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August 4, 2026

Company name: TAUNS Laboratories, Inc.
Representative: Masaki Nonaka, President and CEO
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Notice Concerning Marketing Authorization for the In Vitro Diagnostic

“ImunoAce SARS-CoV-2/Flu II”

TAUNS Laboratories, Inc. hereby announces that it has obtained marketing authorization in Japan on July 31, 2026 for “ImunoAce SARS-CoV-2/Flu II” (generic name: SARS coronavirus antigen kit/influenza virus kit; the “Product”), an in vitro diagnostic antigen test kit for SARS-CoV-2 and influenza virus.

The Product is an improved version of the Company’s existing product, “ImunoAce SARS-CoV-2/Flu.” The time required for result determination has been shortened from 15 minutes to 10 minutes, and the shelf life has been extended from 12 months after manufacture to 22 months after manufacture.

The Company will proceed with the necessary procedures for insurance coverage and preparations for launch in accordance with applicable laws and regulations.

The impact of this matter on the Company’s financial results is currently under review. The Company will promptly disclose any matters that require disclosure, including any revisions to its financial forecasts.

[Product Overview]

Generic name: SARS coronavirus antigen kit/influenza virus kit

Product name: ImunoAce SARS-CoV-2/Flu II

Date of marketing authorization: July 31, 2026

Marketing authorization number: 30800EZX00026000

Marketing authorization holder: TAUNS Laboratories, Inc.

This document has been prepared solely for the purpose of providing information regarding the Company’s business activities and is not intended to solicit investment or provide medical advice. Information contained herein is current as of the date of announcement and is subject to change without notice.

[Contact Information]

For healthcare professionals:

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