○ That using D&B's product is likely to allow a business to have its previously unreported commercial payment experiences added to its credit report;

○ That D&B will actively assist a business in adding its unreported commercial payment experiences to its credit report;

○ That using D&B's product is likely to help a business build or improve its credit report;

○ The ease with which information or payment experiences can be added to a business's credit report; and

○ That D&B's product is needed when it is not, and that a product will enable a prospective customer to have a ''complete'' file.

• Part I also features ancillary relief relating to the challenged conduct by prohibiting misrepresentations relating to what payment experiences customers can add, as well as to D&B's renewal and charging practices.

• Part II provides additional specific relief relating to D&B's renewal and charging practices for products covered under the Proposed Order, to make sure that D&B makes clear disclosures about renewals both before a customer subscribes and during the period of the subscription.

• Parts III and IV require D&B to make certain disclosures to potential customers of CreditBuilder products, so that those potential customers can make better informed decisions about whether to purchase the products.

• Part V sets out specific requirements for D&B to follow when a business disputes information that D&B reports about it. The requirements of this Part V apply generally and are not limited only to D&B customers.

• Part VI requires D&B to offer refunds (or partial refunds) to certain customers and former customers of CreditBuilder products. Refund or partial refund eligibility under the Proposed Order will depend on customers' specific circumstances and how they used or attempted to use their CreditBuilder products.

• Part VII requires D&B to send notices to all current customers of paid products covered under the Proposed Order that automatically renew.

Parts VIII through XII are reporting and compliance provisions. Part VIII mandates that D&B acknowledge receipt of the Proposed Order and, for three years, distribute the Proposed Order to certain employees and agents and secure acknowledgments from recipients of the Proposed Order. Part IX requires D&B to submit compliance reports to the FTC one year after the order's issuance and submit additional reports when certain events occur. Part X requires that, for 10 years, D&B creates certain records and retain them for at least 5 years. Part XI provides for the FTC's continued compliance monitoring of D&B's activity during the Proposed Order's effective dates. Part XII is a provision ''sunsetting'' the Proposed Order after 20 years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the Proposed Order. It is not intended to constitute an official interpretation of the complaint or Proposed Order, or to modify in any way the Proposed Order's terms.

By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2022–00938 Filed 1–18–22; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Patient Safety Organizations: Voluntary Relinquishment for the Theator, Inc. PSO

AGENCY: Agency for Healthcare Research and Quality (AHRQ), Department of Health and Human Services (HHS).

ACTION: Notice of delisting.

SUMMARY: The Patient Safety and Quality Improvement Final Rule (Patient Safety Rule) authorizes AHRQ, on behalf of the Secretary of HHS, to list as a patient safety organization (PSO) an entity that attests that it meets the statutory and regulatory requirements for listing. A PSO can be ''delisted'' by the Secretary if it is found to no longer meet the requirements of the Patient Safety and Quality Improvement Act of 2005 (Patient Safety Act) and Patient Safety Rule, when a PSO chooses to voluntarily relinquish its status as a PSO for any reason, or when a PSO's listing expires. AHRQ accepted a notification of proposed voluntary relinquishment from the Theator, Inc. PSO, PSO number P0218, of its status as a PSO, and has delisted the PSO accordingly.

DATES: The delisting was effective at 12:00 Midnight ET (2400) on December 22, 2021.

ADDRESSES: The directories for both listed and delisted PSOs are ongoing and reviewed weekly by AHRQ. Both directories can be accessed electronically at the following HHS website: *http://www.pso.ahrq.gov/listed.*

FOR FURTHER INFORMATION CONTACT: Cathryn Bach, Center for Quality Improvement and Patient Safety, AHRQ, 5600 Fishers Lane, MS 06N100B, Rockville, MD 20857; Telephone (toll free): (866) 403–3697; Telephone (local): (301) 427–1111; TTY (toll free): (866) 438–7231; TTY (local): (301) 427–1130; Email: *pso@ahrq.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Background

The Patient Safety Act, 42 U.S.C. 299b–21 to 299b–26, and the related Patient Safety Rule, 42 CFR part 3, published in the **Federal Register** on November 21, 2008 (73 FR 70732–70814), establish a framework by which individuals and entities that meet the definition of provider in the Patient Safety Rule may voluntarily report information to PSOs listed by AHRQ, on a privileged and confidential basis, for the aggregation and analysis of patient safety work product.

The Patient Safety Act authorizes the listing of PSOs, which are entities or component organizations whose mission and primary activity are to conduct activities to improve patient safety and the quality of health care delivery.

HHS issued the Patient Safety Rule to implement the Patient Safety Act. AHRQ administers the provisions of the Patient Safety Act and Patient Safety Rule relating to the listing and operation of PSOs. The Patient Safety Rule authorizes AHRQ to list as a PSO an entity that attests that it meets the statutory and regulatory requirements for listing. A PSO can be ''delisted'' if it is found to no longer meet the requirements of the Patient Safety Act and Patient Safety Rule, when a PSO chooses to voluntarily relinquish its status as a PSO for any reason, or when a PSO's listing expires. Section 3.108(d) of the Patient Safety Rule requires AHRQ to provide public notice when it removes an organization from the list of PSOs.

AHRQ has accepted a notification of proposed voluntary relinquishment from the Theator, Inc. PSO to voluntarily relinquish its status as a PSO. Accordingly, the Theator, Inc. PSO, P0218, was delisted effective at 12:00 Midnight ET (2400) on December 22, 2021.

More information on PSOs can be obtained through AHRQ's PSO website at *http://www.pso.ahrq.gov.*

Dated: January 12, 2022.

Marquita Cullom,

Associate Director.

[FR Doc. 2022–00906 Filed 1–18–22; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—DP22–001, Real-World Effectiveness of Structured Lifestyle Interventions in Preventing Type 2 Diabetes.

Date: March 23, 2022.

Time: 10:30 a.m.–6:00 p.m., EDT.

Place: Teleconference.

Agenda: To review and evaluate grant applications.

FOR FURTHER INFORMATION CONTACT: Jaya Raman Ph.D., Scientific Review Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway, Mailstop S107–8, Atlanta, Georgia 30341–3717, Telephone: (770) 488–6511; Email: *JRaman@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022–00868 Filed 1–18–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—DP22–002, Epidemiology of Lupus: Longitudinal Studies in Population-Based Cohorts.

Date: March 17, 2022.

Time: 11:00 a.m.–6:00 p.m., EDT.

Place: Teleconference.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Jaya Raman, Ph.D., Scientific Review Officer, National Center for Chronic Disease and Health Promotion, CDC, 4770 Buford Highway, Mailstop S107–8, Atlanta, Georgia 30341–3717, Telephone: (770) 488–6511, Email: *JRaman@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022–00867 Filed 1–18–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Docket No. CDC–2022–0003]

Draft Policy Statement for Biosafety Level 4 (BSL–4) and Animal BSL–4 (ABSL–4) Laboratory Verification; Notice of Availability

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of availability and comment.

SUMMARY: The Centers for Disease Control and Prevention (CDC) in the Department of Health and Human Services (HHS) announces the opening of a docket to obtain comment on a draft policy statement regarding Biosafety Level 4 (BSL–4)/Animal Biosafety Level 4 (ABSL–4) verification requirements. The policy statement, once finalized, will assist individuals and entities in verifying that the facility design parameters and operational procedures, including heating, ventilation, and air conditioning (HVAC) systems, in BSL–4 and/or ABLS–4 laboratories are functioning as intended to meet the biosafety sufficiency requirement in the HHS/CDC select agent regulations.

DATES: Submit written or electronic comments by March 21, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0003, by either of the following methods.

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Division of Select Agents and Toxins, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H21–7, Atlanta, Georgia 30329.

Instructions: All submissions received must include the Agency name and Docket No. CDC–2022–0003. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. *Do not send comments by email. CDC does not accept comments by email.*

Docket Access: For access to the docket to read background documents or comments received, or to download