

Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), SGIP 2.0, Inc. (“SGIP 2.0”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the standards development organization and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the name and principal place of business of the standards development organization is SGIP 2.0, Inc., c/o Gesmer Updegrave LLP, Boston, MA. The nature and scope of SGIP 2.0’s standards development activities are: SGIP 2.0 is organized exclusively for charitable, religious, educational, literary, and scientific purposes, within the meaning of Section 501(c)(3) of the Internal Revenue Code of 1986, as amended (or the corresponding provision of any subsequent federal tax law), and the regulations currently or hereafter promulgated thereunder. In furtherance of such purposes, SGIP 2.0 is organized and will be operated primarily to continue the work of the unincorporated SmartGrid Interoperability Panel, by supporting the National Institute of Standards and Technology in fulfilling its responsibilities pursuant to the *Energy Independence and Security Act of 2007*, including but not limited to by (a) providing technical guidance and coordination to help facilitate standards development for smart grid interoperability; (b) identifying and specifying testing and certification requirements, including provision of the underlying rationale to assess achievement of interoperability using smart grid standards; (c) informing and educating smart grid industry stakeholders regarding smart grid interoperability and related benefits; (d) liaising with similar organizations in other countries to help establish global smart grid interoperability alignment; and (e) undertaking such other activities as may from time to time be appropriate to further the purposes and achieve the goals set forth above.

Patricia A. Brink,

Director of Civil Enforcement, Antitrust Division.

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DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—3D PDF Consortium, Inc.

Notice is hereby given that, on November 8, 2012, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. § 4301 *et seq.* (“the Act”), 3D Consortium, Inc. (“3D PDF”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Boeing Shared Services Group, Seattle, WA, has been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and 3D PDF intends to file additional written notifications disclosing all changes in membership.

On March 27, 2012, 3D PDF filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on April 20, 2012 (77 FR 23754).

The last notification was filed with the Department on August 20, 2012. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on September 14, 2012 (77 FR 56861).

Patricia A. Brink,

Director of Civil Enforcement, Antitrust Division.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0006]

Agency Information Collection Activities: Proposed Collection; Comments Requested: Application for Individual Manufacturing Quota for a Basic Class of Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine

ACTION: 60-Day Notice.

The Department of Justice (DOJ), Drug Enforcement Administration (DEA), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. Comments are encouraged and will be accepted until February 4, 2013. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments, especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Cathy Gallagher, Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate 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;
- Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and
- 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, e.g., permitting electronic submission of responses.

Overview of Information Collection 1117–0006

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Application for Individual Manufacturing Quota for a Basic Class of Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 189).

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the*