

EXECUTIVE SUMMARY

First half of 2026



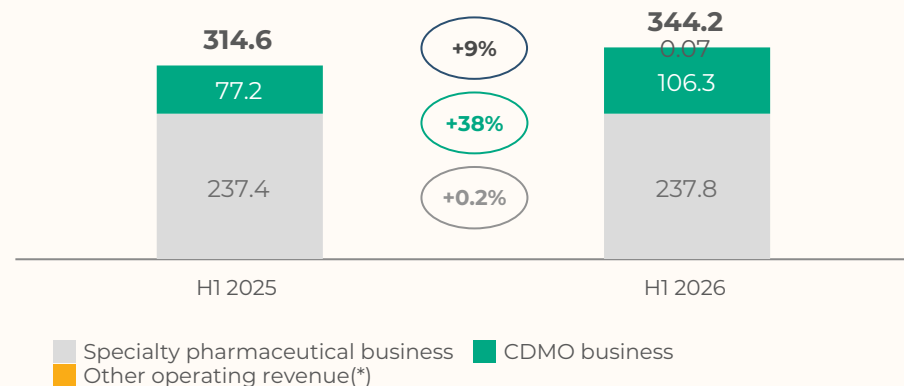
Main figures €Mn

Op. Revenue	EBITDA	EBIT
344.2 (+9%)	121.2 (+85%)	103.2 (+103%)
Net profit	Capex	Net cash
84.4 (+113%)	30.4 (46%)	22.5

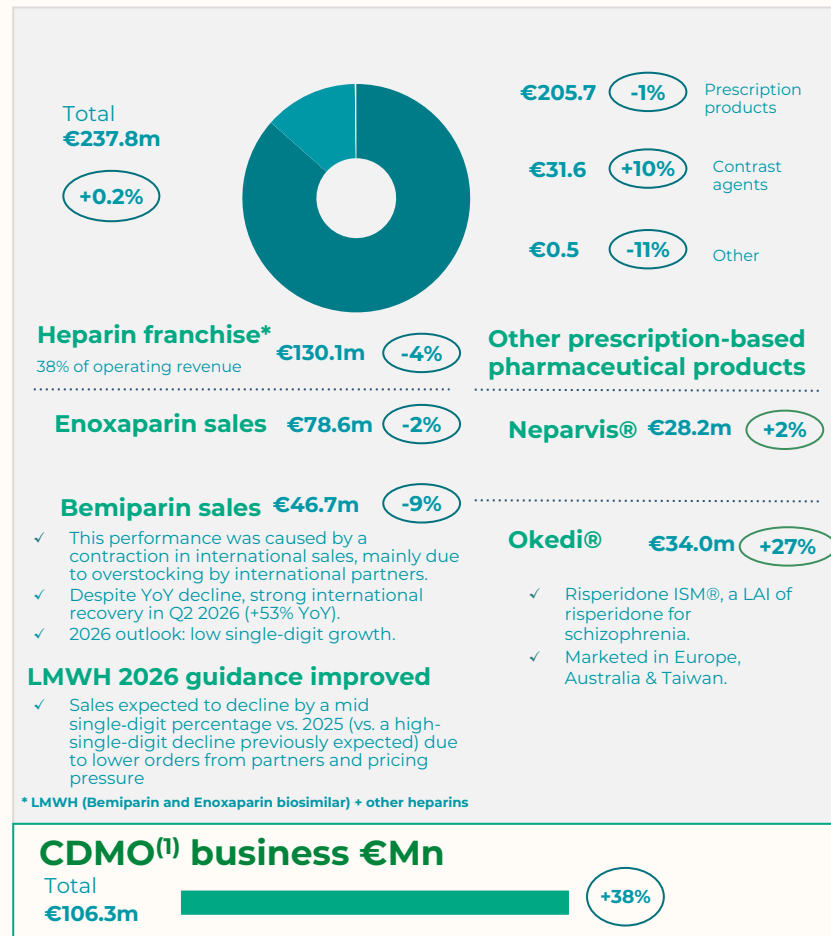
2026 operating revenue growth rate:
Increase by a low- to mid-single-digit percentage vs 2025

- ✓ Phoenix acquisition completed on April 1, 2026. Preliminary purchase price allocation resulted in a €62.4 Mn bargain purchase gain (badwill), recognized as non-recurring income in the consolidated income statement.

Operating revenue €Mn



Specialty pharma business €Mn

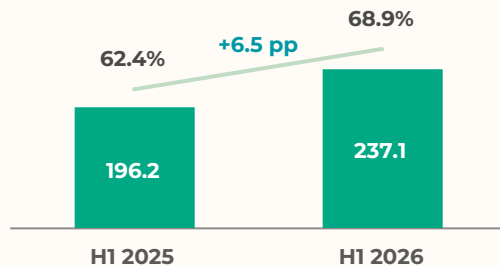


(*) "Other operating income" includes service activities that are not material to the Group.

⁽¹⁾ Contract Development and Manufacturing Organisation

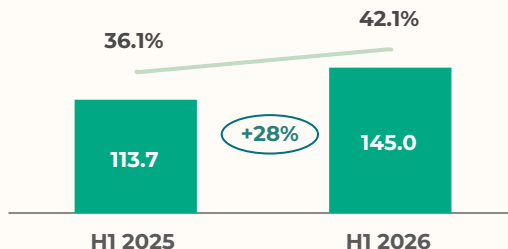
To obtain further information on the alternative performance measures (APMs) and non-IFRS financial indicators used, including the definition thereof and a reconciliation between the applicable management indicators and the financial information set out in the consolidated financial statements prepared under IFRSs, please consult the information included on this subject on page 1 and Appendix 2 (pages 32-36) of the press release on the financial results for the first half of 2026. Said document is available on ROVI's website and may be accessed on the following link <https://www.rovi.es/en/shareholders-investors/financial-business-information>

Gross profit €Mn



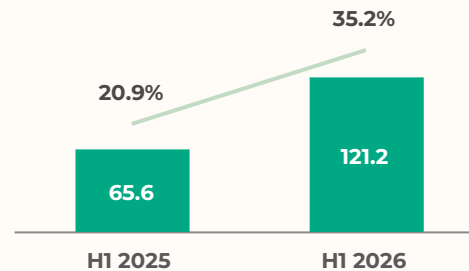
■ Gross profit — Gross margin

SG&A expenses €Mn



■ SG&A — SG&A/Operating revenue

EBITDA €Mn



■ EBITDA — EBITDA Margin

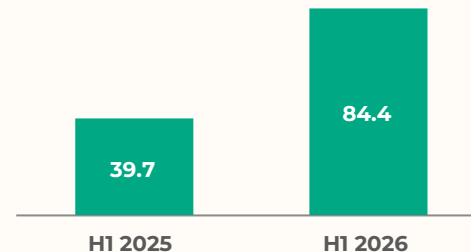
EBITDA Pre-R&D and pre badwill⁽²⁾ €Mn **€92.2m** **+12%**

R&D expenses €Mn €33.3

+98%

Product	Potential Indication	Current Situation	Key Milestones
Risperidone ISM®, monthly	Schizophrenia	Approved	Marketed in Europe, and in Australia & Taiwan
Letrozole ISM®, annual	Breast Cancer	Clinical development on hold	Phase I: Superior oestrogen suppression vs Femara®
Letrozole SIE ⁽¹⁾ , quarterly	Breast Cancer	Completion of phase I	Phase I: positive readout confirms superior estrogen suppression vs Femara® and allows progression to phase III clinical trial
Risperidone ISM®, quarterly	Schizophrenia	Completion of phase I	Phase I: positive readout allows progression to phase III clinical trial
Concentrated on improving posology for already approved compounds, which benefits risk / reward profile Multiple FDA / GMP approved facilities to support the platform			

Net profit €Mn



Closing remarks

- ✓ Strong momentum in CDMO business: consistent and focused execution
- ✓ Value creation from the acquisition of an injectable drug product manufacturing site in Phoenix, Arizona (USA)
- ✓ Vertical integration of the heparin division to enhance cost efficiency and improve competitiveness
- ✓ Sustained growth of the specialty pharmaceutical business, driven by Okedi®
- ✓ Strong R&D investment, with two clinical trials ongoing

(1) Superior Inhibition of Estrogen; (2) calculated excluding badwill in H1 2026 and R&D expenses in H1 2025 and in H1 2026.

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