study, additional clinical updates from the Gemini program, and continued interactions with regulators. If Gemini continues to demonstrate the encouraging biological activity observed thus far and advances successfully through late-stage clinical development, Revelation Biosciences could be positioned to address a significant unmet medical need in acute kidney injury, a condition with limited therapeutic options and substantial healthcare impact. The combination of improving regulatory clarity, encouraging clinical evidence, and a cash runway extending into 2027 provides the company with a stronger foundation as it works toward its next phase of growth.

PROJECTED INCOME STATEMENT & BALANCE SHEET

Revelation Income Statement and Balance Sheet									
	1Q2025A	2Q2025A	3Q2025A	4Q2025A	1Q2026A	2Q2026A	3Q2026E	4Q2026E	2027E
Revenues	0	0	0	0	0	0	0	0	0
Operating Expenses									
Research and Development	858,830	1,317,980	922,857	964,189	1,376,861	1,918,424	1,975,977	2,035,256	5,507,444
General and Administrative	1,236,157	1,143,249	1,020,154	1,607,398	1,715,639	1,743,817	1,813,570	1,886,112	6,862,556
Total Operating Expenses	2,094,987	2,461,229	1,943,011	2,571,587	3,092,500	3,662,241	3,789,546	3,921,368	12,370,000
Gain/(Loss) from Operations	(2,094,987)	(2,461,229)	(1,943,011)	(2,571,587)	(3,092,500)	(3,662,241)	(3,789,546)	(3,921,368)	(12,370,000)
Other Income/(Expense)									
Change in fv of warrant liab.	1,417	44	611	87	-	-	-	-	-
Other income/(expense)	42,285	16,803	35,224	60,493	83,449	338,857	355,800	373,590	333,796
Total other income/(expense)	43,702	16,847	35,835	60,580	83,449	338,857	355,800	373,590	333,796
Net Income/(Loss)	(2,051,285)	(2,444,382)	(1,907,176)	(2,511,007)	(3,009,051)	(3,323,384)	(3,433,747)	(3,547,779)	(12,036,204)
Deemed Dividends	0	(3,181,786)	(2,769,742)	0	(5,681,616)	0	0	0	0
Net loss attributable to Common S/H	(2,051,285)	(5,626,168)	(4,676,918)	(2,511,007)	(8,690,667)	(3,323,384)	(3,433,747)	(3,547,779)	(12,036,204)
Net Income/(Loss) per share	(\$2.11)	(\$7.01)	(\$1.77)	(\$1.65)	(\$2.71)	(\$0.87)	(\$0.82)	(\$0.77)	(\$2.36)
Shares Outstanding-diluted	971,313	802,670	2,644,733	1,524,011	3,207,720	3,829,363	4,212,299	4,633,529	5,096,882
Assets									
Current Assets:									
Cash and equivalents	3,704,611	5,173,871	12,708,489	10,700,331	14,113,415	11,482,792	12,631,071	13,894,178	13,407,744
Other current assets	223,270	186,049	123,934	111,297	92,620	201,724	207,776	214,009	95,399
Total current assets	3,927,881	5,359,920	12,832,423	10,811,628	14,206,035	11,684,516	12,838,847	14,108,187	13,503,143
Property and Equipment and other	42,390	35,170	23,919	771,296	838,247	813,416	821,550	829,766	838,063
Total Assets	3,970,271	5,395,090	12,856,342	11,582,924	15,044,282	12,497,932	13,660,397	14,937,953	14,341,206
Liabilities and SH Equity									
Current Liabilities:									
Accounts payable	625,058	828,894	1,034,951	577,501	828,203	931,093	949,715	968,709	844,767
Accrued expenses	463,632	739,626	819,381	1,397,644	468,082	759,937	775,136	790,638	477,444
Operating lease liability	0	-	-	23,013	120,861	189,370	193,157	197,021	123,278
Warrant liabilities	829	786	175	-	-	-	-	-	-
Total current liabilities	1,089,519	1,569,306	1,854,507	1,998,158	1,417,146	1,880,400	1,918,008	1,956,368	1,445,489
Operating Lease--net of current	-	-	-	723,771	638,183	550,546	523,019	496,868	472,024
Total Liabilities	1,089,519	1,569,306	1,854,507	2,721,929	2,055,329	2,430,946	2,441,027	2,453,236	1,917,513
Shareholder Equity									
Common Stock	964	1,535	3,586	1,584	3,908	3,908	1,535	1,535	1,535
Additional Paid-in Capital	45,436,511	48,825,354	57,906,530	58,278,698	65,413,383	65,814,800	66,472,948	67,137,677	68,684,051
Accumulated Deficit	(42,556,723)	(45,001,105)	(46,908,281)	(49,419,287)	(52,428,338)	(55,751,722)	(55,255,113)	(54,654,495)	(56,717,493)
Total Shareholder Equity	2,880,752	3,825,784	11,001,835	8,860,995	12,988,953	10,066,986	11,219,370	12,484,717	12,423,693
Total Liabilities and SE	3,970,271	5,395,090	12,856,342	11,582,924	15,044,282	12,497,932	13,660,397	14,937,953	14,341,206

HISTORICAL STOCK PRICE



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