

cert. denied, 486 U.S. 1014–15 (1988); *see also FTC v. Gem Merchandising Corp.*, 87 F.3d 466, 470 (11th Cir. 1996).

The Commission is sensitive to the interest in avoiding duplicative recoveries by injured persons or “excessive” multiple payments by defendants for the same injury. Thus, although a particular illegal practice may give rise both to monetary equitable remedies and to damages under the antitrust laws, when an injured person obtains damages sufficient to erase an injury, we do not believe that equity warrants restitution to that person. We will take pains to ensure that injured persons who recover losses through private damage actions under the Clayton Act not recover doubly for the same losses via FTC-obtained restitution. Similarly, in cases involving both disgorgement and restitution, we would apply any available disgorged funds toward restitution and credit any funds paid for restitution against the amount of disgorgement.

We do not, however, consider it appropriate to offset a civil penalty assessment against disgorgement or restitution. As noted above, disgorgement is an equitable remedy whose purpose is simply to remove the unjust gain of the violator. Penalties are intended to punish the violator and reflect a different, additional calculation of the amount that will serve society’s interest in optimal deterrence, retribution, and perhaps other interests. A penalty award would have no punitive effect if it were simply offset against these equitable remedies. It is not the Commission’s intent, therefore, to allow its monetary relief proceedings to dilute the effectiveness of a civil penalty.

When the same conduct gives rise to two different causes of action, moreover, the imposition of remedies for each cause of action does not necessarily mean the resulting sanctions are “excessive.” *See e.g., California v. ARC America Corp.*, 490 U.S. 93 (1989); *Loeb Industries, Inc. v. Sumitomo Corp.*, 306 F.3d 469, 492 (7th Cir. 2002), *cert. denied*, 123 S. Ct. 2247 (2003); *In Re Lorazepam & Clorazepate Antitrust Litigation*, MDL Dkt. No. 1290 (D.D.C.) (denial of motion to dismiss, July 2, 2001) Mem. Order at 15–16. Ultimately, we believe that courts considering equitable remedies have sufficient flexibility to craft orders to avoid unjust results.¹⁶ We have not yet encountered any such complications.

As a procedural matter, in the Commission’s two recent cases in which disgorgement was approved, claims administration procedures were being developed in parallel state and private litigation. To simplify the process and avoid any appearance of duplicative payments, in each of those cases the funds recovered by the Commission were combined with other recoveries and a single claims administration process handled the administration of all the funds. In future cases, the Commission could also consider the suggestion of several commentors to set up an escrow fund, to seek appointment of a special master or claims administrator to determine the appropriate allocation of funds collected, or to seek to coordinate parallel actions.

By direction of the Commission

Donal S. Clark,
Secretary.

[FR Doc. 03–19722 Filed 8–1–03; 8:45 am]

BILLING CODE 6750–01–M

HARRY S. TRUMAN SCHOLARSHIP FOUNDATION

Notice of Intent To Extend an Information Collection

AGENCY: Harry S. Truman Scholarship Foundation.

ACTION: Notice and request for comments.

SUMMARY: In compliance with the requirements of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Harry S. Truman Scholarship Foundation [Foundation] will publish periodic summaries of proposed projects.

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 or the forms of information technology.

(1997); *SEC v. Penn Cent. Co.*, 425 F. Supp. 593, 599 (E.D. Pa. 1976); *see also SEC v. Texas Gulf Sulphur Co.*, 446 F.2d 1301, 1307 (2d Cir.) (establishing escrow fund to prevent “double liability”), *cert denied*, 404 U.S. 1005 (1971).

DATES: Written comments on this notice must be received by October 3, 2003 to be assured of consideration. Comments received after that date will be considered to the extent practicable.

FOR FURTHER INFORMATION CONTACT:

Contact Louis H. Blair, Executive Secretary, Harry S. Truman Scholarship Foundation, 712 Jackson Place, NW., Washington, DC 20006; telephone 202–395–4831; or send e-mail to lblair@truman.gov. You also may obtain a copy of the data collection instrument and instructions from Mr. Blair.

SUPPLEMENTARY INFORMATION:

Title of Collection: Truman Scholar Payment Request Form.

OMB Approval Number: 3200–0005.

Expiration Date of Approval: September 30, 2003.

Type of Request: Intent to seek approval to extend an information collection for three years.

Proposed Project: The Foundation has been providing scholarships since 1977 in compliance with Public Law 93–642. This data collection instrument is used to collect essential information to enable the Truman Scholarship Foundation to determine the amount of financial support to which each Truman Scholar is eligible and then to make the payment. A total response rate of 100% was provided by the 315 Truman Scholars who received support in FY 2002.

Estimate of Burden: The Foundation estimates that, on average, 0.5 hours per Scholar applying for funds will be required to complete the Payment Request Form, for a total annual burden of 157.5 hours for all applicants.

Respondents: Individuals.

Estimated Number of Responses: 215.

Estimated Total Annual Burden on Respondents: 157.5 hours.

Dated: July 30, 2003.

Louis H. Blair,

Executive Secretary.

[FR Doc. 03–19777 Filed 8–1–03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–59–03]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and

¹⁶Courts routinely allows “set-offs” and credits, for example, to avoid duplicative payments. *See, e.g., SEC v. First Jersey Sec., Inc.*, 101 F. 3d 1450, 1475 (2d Cir. 1996), *cert. denied*, 552 U.S. 812

Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498-1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503 or by fax to (202) 395-6974. Written comments should be received within 30 days of this notice.

Proposed Project

Emergency Epidemic Investigations (0920-0008)—Extension—(Epidemiology Program Office, EPO)—One of the objectives of CDC’s epidemic services is to provide for the prevention and control of epidemics and protect the population from public health crises such as man made or natural biological disasters and chemical emergencies. This is carried out, in part, by training investigators, maintaining laboratory capabilities for identifying potential problems, collecting and analyzing data, and recommending appropriate actions to protect the public’s health. When state, local, or foreign health authorities request help in controlling an epidemic or solving other health problems, CDC dispatches skilled epidemiologists from the Epidemic Intelligence Service (EIS) to investigate and resolve the problem. Resolving public health problems rapidly ensures costs effective health care and enhances health promotion and disease prevention. Annually, the EIS Program coordinates 400 Epidemic Assistance Investigations (Epi-Aids) and state-based field investigations.

Epidemics are prevented and controlled by mobilizing and deploying CDC staff, primarily EIS officers to respond rapidly to disease outbreaks and disaster situations. At the request of public health officials—at the state, national, or international level—CDC provides assistance by participating in epidemiologic field investigations.

The purpose of the Emergency Epidemic Investigation surveillance is to collect data on the conditions surrounding and preceding the onset of a problem. The data must be collected in a timely fashion so that information can be used to develop prevention and control techniques, to interrupt disease transmission and to help identify the cause of an outbreak. Since the events necessitating the collections of information are of an emergency nature, most data collection is done by direct interview or written questionnaire and are one-time efforts related to a specific outbreak or circumstance. If during the emergency investigation, the need for further study is recognized, a project is designed and separate OMB clearance is required. Interviews are conducted to be as unobtrusive as possible and only the minimal information necessary is collected. The Emergency Epidemic Investigations is the principal source of data on outbreaks of infectious and noninfectious diseases, injuries, nutrition, environmental health and occupational problems.

Each investigation does contribute to the general knowledge about a particular type of problem or

emergency, so that data collections are designed taking into account similar situations in the past. Some questionnaires have been standardized, such as investigations of outbreaks aboard aircraft or cruise vessels.

The Emergency Epidemic Investigations provides a range of data on the characteristics of outbreaks and those affected by them. Data collected include demographic characteristics, exposure to the causative agent(s), transmission patterns and severity of the outbreak on the affected population. These data, together with trend data, may be used to monitor the effects of change in the health care system, planning of health services, improving the availability of medical services and assessing the health status of the population.

Users of the Emergency Epidemic Investigations data include, but are not limited to EIS Officers in investigating the patterns of disease or injury, investigating the level of risky behaviors, identifying the causative agent and identifying the transmission of the condition and the impact of interventions.

It is difficult to predict the number of epidemic investigations which might occur in any given year. The previous three years’ experience shows an annualized burden of 3,000 hours and respondent total of 12,000. For this clearance, we are requesting 3,750 total burden hours. This is due to the increase in the number of investigations performed over the past two years.

Respondents	Number of respondents	Number of responses/respondent	Avg. burden per response (in hrs.)
Total Respondents	15,000	1	15/60

Dated: July 29, 2003.
Thomas A. Bartenfeld,
Acting Associate Director for Policy, Planning, and Evaluation, Centers for Disease Control and Prevention (CDC).
 [FR Doc. 03-19685 Filed 8-1-03; 8:45 am]
BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-55-03]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498-1210. Send written

comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503 or by fax to (202) 395-6974. Written comments should be received within 30 days of this notice.

Proposed Project

Congenital Syphilis (CS) Case Investigation and Report Form (OMB Control No. 0920-0128)—Revision—National Center for HIV, STD, and TB Prevention (NCHSTP), Centers for Disease Control and Prevention (CDC). CDC proposes to continue data collection for congenital syphilis case investigations under the “Congenital Syphilis Case Investigation and Report Form” (CDC73.126 REV 11-98); this form is currently approved under OMB