1,2,4-Trichlorobenzene (as the metabolite 2,4,5-trichlorophenol)

Organophosphate Pesticides

Chlorpyrifos (as the metabolite 3,5,6-trichloro-2-pyridinol)

Phenols

ortho-Phenylphenol

N-Methyl Carbamates

Carbaryl (as the metabolite 1-naphthol)
Carbofuran (as the metabolite carbofuranphenol)
Propoxur (as the metabolite 2-isopropoxyphenol)
Alachlor (as the metabolite alachlor mercapturate)

Triazines

Atrazine (as the metabolite atrazine mercapturate)

Other Herbicides

2,4-Dichlorophenoxyacetic acid (2,4-D)

Polychlorinated Biphenyls (PCBs)

PCB numbers 52, 66, 81, 99, 101, 105, 118, 126, 128, 138, 146, 153, 156, 157, 167, 169, 170, 180, and 183

Nominated Chemicals or Chemical Categories Already Planned for Inclusion in Future ''Reports''

Metals

Arsenic (speciated and total)
Methyl mercury

Volatile Organic Compounds (VOCs)

Benzene
1,1,1-Trichloroethane
1,2-Dichlorobenzene
Carbon tetrachloride
Chloroform
Ethylene dichloride
Perchloroethylene
Toluene
Xylenes (o, m, p-isomers)

Polybrominated Compounds

Polybrominated biphenyls (PBBs)
Polybrominated diphenyl ethers (PBDEs)
Tetrabromodiphenyl ether (TeBDE)
Pentabromodiphenyl ether (PeBDE)
Heptabromodiphenyl ether (HpBDE)
Decabromodiphenyl ether (DeBDE)
Hexabromocyclododecane (HBCD)
Tetrabromobisphenol A (TBBP-A)

Organochlorine Pesticides

alpha Hexachlorocyclohexane (HCH)
Aldrin (as the metabolite endrin)
4,4′-DDD (p,p′-DDD; DDD)
Dieldrin
Endosulfan
Endrin
Methoxychlor
Octachlorostyrene
Pentachlorobenzene (as the metabolite pentachlorophenol)
Toxaphene

Chloroacetanilides

Acetochlor (as the metabolite acetochlor mercapturate)
Metolachlor (as the metabolite metolachlor mercapturate)

Phenols

Bisphenol A
Nonylphenol
Octylphenol

Dithiocarbamates

Ethylenethiourea

Pyrethrins and Pyrethroids

Cyfluthrin
Cypermethrin
Deltamethrin
Esfenvalerate
Fenvalerate
Permethrin

Perfluorinated Compounds

Perfluorohexanoic sulfonic acid (PFHS)
Perfluorooctane sulfonate (PFOS), including nominated salt forms
Perfluorooctanoic acid (PFOA), including nominated salt forms

Organophosphate Pesticides

Acephate
Azinophos-methyl (also as dialkyl phosphate metabolites)
Coumaphos (also as dialkyl phosphate metabolites)
Methamidophos

Polychlorinated Biphenyls (PCBs)

PCB numbers 77, 87, 151, 158, 189, 194, 195, 169, 203, 206, and 209

ADDRESSES: Address all correspondence related to this notice to Dorothy Sussman, CDC, National Center for Environmental Health, Division of Laboratory Sciences, Mail Stop F–20, 4770 Buford Highway, Atlanta, Georgia 30341.

SUPPLEMENTARY INFORMATION: CDC publishes the ''Report'' under the authorities 42 U.S.C. 241 and 42 U.S.C. 242k. The ''Report'' provides an ongoing assessment using biomonitoring of the exposure of the noninstitutionalized, U.S. civilian population to environmental chemicals. Biomonitoring assesses human exposure to chemicals by measuring the chemicals or their breakdown products in human specimens such as blood or urine. For the ''Report,'' an environmental chemical means a chemical compound or chemical element present in air, water, soil, dust, food, or other environmental medium. The ''Report'' provides exposure information about participants in an ongoing national survey known as the National Health and Nutrition Examination Survey (NHANES). This survey is conducted by CDC's National Center for Health Statistics; biomonitoring measurements are conducted by CDC's National Center for Environmental Health. The first ''Report,'' published in March 2001, gave information about levels of 27 chemicals in the U.S. population. The second ''Report,'' published in January 2003, provided data on 116 chemicals, including expanded data on the 27 in the first ''Report,'' and was the most extensive assessment ever of the exposure of the U.S. population to environmental chemicals. The ''Report'' can be obtained in the following ways: Access *http://www.cdc.gov/exposurereport;* e-mail *ncehdls@cdc.gov;* or telephone 1–866–670–6052.

Current plans are to release future reports of exposure of the U.S. population that cover 2-year periods (*e.g.*, 2001–2002; 2003–2004; 2005–2006) and that will include data on more chemicals than the 116 listed in the second ''Report.'' Over time, CDC will be able to track trends in exposure levels. Future releases also may include additional exposure information for special exposure populations (*e.g.*, children, women of childbearing age, elderly people) from studies examining localized or point sources and from studies of adverse health effects resulting from exposure to varying levels of environmental chemicals.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both the CDC and ATSDR.

Dated: September 24, 2003.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 03–24671 Filed 9–29–03; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2000N–1530]

Richard L. Borison; Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (the act) debarring Dr. Richard L. Borison for 10 years from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Dr. Borison was convicted of felonies under Georgia State law for racketeering, theft, and false statements and representations,

and was a material participant in offenses for which another person is being debarred. Dr. Borison has failed to request a hearing and, therefore, has waived his opportunity for a hearing concerning this action.

DATES: This order is effective September 30, 2003.

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm., 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Mary Catchings, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION:

I. Background

On October 8, 1998, the Superior Court for the County of Richmond, State of Georgia, accepted Dr. Borison's plea of guilty and entered judgment against him for 36 counts of criminal offenses under Georgia State law for racketeering, theft, and false statements and representations.

As a result of this conviction, FDA served Dr. Borison by certified mail on December 5, 2002, a notice proposing to debar him for 10 years from providing services in any capacity to a person that has an approved or pending drug product application. The proposal also offered Dr. Borison an opportunity for a hearing on the proposal. The debarment proposal was based on findings: (1) Under section 306(b)(2)(B)(ii) of the act (21 U.S.C. 335a(b)(2)(B)(ii) that Dr. Borison was convicted of felonies under State law for racketeering, theft, and false statements and representations; and (2) under section 306(b)(2)(B)(iii) of the act that Dr. Borison was a material participant in offenses leading to the conviction and debarment of another individual. Dr. Borison was provided 30 days to file objections and to request a hearing. Dr. Borison did not request a hearing. His failure to request a hearing constitutes a waiver of his opportunity for a hearing and a waiver of any contentions concerning his debarment.

II. Findings and Order

Therefore, the Director, Center for Drug Evaluation and Research, under section 306(b)(2)(B) of the act, and under authority delegated to her (21 CFR 5.34), finds that Dr. Richard L. Borison: (1) Has been convicted of a felony under State law for racketeering, theft, and false statements and representations; and (2) was a material participant in offenses leading to the conviction of another individual.

As a result of the foregoing findings, Dr. Richard L. Borison is debarred for 10 years (two periods of 5 years, to run consecutively, based on his conviction for State felonies and his role as a material participant in the offenses leading to the conviction and debarment of another individual) from providing services in any capacity to a person that has an approved or pending drug product application under section 505, 512, or 802 of the act (21 U.S.C. 355, 360b, or 382) or under section 351 of the Public Health Service Act (42 U.S.C. 262)(see sections 306(c)(1)(B) and (c)(2)(A)(iii) and 201(dd) of the act (21 U.S.C. 321(dd))). Any person with an approved or pending drug product application who knowingly uses the services of Dr. Borison in any capacity during his period of debarment will be subject to civil money penalties. If Dr. Borison, during his period of debarment, provides services in any capacity to a person with an approved or pending drug product application, he will be subject to civil money penalties. In addition, FDA will not accept or review any abbreviated new drug applications submitted by or with the assistance of Dr. Borison during his period of debarment.

Any application by Dr. Borison for termination of debarment under section 306(d)(4) of the act should be identified with Docket No. 2000N–1530 and sent to the Division of Dockets Management (see **ADDRESSES**). All such submissions are to be filed in four copies. The public availability of information in these submissions is governed by 21 CFR 10.20(j). Publicly available submissions may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: September 4, 2003.

Janet Woodcock,

Director, Center for Drug Evaluation and Research.

[FR Doc. 03–24656 Filed 9–29–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2000N–1428]

Suhas V. Sardesai; Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (the act) permanently debarring Suhas V. Sardesai from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Mr. Sardesai was convicted of a felony under Federal law for conduct relating to the regulation of a drug product under the act. Mr. Sardesai failed to request a hearing and, therefore, has waived his opportunity for a hearing concerning this action.

DATES: This order is effective September 30, 2003.

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Mary Catchings, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION:

I. Background

On October 28, 1998, the U.S. District Court for the District of Maryland accepted Mr. Suhas V. Sardesai's plea of guilty to one count of distributing an adulterated drug into interstate commerce, a Federal felony offense under section 501(a)(2)(B) of the act (21 U.S.C. 351(a)(2)(B)).

As a result of this conviction, FDA served Mr. Sardesai by certified mail on July 24, 2002, a notice proposing to permanently debar him from providing services in any capacity to a person that has an approved or pending drug product application. The proposal also offered Mr. Sardesai an opportunity for a hearing on the proposal. The proposal was based on a finding, under section 306(a)(2)(B) of the act (21 U.S.C. 335a(a)(2)(B)), that Mr. Sardesai was convicted of a felony under Federal law for conduct relating to the regulation of a drug product under the act. Mr. Sardesai was provided 30 days to file objections and request a hearing. Mr. Sardesai did not request a hearing. His failure to request a hearing constitutes a waiver of his opportunity for a hearing and a waiver of any contentions concerning his debarment.

II. Findings and Order

Therefore, the Director, Center for Drug Evaluation and Research, under section 306(a)(2)(B) of the act, and under authority delegated to her (21 CFR 5.34), finds that Mr. Suhas V.