

Determinations of the Investigations of Certain Durum Wheat and Hard Red Spring Wheat from Canada” from Jeffrey May, Deputy Assistant Secretary, Import Administration, to James J. Jochum, Assistant Secretary for Import Administration, dated August 28, 2003, (“*Decision Memorandum*”), which is hereby adopted by this notice. Attached to this notice as an Appendix is a list of the issues which parties have raised and to which we have responded in the *Decision Memorandum*. Parties can find

a complete discussion of all issues raised in these investigations and the corresponding recommendations in this public memorandum which is on file in the CRU. In addition, a complete version of the Decision Memorandum can be accessed directly on the Internet at <http://www.ia.ita.doc.gov/frn/summary/list.htm> under the heading “Canada.” The paper copy and electronic version of the Decision Memorandum are identical in content.

**Suspension of Liquidation**

In accordance with section 705(c)(1)(B)(i) of the Act, we have calculated an individual net subsidy rate for each manufacturer of the subject merchandise. In accordance with sections 777A(e)(2)(B) and 705(c)(5)(A) of the Act, we have set the “all others” rate as CWB’s rate, because it is the only exporter/manufacturer investigated. We determine the total estimated net subsidy rate for the CWB and “all others” to be:

Exporter/Manufacturer	Net Subsidy Rate (Hard Red Spring Wheat)	Net Subsidy Rate (Durum Wheat)
Canadian Wheat Board .....	5.29 percent	5.29 percent
All Others .....	5.29 percent	5.29 percent

As a result of our *Preliminary Determinations* and pursuant to section 705(c)(1)(B)(ii) of the Act, we instructed the U.S. Bureau of Customs and Border Protection (“BCBP”) to suspend liquidation of all entries of durum wheat and hard red spring wheat from Canada which were entered or withdrawn from warehouse, for consumption on or after March 10, 2003, the date of the publication of the *Preliminary Determinations* in the **Federal Register**. In accordance with section 703(d) of the Act, we instructed the BCBP to discontinue the suspension of liquidation for subject merchandise for countervailing duty purposes entered on or after July 8, 2003, but to continue the suspension of liquidation of entries made from March 10, 2003, through July 7, 2003.

We will issue countervailing duty orders and reinstate the suspension of liquidation under section 706(a) of the Act if ITC issues final affirmative injury determinations, and will require a cash deposit of estimated countervailing duties for such entries of merchandise in the amounts indicated above. If the ITC determines that material injury, or threat of material injury, does not exist, these proceedings will be terminated and all estimated duties deposited or securities posted as a result of the suspension of liquidation will be refunded or canceled.

**ITC Notification**

In accordance with section 705(d) of the Act, we have notified the ITC of our determinations. In addition, we are making available to the ITC all non-privileged and non-proprietary information related to these investigations. We will allow the ITC access to all privileged and business proprietary information in our files,

provided the ITC confirms that it will not disclose such information, either publicly or under an Administrative Protective Order (“APO”), without the written consent of the Assistant Secretary for Import Administration.

**Return or Destruction of Proprietary Information**

In the event that the ITC issues final negative injury determination, this notice will serve as the only reminder to parties subject to an APO of their responsibility concerning the destruction of proprietary information disclosed under APO in accordance with 19 CFR 351.305(a)(3). Failure to comply is a violation of the APO.

These determinations are published pursuant to sections 703(f) and 777(i) of the Act.

Dated: August 28, 2003.

**James J. Jochum,**  
*Assistant Secretary for Import Administration.*

**APPENDIX**

**List of Comments and Issues in the Decision Memorandum**

*Comment 1:* The Department Should Treat the Government-Leased Railcars Differently from the Government-Owned Railcars.

*Comment 2:* The Provision of Government-Owned and Leased Railcars is Tied to Non-U.S. Markets.

*Comment 3:* The Provision of Rail Cars Does Not Result in an Indirect Subsidy to the CWB.

*Comment 4:* Countervailability of Subsidies Given to Third Party Service Providers.

*Comment 5:* The Governments’ Entrustment or Direction of the Railways to Provide Rail Service.

*Comment 6:* The Provision of Government-Owned and Leased Railcar Confers No Benefit.

*Comment 7:* Measurement of Benefit from the Government-Provided Railcars.

*Comment 8:* The Revenue Cap Does Confer a Benefit.

*Comment 9:* The Rail Freight Revenue Cap Does Not Provide a Financial Contribution.

*Comment 10:* The Department Should Determine That the Revenue Cap Does Not Provide a Financial Contribution Because It is Consistent With Market Principles.

*Comment 11:* The Benefit of the Revenue Cap Extends to All CWB Shipments, Including Shipments to the United States.

*Comment 12:* The Closure Fee for Grain Dependent Branch Lines Confers a Financial Contribution.

*Comment 13:* Impact of the Lending and Initial Payment Guarantees on the CWB’s Cost of Borrowing.

*Comment 14:* The Benchmark.

*Comment 15:* The Borrowing Guarantee is Tied to Non-U.S. Markets.

*Comment 16:* The Department’s Analysis of the Initial Payment Guarantee is Based on Incomplete and Inaccurate Data.

[FR Doc. 03-22662 Filed 9-4-03; 8:45 am]

**BILLING CODE 3510-DS-S**

**DEPARTMENT OF COMMERCE**

**Patent and Trademark Office**

**Deposit of Biological Materials**

**ACTION:** Proposed collection; comment request.

**SUMMARY:** The United States Patent and Trademark Office (USPTO), 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 the continuing and proposed information collection, as required by the Paperwork Reduction Act of 1995, Public Law 104-13 (44 U.S.C. 3506(c)(2)(A)).

**DATES:** Written comments must be submitted on or before November 4, 2003.

**ADDRESSES:** Direct all written comments to Susan K. Brown, Records Officer, Office of the Chief Information Officer, Office of Data Architecture and Services, Data Administration Division, 703-308-7400, U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, Virginia 22313, Attn: CPK 3 Suite 310; by e-mail at [susan.brown@uspto.gov](mailto:susan.brown@uspto.gov); or by facsimile at 703-308-7407.

**FOR FURTHER INFORMATION CONTACT:** Requests for additional information should be directed to the attention of Robert J. Spar, Director, Office of Patent Legal Administration, United States Patent and Trademark Office (USPTO), P.O. Box 1450, Alexandria, VA 22313-1450; by telephone at 703-308-5107; or by e-mail at [bob.spar@uspto.gov](mailto:bob.spar@uspto.gov).

**SUPPLEMENTARY INFORMATION:**

**I. Abstract**

The deposit of biological materials as part of a patent application is required by 35 U.S.C. 2(b)(2) and outlined in 37 CFR chapter 1, subpart G, 1.801-809. Every patent must contain a description of the invention sufficient to enable a person (knowledgeable in the relevant science) to make and use the invention

as specified by 35 U.S.C. 112. The term biological includes material that is capable of self-replication either directly or indirectly. When the invention involves a biological material, sometimes words alone cannot sufficiently describe how to make and use the invention in a reproducible or repeatable manner. In such cases, the required biological material must either be known and readily (and continually) available, or be deposited in a suitable depository to meet the enablement and written description requirements of 35 U.S.C. 112.

In cases where a novel microorganism is involved, the USPTO traditionally requires the deposit of a sample with a recognized patent depository in order to meet the above disclosure requirements. When a deposit is necessary, the USPTO collects information to determine whether the depositor is in compliance with the patent statute. This includes a statement proving notification to the interested public on where to obtain samples of the deposits. A viability statement showing that the biological material was tested by the depository, and is a viable or acceptable deposit, must also be submitted to the USPTO.

In order to meet and satisfy requirements for international patenting, all countries signing the Budapest Treaty must recognize the deposit of biological material with any International Depository Authority (IDA).

**II. Method of Collection**

By mail, facsimile, or hand delivery to the USPTO when the applicant or agent

files a patent application with the USPTO or submits subsequent papers during the prosecution of the application to the USPTO.

**III. Data**

*OMB Number:* 0651-0022.

*Form Number(s):* None.

*Type of Review:* Extension of a currently approved collection.

*Affected Public:* Individuals or households; business or other for-profit; not-for-profit institutions; and the Federal Government.

*Estimated Number of Respondents:* 3,500 responses per year for deposited materials and 0.25 per year for depository approval.

*Estimated Time Per Response:* The USPTO estimates that it will take approximately 1 hour per application for deposited materials and 5 hours per application for depository approval.

*Estimated Total Annual Respondent Burden Hours:* 3,501 hours per year.

*Estimated Total Annual Respondent Cost Burden:* \$105,315 per year to submit the information to the USPTO. Using the professional hourly rate of \$30 for a senior administrative assistant, the USPTO estimates \$105,000 per year for salary costs associated with collecting and submitting the necessary deposit information. Using the professional hourly rate of \$252 for associate attorneys in private firms, the USPTO estimates \$315 per year for salary costs associated with the average depository seeking approval to store biological material.

Item	Estimated time for response	Estimated annual responses	Estimated annual burden hours
Deposited Materials .....	1 hour .....	3,500	3,500
Depository Approval .....	5 hours .....	0.25	1.25
Total .....	.....	3,500	3,501

*Estimated Total Annual Non-hour Respondent Cost Burden:* \$979,010. There are no maintenance costs or filing fees associated with this information collection. There are, however, capital start-up and postage costs.

Depositories charge fees to depositors; all depositories charge about the same rates for their services. For example, the American Type Culture Collection (ATCC), one of the world's leading biological supply houses and recognized patent depositories, charges a one-time fee of \$1,150 per deposit for basic storage and informing, and a minimum of \$160 per deposit for viability testing,

depending upon the type of deposit being tested. Most deposits received from outside the United States require an import permit from the U.S. Department of Agriculture (USDA). Also required is a public Health Service (PHS) permit, available from the Centers for Disease Control and Prevention (CDC), for importation of agents infectious to humans. This permit application processing fee is \$150. The USPTO estimates that the total non-hour respondent cost burden in the form of capital start-up costs amounts to \$1,460.

In addition, this collection does have postage costs. Biological deposits are

generally shipped to the depository Domestic Overnight by Federal Express (FedEx) and, since depositors are urged to supply frozen or freeze dried material, it must be packed in dry ice, according to a representative from the Patent Department at ATCC. Dry ice itself is considered dangerous goods and requires special packaging. Additional FedEx special handling charges of \$60 per shipment apply for temperature-sensitive biological material and also for the dry ice. An average cost for shipping by FedEx Domestic Overnight is estimated to be \$75. If the shipment requires pick-up by FedEx, there is an

additional charge of \$2.50. Special packaging is also required for these shipments. According to DG Supplies Inc., a supplier of infectious and diagnostic goods packaging, frozen infectious shippers are estimated to cost \$141.80 per package for specimen

shipments requiring refrigeration or dry ice. Therefore, postage costs average \$279.30 per shipment, for a total cost to all the respondents of \$977,550. The postage cost for a depository seeking recognition is estimated to be \$3.85, sent to the USPTO by priority mail

through the United States Postal Service. Therefore, the USPTO estimates that the total non-hour respondent cost burden in the form of postage costs amounts to \$977,551.

Item	Responses (a)	Postage costs (\$) (b)	Total non-hour cost burden (a) × (b)
Deposited Materials .....	3,500	\$279.30	\$977,550.00
Depository Approval .....	0.25	3.85	1.00
Total .....	3,501	.....	977,551.00

The USPTO estimates that the total non-hour respondent cost burden for this collection in the form of capital start-up costs (\$1,460) and postage costs (\$977,551) amounts to \$979,011.

**IV. Request for 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 (including hours and cost) of the proposed collection of information; (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 respondents, including through the use of automated collection techniques or other forms of information technology.

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

Dated: August 29, 2003.

**Susan K. Brown,**

*Records Officer, USPTO, Office of the Chief Information Officer, Office of Data Architecture and Services, Data Administration Division.*

[FR Doc. 03-22612 Filed 9-4-03; 8:45 am]

**BILLING CODE 3510-16-P**

**COMMITTEE FOR THE IMPLEMENTATION OF TEXTILE AGREEMENTS**

**Adjustment of Import Limits for Certain Cotton, Wool and Man-Made Fiber Textile Products Produced or Manufactured in Malaysia**

August 29, 2003.

**AGENCY:** Committee for the Implementation of Textile Agreements (CITA).

**ACTION:** Issuing a directive to the Commissioner, Bureau of Customs and Border Protection adjusting limits.

**EFFECTIVE DATE:** September 5, 2003.

**FOR FURTHER INFORMATION CONTACT:** Ross Arnold, International Trade Specialist, Office of Textiles and Apparel, U.S. Department of Commerce, (202) 482-4212. For information on the quota status of these limits, refer to the Quota Status Reports posted on the bulletin boards of each Customs port, call (202) 927-5850, or refer to the Bureau of Customs and Border Protection website at <http://www.customs.gov>. For information on embargoes and quota reopenings, refer to the Office of Textiles and Apparel website at <http://otexa.ita.doc.gov>.

**SUPPLEMENTARY INFORMATION:**

**Authority:** Section 204 of the Agricultural Act of 1956, as amended (7 U.S.C. 1854); Executive Order 11651 of March 3, 1972, as amended.

The current limits for certain categories are being adjusted, variously, for swing, special shift and carryover.

A description of the textile and apparel categories in terms of HTS numbers is available in the **CORRELATION:** Textile and Apparel Categories with the Harmonized Tariff Schedule of the United States (see **Federal Register** notice 68 FR 1599, published on January 13, 2003). Also

see 67 FR 63896, published on October 16 2002.

**James C. Leonard III,**

*Chairman, Committee for the Implementation of Textile Agreements.*

Committee for the Implementation of Textile Agreements

August 29, 2003.

Commissioner,  
*Bureau of Customs and Border Protection, Washington, DC 20229.*

Dear Commissioner: This directive amends, but does not cancel, the directive issued to you on October 9, 2002, by the Chairman, Committee for the Implementation of Textile Agreements. That directive concerns imports of certain cotton, wool and man-made fiber textiles and textile products and silk blend and other vegetable fiber apparel, produced or manufactured in Malaysia and exported during the twelve-month period which began on January 1, 2003 and extends through December 31, 2003.

Effective on September 5, 2003, you are directed to adjust the limits for the following categories, as provided for under the Uruguay Round Agreement on Textiles and Clothing:

Category	Adjusted twelve-month limit <sup>1</sup>
Other specific limits	
333/334/335 .....	506,699 dozen of which not more than 263,719 dozen shall be in Category 333.
342/642 .....	762,634 dozen.
345 .....	296,802 dozen.
634/635 .....	1,451,092 dozen.
645/646 .....	513,885 dozen.
647/648 .....	3,039,496 dozen of which not more than 2,212,459 dozen shall be in Category 647-K <sup>2</sup> and not more than 2,212,459 dozen shall be in Category 648-K <sup>3</sup>

<sup>1</sup> The limits have not been adjusted to account for any imports exported after December 31, 2002.