



Acquisition of pacritinib Royalty and CTI BioPharma Secured Loan

August 25, 2021

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TRANSACTION HIGHLIGHTS

- Medically necessary product / addresses unmet medical need
- Strong and long-lasting intellectual property
- Extends weighted average duration of royalty portfolio (to over 9 years)
- Within target deal size of US\$25M to US\$150M
- Synthetic royalty stream aligns with stated growth strategy

TRANSACTION HIGHLIGHTS

- Transaction demonstrates ability of DHT to acquire quality assets with “win-win” solutions through bespoke transaction structures that meet objectives of counterparties
- Immediate deployment of US\$50 million via secured loan, with overall deal size of US\$110-US\$135 million subject to FDA approval and achievement of certain pacritinib sales thresholds
- Pacritinib is expected to become the first drug to be approved for treatment of Myelofibrosis with severe thrombocytopenia
- Immediate cash flow from interest income on loan commencing Q3 2021 with first royalty payment expected in Q1 2022

TRANSACTION TERMS

Secured long-term royalty stream on high quality product, with immediate contribution to cash flow



- Senior secured loan of US\$50 million to CTI BioPharma Corp. on closing
 - Interest rate of LIBOR plus 825 bps (subject to minimum 10.0%)
 - 5-year term
- Purchase of royalty on pacritinib on sales in the United States for US\$60 million upon FDA approval of pacritinib (expected decision date of November 30, 2021)
 - Royalty entitlement of 9.60% on first US\$125 million of sales, 4.50% on next US\$50 million of sales, 0.50% on the next US\$225 million of sales, and a cap at US\$400 million of sales
- Up to US\$25 million in milestone payments to CTI upon meeting sales thresholds by Q3 2023

MYELOFIBROSIS w/ SEVERE THROMBOCYTOPENIA

- Myelofibrosis (MF) is a form of bone marrow cancer that can lead to thrombocytopenia and anemia, weakness, fatigue and enlarged spleen and liver, with a median survival range of 2.25 – 11.25 years⁽¹⁾
- Thrombocytopenia (low blood platelet count) is an adverse prognostic variable that increases in prevalence with disease progression
- Patients suffering from severe thrombocytopenia (platelet counts less than $50 \times 10^9/L$) typically have more advanced disease, a higher risk of bleeding, increased risk of leukemic transformation, and a shorter overall survival, shown to be just 15 months⁽²⁾⁽³⁾
- While stem cell transplant is used in a small number of MF patients, there are currently no indicated therapies for MF patients with severe thrombocytopenia

PACRITINIB

- An investigational oral kinase inhibitor developed by CTI BioPharma Inc., with specificity for JAK2, IRAK1, and CSF1R
 - Mutations in these kinases have been shown to be directly related to the development of a variety of blood-related cancers, including myeloproliferative neoplasms, leukemia, and lymphoma
- The only therapy that has demonstrated evidence in reduction of spleen volumes in severely thrombocytopenic MF patients as evaluated in two prospective randomized phase III clinical trials
- A phase III confirmatory study (PACIFICA) is currently ongoing in MF patients with platelets $<50 \times 10^9/L$ with a top line readout anticipated in 2022

SUMMARY AND OUTLOOK

Transaction demonstrates our commitment and ability to build DRI's asset base

- On track to deliver on growth targets
 - US\$650 to US\$750 million five-year aggregate acquisition target
- Target sustainable growth in cash royalty receipts
- Acquisition pipeline remains strong and growing; transactions advancing with continual expansion of opportunities