



**3R MATRIX**

	+	=	-
Right Sector (RS)	✓	■	■
Right Quality (RQ)	✓	■	■
Right Valuation (RV)	✓	■	■

+ Positive = Neutral - Negative

**What has changed in 3R MATRIX**

	Old		New
RS	■	↔	■
RQ	■	↔	■
RV	■	↔	■

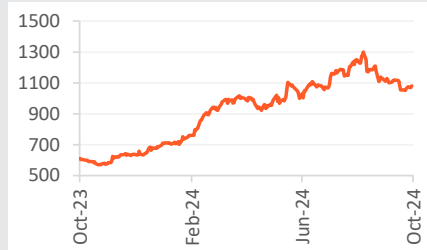
**Company details**

Market cap:	Rs. 98,497 cr
52-week high/low:	Rs. 1324/568
NSE volume: (No of shares)	19.66 lakh
BSE code:	532321
NSE code:	ZYDUSLIFE
Free float: (No of shares)	25.6 cr

**Shareholding (%)**

Promoters	75.0
FII	5.6
DII	12.6
Others	6.9

**Price chart**



**Price performance**

(%)	1m	3m	6m	12m
Absolute	-12.7	-9.7	-3.0	58.7
Relative to Sensex	-0.1	-0.5	-0.4	48.7

Sharekhan Research, Bloomberg

**Zydu Lifesciences Ltd**  
R&D advancements to drive growth

<b>Pharmaceuticals</b>	<b>Sharekhan code: ZYDUSLIFE</b>		
<b>Reco/View: Buy</b>	↔	<b>CMP: Rs. 1,066</b>	<b>Price Target: Rs. 1,268</b>
↑ Upgrade	↔ Maintain	↓ Downgrade	

**Summary**

- Zydu Lifesciences Ltd has partnered with the Central Drug Research Institute (CDRI) to develop an oral treatment for osteoporosis related to chronic kidney disease (CKD) and postmenopausal conditions, focusing on small-molecule inhibitors of the protein sclerostin.
- The company has launched a Phase-1 trial for the ZY19489-Ferroquine combination, now actively recruiting 36 patients to assess safety, tolerability, and pharmacokinetics, with a projected completion date of February 1, 2025.
- Zydu has received a tentative approval from the USFDA to manufacture Enzalutamide Tablets for prostate cancer treatment, with production set to occur at its SEZ facility in Ahmedabad; these tablets generated annual sales of approximately \$1.42 billion in the US as of July 2024.
- Clinical pipeline includes ZYL1, currently in Phase 1 and Phase 2 trials for Amyotrophic Lateral Sclerosis (ALS) and undergoing a Phase 2 trial for mild to moderately active ulcerative colitis in patients resistant or intolerant to oral aminosaliclates.

Zydu Lifesciences Ltd has partnered with the Central Drug Research Institute (CDRI) to develop an oral treatment for osteoporosis linked to chronic kidney disease (CKD) and post-menopausal conditions, focusing on small-molecule inhibitors of the protein sclerostin, which disrupts bone metabolism in advanced CKD patients. Collaboration will involve joint pre-clinical studies, with Zydu leading further drug development to address needs of over 10% of the global CKD-affected population. Additionally, Zydu has launched a Phase 1 trial for the ZY19489-Ferroquine combination, which is actively recruiting 36 patients to evaluate safety, tolerability, and pharmacokinetics, with an anticipated completion date of February 1, 2025. The company has also received tentative USFDA approval to manufacture Enzalutamide Tablets for treating castration-resistant and metastatic castration-sensitive prostate cancer, with production at its SEZ facility in Ahmedabad, and these tablets have generated approximately USD 1.42 billion in annual sales in the US as of July 2024. Furthermore, Zydu is advancing its clinical pipeline with ZYL1, currently undergoing Phase 1 and Phase 2 trials for Amyotrophic Lateral Sclerosis (ALS) and a Phase 2 trial for mild to moderately active ulcerative colitis, targeting patients intolerant to oral aminosaliclates.

**Partnership with Zydu:** Zydu Lifesciences Ltd has teamed up with the CDRI to develop an oral treatment for osteoporosis associated with chronic kidney disease (CKD) and post-menopausal conditions. This collaboration focuses on identifying small-molecule inhibitors of sclerostin, a protein that disrupts bone metabolism, particularly in advanced CKD patients. CDRI's research has highlighted sclerostin as a key target for drug development, aiming to create therapies that reduce fracture risk while preserving renal function. This initiative seeks to meet the healthcare needs of over 10% of the global population suffering from CKD, underscoring the demand for safer osteoporosis treatments for this vulnerable group.

**Phase 1 Trial for ZY19489-ferroquine combination launched:** As of August 30, 2024, Zydu Lifesciences Limited has launched a Phase 1 trial for the ZY19489-Ferroquine combination, which is currently "actively recruiting" participants. This trial will be conducted at a single site with the goal of enrolling 36 patients to assess the safety, tolerability, and pharmacokinetics of the treatment. The primary objective is to develop a simplified treatment regimen for malaria, with an expected completion date for the primary phase set for February 1, 2025.

**Tentative USFDA nod for enzalutamide tablets received:** Zydu Lifesciences Limited has received tentative approval from the USFDA to produce Enzalutamide Tablets in 40 mg and 80 mg dosages for the treatment of castration-resistant and metastatic castration-sensitive prostate cancer. These tablets will be manufactured at Zydu's facility located in the SEZ, Ahmedabad. As of July 2024, Enzalutamide tablets have generated approximately USD 1.42 billion in annual sales in the United States, underscoring their substantial market potential.

**Additional active programs in clinical pipeline:** Zydu Lifesciences Limited is advancing its clinical pipeline, particularly with ZYL1, which is currently undergoing Phase 1 and Phase 2 trials for Amyotrophic Lateral Sclerosis (ALS). These trials aim to evaluate the efficacy, safety, and pharmacokinetics of ZYL1, with the most advanced trial expected to conclude by August 25, 2024. Additionally, ZYL1 is in a Phase 2 trial for treating mild to moderately active ulcerative colitis in patients who are resistant to or intolerant of oral aminosaliclates.

**Our Call**

**Valuation – Maintain Buy with a revised PT of Rs. 1,268:** Zydu Lifesciences Ltd. is well-positioned for significant growth, supported by its strategic partnerships and active clinical programs. Collaboration with the CDRI to develop an oral treatment for osteoporosis associated with chronic kidney disease (CKD) targets a critical healthcare need, potentially addressing the concerns of over 10% of the global CKD population. Additionally, the company's Phase 1 trial for the ZY19489-Ferroquine combination showcases its commitment to innovating malaria treatments, with a focus on safety and tolerability. Zydu's recent tentative approval from the USFDA to manufacture Enzalutamide Tablets further underscores its market potential, particularly as these tablets generated approximately \$1.42 billion in annual sales in the U.S. as of July 2024. Furthermore, Zydu is advancing its clinical pipeline with ZYL1 for ALS and ulcerative colitis, enhancing its therapeutic portfolio. Collectively, these initiatives not only strengthen Zydu's position in the pharmaceutical market but also suggest a robust valuation trajectory driven by diverse revenue streams and innovative product development. is currently trading at 23x and 22x its estimated EPS for FY26 and FY27, respectively, with projected EPS of Rs. 45.56 and Rs. 48.75.

**Key Risks**

- Price erosion in the US generics business could hurt performance.
- Forex volatility could affect earnings

**Valuation (Consolidated)**

Particulars	FY23	FY24	FY25E	FY26E	FY27E
Net Sales	1724	1955	2228	2518	2820
Growth %	13%	13%	14%	13%	12%
EBITDA	378	538	624	692	761
Margin	22%	28%	28%	28%	27%
Adjusted PAT	189	384	415	458	490
EPS	18.67	38.14	41.29	45.56	48.75
PE	57x	28x	26x	23x	22x
P/BV	6x	5x	5x	4x	3x
EV/EBITDA	29x	20x	17x	16x	14x
ROE (%)	11%	19%	18%	17%	16%
ROCE (%)	15%	19%	19%	19%	18%

Source: Company; Sharekhan estimates

## Outlook and Valuation

### ■ Sector view - Input cost easing with companies focusing on complex product launches

Over the years, Indian pharmaceutical companies have established themselves as a dependable source for global peers. A confluence of other factors, including a focus on specialty/complex products in addition to emerging opportunities in the API space, would be key growth drivers over the long term. The sector is seeing an easing of input costs – raw material, freight and power, which aids the sector in expanding margins. The sector is also witnessing an easing of price erosion followed by increasing contribution from product launches. We believe the sector is in a sweet spot, where it is experiencing healthy product mix and cost rationalization, which increases operational profit of companies. The sector is mainly a low-debt sector and increasing operational profit followed by experiencing advantage of a low tax rate due to its operations in the SEZ sector, hence overall, we stay positive on the sector.

### ■ Company outlook - Healthy growth prospects

Over the long term, both the US and India have a healthy growth outlook. The US business is on a strong footing, helped by a sturdy new product pipeline and ramp-up of recent product launches, which would be long-term growth drivers. However, in the near term, high price erosion would act as dampeners. Efforts to build a presence in the injectables space would also add to growth albeit over the medium to long term. The India business has a robust growth outlook, backed by pick-up in chronic as well as acute therapies and a few substantial high-value launches lined up. Over the long term, product launches such as Saroglitazar, gRevlimid, and Desidustat offer substantial growth potential. With a sharp reduction in debt, Zydus Lifesciences has strengthened its balance sheet. The management looks to keep an eye on debt reduction going ahead as well. This augurs well and would strengthen the company's financial muscle. Strong earnings prospects, healthy return ratios, and strengthening balance sheet are key positives for Cadila. In the near term, US market's growth is expected to moderate, while India and other geographies are likely to stage double-digit growth.

### ■ Valuation - Maintain Buy with a revised PT of Rs. 1,268

Zydus Lifesciences Ltd. is well-positioned for significant growth, supported by its strategic partnerships and active clinical programs. Collaboration with the CDRI to develop an oral treatment for osteoporosis associated with chronic kidney disease (CKD) targets a critical healthcare need, potentially addressing the concerns of over 10% of the global CKD population. Additionally, the company's Phase 1 trial for the ZY19489-Ferroquine combination showcases its commitment to innovating malaria treatments, with a focus on safety and tolerability. Zydus's recent tentative approval from the USFDA to manufacture Enzalutamide Tablets further underscores its market potential, particularly as these tablets generated approximately \$1.42 billion in annual sales in the U.S. as of July 2024. Furthermore, Zydus is advancing its clinical pipeline with ZYIL1 for ALS and ulcerative colitis, enhancing its therapeutic portfolio. Collectively, these initiatives not only strengthen Zydus's position in the pharmaceutical market but also suggest a robust valuation trajectory driven by diverse revenue streams and innovative product development. is currently trading at 23x and 22x its estimated EPS for FY26 and FY27, respectively, with projected EPS of Rs. 45.56 and Rs. 48.75.

## About company

Zydu Lifescience is one of the leading pharmaceutical companies in India. The company is present across the pharmaceutical value chain of research, development, manufacturing, marketing, and selling of finished dosage human formulations (generics, branded generics, and specialty formulations, including biosimilars and vaccines), active pharmaceutical ingredients (APIs), animal healthcare products, and consumer wellness products. The company has a global presence and sells its products in the US, India, Europe, and emerging markets, including countries in Latin America, Asia Pacific region, and Africa. The company is also engaged in research and development activities focused across the value chain of API process development, generics development for simple as well as differentiated dosage forms such as modified release oral solids, transdermal, topicals and nasals, biologics, vaccines, and new chemical entities (NCE).

## Investment theme

Zydu Lifescience is favorably progressing in its efforts to build an alternative growth platform (NCE, biologics, and vaccines) that should start delivering over the medium to long term and reduce the company's dependence on limited competition assets in the US for its earnings. India business, including the consumer wellness segment, is likely to grow at a healthy pace, albeit over the medium to long term. Zydu Lifesciences is in a sweet spot, wherein both its geographies have an improved growth outlook. Easing pricing pressures, sturdy new product pipeline, and ramp-up in the recent product launches would be key growth drivers for the US business. The efforts to build up presence in the injectables space would also add to growth albeit over the medium to long term. India business is also showing signs of a pick-up in growth momentum, led by a solid presence in the chronic and sub-chronic segments and an improving outlook for the acute segment. Further, COVID-19 related opportunities would add to the company's growth momentum.

## Key Risks

1) Regulatory compliance risk; 2) delay in product approvals; 3) currency risk; and 4) risk concentration in the US portfolio.

## Additional Data

### Key management personnel

Pankaj R. Patel	Chairman
Dr. Sharvil P. Patel	Managing Director
Mr. Ganesh Nayak	COO & Executive Director
Mr. Nitin Parekh	CFO

Source: Company

### Top 10 shareholders

Sr. No.	Holder Name	Holding (%)
1	Life Insurance Corp of India	3.16
2	Kotak Mahindra Asset Management Co	1.51
3	ICICI Prudential Asset Management	1.16
4	Vanguard Group Inc/The	0.98
5	Blackrock Finance Inc	0.86
6	PPFAS Asset Management	0.77
7	Nippon Life India Asset Management	0.46
8	Axis Asset Management Co Ltd/India	0.45
9	Norges Bank	0.43
10	UTI Asset Management Co Ltd	0.31

Source: Bloomberg

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## Understanding the Sharekhan 3R Matrix

Right Sector	
<b>Positive</b>	Strong industry fundamentals (favorable demand-supply scenario, consistent industry growth), increasing investments, higher entry barrier, and favorable government policies
<b>Neutral</b>	Stagnancy in the industry growth due to macro factors and lower incremental investments by Government/private companies
<b>Negative</b>	Unable to recover from low in the stable economic environment, adverse government policies affecting the business fundamentals and global challenges (currency headwinds and unfavorable policies implemented by global industrial institutions) and any significant increase in commodity prices affecting profitability.
Right Quality	
<b>Positive</b>	Sector leader, Strong management bandwidth, Strong financial track-record, Healthy Balance sheet/cash flows, differentiated product/service portfolio and Good corporate governance.
<b>Neutral</b>	Macro slowdown affecting near term growth profile, Untoward events such as natural calamities resulting in near term uncertainty, Company specific events such as factory shutdown, lack of positive triggers/events in near term, raw material price movement turning unfavourable
<b>Negative</b>	Weakening growth trend led by led by external/internal factors, reshuffling of key management personal, questionable corporate governance, high commodity prices/weak realisation environment resulting in margin pressure and deteriorating balance sheet
Right Valuation	
<b>Positive</b>	Strong earnings growth expectation and improving return ratios but valuations are trading at discount to industry leaders/historical average multiples, Expansion in valuation multiple due to expected outperformance amongst its peers and Industry up-cycle with conducive business environment.
<b>Neutral</b>	Trading at par to historical valuations and having limited scope of expansion in valuation multiples.
<b>Negative</b>	Trading at premium valuations but earnings outlook are weak; Emergence of roadblocks such as corporate governance issue, adverse government policies and bleak global macro environment etc warranting for lower than historical valuation multiple.

Source: Sharekhan Research

# Sharekhan

by BNP PARIBAS

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