UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
of the Securities Exchange Act of 1934

Date of Report: December 14, 2020

Commission File Number: 001-39307

Legend Biotech Corporation
(Exact Name of Registrant as Specified in its Charter)

2101 Cottontail Lane
Somerset, New Jersey 08873
(Address of principal executive office)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F:

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

Legend Biotech Announcement Regarding IND for LB1901

On December 14, 2020, Legend Biotech Corporation issued a press release announcing that the U.S. Food and Drug Administration (FDA) has cleared its Investigational New Drug (IND) application to evaluate LB1901 for the treatment of adults with relapsed or refractory T-cell lymphoma, which is attached to this Form 6-K as Exhibit 99.1.

EXHIBIT INDEX

Exhibit	Title
99.1	Press Release, dated December 14, 2020.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

LEGEND BIOTECH CORPORATION
(Registrant)

December 14, 2020

By: /s/ Ying Huang
Ying Huang, Ph.D.
Chief Executive Officer and Chief Financial Officer

Exhibit 99.1

Legend Biotech Announces FDA Clearance of the IND for LB1901, an Investigational Autologous Anti-CD4 CAR-T Therapy for Relapsed or Refractory T-Cell Lymphoma

SOMERSET, N.J., December 14, 2020 – Legend Biotech Corporation (NASDAQ: LEGN) ("Legend Biotech"), a global clinical-stage biopharmaceutical company engaged in the discovery and development of novel cell therapies for oncology and other indications, today announced that the U.S. Food and Drug Administration (FDA) has cleared its Investigational New Drug (IND) application to evaluate LB1901, the company's investigational autologous chimeric antigen receptor T-cell (CAR-T) therapy, for the treatment of adults with relapsed or refractory T-cell lymphoma (TCL). Under this IND, Legend will initiate a Phase 1 clinical study for LB1901 in the United States.

LB1901 is an investigational CAR-T product targeting CD4, which is a surface membrane glycoprotein uniformly expressed in a majority of TCL subtypes. A Phase 1, first-in-human, open-label, multicenter, multicohort clinical study will enroll patients with relapsed or refractory peripheral T-cell lymphoma (PTCL) or cutaneous T-cell lymphoma (CTCL) in the United States. The primary objectives of the study are to characterize the safety and tolerability of LB1901 and to determine the recommended Phase 2 dose.

TCL is a heterogeneous group of disorders accounting for less than 15 percent of Non-Hodgkin lymphoma cases in the US.[1,2] PTCL comprises subtypes which are uncommon and often aggressive, with a 5-year overall survival of 39% that varies by subtype.[3,4] Cutaneous T-cell lymphomas are a group of T-cell malignancies, which occur primarily in the skin.[5] Despite current treatment options, a substantial proportion of patients with PTCL or CTCL experiences relapse. A high unmet medical need remains for patients with relapsed or refractory PTCL and CTCL.

"The FDA's clearance of Legend's IND application for LB1901 is a milestone representative of the company's scientific expertise in cell therapy innovation," said Ying Huang, PhD, Chief Executive Officer and Chief Financial Officer of Legend Biotech. "We look forward to working with the investigators as we explore its potential in meeting the great unmet medical needs in the TCL population."

About Legend Biotech

Legend Biotech is a global clinical-stage biopharmaceutical company engaged in the discovery and development of novel cell therapies for oncology and other indications. Our team of more than 800 employees across the United States, China and Europe, along with our differentiated technology, global development, and manufacturing strategies and expertise, provide us with the strong potential to discover, develop, and manufacture cutting edge cell therapies for patients in need.

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Cautions Concerning Forward-Looking Statements

This information constitutes forward-looking statements relating to the business of Legend, including express or implied discussions regarding the clinical development of its product candidates and potential attributes and benefits of such product candidates. Such forward-looking statements reflect the current views of Legend's management regarding future events, and involve known and unknown risks, uncertainties and other factors that may cause actual results to be materially different from any future results, performance or achievements expressed or implied by such statements. In particular, Legend's expectations could be affected by, among other things, uncertainties involved in the development of new pharmaceutical products; unexpected clinical trial

results, including additional analysis of existing clinical data or unexpected new clinical data; unexpected regulatory actions or delays or government regulation generally; Legend's ability to obtain or maintain patent or other proprietary intellectual property protection; competition in general; government, industry, and general public pricing and other political pressures. Should one or more of these risks or uncertainties materialize, or should underlying assumptions prove incorrect, actual results may vary materially from those described herein as anticipated, believed, estimated or expected.

The safety and efficacy of the product candidates and/or uses under investigation have not been established. There is no guarantee that the product candidates will receive health authority approval or become commercially available in any country for the uses being investigated.

The information in this press release speaks only as of the date hereof. Legend assumes no duty to update the information to reflect subsequent developments. Readers should not rely upon the information on this page as current or accurate after its publication date.

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1 Scherer LD, Brenner MK, Mamonkln M. Chimeric antigen receptors for T-cell malignancies. *Frontiers in Oncology*. 2019 March;9(article 126):1-10.
2 American Cancer Society. Types of T-cell Lymphoma. Available at: https://www.cancer.org/cancer/non-hodgkin-lymphoma/about/t-cell-lymphoma.html. Accessed December 2020.
3 Casulo C, O'Connor O, Shustov A, Fanale M, Friedberg JW, Leonard JP, et al. T-cell lymphoma: Recent advances in characterization and new opportunities for treatment. *J Natl Cancer Inst*. 2017;109(2):1-9.
4 Abouyabis AN, Shenoy PJ, Sinha R, Flowers CR, Lechowicz MJ. A systematic review and meta-analysis of front-line anthracycline based chemotherapy regimens for peripheral T-cell lymphoma. *ISRN Hematol*. 2011;2011:623924.
5 Scarfo I, Frigault M, Maus M. CAR-based approaches to cutaneous T-cell lymphoma. *Frontiers in Oncology*. 2019;9(article 259):1-6.