

to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

(Authority: 43 CFR 46.435 and 43 CFR 1610.2)

Kristina Kirby,
State Director (Acting).

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INTERNATIONAL TRADE COMMISSION

Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Mobile Cellular Communications Devices, DN 3835*; the Commission is soliciting comments on any public interest issues raised by the complaint or complainant's filing pursuant to the Commission's Rules of Practice and Procedure.

FOR FURTHER INFORMATION CONTACT: Lisa R. Barton, Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–2000. The public version of the complaint can be accessed on the Commission's Electronic Document Information System (EDIS) at <https://edis.usitc.gov>. For help accessing EDIS, please email EDIS3Help@usitc.gov.

General information concerning the Commission may also be obtained by accessing its internet server at United States International Trade Commission (USITC) at <https://www.usitc.gov>. The public record for this investigation may be viewed on the Commission's Electronic Document Information System (EDIS) at <https://edis.usitc.gov>. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Pantech Corporation on July 3, 2025. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation,

and the sale within the United States after importation of certain mobile cellular communications devices. The complaint names as respondents: OnePlus Technology (Shenzhen) Co., Ltd. of China; OnePlus USA Corp. of Irving, TX; Lenovo Group Ltd. of China; Lenovo (United States) Inc. of Morrisville, NC; Motorola Mobility LLC of Libertyville, IL; TCL Industries Holdings Co., Ltd. of China; TCL Electronics Holdings Ltd. of Hong Kong; TCL Communication Ltd. of Hong Kong; TCL Communication Technology Holdings Ltd. of China; TCL Mobile International Ltd. of Hong Kong; Huizhou TCL Mobile Communication Co., Ltd. of China; TCL Mobile Communication (HK) Company Ltd. of Hong Kong; Tinno USA, Inc. of Plano, TX; Shenzhen Tinno Mobile Technology Corp. of China; HMD Global of Finland; HMD Global OY of Finland; and HMD America, Inc. of Miami, FL. The complainant requests that the Commission issue a limited exclusion order, cease and desist orders and impose a bond upon respondents' alleged infringing articles during the 60-day Presidential review period pursuant to 19 U.S.C. 1337(j).

Proposed respondents, other interested parties, members of the public, and interested government agencies are invited to file comments on any public interest issues raised by the complaint or § 210.8(b) filing. Comments should address whether issuance of the relief specifically requested by the complainant in this investigation would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) explain how the articles potentially subject to the requested remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the requested remedial orders;

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and

desist order within a commercially reasonable time; and

(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business, eight calendar days after publication of this notice in the **Federal Register**. Complainant may file replies to any written submissions no later than three calendar days after the date on which any initial submissions were due, notwithstanding § 201.14(a) of the Commission's Rules of Practice and Procedure. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number ("Docket No. 3835") in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures¹). Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>.) No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice. Persons with questions regarding filing should contact the Secretary at EDIS3Help@usitc.gov.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which

¹ Handbook for Electronic Filing Procedures: https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf

confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel², solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.³

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.

Issued: July 3, 2025.

Lisa Barton,

Secretary to the Commission.

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1022 (Fourth Review)]

Refined Brown Aluminum Oxide From China

Determination

On the basis of the record¹ developed in the subject five-year review, the United States International Trade Commission (“Commission”) determines, pursuant to the Tariff Act of 1930 (“the Act”), that revocation of the antidumping duty order on refined brown aluminum oxide from China would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.

Background

The Commission instituted this review on February 3, 2025 (90 FR 8812, February 3, 2025) and determined on May 9, 2025, that it would conduct an

expedited review (90 FR 22113, May 23, 2025).

The Commission made this determination pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determination in this review on July 3, 2025. The views of the Commission are contained in USITC Publication 5645 (July 2025), entitled *Refined Brown Aluminum Oxide from China: Investigation No. 731–TA–1022 (Fourth Review)*.

By order of the Commission.

Issued: July 3, 2025.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2025–12665 Filed 7–7–25; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

William Washington, M.D.; Decision and Order

On January 22, 2025, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause (OSC) to William Washington, M.D., of Bellevue, Washington (Registrant). Request for Final Agency Action (RFAA), Exhibit (RFAAX) A, at 1, 3. The OSC proposed the revocation of Registrant's Certificate of Registration No. FW5625213, alleging that Registrant is “currently without authority to handle controlled substances in the State of Washington, the state in which [he is] registered with DEA.” *Id.* at 2 (citing 21 U.S.C. 824(a)(3)).

The OSC notified Registrant of his right to file a written request for hearing, and that if he failed to file such a request, he would be deemed to have waived his right to a hearing and be in default. *Id.* at 2 (citing 21 CFR 1301.43). Here, Registrant did not request a hearing. RFAA, at 2.¹ “A default, unless excused, shall be deemed to constitute a waiver of the registrant's/applicant's right to a hearing and an admission of the factual allegations of the [OSC].” 21 CFR 1301.43(e).

Further, “[i]n the event that a registrant . . . is deemed to be in default . . . DEA may then file a request for final agency action with the Administrator, along with a record to support its request. In such circumstances, the Administrator may

enter a default final order pursuant to [21 CFR] 1316.67.” *Id.* 1301.43(f)(1). Here, the Government has requested final agency action based on Registrant's default pursuant to 21 CFR 1301.43(c), (f), 1301.46. RFAA, at 1–2; *see also* 21 CFR 1316.67.

Findings of Fact

The Agency finds that, in light of Registrant's default, the factual allegations in the OSC are admitted. According to the OSC, on November 26, 2024, the Washington Medical Commission permanently revoked Registrant's Washington medical license. RFAAX A, at 1. According to Washington online records, of which the Agency takes official notice,² Registrant's Washington medical license remains revoked. Washington State Department of Health Provider Credential Search, <https://fortress.wa.gov/doh/providercredentialsearch/default.aspx> (last visited date of signature of this Order). Accordingly, the Agency finds that Registrant is not licensed to practice medicine in Washington, the state in which he is registered with DEA.³

Discussion

Pursuant to 21 U.S.C. 824(a)(3), the Attorney General may suspend or revoke a registration issued under 21 U.S.C. 823 “upon a finding that the registrant . . . has had his State license or registration suspended . . . [or] revoked . . . by competent State authority and is no longer authorized by State law to engage in the . . . dispensing of controlled substances.”

With respect to a practitioner, DEA has also long held that the possession of authority to dispense controlled substances under the laws of the state in which a practitioner engages in professional practice is a fundamental condition for obtaining and maintaining a practitioner's registration. *Gonzales v.*

² Under the Administrative Procedure Act, an agency “may take official notice of facts at any stage in a proceeding—even in the final decision.” United States Department of Justice, Attorney General's Manual on the Administrative Procedure Act 80 (1947) (Wm. W. Gaunt & Sons, Inc., Reprint 1979).

³ Pursuant to 5 U.S.C. 556(e), “[w]hen an agency decision rests on official notice of a material fact not appearing in the evidence in the record, a party is entitled, on timely request, to an opportunity to show the contrary.” The material fact here is that Registrant, as of the date of this decision, is not licensed to practice medicine in Washington. Accordingly, Registrant may dispute the Agency's finding by filing a properly supported motion for reconsideration of findings of fact within fifteen calendar days of the date of this Order. Any such motion and response shall be filed and served by email to the other party and to the DEA Office of the Administrator, Drug Enforcement Administration, at dea.addo.attorneys@dea.gov.

² All contract personnel will sign appropriate nondisclosure agreements.

³ Electronic Document Information System (EDIS): <https://edis.usitc.gov>

¹ The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

¹ Based on the Government's submissions in its RFAA dated February 24, 2025, the Agency finds that service of the OSC on Registrant was adequate. Specifically, a Form DEA–12, Receipt for Cash or Other Items, indicates that a DEA Diversion Investigator personally served Registrant with the OSC on January 22, 2025. RFAAX B, at 1.