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Dated at King of Prussia, Pennsylvania this 24th day of January, 2005.

For the Nuclear Regulatory Commission.

James P. Dwyer,

Chief, Commercial and R&D Branch, Division of Nuclear Materials Safety, Region I.

[FR Doc. 05-1685 Filed 1-28-05; 8:45 am]

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NUCLEAR REGULATORY COMMISSION

[Docket No. 030-36602]

Notice of Availability of Environmental Assessment and Finding of No Significant Impact for License Amendment for SWATCH Group(U.S.), Inc.'s Facility in Lancaster, PA

AGENCY: Nuclear Regulatory Commission.

ACTION: Notice of availability.

FOR FURTHER INFORMATION CONTACT:

Marjorie McLaughlin, Decommissioning Branch, Division of Nuclear Materials Safety, Region I, 475 Allendale Road, King of Prussia, Pennsylvania, 19406, telephone (610) 337-5240, fax (610) 337-5269; or by email: mmm3@nrc.gov.

SUPPLEMENTARY INFORMATION:

I. Introduction

The Nuclear Regulatory Commission (NRC) is issuing a license amendment to Swatch Group (U.S.), Inc. for Materials License No. 29-30923-01, to authorize release of its facility in Lancaster, Pennsylvania for unrestricted use. NRC has prepared an Environmental Assessment (EA) in support of this action in accordance with the requirements of 10 CFR Part 51. Based on the EA, the NRC has concluded that a Finding of No Significant Impact (FONSI) is appropriate. The amendment will be issued following the publication of this Notice.

II. EA Summary

The purpose of the action is to authorize the release of the licensee's Lancaster, Pennsylvania, facility for unrestricted use. Swatch Group (U.S.), Inc. was authorized by NRC from August, 1986, to use radioactive materials for manufacturing and

distribution purposes at the site. On August 16, 2004, Swatch Group (U.S.), Inc. requested that NRC release the facility for unrestricted use. Swatch Group (U.S.), Inc. has conducted surveys of the facility and provided information to the NRC to demonstrate that the site meets the license termination criteria in Subpart E of 10 CFR Part 20 for unrestricted use.

The NRC staff has prepared an EA in support of the license amendment. The facility was remediated and surveyed prior to the licensee requesting the license amendment. The NRC staff has reviewed the information and final status survey submitted by Swatch Group (U.S.), Inc. Based on its review, the staff has determined that there are no additional remediation activities necessary to complete the proposed action. Therefore, the staff considered the impact of the residual radioactivity at the facility and concluded that since the residual radioactivity meets the requirements in Subpart E of 10 CFR Part 20, a Finding of No Significant Impact is appropriate.

III. Finding of No Significant Impact

The staff has prepared the EA (summarized above) in support of the license amendment to release the facility for unrestricted use. The NRC staff has evaluated Swatch Group (U.S.), Inc.'s request and the results of the surveys and has concluded that the completed action complies with the criteria in Subpart E of 10 CFR Part 20. The staff has found that the environmental impacts from the action are bounded by the impacts evaluated by NUREG-1496, Volumes 1-3, "Generic Environmental Impact Statement in Support of Rulemaking on Radiological Criteria for License Termination of NRC-Licensed Facilities" (ML042310492, ML042320379, and ML042330385). On the basis of the EA, the NRC has concluded that the environmental impacts from the action are expected to be insignificant and has determined not to prepare an environmental impact statement for the action.

IV. Further Information

Documents related to this action, including the application for the license amendment and supporting documentation, are available electronically at the NRC's Electronic Reading Room at <http://www.nrc.gov/reading-rm/adams.html>. From this site, you can access the NRC's Agencywide Document Access and Management System (ADAMS), which provides text and image files of NRC's public documents. The ADAMS accession

numbers for the documents related to this Notice are: The Environmental Assessment (ML043410211), the letter dated August 16, 2004, requesting amendment of the license (ML042680179), the Final Status Survey, dated September 9, 2004 (ML042670407), additional information submitted October 19, 2004 containing survey maps (ML043010357), and a facsimile dated November 15, 2004 containing radwaste shipping papers (ML043340152). Please note that on October 25, 2004, the NRC terminated public access to ADAMS and initiated an additional security review of publicly available documents to ensure that potentially sensitive information is removed from the ADAMS database accessible through the NRC's web site. Interested members of the public may obtain copies of the referenced documents for review and/or copying by contacting the Public Document Room pending resumption of public access to ADAMS. The NRC Public Documents Room is located at NRC Headquarters in Rockville, MD, and can be contacted at (800) 397-4209 or (301) 415-4737, or by e-mail to pdr@nrc.gov. The PDR reproduction contractor will copy documents for a fee. The PDR is open from 7:45 a.m. to 4:15 p.m., Monday through Friday, except on Federal holidays.

Dated at King of Prussia, Pennsylvania this 24th day of January, 2005.

For the Nuclear Regulatory Commission.

James Dwyer,

Chief, Commercial & R&D Branch, Division of Nuclear Materials Safety, Region I.

[FR Doc. 05-1686 Filed 1-28-05; 8:45 am]

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NUCLEAR REGULATORY COMMISSION

Advisory Committee on Reactor Safeguards; Subcommittee Meeting on Thermal-Hydraulic Phenomena; Notice of Meeting

The ACRS Subcommittee on Thermal-Hydraulic Phenomena will hold a meeting on February 15, 2005, Room T-2B3, 11545 Rockville Pike, Rockville, Maryland.

The agenda for the subject meeting shall be as follows:

Tuesday, February 15, 2005-8:30 a.m. until the conclusion of business.

The Subcommittee will continue review of the development of the TRACE thermal-hydraulic computer code. The Subcommittee will hear presentations by and hold discussions with representatives of the NRC staff and their contractors regarding this

matter. The Subcommittee will gather information, analyze relevant issues and facts, and formulate proposed positions and actions, as appropriate, for deliberation by the full Committee.

Members of the public desiring to provide oral statements and/or written comments should notify the Designated Federal Official, Mr. Ralph Caruso (Telephone: 301-415-8065) five days prior to the meeting, if possible, so that appropriate arrangements can be made. Electronic recordings will be permitted.

Further information regarding this meeting can be obtained by contacting the Designated Federal Official between 7:30 a.m. and 4:15 p.m. (e.t.). Persons planning to attend this meeting are urged to contact the above named individual at least two working days prior to the meeting to be advised of any potential changes to the agenda.

Dated: January 25, 2005.

John H. Flack,

Acting Branch Chief, ACRS/ACNW.

[FR Doc. 05-1688 Filed 1-28-05; 8:45 am]

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RAILROAD RETIREMENT BOARD

Proposed Collection; Comment Request

SUMMARY: In accordance with the requirement of section 3506 (c)(2)(A) of the Paperwork Reduction Act of 1995 which provides opportunity for public comment on new or revised data collections, the Railroad Retirement Board (RRB) will publish periodic summaries of proposed data collections.

Comments are invited on: (a) Whether the proposed information collection is necessary for the proper performance of the functions of the agency, including whether the information has practical utility; (b) the accuracy of the RRB's estimate of the burden of the collection of the information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden related to the collection of information on

respondents, including the use of automated collection techniques or other forms of information technology.

Title and Purpose of Information Collection: Medical Reports; OMB 3220-0038.

Under Sections 2(a)(1)(iv), 2(a)(2) and 2(a)(3) of the Railroad Retirement Act (RRA), annuities are payable to qualified railroad employees whose physical or mental condition is such that they are unable to (1) work in their regular occupation (occupational disability); or (2) work at all (permanent total disability). The requirements for establishment of disability and proof of continuance of disability are prescribed in 20 CFR part 220.

Under sections 2(c)(1)(ii)(c) and 2(d)(1)(ii) of the RRA, annuities are also payable to qualified spouses and widow(ers), respectively, who have a qualified child who is under a disability which began before age 22. Annuities are also payable to surviving children on the basis of disability under section 2(d)(1)(iii)(C) if the child's disability began before age 22 and to widow(ers) on the basis of disability under section 2(d)(1) (i)(B). To meet the disability standard, the RRA provides that individuals must have a permanent physical or mental condition such that they are unable to engage in any regular employment.

Under section 2(d)(1)(v) of the RRA, annuities are also payable to remarried and surviving divorced spouses on the basis of, *inter alia*, disability or having a qualified disabled child in care. However, the disability standard in these cases is that found in the Social Security Act. That is, individuals must be able to engage in any substantial gainful activity by reason of any medically determinable physical or mental impairment. The RRB also determines entitlement to a period of early disability and early Medicare entitlement for qualified claimants in accordance with Section 216 of the Social Security Act.

When making disability determinations, the RRB needs evidence

from acceptable medical sources. The RRB currently utilizes Forms G-3EMP, Report of Medical Condition by Employer; G-250, Medical Assessment; G-250a, Medical Assessment of Residual Functional Capacity; G-260, Report of Seizure Disorder; RL-11b, Disclosure of Hospital Medical Records; and RL-11d, Disclosure of Medical Records from a State Agency; to obtain the necessary medical evidence.

The RRB proposes significant changes to the information collection. The primary change is to add proposed Form G-197, Authorization to Release Medical Information, to the information collection. Proposed Form G-197 will be the standard Health Insurance Portability and Accountability Act (HIPAA) compliant release form used by the RRB to obtain consent for the release of medical evidence under the RRA. The RRB also proposes to revise, renumber, and rename current Form G-250 to proposed Form RL-250, Request for Medical Assessment. Currently, Form G-250, requests a narrative report, copies of office records of the claimant's treatment, requests completion of Form G-250a, and includes a consent statement that the claimant must sign. Proposed Form RL-250 will not request the narrative report nor contain a consent statement. A new Form G-250, titled Medical Assessment is proposed. It is intended to provide more complete information while being more user-friendly by formatting responses into a question and answer format by body system being evaluated. Forms G-3EMP, and RL-11b are being revised to delete the consent portions from the versions currently in use. Minor editorial changes are proposed to Form G-250a which will continue to be used by agency hearings officers as a means to clarify, when necessary, information previously received or to obtain precise information needed to make a residual functional capacity determination. No changes are proposed to Form G-260. Completion is voluntary. One response is requested of each respondent.

ESTIMATE OF RESPONDENT BURDEN

Form No.	Annual response	Time (min)	Burden (hours)
G-3EMP	600	10	100
G-197	6,000	10	1,000
G-250	11,950	30	5,975
G-250a	50	20	17
G-260	100	25	42
RL-11b	5,000	10	833
RL-250	11,950	10	1,992
Total	35,900	9,501