ROYALTY PHARMA

IMDELLTRA royalty acquisition

August 2025

Forward Looking Statements

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Key messages

1

Imdelltra addresses high unmet need in SCLC

Amgen's Imdelltra, a first-in class targeted immunotherapy, received accelerated approval (May 2024) for small cell lung cancer (SCLC)

Significant unmet need in SCLC; median survival of ~12 months following initial therapy and 5-year survival of 7% across all stages⁽¹⁾

Treatment options in SCLC limited and provide only modest benefit beyond chemotherapy

2

Transformative Phase 3 clinical results

Imdelltra reduced risk of death by 40% vs. standard-of-care chemotherapy in 2L+ SCLC

Imdelltra seeing broad physician uptake as comfort with safety profile grows⁽²⁾

Additional studies ongoing in first-line SCLC and other DLL3-expressing tumors

3

Rapid uptake and multiblockbuster potential

Rapid launch driven by Amgen with strong first-mover advantage and aggressive clinical development strategy

Multi-blockbuster potential with consensus sales of ~\$520m in FY2025 projected to grow to >\$2.8bn in FY 2035⁽³⁾

Royalty duration expected until 2038-2041

Transaction expected to deliver unlevered IRR in the low-double digits

²L: second-line; DLL3: Delta-like ligand 3

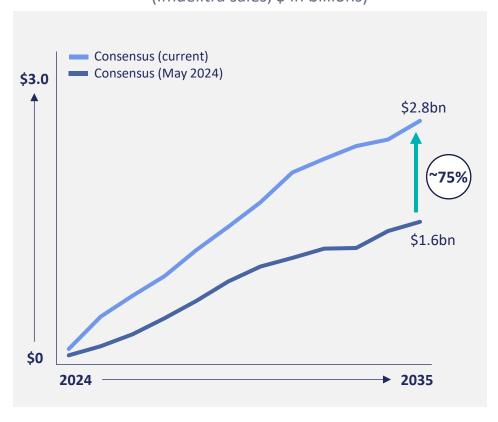
^{1.} Amgen presentation, May 2024.

^{2.} Cytokine release syndrome (CRS) and immune effector cell-associated neurotoxicity syndrome (ICANS) with Imdelltra was primarily low grade and manageable with supportive care.

Acquired royalty on Amgen's Imdelltra for small cell lung cancer

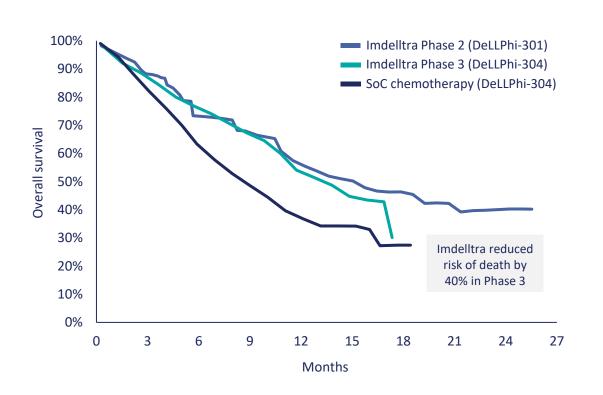
- Up to \$950m for portion of BeOne's approved, uncapped royalty in Amgen's Imdelltra for extensive-stage small cell lung cancer
 - \$885m upfront payment
 - Approximately 7% royalty on worldwide net sales⁽²⁾
 - Royalty duration expected through 2038-2041
- Option structure provides BeOne added flexibility
 - BeOne can sell additional portion of Imdelltra royalty to Royalty Pharma for up to \$65m over next 12 months
- Transaction expected to deliver unlevered IRR in the low-double digits

Amgen's launch has outperformed consensus⁽¹⁾ (Imdelltra sales; \$ in billions)

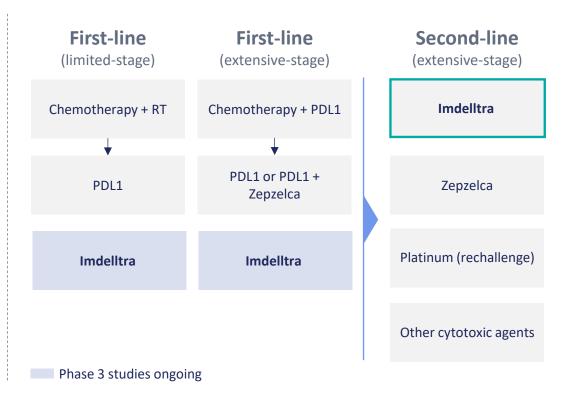


Imdelltra is transforming the SCLC treatment paradigm

Imdelltra OS benefit has impressive durability⁽¹⁾



SCLC treatment paradigm



Positive DelLphi-304 confirmatory results to support full U.S. approval and geographic expansion outside the U.S.

Royalty Pharma sees multi-blockbuster potential

RP sees significant commercial opportunity in the approved 2L+ SCLC indication with high conviction in first-line label expansion

Extensive-stage 2L+ SCLC

(Imdelltra approved)

>8,000-10,000 U.S. patients/year⁽¹⁾

- ☑ DelLphi-304 (Phase 3)
- ☑ DelLphi-301 (Phase 2)

First-line SCLC

(Phase 3 studies ongoing)

~27,000 U.S. patients/year(1)

Dellphi-305 (ES - maintenance)
Dellphi-312 (ES - induction)
Dellphi-306 (LS - maintenance)

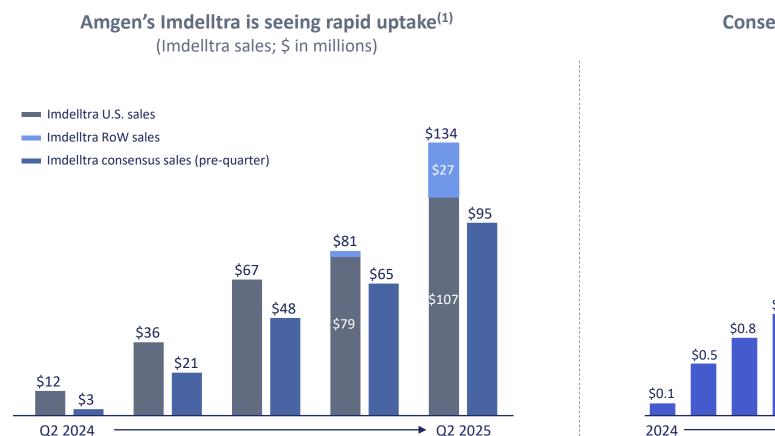
Additional opportunities

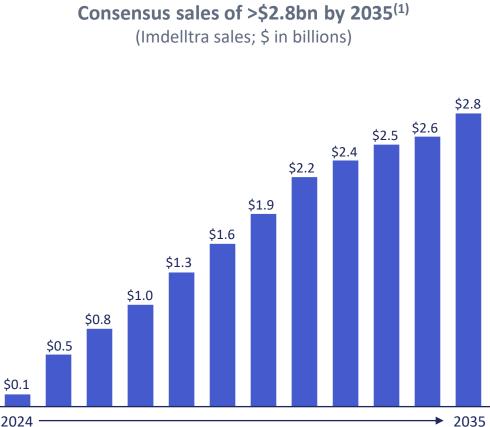
(Clinical studies ongoing)

Other DLL3-expressing tumors

TARLANEC (Phase 3 IIT)(2)

Imdelltra to enhance RP long-term growth and diversification





Amgen is driving rapid uptake of Imdelltra in the U.S., which is also off to a strong launch internationally