

companies that enter the market each year from 1,000 to 1,985. This, too, is based on 1999 Statistical Abstract data. Thus, the current total of affected firms consists of approximately 47,904 established and new companies.

Accordingly, staff estimates total industry hours to comply with the MTOR is 2,752,500 hours [(45,919 X 50 hours) + (1,985 X 230 hours)].

This is a conservative estimate. Arguably much of the estimated time burden for disclosure-related compliance would be incurred even absent the Rule. Representatives from industry trade associations and other knowledgeable individuals have consistently stated that compliance with the Rule is widely regarded by direct marketers as being good business practice. The Rule's notification requirements would be voluntarily initiated by most merchants to meet consumer expectations regarding timely shipment, notification of delay, and prompt and full refunds. Providing consumers with notice about the status of their orders fosters consumer loyalty and encourages repeat purchases, which are important to direct marketers' success. Thus, it appears that much of the time and expense associated with Rule compliance may not constitute "burden" under the PRA³ although the above estimates account for it as such.

In estimating PRA burden, staff considered "the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency." 5 CFR 1320.3(b)(1). This includes "developing, acquiring, installing, and utilizing technology and systems for the purpose of disclosing and providing information." 5 CFR 1320.3(b)(1)(iv). Although not expressly stated in the OMB regulation implementing the PRA, the definition of burden arguably includes upgrading and maintaining computer and other systems used to comply with a rule's requirements. Conversely, to the extent that these systems are used in the ordinary course of business independent of the Rule, their associated upkeep would fall outside the realm of PRA "burden."

The mail order industry has been subject to the basic provisions of the Rule since 1976 and the telephone order industry since 1994. Thus, businesses have had several years (and some have had decades) to integrate compliance systems into their business procedures. Since 1997 many businesses have

upgraded the information management systems they need, in part, to comply with the Rule, and to more effectively track orders. These upgrades, however, mostly were needed to deal with growing consumer demand for merchandise resulting, in part, from increased public acceptance of making purchases over the telephone and, more recently, the Internet.

Accordingly, most companies now maintain records and provide updated order information of the kind required by the Rule in their ordinary course of business. Nevertheless, staff conservatively assumes that the time existing and new companies devote to compliance with the Rule remains the same as in 1997.

Estimated labor costs: \$31,136,000, rounded to the nearest thousand.

Labor costs are derived by applying appropriate hourly cost figures to the burden hours described above. According to the 1999 Statistical Abstract, average payroll for "non-store catalogue and mail order houses" and "non-store direct selling establishments" rose \$0.322 per hour per year between 1991 and 1996. In 1996, average payroll was \$10.34 per hour. Assuming average payroll continued to increase \$0.322 per hour per year, in 1999 average payroll would have reached \$11.31 per hour. Because the bulk of the burden of complying with the MTOR is borne by clerical personnel, staff believes that the average hourly payroll figure for non-store catalogue and mail order houses and non-store direct selling establishments is an appropriate measure of a direct marketer's average labor cost to comply with the Rule. Thus, the total annual labor cost to new and established businesses in 1999 for Rule compliance is approximately \$31,136,000 (2,753,000 hours x \$11.31/hr.). Relative to direct industry sales, this total is negligible.⁴

Estimated annual non-labor cost burden: \$0 or minimal.

The applicable requirements impose minimal start-up costs, as businesses subject to the Rule generally have or obtain necessary equipment for other business purposes, i.e., inventory and order management, and customer relations. For the same reason, staff anticipates printing and copying costs to be minimal, especially given that telephone order merchants have increasingly turned to electronic

communications to notify consumers of delay and to provide cancellation options. Staff believes that the above requirements necessitate ongoing, regular training so that covered entities stay current and have a clear understanding of federal mandates. This training, however, would be a small portion of and subsumed within the ordinary training that employees receive apart from that associated with the information collected under the Rule.

John D. Graubert,

Acting General Counsel.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control And Prevention

[60Day-01-07]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506 (c)(2)(A) of the Paperwork reduction Act of 1995, the Center for Disease Control and Prevention is providing opportunity for public comment on proposed data collection projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639-7090.

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 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 for other forms of information technology. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project

National Exposure Registry—
Extension—(OMB No. 0923-0006)

³ Under the OMB regulation implementing the PRA, burden is defined to exclude any effort that would be expended regardless of any regulatory requirement. 5 CFR 1320.3(b)(2).

⁴ Projecting sales for "non-store catalogue and mail order houses" and "non-store direct selling establishments" (according to the 1999 Statistical Abstract) to all merchants subject to the MTOR, staff estimates that direct sales to consumers in 1999 would have been \$109.45 billion. Thus, the labor cost of compliance by existing and new businesses in 1999 would have amounted to less than .03% of sales.

Agency for Toxic Substances and Disease Registry (ATSDR), Centers for Disease Control and Prevention (CDC). The Agency for Toxic Substances and Disease Registry (ATSDR) is mandated pursuant to the 1980 Comprehensive Environmental Response Compensation and Liability Act (CERCLA) and its 1986 Amendments, the Superfund Amendments and Re-authorization Act (SARA), to establish and maintain a national registry of persons who have been exposed to hazardous substances in the environment and a national registry of persons with illnesses or health problems resulting from such exposure. ATSDR created the National Exposure Registry (NER) as a result of this legislation in an effort to provide scientific information about potential adverse health effects people develop as a result of low-level, long-term exposure to hazardous substances.

The National Exposure Registry is a program that collects, maintains, and

analyzes information obtained from participants (called registrants) whose exposure to selected toxic substances at specific geographic areas in the United States was documented. Relevant health data and demographic information are also included in the NER database. The NER databases furnish the information needed to generate appropriate and valid hypothesis for future activities such as epidemiologic studies. The NER also serves as a mechanism for longitudinal health investigations that follow registrants over time to ascertain adverse health effects and latency periods.

The NER is currently composed of four sub-registries of persons known to have been exposed to specific chemicals: 1,1,1-Trichloroethane (TCA), Trichloroethylene (TCE), 2,3,7,8-tetrachlorodibenzo-p-dioxin (dioxin), and benzene. In 2001, the NER will establish a new asbestos subregistry.

Participants in each subregistry are interviewed initially with a baseline questionnaire. An identical follow-up telephone questionnaire is administered to participants every three years until the criteria for terminating a specific subregistry have been met. The annual number of participants varies greatly from year to year. Two factors influencing the number of respondents per year are the number of subregistry updates that are scheduled and whether a new subregistry will be established. The addition of the new asbestos subregistry is expected to add approximately 6,000 persons to the NER. This increase is reflected in the following estimated burden table.

The following table is annualized to reflect one new subregistry (asbestos) and five updates for the requested three-year extension of OMB No. 0923-0006. There is no cost to registrants.

Respondents	Number of respondents	Responses per respondent	Average burden per response (in hrs.)	Total annualized burden (in hrs.)
One New Subregistry	2,000	1	0.50	1,000
Five updates	4,927	1	0.42	2,069
Total	3,069

Dated: December 1, 2000.

John Moore,

Acting Associate Director for Policy, Planning, and Evaluation, Centers for Disease Control and Prevention (CDC).

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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Proposed Data Collections Submitted for Public Comment and Recommendations

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Proposed Project

NIOSH Research Study for the Prevention of Work-related Musculoskeletal Disorders (MSDs)—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC). The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all

people through research and prevention. There is evidence of causal relationships between physical job stressors (e.g., repetitive or static exertion, forcefulness, awkward postures) and MSDs, and some quantitative information is available on how much rates of MSDs change at varying levels of exposure to each stressor and combination of stressors (exposure-response relationships). Additional information would foster the further development of effective strategies for prevention.

A research project is proposed to conduct a prospective cohort study to quantify the risk for upper limb and low back MSDs at varying levels of exposure to physical job stressors (repetitive, forceful exertion, awkward postures, vibration, manual handling, etc.). This research will involve multiple work sites from the service and manufacturing industries with job tasks that represent a range of exposures to physical job stressors that can result in musculoskeletal disorders of the upper limb (e.g., carpal tunnel syndrome, hand-wrist tendinitis, medial and lateral epicondylitis, hand-arm vibration syndrome (HAVS)) and low back disorders. Because of the limitations of