



Our STN: BL 125643/725

**SUPPLEMENT APPROVAL**

June 26, 2025

Kite Pharma, Inc.  
Attention: Arjana Pradhan, PhD  
2400 Broadway  
Santa Monica, CA 90404

Dear Dr. Pradhan:

We have approved your request received March 21, 2025, to supplement your Biologics License Application (BLA) submitted under section 351(a) of the Public Health Service Act for axicabtagene ciloleucel [YESCARTA] to remove references to Risk Evaluation and Mitigation Strategies (REMS) from the United States Prescribing Information (USPI) and to revise Sections 2 (Dosage and Administration), 5 (Warnings and Precautions), 17 (Patient Counseling Information) of the US Prescribing Information and Medication Guide, to align with REMS elimination and streamline monitoring for patients. This supplement is in response to our March 7, 2025, REMS Modification Notification letter.

## **LABELING**

We hereby approve the draft content of labeling for the Package Insert submitted under amendment 1, dated June 9, 2025

## **CONTENT OF LABELING**

As soon as possible, but no later than 14 days from the date of this letter, please submit the final content of labeling (21 CFR 601.14) in Structured Product Labeling (SPL) format via the FDA automated drug registration and listing system, (eLIST) as described at <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>. Content of labeling must be identical to the Package Insert submitted on June 9, 2025. Information on submitting SPL files using eLIST may be found in the guidance for industry *SPL Standard for Content of Labeling Technical Qs and As* at <http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM072392.pdf>.

The SPL will be accessible via publicly available labeling repositories.

We request that the labeling approved today be available on your website within 10 days of receipt of this letter.

All final labeling should be submitted as Product Correspondence to this BLA, STN BL 125643/0 at the time of use and include implementation information on Form FDA 356h.

## **ADVERTISING AND PROMOTIONAL LABELING**

You may submit two draft copies of the proposed introductory advertising and promotional labeling with Form FDA 2253 to the Advertising and Promotional Labeling Branch at the following address:

Food and Drug Administration  
Center for Biologics Evaluation and Research  
Document Control Center  
10903 New Hampshire Ave.  
WO71–G112  
Silver Spring, MD 20993-0002

You must submit copies of your final advertising and promotional labeling at the time of initial dissemination or publication, accompanied by Form FDA 2253 (21 CFR 601.12(f)(4)).

All promotional claims must be consistent with and not contrary to approved labeling. You should not make a comparative promotional claim or claim of superiority over other products unless you have substantial evidence or substantial clinical experience to support such claims (21 CFR 202.1(e)(6)).

For each pending supplemental application for this BLA that includes proposed revised labeling, please submit an amendment to update the proposed revised labeling with the changes approved today.

We will include information contained in the above-referenced supplement in your BLA file.

Sincerely,

Asha Das, MD  
Acting Director  
Office of Clinical Evaluation  
Office of Therapeutic Products  
Center for Biologics Evaluation and Research