



COMPANY OVERVIEW

Apricus Biosciences, Inc. (APRI) is a biopharmaceutical company advancing innovative medicines in urology and rheumatology. Apricus has two product candidates currently in development. Vitaros™ is a product candidate in the United States for the treatment of erectile dysfunction, which Apricus in-licensed from Warner Chilcott Inc., now a subsidiary of Allergan. RayVa™ is our product candidate in Phase 2 development for the treatment of circulatory disorder Raynaud's Phenomenon, secondary to scleroderma, which Apricus owns worldwide.

EQUITY OVERVIEW

NSDQ: APRI

Market Cap: \$26.0m[†]

52 Week Range: \$0.86 - \$4.07[†]

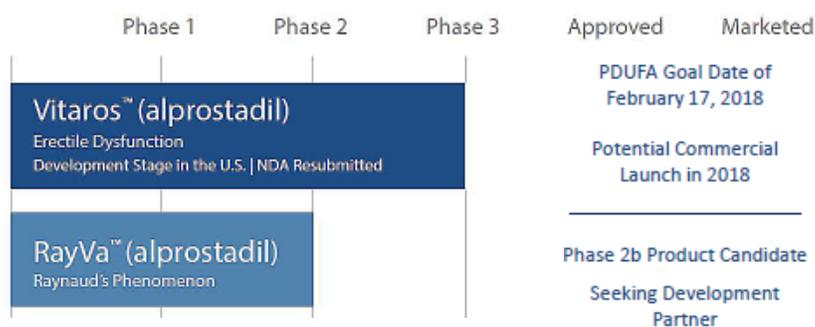
3 Month Average Volume: 426,406[†]

Common Shares Outstanding: 15.03m[†]

Cash Balance: \$7.8m^{††}

[†] As of 9/25/2017 ^{††} As of 6/30/2017. Does not include \$3.7m PIPE on 9/13/2017

DRUG DEVELOPMENT PIPELINE



ANALYST COVERAGE

H.C. Wainwright
 Roth Capital Partners
 See ThruEquity, LLC

POTENTIAL MILESTONES

- Vitaros NDA approval decision on or around February 17, 2018
- RayVa Partnership
- Vitaros Commercialization partnership in H1 2018
 - Allergan Opt-In
 - Out-License
 - Other Strategic Transaction
- Vitaros U.S. launch as early as H2 2018

INVESTOR CONTACT

Matthew Beck
 The Trout Group, LLC
 (646) 378-2933

INVESTMENT HIGHLIGHTS

Company:

- Specialty pharmaceutical company focused on unmet need in urology and rheumatology
- Novel, capital efficient pipeline with multiple potential value-creating milestones
- Balance sheet supports operations through 2018
- No debt

Vitaros:

- First and only topical treatment for erectile dysfunction
- Rapid onset of action (5-30 minutes)
- Studied in over 3,300 patients with favorable safety and efficacy profile
- Approved in 29 countries outside of US, where it is being commercialized by Ferring and its licensees
- US approval decision expected on or around February 17, 2018
- Allergan holds commercialization option in US
- Company estimates peak US revenue at \$350 million
- Broad IP position with potential exclusivity to 2032

RayVa:

- First in class topical treatment for Raynaud's phenomenon secondary to scleroderma
- No approved Raynaud's treatments in US
- Completed Phase 2a clinical trial
- Broad IP position with potential exclusivity to 2032
- May be eligible for priority review in the US