

device does not perform as it should; the sooner they report it to the FDA, the faster the agency can analyze the problem and take corrective action," said FDA Commissioner Dr. David A. Kessler.

An important part of MEDWatch is a simplified reporting form—a self-addressed, one-page form that can be folded and mailed postage-free. In addition, the agency has established a special 24-hour toll-free telephone line, (800) FDA-1088, for adverse event information. Health professionals also will be able to file reports on a computer.

Joint Commission to Make Hospital Performance Information Available to Public

Starting next year, the Joint Commission on Accreditation of Healthcare Organizations for the first time will release information to the public on how hospitals meet specific performance standards. Commission President Dennis S. O'Leary said, "This is a landmark issue for us ... it is very much in line with the reform environment. The change in the Joint Commission's confidentiality and disclosure policy, approved recently by the commissions board, recognizes the accrediting agency's obligation to share information with patients, purchasers, and other stakeholders in healthcare delivery systems."

Under the new policy, standards-compliance ratings in such areas as nursing care, infection control, patient rights, and life safety will be provided on a comparative basis. Also available for comparison will be facilities' overall summary grid scores, which are the basis for accreditation decisions.

Healthcare organizations will be able to review and comment on the data in the Joint Commission performance reports prior to their publication, but official survey reports will remain confidential.

FROM: Trustee magazine. American Hospital Association, July 1993.

CDC Clarifies Recommendation for Use of Bleach to Disinfect Needles

The CDC clarified its April 19, 1993, joint bulletin regarding the use of bleach for disinfecting needles and syringes, including the recommendation for the use of full-strength household bleach. In a letter signed by Dr. James Curran, Associate Director for HIV/AIDS, the CDC explained that it recommended full-strength bleach to disinfect needles and syringes because of the difficulty of cleaning the interior of needles and syringes and the use of needles and syringes for parenteral injection.

This recommendation does not change the recommendations for situations encountered in healthcare settings, such as contamination of environmental

surfaces with blood. The CDC recommendation for disinfecting environmental surfaces continues to include the use of a 1:100 dilution of household bleach (or 1/4 cup bleach to 1 gallon tap water) (See *MMWR* 1993;46:1-18S; August 21, 1987.)

Lifetime AIDS Treatment Cost: \$119,274

The lifetime cost of treating a person with HIV disease, assuming treatment begins immediately after infection and diagnosis, averages \$119,274, Dr. Fred Hellinger of the Agency for Health Care Policy and Research recently reported.

The figure was calculated by multiplying the average monthly cost of care in each of four stages of HIV disease by the average number of months the infected person spends in each stage. Dr. Hellinger's study was based on self-reported interviews and data every three months from 1,164 respondents from March 1991 through August 1992 at 26 sites in 10 cities in the United States. Costs included inpatient care, outpatient visits, home healthcare, long-term care, and drugs, both self-paid and covered by insurance. The survey assumed an average of 10.3 years from infection until progression to AIDS and a 25-month survival time from then until death.

Hellinger said the study was not random and thus the lifetime costs represented the "upper bounds" for the cost of treatment. The study found that the cost of treating a person with AIDS, which has risen rapidly in the past, has fallen as a result of reducing the use of inpatient hospital services.

FROM: Hellinger FJ. The lifetime cost of treating a person with HIV infection. *JAMA* 1993;270:474-478.

Government Considers Relaxing CLIA Regulations

The federal government now is looking at proposals for relaxing physician office laboratory regulations under the Clinical Laboratory Improvement Acts (CLIA). Although the Clinton administration vowed to ease some of the more burdensome requirements, it appears that the proposals for easing the rules are driven by the lack of money rather than a desire to limit the burden on physicians. The current plan is to let "low-volume" and "low-risk" labs perform self-assessments rather than submit to on-site surveys by government inspectors.

CLIA set up four categories of labs based on test complexity. It is unclear how the government would define low-risk labs for the purpose of this new proposal. However, it is assumed that the tests they perform would fall into CLIA's "moderately complex" category.

Labs doing high-complexity testing still would be