# **DEPARTMENT OF JUSTICE**

### **Drug Enforcement Administration**

[Docket No. DEA-1045]

Importer of Controlled Substances Application: Biopharmaceutical Research Company LLC

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

SUMMARY: Biopharmaceutical Research Company LLC has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to SUPPLEMENTARY INFORMATION listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before August 29, 2022. Such persons may also file a written request for a hearing on the application on or before August 29, 2022.

**ADDRESSES:** The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to https://www.regulations.gov and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on https://www.regulations.gov. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on June 8, 2022, Biopharmaceutical Research Company LLC, 11045 Commercial Parkway, Castroville, California 95012–3209,

applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana	7360	1

The company plans to import narcotic raw material for bulk manufacture and analytical purposes. This notice does not constitute an evaluation or determination of the merits of the company's application. No other activity for this drug code is authorized for this registration. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

#### Kristi O'Malley,

Assistant Administrator.

[FR Doc. 2022–16204 Filed 7–27–22; 8:45 am]

BILLING CODE P

### **DEPARTMENT OF JUSTICE**

# **Drug Enforcement Administration**

[Docket No. DEA-1039]

Importer of Controlled Substances Application: AndersonBrecon Inc.

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

**SUMMARY:** AndersonBrecon Inc. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before August 29, 2022. Such persons may also file a written request for a hearing on the application on or before August 29, 2022.

Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <a href="https://www.regulations.gov">https://www.regulations.gov</a> and follow the online instructions at that site for submitting comments. Upon submission

of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on https://www.regulations.gov. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 6, 2022, AndersonBrecon Inc., 5775 Logistics Parkway, Rockford, Illinois 61109–3608, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	1

The company plans to import the listed controlled substance for clinical trial studies only. No other activity for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

# Kristi O'Malley,

Assistant Administrator.

[FR Doc. 2022–16205 Filed 7–27–22; 8:45 am] **BILLING CODE P** 

### **DEPARTMENT OF JUSTICE**

### **Drug Enforcement Administration**

[Docket No. DEA-1040]

Bulk Manufacturer of Controlled Substances Application: AMPAC Fine Chemicals Virginia, LLC

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.