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 will have practical utility; (b) the accuracy of the Agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology. Comments may be sent to Jeanne Jacobs, Regulations and Paperwork Management Branch, U.S. Department of Agriculture, Rural Development, STOP 0742, 1400 Independence Ave., SW., Washington, DC 20250–0742. All responses to this notice will be summarized and included in the request for OMB approval. All comments will also become a matter of public record.

Dated: June 7, 2010.

Tammye Treviño,

Administrator, Rural Housing Service.

[FR Doc. 2010–15063 Filed 6–21–10; 8:45 am]

BILLING CODE 3410–XV–P

DEPARTMENT OF AGRICULTURE

Agricultural Research Service

Notice of Intent To Grant Partially Exclusive License

AGENCY: Agricultural Research Service, USDA.

ACTION: Notice of intent.

SUMMARY: Notice is hereby given that the U.S. Department of Agriculture, Agricultural Research Service, intends to grant to KGK Synergize Inc. of London, Ontario, Canada, a partially exclusive license to U.S. Patent No. 6,987,125, "Compositions and Methods of Treating, Reducing and Preventing Cardiovascular Diseases and Disorders with Polymethoxyflavones," issued on January 17, 2006. This will be the second license granted for this invention. The Agricultural Research Service intends to grant no additional licenses.

DATES: Comments must be received on or before July 22, 2010.

ADDRESSES: Send comments to: USDA, ARS, Office of Technology Transfer, 5601 Sunnyside Avenue, Rm. 4–1174, Beltsville, Maryland 20705–5131.

FOR FURTHER INFORMATION CONTACT: June Blalock of the Office of Technology Transfer at the Beltsville address given above; telephone: 301–504–5989.

SUPPLEMENTARY INFORMATION: The Federal Government's patent rights in this invention are assigned to the United States of America, as represented by the Secretary of Agriculture. It is in the public interest to so license this invention as KGK Synergize Inc. of London, Ontario, Canada, has submitted a complete and sufficient application for a license. The prospective partially exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective partially exclusive license may be granted unless, within thirty (30) days from the date of this published notice, the Agricultural Research Service receives written evidence and argument which establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Richard J. Brenner,

Assistant Administrator.

[FR Doc. 2010–15049 Filed 6–21–10; 8:45 am]

BILLING CODE 3410–03–P

ARCHITECTURAL AND TRANSPORTATION BARRIERS COMPLIANCE BOARD

Medical Diagnostic Equipment Accessibility Standards

AGENCY: Architectural and Transportation Barriers Compliance Board.

ACTION: Notice of public information meeting.

SUMMARY: Section 4203 of the Patient Protection and Affordable Care Act (Pub. L. 111–148, 124 Stat. L. 119) amended the Rehabilitation Act of 1973 by adding Section 510 to the Rehabilitation Act. Section 510 of the Rehabilitation Act requires the Architectural and Transportation Barriers Compliance Board (Access Board), in consultation with the Food and Drug Administration, to issue accessibility standards for medical diagnostic equipment to ensure that such equipment is accessible to, and usable by, individuals with disabilities to the maximum extent possible. The Access Board will hold a public information meeting to discuss the accessibility needs of individuals with disabilities with respect to medical diagnostic equipment and existing guidance for designing accessible medical diagnostic equipment. The meeting will provide an opportunity for individuals with disabilities, health care providers, and medical diagnostic equipment manufacturers to provide information to assist the Access Board in establishing accessibility standards for medical diagnostic equipment.

DATES: The information meeting will be on Thursday, July 29, 2010 from 9 a.m. until 5 p.m.

ADDRESSES: The information meeting will be held at the Access Board's conference space, 1331 F Street, NW., suite 800, Washington, DC 20004–1111.

FOR FURTHER INFORMATION CONTACT: David Baquis, Office of Technical and Information Services, Architectural and Transportation Barriers Compliance Board, 1331 F Street, NW., suite 1000, Washington, DC 20004–1111. Telephone number: 202–272–0013 (voice); 202–272–0082 (TTY). Electronic mail address: *baquis@access-board.gov.*

SUPPLEMENTARY INFORMATION: Section 4203 of the Patient Protection and Affordable Care Act (Pub. L. 111–148, 124 Stat. L. 119) amended the Rehabilitation Act of 1973 by adding Section 510 to the Rehabilitation Act. Section 510 of the Rehabilitation Act requires the Architectural and Transportation Barriers Compliance Board (Access Board), in consultation with the Food and Drug Administration, to issue accessibility standards for medical diagnostic equipment to ensure that such equipment is accessible to, and usable by, individuals with disabilities to the maximum extent possible. The standards will address equipment used by health care professionals in, or in conjunction with, physician's offices, clinics, emergency rooms, hospitals, and other medical settings for diagnostic purposes. Examination tables and chairs, mammography equipment, x-ray machines and other radiological equipment, and weight scales are examples of the types of equipment that the accessibility standards will address. Section 510 of the Rehabilitation Act requires the Access Board to issue the standards by March 22, 2012, and to periodically review and update the standards.

The Access Board will hold a public information meeting on Thursday, July 29, 2010 to discuss the accessibility needs of individuals with disabilities with respect to medical diagnostic equipment and existing guidance for designing accessible medical diagnostic equipment. The meeting will provide an opportunity for individuals with