

Company Update

bluebird bio **Biotechnology**

years

US Equity Research

8 December 2020

Rating Price Target **BUY** US\$86.00 unchanged unchanged Price **BLUE-NASDAO** US\$45.40

Market Data

| 52-Week Range (US\$): | 38.95 - 99.36 |
|-----------------------|---------------|
| Market Cap (US\$M): | 3,114.2 |
| Shares Out. (M): | 66.4 |

| FYE Dec | 2018A | 2019A | 2020E | 2021E |
|-----------------------------|---------|---------|---------|--------|
| Revenue (US\$M) | 54.6 | 44.7 | 249.0 | 59.6 |
| EPS Adj&Dil (US\$) | (10.68) | (14.31) | (9.62) | (8.89) |
| Quarterly Revenue | Q1 | Q2 | Q3 | Q4 |
| 2018A | 16.0 | 7.9 | 11.5 | 19.2 |
| 2019A | 12.5 | 13.3 | 8.9 | 10.0 |
| 2020E | 21.9A | 198.9A | 19.3A | 8.9 |
| 2021E | 10.3 | 12.6 | 15.2 | 21.5 |
| Quarterly EPS Adj&Dil | Q1 | Q2 | Q3 | Q4 |
| 2018A | (2.31) | (2.91) | (2.73) | (2.72) |
| 2019A | (2.99) | (3.55) | (3.73) | (4.04) |
| 2020E | (3.64)A | (0.36)A | (2.94)A | (2.69) |
| 2021E | (2.27) | (2.20) | (2.21) | (2.21) |



NYSE Arca Biotechnology Index (BTK) (rebased)

Priced as of close of business 7 December 2020

Severe vaso-occlusive event resolution for up to two

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Key takeaways: 100% resolution of severe vaso-occlusive events (VOEs) between months 6 and 24 of follow-up was reported in Sickle Cell Disease patients treated with LentiGlobin (n=19). Additionally, patient-reported outcomes show a reduction in pain intensity after treatment, improving quality of life.

Resolution of severe VOEs for up to two years

Data as of August 20, 2020 show 100% resolution of severe vaso-occlusive events (VOEs) between months 6 and 24 of follow-up (n=19). Prior to treatment, these patients experienced a median of 3 severe VOEs, and resolution of these VOEs represents an important clinical benefit. After 6 or more months post-LentiGlobin treatment, median total hemoglobin is consistently >= 11 g/dL, and gene therapyderived hemoglobin levels stabilizes at median >= 40% with near pancellular expression (n=22). Hemolysis markers trended towards normalization after treatment, and all patients were able to stop regular blood transfusions by three months post-treatment and remain off transfusions as of the data cut-off. Safety remains generally consistent with known side effects of busulfan conditioning as well as underlying SCD.

Patient-reported outcomes show reduction in pain intensity

BLUE reported reduction in pain intensity at 12 months after treatment with LentiGlobin for SCD, using validated PROMIS-57 (0-10 scale) to collect patient-reported clinical outcomes. For five patients with baseline pain worse than population average (2.6), their mean score improved from 6 at baseline to 2.4 at month 12. For four patients with baseline pain better than or near population average at baseline, two reported improvement and two remained stable with a mean score of 2.3 at baseline and 0.8 at Month 12. These results support improvement in quality of life.

Maintain BUY, \$86 price target

We maintain our BUY rating and \$86 price target for bluebird bio. We are encouraged by the long-term follow-up data in SCD, and we also continue to expect FDA approval for LentiGlobin in B-thalassemia. Additionally, we look for revenues to ramp for ZYNTEGLO in Europe for B-thalassemia, beginning 2021.