

FEATURE · COMPLEX GENERICS & PE

The \$1.5 Billion Question in Hyderabad: Orbicular, Peptides and the PE Re-Rating of Indian Complex Generics

Warburg Pincus and CVC have begun circling a specialty developer whose value now rests less on a 90-ANDA filing machine than on two GLP-1 peptides, a pen device built in India, and a partnership model that already ships generic Ozempic in Canada.

By Anup Soans · MedicinMan · 15 September 2026

Private equity does not pay a billion-dollar multiple for a Hyderabad research shop because the PowerPoint says “complex generics.” It pays when a laboratory has converted scientific difficulty into a file that two G7 regulators will accept — and when that file is attached to the most valuable peptide class of the decade.

That is the story behind this morning’s Mint report: Orbicular Pharmaceutical Technologies is exploring a stake sale, Jefferies is mandated, Tata Capital Healthcare Fund may take a partial or full exit, and Warburg Pincus and CVC Capital Partners are already spending time on the asset. People familiar with the matter put the valuation band at \$1 billion to \$1.5 billion. The process is early. The official launch is expected next quarter. Other PE firms and strategics are expected to look. Nothing is signed.

The number, however, is not early. It is a verdict on what Indian specialty development is worth when the product is no longer a cream or a nasal spray, but a synthetic incretin in a multi-dose pen.

The Deal, as It Stands

Mint, citing three people familiar with the matter, reported on 15 September 2026 that Orbicular has attracted early interest from Warburg Pincus and CVC. Jefferies is the sell-side advisor. Tata Capital Healthcare Fund II, which invested approximately \$20 million in May 2024 for an undisclosed stake, is likely to explore a part or full exit. The exact stake size and valuation “may evolve depending on the incoming investor.” Orbicular, CVC and Tata Capital did not comment; Jefferies and Warburg declined.

Moneycontrol had flagged a \$1 billion-plus process as early as 11 August 2026. Together, the two reports describe a familiar PE choreography: a growth investor that came in two years ago, a company whose earnings have just doubled, a category (GLP-1 peptides) that global capital already understands, and a formal auction still weeks away.

The structure will likely mix secondary (Tata Capital and, possibly, some promoter liquidity) with primary capital to fund the next wave of peptide filings and scale-up. That is the standard mid-life transaction for an Indian specialty developer that has outgrown its first institutional cheque.

Who Orbicular Actually Is

Orbicular Pharmaceutical Technologies Private Limited (CIN U24233TG2008PTCo60606) was incorporated in 2008 and commenced operations in 2010. It is based at Plot 53, ALEAP Industrial Estate, Kukatpally / Bachupally, Hyderabad. The company describes a research campus of more than 100,000 square feet and a scientific team of roughly 250 to 300 people.

Founders Dr M. S. Mohan (Managing Director) and Dr Hiren Patel (CEO / President) built the firm as a partnership-led developer: select a hard product, take it through formulation, IP, analytics, PK/clinical and regulatory packaging, then license it late to a company that already owns the market. They call the domain CI-TOP SPIN — complex injectables, topicals, ophthalmics, parenterals, nasal sprays and inhalations. The acronym isn't for branding. It is product selection: complexity, entry barriers, limited competition.

The model is older than peptides. In 2016, Alembic took 60 per cent of Aleor Dermaceuticals, a dermatology JV with Orbicular. In 2023, Strides partnered with Orbicular to develop four nasal sprays (combined IQVIA market then put above \$400 million), to be made at Strides' Chestnut Ridge, New York site and sold through Strides' US and European engine. Those deals established the template PE is now underwriting at a different scale: Orbicular does the science; a larger partner does the plant, the ANDA nameplate and the sales force.

The board today includes promoters (Dr Mohan, Dr Patel, Meena Mohan) and Tata Capital nominee Visalakshi Chandramouli, managing partner of the healthcare fund that wrote the 2024 cheque.

From R&D Consultancy to Peptide Specialist

ICRA, cited by Mint, draws the arc cleanly. Orbicular began as an R&D consultancy and became a developer of complex formulations with a focus on niche peptide molecules. In FY25 it began commercial production of liraglutide formulations. It now has two molecules in commercial production: liraglutide and semaglutide.

As of March 2026, ICRA said the company had filed more than 90 abbreviated new drug applications — some on its own, some with partners — and received approvals for more than 66 products. That filing machine is the old Orbicular. The new Orbicular is the one ICRA flagged as one of three players with an approved semaglutide molecule in Canada, supporting a strong order book at 31 March 2026.

The financial step-change matches the product step-change. Operating income rose from ₹265.2 crore in FY25 to ₹525 crore in FY26. Net profit rose from ₹28.7 crore to ₹53.3 crore. Inc42 had earlier placed FY25 revenue in the ₹262–272 crore range. Doubling sales in a year is not a consulting-shop trajectory. It is what happens when a peptide leaves the lab.

The Peptide Pipeline: What Is Proven, What Is Promised

Semaglutide is the franchise

Semaglutide is not “in development.” It is a multi-market regulatory asset, chemically synthesised to match a recombinant originator — the hardest form of peptide generic work, because impurity profile, potency, device compatibility and stability must satisfy regulators who know the reference product was made in a different way.

The architecture is a three-party stack. Orbicular led product development and the technical programme, and has claimed that API, pen device, formulation, manufacturing and analytical technology were developed end-to-end in India. Apotex is the ANDA / ANDS applicant and commercialiser in the United States and Canada. OneSource Specialty Pharma is the CDMO; commercial supply is planned from its USFDA-approved flagship site in Bengaluru.

Table 1. Semaglutide programme — disclosed status (as of September 2026)

Market	Status	What it means
United States	FDA tentative ANDA for generic Ozempic (Apr 2026)	Science accepted; launch blocked until patents/exclusivity lapse. Reports cite ~2030 or a main compound cliff around 2032. A parked, validated file — not FY27 US sales.
Canada — T2D	Health Canada NOC; Apo-Semaglutide Injection. Strengths: 2 mg/3 mL (0.68 mg/mL) and 4 mg/3 mL (1.34 mg/mL)	Canada’s originator protections lapsed 5 January 2026 — first G7 market open to generics. Apotex shipped mid-May. List price reported near C\$78 vs ~C\$228–240 brand per four weeks.
Canada — obesity	SEVMIA (Apotex + Orbicular): first generic Wegovy equivalent. 4 mg/3 mL pen. Approved ~25–29 June 2026	Weight-management indication, including adolescents 12+ on the Health Canada label. Third generic semaglutide product authorised in Canada; first for chronic weight management.
European Union	EU DCP filing announced February 2026 (diabetes and weight management)	Execution signal, not an approval. EU timing will follow both assessment and originator IP.

Dr Mohan called the Health Canada notice of compliance, after the FDA tentative approval, validation of an integrated approach across formulation, analytics and regulatory strategy. He also said the quiet thing PE needs to hear: “Peptides remain a strategic growth area for Orbicular, with Semaglutide the first of several programs in our pipeline targeted at global regulated markets.”

Canada has already tested the commercial stack. By late July 2026, after manufacturing trouble at an Indian plant sidelined at least two competitors, Apotex was briefly the only generic Ozempic on Canadian shelves. The Globe and Mail explicitly tied key ingredients to Orbicular. That is both a capability proof and a concentration risk: few sites are qualified to make this product at this quality.

Early sales remain modest against brand. One secondary compilation put Apotex generic sales at about C\$15 million by 30 June 2026 versus a still-dominant branded franchise. First-mover rent in Canada will compress as Dr Reddy’s, Aspen and others fill the channel. The PE case is not that Canada alone is worth \$1.5 billion. It is that Canada is the live demonstration of a file that can travel.

Liraglutide is commercial — and quieter

ICRA names liraglutide as the first GLP-1 Orbicular put into commercial production, in FY25. Unlike semaglutide, the partner, market and label have not been as publicly documented. The honest reading is earlier-generation GLP-1 (Victoza / Saxenda analogues) already shipping into markets where intellectual property has expired. It matters because it shows the peptide plant and the tech-transfer muscle were working before the Ozempic file became a headline.

What is not on the board

Orbicular's public language includes "complex peptide combination products," device-based peptide injections, long-acting polymeric suspensions, liposomes and nano-systems, and some 505(b)(2) work under the complex-injectable vertical led by Dr Bhavesh Patel. None of that has been reduced to a named next molecule. There is no public confirmation of tirzepatide, CagriSema, oral semaglutide or a dual/triple agonist. Those are the logical next bets if the firm stays inside incretins. Treating them as in-house would be invention, not analysis.

So the disclosed peptide pipeline, stripped of brochure language, is this: two commercial molecules (liraglutide and semaglutide); two Canadian labels on semaglutide (diabetes and obesity); one FDA-tentative US Ozempic file; one EU DCP filing; and an undisclosed follow-on peptide slate that management has chosen to describe but not name.

Why the Multiple Looks Aggressive — and Why It May Not Be

At \$1.0–1.5 billion against FY26 operating income of ₹525 crore (about \$60 million at prevailing rates), the implied sales multiple is roughly 16–24 times trailing revenue. That is not a generic-Indian-formulations multiple. It is a scarce-capability multiple.

Four things support it. First, the science is no longer theoretical: FDA tentative approval plus two Health Canada authorisations on a synthetic-versus-biologic incretin is a high bar. Second, cash has started: Canada is open; FY26 already shows the inflection. Third, the US file is a 2030s call option on the largest semaglutide market in the world, sitting inside a partnership with a commercialiser that knows North America. Fourth, the platform argument — "first of several programs" — is what growth PE is paid to underwrite, provided diligence confirms the next files exist.

Four things cut against it. Named pipeline visibility after semaglutide is thin. US economics are back-ended and patent-sensitive; published dates do not even agree. Partner contracts with Apotex and OneSource — royalties, supply price, capacity reservation, change-of-control — are not public and will make or break the model. And Canada is filling up. A valuation that assumes durable Canadian rents will be marked down in the data room.

The Encube Ethicals process, where a Warburg–Mubadala consortium was reported to be leading a majority bid around \$1.8 billion for a scaled topical CDMO, is the nearest comparable in the current Indian PE cycle. Encube is a plant-and-brand story. Orbicular is an IP-and-file story. Same buyer universe; different asset.

What This Deal Says About Indian Pharma

India has spent two decades proving it can make high-volume oral solids. The next proof is whether it can industrialise complexity: peptides, devices, long-acting injectables, products that fail if the analytical method is merely good rather than forensic. Orbicular is a small company in that argument, but it is a clean exhibit. The API, the pen and the file were claimed as Indian work. Canada was the first G7 commercial market. The first US regulatory handshake was tentative approval, not a refuse-to-receive.

It also says something about capital. Tata Capital's \$20 million in 2024 looked like a specialist growth cheque. Twenty-eight months later, the same asset is being discussed at fifty to seventy-five times that cheque. That is what happens when a category (GLP-1) re-prices an entire skill set. Warburg and CVC do not need to be taught what semaglutide is. They need to be convinced that Orbicular can repeat the file.

Leadership, in this business, is still measured the old way: results that improve the organisation's performance by improving the capabilities of the people who do the work. Capability here is not a poster on the wall. It is a peptide assay, a pen specification, a Health Canada notice of compliance, and a plant in Bengaluru that kept shipping when another Indian site could not.

What to Watch Next

1. Process timing. Formal launch next quarter, as Mint's sources described it. Binding offers and exclusivity will tell us whether \$1.5 billion was aspiration or clearing price.
2. Stake mix. How much is Tata Capital exiting, how much is primary, and whether promoters remain controlling partners — the Encube pattern of a junior promoter residual is one template, not a law.
3. The next named peptide. Until a second incretin or combination is disclosed with a partner and a filing date, the "several programs" claim is a diligence item, not a portfolio.
4. Canada share and price. If generic semaglutide pricing collapses faster than volume rises, FY27 earnings quality will disappoint anyone who annualised May–July.
5. US patent outcome. Tentative approval is a scientific trophy until the last relevant patent or exclusivity falls. The difference between 2030 and 2032 is two years of option value.

Orbicular is not yet a household name in Indian pharma. It does not need to be. The firms that change the industry's centre of gravity are often the ones that spent a decade on products other people found too difficult, then arrived in the market just as the world decided those products were indispensable. Peptides are that moment. The PE process is simply the market putting a number on it.

Anup Soans is Editor of MedicinMan and a consultant to the pharmaceutical industry. This article is based on published regulatory announcements, company and partner statements, and contemporaneous business reporting. It is not investment advice.

Appendix: Sources

All facts in the article are drawn from the sources below. Where reports disagree (notably US patent timing for semaglutide), both versions are flagged in the text. URLs are current as of 15 September 2026.

A. The stake-sale process

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2. Moneycontrol, "Mega pharma deal brewing! Stake sale in the works at Orbicular Pharmaceutical Technologies; firm seeking \$1 bn plus valuation" (Ashwin Mohan), 11 August 2026. <https://x.com/moneycontrolcom/status/2087185044465168448>

B. Company identity, funding and financials

3. Tata Capital Healthcare Fund II investment of ~\$20 million: Economic Times, 31 May / 1 June 2024; Entrepreneur India, 3 June 2024; Indian Startup News; BW Healthcare World; Digital Health News. Ambit advised Orbicular.
4. PitchBook company profile, Orbicular Pharmaceutical Technologies (founded 2010; PE-backed; ~\$20m raised). <https://pitchbook.com/profiles/company/234202-42>
5. CB Insights company profile. <https://www.cbinsights.com/company/orbicular-pharmaceutical-technologies>
6. Inc42 company page (FY25 revenue ₹261.5–272 crore; last funding 30/31 May 2024). <https://inc42.com/company/orbicular-pharmaceutical-technologies/>
7. Tofter company dossier: CIN U24233TG2008PTC060606; directors including M.S. Mohan, Hiren Patel, Meena Mohan, Visalakshi Chandramouli (Tata Capital nominee). <https://www.tofter.in/orbicular-pharmaceutical-technologies-private-limited/company/U24233TG2008PTC060606>
8. ICRA note as cited in Mint (August/September 2026): 90+ ANDAs filed, 66+ approvals as of March 2026; liraglutide commercial from FY25; semaglutide also in commercial production; one of three Canadian-approved semaglutide players; FY26 operating income ₹525 crore vs ₹265.2 crore; PAT ₹53.3 crore vs ₹28.7 crore.

C. Company self-description and platform

9. Orbicular website — About, Business, Team, Infrastructure. CI-TOP SPIN; high-end peptides; device-based peptide injections. <https://www.orbicular.co.in/>
10. Orbicular LinkedIn company page (specialties: niche injectables, nasal sprays, peptides, drug-device combinations, long-acting complex injectables). <https://in.linkedin.com/company/orbicularpharma>

D. Semaglutide — United States

11. Orbicular / Apotex: USFDA tentative approval for generic Ozempic (semaglutide injection). The Hindu Business Line, 11 April 2026; Indian Pharma Post, 11 April 2026; ET Pharma, 15 April 2026. Apotex Corp. is ANDA applicant and US commercialiser.
12. OneSource Specialty Pharma exchange / press announcement, 21 April 2026: Orbicular + US front-end partner received FDA tentative ANDA; OneSource is CDMO from Bengaluru USFDA site. https://www.onesourcecdmo.com/wp-content/uploads/2026/04/Press_Release_Orbiculars-Partner-secures-tentative-U.S.-ANDA-approval-for-a-generic-version-of-Ozempic%C2%AE-.pdf
13. Express Pharma; Financial Express; PharmaSource; CNBC-TV18; Business Standard market notes on the same 21 April 2026 OneSource disclosure.
14. The Globe and Mail, 12–14 April 2026: Apotex tentative FDA nod; product developed with Orbicular; discussion of US patent timing vs Canadian expiry.
15. ET Pharma, 15 April 2026: notes molecule exclusivity "expected to lapse in 2030." Contrast with Globe and Mail / GLP1Prices commentary citing a main US patent around 2032. Both recorded; not reconciled.

E. Semaglutide — Canada (diabetes and obesity)

16. Orbicular press release via The Hindu BrandHub, 5 May 2026: Health Canada NOC for generic Ozempic with Apotex; presentations 2 mg/3 mL and 4 mg/3 mL; “second G7 approval” after FDA TA; claim of end-to-end Indian development of API, pen, formulation and analytics. <https://www.thehindu.com/brandhub/pr-release/orbicular-announces-health-canada-approval-for-a-g-ozempic-semaglutide-injection-in-partnership-with-apotex/article70942553.ece>
17. Health Canada, “Canada approves second generic semaglutide, the first G7 country to do so,” 1 May 2026 (Apotex ANDS for Ozempic-equivalent; Dr Reddy’s authorised days earlier).
18. The Globe and Mail, 1 May 2026 and subsequent availability explainers (4 June 2026): Apotex second approval, first with Canadian manufacturing presence; shipping from 19 May; price comparisons.
19. OneSource, 4 May 2026: “second generic semaglutide approval in Canada”; commercial supply from Bengaluru.
20. Apotex Health Corp. / CNW, 30 June 2026: first Health Canada approval for generic Wegovy-equivalent, SEVMIA, developed with Orbicular; 4 mg/3 mL (1.34 mg/mL) pen.
21. Health Canada, “Canada approves first generic semaglutide for weight loss,” 29 June 2026; Regulatory Decision Summary for Sevmia (synthetic product comparable to biologic CRP Wegovy).
22. Pearce IP / Lexology, 29 June 2026; Globe and Mail, 30 June 2026.
23. The Globe and Mail, 28–29 July 2026: Apotex briefly sole generic Ozempic supplier after Indian-plant issues; key ingredients via Orbicular.
24. BBC News, 4 June 2026: Canada first G7 generic semaglutide market; Apotex and Dr Reddy’s.

F. Semaglutide — Europe

25. Orbicular Pharmaceutical Technologies LinkedIn post, February 2026: EU DCP filing for generic semaglutide, diabetes and weight-management indications. https://www.linkedin.com/posts/orbicularpharma_semaglutide-glp1-peptidgenerics-activity-7428085768658821120-8Gp8

G. Earlier partnership history (non-peptide platform)

26. Mint / Business Standard, 21 April 2016: Alembic 60% / Orbicular 40% JV, Aleor Dermaceuticals, for global dermatology.
27. Strides–Orbicular nasal-spray partnership, 3 May 2023: four sprays, IQVIA market >\$400 million; manufacture at Chestnut Ridge, NY. Medical Dialogues; OINDP News.

H. Comparable PE process

28. Economic Times: Warburg–Mubadala leading race for majority in Encube Ethicals at around ₹16,500 crore (~\$1.8 billion). Context only — different business mix (topical CDMO / brands) from Orbicular’s development-and-license model.