Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Maria E. Davila-Bloom, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 758, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–7637, *davila-bloomm@extra.niddk.nih.gov.*

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; T2D Outcomes Ancillary Study.

Date: August 6, 2015.

Time: 11:00 a.m. to 1:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Robert Wellner, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 706, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, 301–594–4721, *rw175w@nih.gov.*

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Reengineering Organs P01.

Date: August 10, 2015.

Time: 11:00 a.m. to 1:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Thomas A. Tatham, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 760, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–3993, *tathamt@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: June 30, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–16503 Filed 7–6–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: Treatment of Acute and Chronic Neurological Injuries Involving Axonal Regeneration

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: This is notice, in accordance with 35 U.S.C. 209 and 37 CFR part 404, that the National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive patent license to practice the following inventions embodied in the following patent applications:

HHS Ref. No.: E–214–2012/0

Titled: ''Compositions and Methods for the Treatment of Central Nervous System Injury''

1. US Provisional Patent Application No.: 61/705,555 HHS Ref. No.: E–214–2012/0–US–01 Filing *Date:* September 25, 2012
2. PCT Patent Application No.: PCT/US2013/061693 HHS Ref. No.: E–214–2012/0–PCT–02 Filing *Date:* September 25, 2013
3. Australian Patent Application No.: 2013–32367 HHS Ref No.: E–214–2012/0–AU–03 Filing *Date:* September 25, 2013
4. European Patent Application No.: 13771750 HHS Ref. No.: E–214–2012/0–EP–04 Filing *Date:* September 25, 2013
5. U.S. Patent Application No.: 14/430,850 HHS Ref. No.: E–214–2012/0–US–06 Filing *Date:* September 25, 2013

to BioAxone Biosciences Incorporated (''BioAxone''), a company incorporated under the laws of the State of Delaware having an office in at least Cambridge, Massachusetts, U.S.A. The patent rights in these inventions have been assigned to the United States of America. BioAxone is seeking all worldwide territories for this license. The field of use may be limited to ''Treatment of human acute and chronic neurological injuries involving axonal regeneration, as monotherapy or in combination with other therapeutic drugs or medical devices''.

DATES: Only written comments and/or applications for a license which are received by the NIH Office of Technology Transfer on or before August 6, 2015 will be considered.

ADDRESSES: Requests for copies of the patent application, patents, inquiries, comments, and other materials relating to the contemplated exclusive license should be directed to: Cristina Thalhammer-Reyero, Ph.D., M.B.A., Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852–3804; Telephone: (301) 435–4507; Facsimile: (301) 402–0220; Email: *thalhamc@mail.nih.gov.* A signed confidentiality nondisclosure agreement will be required to receive copies of any patent applications or patents that have not been published or issued by the United States Patent and Trademark Office or the World Intellectual Property Organization.

SUPPLEMENTARY INFORMATION: This technology, and its corresponding patent applications, is directed to methods of treating or preventing spinal cord injury or a glial scar by administering an agent that reduces the amount or activity of a 4-sulfacted GalNAc in a chondroitin glycosaminoglycan chain, wherein said agent includes human enzyme, arylsulfatase B (ARSB). This technology, and its corresponding patent applications, is also directed to methods of increasing neuron growth, proliferation, or migration by administering an agent that reduces the amount or activity of a 4-sulfacted GalNAc in a chondroitin glycosaminoglycan chain, wherein said agent includes ARSB. This technology may be useful as a means to treat paralysis and motor defects induced by spinal cord injury, such as by promoting axonal regrowth.

The prospective exclusive license will be royalty bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR part 404. The prospective exclusive license may be granted unless within thirty (30) days from the date of this published notice, the NIH receives written evidence and argument that establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR part 404.

Properly filed competing applications for a license in response to this notice will be treated as objections to the contemplated license. Comments and objections submitted in response to this notice will not be made available for public inspection and, to the extent permitted by law, will not be released under the Freedom of Information Act, 5 U.S.C. 552.

Dated: June 30, 2015.

Richard U. Rodriguez,

Acting Director, Office of Technology Transfer, National Institutes of Health.

[FR Doc. 2015–16500 Filed 7–6–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Generic Clearance To Conduct Voluntary Customer/Partner Surveys (NLM)

SUMMARY: Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget

(OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on April 22, 2015, page 22542 and allowed 60-days for public comment. One public comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Library of Medicine (NLM), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments or request more information on the proposed project contact: David Sharlip, Office of Administrative and Management Analysis Services, National Library of Medicine, Building 38A, Room B2N12, 8600 Rockville Pike, Bethesda, MD 20894, or call non-toll-free number (301) 402–9680, or Email your request, including your address to: *sharlipd@mail.nih.gov* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Generic Clearance to Conduct Voluntary Customer/Partner Surveys (NLM), 0925–0476, Expiration Date 07/31/2015, EXTENSION, National Library of Medicine (NLM), National Institutes of Health (NIH).

Need and Use of Information Collection: E.O. 12962 directed agencies that provide significant services directly to the public to survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services. Additionally, since 1994, the NLM has been a ''Federal Reinvention Laboratory'' with a goal of improving its methods of delivering information to the public. An essential strategy in accomplishing reinvention goals is the ability to periodically receive input and feedback from customers about the design and quality of the services they receive. The NLM provides significant services directly to the public including health providers, researchers, universities, other federal agencies, state and local governments, and to others through a range of mechanisms, including publications, technical assistance, and Web sites. These services are primarily focused on health and medical information dissemination activities. The purpose of this submission is to obtain OMB's generic approval to continue to conduct satisfaction surveys of NLM's customers. The NLM will use the information provided by individuals and institutions to identify strengths and weaknesses in current services and to make improvements where feasible. The ability to periodically survey NLM's customers is essential to continually update and upgrade methods of providing high quality service.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 750.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Frequency of response	Average time per response (minutes/hour)	Total burden hours
General Public	1,000	1	20/60	333
Health Professionals	500	1	15/60	125
Librarians	500	1	20/60	167
Health Educators	500	1	15/60	125

Dated: June 29, 2015.

David Sharlip,

Project Clearance Liaison, NLM, NIH.

[FR Doc. 2015–16633 Filed 7–6–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health and Human Development; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in section 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. 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: National Institute of Child Health and Human Development Special Emphasis Panel.

Date: August 7, 2015.

Time: 2:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6100 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Sherry L. Dupere, Ph.D., Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, NIH, 6100 Executive Boulevard, Room 5B01, Bethesda, MD 20892–9304, (301) 451–3415, *duperes@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: June 30, 2015.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–16528 Filed 7–6–15; 8:45 am]

BILLING CODE 4140–01–P