

## Update Report

December 13, 2017

**RedHill Biopharma Ltd.: RedHill gives up migraine product (RHB-103). Upside remains, but the price target is decreased to NIS 2.27.**

Primary exchange: TASE

Secondary exchange (ADS/share 1:10):  
NASDAQ

Symbol: TASE, NASDAQ: RDHL

Sector: Biotechnology

Sub-sector: Drug Development

Stock price target: NIS 2.27

As of December 11, 2017

(source: TASE):

Closing price: NIS 1.59

Market cap: NIS 338.9M

# of shares: 212.7M

Stock performance (YTD): -60%

Daily-trading-vol. (12 months):  
NIS 1.3M

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### Company Overview

RedHill Biopharma Ltd. ("RedHill") is an Israeli publicly-traded specialty biopharmaceutical company focused on the development and commercialization of late clinical-stage drugs candidates. The company's primary focus is advanced clinical development and commercialization in the US of orally-administered, proprietary, small molecule drugs for the treatment of gastrointestinal and inflammatory diseases and cancer.

RedHill is currently promoting two gastrointestinal products and is advancing multiple clinical programs: three Phase III for gastrointestinal and inflammation indications; and four Phase II for multiple indications, including multiple myeloma, hepatocellular carcinoma, pancreatic cancer, and irritable bowel syndrome with diarrhea.

### Highlights

RedHill announced a 2017 year-end summary on December 5, 2017, with its main activities and key milestones expected in 2018:

- **Cost reduction plan.** Cash balance at the end of 2017 is expected to be approximately \$45 million, with no debt. A cost reduction plan is in place to gradually reduce the average quarterly cash burn rate in 2018 to approximately \$8.5 million.
- **Increasing resource optimization and focus on GI, including termination of RIZAPORT® license.** Given the Company's increasing focus on gastrointestinal (GI) diseases and in particular its two key Phase III GI programs with near-term data points and blockbuster potential for Crohn's disease and H. pylori infection, a notice has been provided to IntelGenx Corp. that **RedHill will terminate, effective January 6, 2018, its co-development and commercialization agreement for the non-core migraine drug product candidate, RIZAPORT®, for which a recent Incomplete Response Letter has been received from the FDA.**
- **Top-line results from the ongoing Phase III study with RHB-104 for Crohn's disease expected mid-2018.** Enrollment of all 331 subjects in the MAP US study has been completed and the last patient to reach the primary endpoint assessment (remission at week 26) is expected by May 2018.
- **Top-line results from the ongoing confirmatory Phase III study with TALICIA™ (RHB-105)2 for H. pylori infection are expected in H2/2018.** To date, 136 patients out of a planned total of 444 subjects have been enrolled. TALICIA™ was previously granted QIDP fast track designation from FDA.

### Analysis

- **RedHill's termination of its co-development and commercialization agreement for the non-core migraine drug product candidate, RIZAPORT® decreases company value.**
- This act will increase resource optimization and focus on GI sales platform and further development of other drugs in the pipeline.
- RHB-104 Crohn's disease - We assume that Redhill will achieve statistically significant results and achieve faster time-to-market. Top-line results from the ongoing Phase III study expected mid-2018.
- **Thus, we decrease company's equity value to \$137.0M/NIS 483.6M (previously, at \$145.4 million/NIS 511.8M in our Q3 2017 report) and the price target to the range of NIS 2.22-NIS 2.33- a mean of NIS 2.27, higher than current stock price.**

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