

NDA 208799/S-026

## SUPPLEMENT APPROVAL

Teva Pharmaceutical Industries Ltd. Attention: Lisa Carle Director Regulatory Affairs Labeling 145 Brandywine Parkway West Chester, PA 19380

Dear Ms. Carle:

Please refer to your Supplemental New Drug Application (sNDA) dated and received May 24, 2023, and your amendment, pursuant to section 505(b)(2) of the Federal Food, Drug, and Cosmetic Act (FDCA) for AIRDUO DIGIHALER (fluticasone propionate and salmeterol) inhalation powder.

This Prior Approval supplemental new drug application provides for revised carton for AirDuo Digihaler to improve understanding and to increase awareness around connecting to the Digihaler app.

## **APPROVAL & LABELING**

We have completed our review of this supplemental application, as amended. It is approved, effective on the date of this letter, for use as recommended in the enclosed agreed-upon labeling.

## **CARTON AND CONTAINER LABELING**

Submit final printed carton and container labeling that are identical to the enclosed carton and container labeling, as soon as they are available, but no more than 30 days after they are printed. Please submit these labeling electronically according to the guidance for industry *Providing Regulatory Submissions in Electronic Format* — Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications. For administrative purposes, designate this submission "**Product Correspondence – Final Printed Carton and Container Labeling for approved NDA 208799/S-026**." Approval of this submission by FDA is not required before the labeling is used.

## **REPORTING REQUIREMENTS**

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

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If you have any questions, call Emma Gimose, PharmD, BCPS, Regulatory Business Process Manager, at (240) 402 - 1681.

Sincerely,

{See appended electronic signature page}

David Lewis, Ph.D. Branch Chief, Branch II Division of Post-Marketing Activities I Office of Lifecycle Drug Products Office of Pharmaceutical Quality Center for Drug Evaluation and Research

Enclosure(s): Carton Labeling



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Digitally signed by David Lewis Date: 11/16/2023 05:12:45PM GUID: 508da72000029f287fa31e664741b577