

M.D., 76 FR 71,371 (2011), *pet. for rev. denied*, 481 F. App'x 826 (4th Cir. 2012); *Frederick Marsh Blanton, M.D.*, 43 FR 27,616, 27,617 (1978).

This rule derives from the text of two provisions of the CSA. First, Congress defined the term “practitioner” to mean “a physician . . . or other person licensed, registered, or otherwise permitted, by . . . the jurisdiction in which he practices . . . , to distribute, dispense, . . . [or] administer . . . a controlled substance in the course of professional practice.” 21 U.S.C. 802(21). Second, in setting the requirements for obtaining a practitioner’s registration, Congress directed that “[t]he Attorney General shall register practitioners . . . if the applicant is authorized to dispense . . . controlled substances under the laws of the State in which he practices.” 21 U.S.C. 823(f). Because Congress has clearly mandated that a practitioner possess state authority in order to be deemed a practitioner under the CSA, DEA has held repeatedly that revocation of a practitioner’s registration is the appropriate sanction whenever he is no longer authorized to dispense controlled substances under the laws of the state in which he practices. *See, e.g., James L. Hooper, M.D.*, 76 FR at 71,371–72; *Sheran Arden Yeates, M.D.*, 71 FR 39,130, 39,131 (2006); *Dominick A. Ricci, M.D.*, 58 FR 51,104, 51,105 (1993); *Bobby Watts, M.D.*, 53 FR 11,919, 11,920 (1988); *Frederick Marsh Blanton, M.D.*, 43 FR at 27,617.

According to New Mexico statute, “A person who . . . dispenses a controlled substance or who proposes to engage in the . . . dispensing of a controlled substance shall obtain a registration issued by the board in accordance with its regulations.” N.M. Stat. Ann. § 30–31–12(A) (West, current with 2020 Regular Session laws in effect through May 20, 2020). In turn, “dispense” means “to deliver a controlled substance to an ultimate user . . . pursuant to the lawful order of a practitioner.” N.M. Stat. Ann. § 30–31–2(H) (West, current with 2020 Regular Session laws in effect through May 20, 2020). Further, “practitioner” means a “physician . . . licensed or certified to prescribe and administer drugs that are subject to the Controlled Substances Act.” N.M. Stat. Ann. § 30–31–2(S) (West, current with 2020 Regular Session laws in effect through May 20, 2020).

Here, the undisputed evidence in the record is that Registrant’s license to practice medicine is revoked. As such, he is not a “practitioner,” a physician licensed or certified to prescribe a controlled substance according to New

Mexico law. Further, under New Mexico law, a person who dispenses a controlled substance in New Mexico must be registered. The undisputed record evidence is that Registrant’s New Mexico controlled substance license is expired.

For all of these reasons, Registrant lacks authority to practice medicine and prescribe controlled substances in New Mexico. Accordingly, I will order that Registrant’s DEA registration be revoked.

**Order**

Pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 824(a), I hereby revoke DEA Certificate of Registration No. FB0178194 issued to Mark D. Beale, M.D. This Order is effective September 10, 2020.

**Timothy J. Shea,**

*Acting Administrator.*

[FR Doc. 2020–17448 Filed 8–10–20; 8:45 am]

**BILLING CODE 4410–09–P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA–693]

**Bulk Manufacturer of Controlled Substances Application: National Center for Natural Products Research NIDA MPROJECT**

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

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 13, 2020.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on July 7, 2020, National Center for Natural Products Research National Institute of Drug Abuse (NIDA) MPROJECT, University of Mississippi, 135 Coy Waller Complex, P.O. Box 1848, University, Mississippi 36877–1848, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Marijuana Extract .....	7350	I

Controlled substance	Drug code	Schedule
Marijuana .....	7360	I
Tetrahydrocannabinols ..	7370	I

The company plans to bulk manufacture the above-listed controlled substances to make a supply of marihuana available to the National Institute of Drug Abuse (NIDA) for distribution to research investigators in support of the national research program needs. No other activities for these drug codes are authorized for this registration.

**William T. McDermott,**

*Assistant Administrator.*

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**BILLING CODE P**

**NATIONAL AERONAUTICS AND SPACE ADMINISTRATION**

[NOTICE: (20–067)]

**Name of Information Collection: NASA Enterprise Salesforce COVID–19 Contact Tracing**

**AGENCY:** National Aeronautics and Space Administration (NASA).

**ACTION:** Notice of information collection.

**SUMMARY:** The National Aeronautics and Space Administration, as part of its continuing effort to reduce paperwork and respondent burden, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections.

**DATES:** Comments are due by Monday, October 4, 2020.

**ADDRESSES:** All comments should be addressed to Roger Kantz, National Aeronautics and Space Administration, 300 E Street SW, Washington, DC 20546–0001.

**FOR FURTHER INFORMATION CONTACT:** Requests for additional information or copies of the information collection instrument(s) and instructions should be directed to Roger Kantz, NASA Clearance Officer, NASA Headquarters, 300 E Street SW, JF0000, Washington, DC 20546, 281–792–7885 or email [Travis.Kantz@nasa.gov](mailto:Travis.Kantz@nasa.gov).

**SUPPLEMENTARY INFORMATION:**

**I. Abstract**

The information will be used to determine whether NASA personnel have been exposed to the COVID–19 virus and to track and trace their interactions across the NASA community for identifying possible points of exposure.

Those individuals that volunteer, will be contacted by a NASA Contact Tracer, a to-be-designated NASA healthcare employee, and will be first read the privacy act, to understand their rights and what this information will be used for. Then they will be asked, orally, to confirm if they have symptoms or not (yes/no question). The Tracer will then enter that information, as well as the names, phone numbers, and emails of those they have been in contact with into the newly developed tracking and tracing digital application on NASA's enterprise solution, Salesforce.

While participation is voluntary, it is strongly encouraged as failure to provide the requested information may result in potential increased exposure of personnel to the virus.

NASA may share this information for authorized purposes with (1) private or other government health care providers or agencies for referral or special program responsibilities, and (2) other entities outlined under standard routine uses for all NASA systems of records.

## II. Methods of Collection

The voluntary data is collected orally by a NASA Contact Tracer, a to-be-designated NASA healthcare employee, who will then enter all the information into the newly developed tracking and tracing digital application on NASA's enterprise solution, Salesforce.

The ability for the Tracer to keep records through this electronic method will ensure higher rate of inclusion and assists in the efficiency of the stages of report processing by human subject matter analysts.

## III. Data

*Title:* NASA Enterprise Salesforce COVID-19 Contact Tracing.

*OMB Number:* 2700-0178.

*Type of Review:* New.

*Affected Public:* Individuals.

*Estimated Annual Number of Activities:* 5,400.

*Estimated Number of Respondents per Activity:* 1.

*Annual Responses:* 5,400.

*Estimated Time per Response:* 8 hours.

*Estimated Total Annual Burden Hours:* 43,200 hours.

*Estimated Total Annual Cost:* \$1,900,800.

## IV. Request for Comments

*Comments are invited on:* (1) Whether the proposed collection of information is necessary for the proper performance of the functions of NASA, including whether the information collected has practical utility; (2) the accuracy of NASA's estimate of the burden

(including hours and cost) of the proposed collection of information; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including automated collection techniques or the use of other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the request for OMB approval of this information collection. They will also become a matter of public record.

**Roger Kantz,**

*NASA PRA Clearance Officer.*

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**BILLING CODE 7510-13-P**

## NATIONAL SCIENCE FOUNDATION

### Agency Information Collection Activities: Comment Request; NSF Surveys To Measure Customer Service Satisfaction

**AGENCY:** National Science Foundation.

**ACTION:** Submission for OMB Review; Comment Request.

**SUMMARY:** The National Science Foundation (NSF) has submitted the following information collection requirement to OMB for review and clearance under the Paperwork Reduction Act of 1995. This is the second notice for public comment; the first was published in the **Federal Register**, and no comments were received. NSF is forwarding the proposed submission to the Office of Management and Budget (OMB) for clearance simultaneously with the publication of this second notice.

**DATES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAmain](http://www.reginfo.gov/public/do/PRAmain). Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

**FOR FURTHER INFORMATION CONTACT:** Suzanne H. Plimpton, Reports Clearance Officer, National Science Foundation, 2415 Eisenhower Avenue, Alexandria, VA 22314, or send email to [splimpto@nsf.gov](mailto:splimpto@nsf.gov). Individuals who use a telecommunications device for the deaf (TDD) may call the Federal Information Relay Service (FIRS) at 1-800-877-8339, which is accessible 24 hours a day, 7 days a week, 365 days a year (including federal holidays).

Copies of the submission may be obtained by calling 703-292-7556.

**SUPPLEMENTARY INFORMATION:** NSF may not conduct or sponsor a collection of information unless the collection of information displays a currently valid OMB control number and the agency informs potential persons who are to respond to the collection of information that such persons are not required to respond to the collection of information unless it displays a currently valid OMB control number.

*Title of Collection:* NSF Surveys to Measure Customer Service Satisfaction.

*OMB Number:* 3145-0157.

*Type of Request:* Renewal without change of a new information collection.

*Proposed project:* On September 11, 1993, President Clinton issued Executive Order 12862, "Setting Customer Service Standards," which calls for Federal agencies to provide service that matches or exceeds the best service available in the private sector. Section 1(b) of that order requires agencies to "survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services." The National Science Foundation (NSF) has an ongoing need to collect information from its customer community (primarily individuals and organizations engaged in science and engineering research and education) about the quality and kind of services it provides and use that information to help improve agency operations and services.

*Estimate of Burden:* The burden on the public will change according to the needs of each individual customer satisfaction survey; however, each survey is estimated to take approximately 30 minutes per response.

*Respondents:* Will vary among individuals or households; business or other for-profit; not-for-profit institutions; farms; federal government; state, local or tribal governments.

*Estimated Number of Responses per Survey:* This will vary by survey.

*Comments:* Comments are invited on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information shall have practical utility; (b) the accuracy of the Agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information on respondents, including through the use of automated collection techniques or other forms of information technology; and (d) ways to minimize the burden of the collection of information on those who are to