

LEGISLATIVE HISTORY
of
LEAD-BASED PAINT POISONING PREVENTION ACT
(Public Law 91-695)

Early in the 91st Congress (March, 1969), Rep. William F. Ryan (D-N.Y.) introduced the first bill to provide Federal financial assistance to States and their subdivisions for projects involving the detection and treatment of lead poisoning in children and for the rehabilitation of areas found to be seriously contaminated by old lead-based paints. Subsequently, approximately 50 members of both the House and Senate demonstrated a recognition of this serious problem and strong bi-partisan support by introducing similar legislative proposals or joining as sponsor thereof. One of these bills (S. 3941) -- introduced by Senator Richard S. Schweiker (R - Pa.) -- contained a provision making it unlawful "for any person to apply or otherwise use, after the effective date of the Act, any paint with an excess of 1 per centum lead pigments or lead additives on the interior of any dwelling..."

The first public hearings on this legislation were not held until July 30, 1970 when the Housing Subcommittee of the House Banking and Currency Committee heard witnesses testify on H.R. 17260, the bill introduced by Rep. William A. Barrett (D - Pa.), Chairman of the Housing Subcommittee. NPVLA submitted a written statement in order to present additional facts relevant to the Subcommittee's review of the bill and to attempt to clarify certain erroneous matters and possible misconceptions on the subject which developed during examination of witnesses at the hearings.

Notwithstanding the testimony of medical witnesses at the hearings and the detailed information provided by NPVLA to emphasize that the lead poisoning of children was being caused by ingestion of flakes or chips of old lead-based paint from the walls and ceilings of poorly-maintained pre-World War II dwellings, the House Committee reported out a bill (H.R. 19172) on September 14, 1970 which contained a new provision prohibiting "the use of lead-based paint in all future Federal construction and rehabilitation of any building or structure with Federal assistance in any form."

At this time, NPVLA representatives met with Rep. Barrett and pointed out that his modified bill would deny the proper uses of lead-based paints in industrial areas where its rust-inhibiting qualities were of great importance and where no possible hazard to children now or in the future would exist. Rep. Barrett assured that his Committee did not intend to extend the coverage of his bill to that extent. However, the bill passed the House without further amendment on October 5, 1970, and was referred to the Senate Committee on Labor and Public Welfare.

With this development, NPVLA addressed a letter to the Chairman of the Senate Committee and requested permission to testify "in support of amendments deemed vital to make the bill an effective law responsive to the public need." The issue became further clouded by politics about this time, and a provision in the Senate "Omnibus Housing Bill" (S. 4368) -- limiting the restrictions on use of lead-based paint, as defined, to the interior of dwellings subject to the Act -- was not agreed to by conferees from the House and Senate Banking and Currency Committees. As finally approved by the Congress, the "Omnibus Housing Bill" (Housing and Urban Development Act of 1970) contained no provisions relating to the use of lead-based paint.

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grams to insure that identified cases of lead-based paint poisoning are protected against further exposure to lead-based paints in their living environment and

(4) any other actions which will reduce or eliminate lead-based paint poisoning.

(d) Each local program shall afford opportunities for employing the residents of communities or neighborhoods affected by lead-based paint poisoning, and for providing appropriate training, education, and any information which may be necessary to inform such residents of opportunities for employment in lead-based paint poisoning elimination programs.

TITLE II -- GRANTS FOR THE ELIMINATION OF LEAD-BASED PAINT POISONING

SEC. 301. The Secretary of Health, Education, and Welfare is authorized to make grants to units of general local government in any State for the purpose of assisting such units in developing and carrying out programs that identify those areas that present a high risk to the health of residents because of the presence of lead-based paints on interior surfaces, and then to develop and carry out programs to eliminate the hazards of lead-based paint poisoning.

(a) A local program should include:

(1) development and carrying out of comprehensive testing programs to detect the presence of lead-based paints on surfaces of residential housing;

(2) the development and carrying out of a comprehensive program requiring the prompt elimination of lead-based paints from all interior surfaces, porches, and exterior surfaces to which children may be commonly exposed, of residential housing on which lead-based paints have been used as a surface covering, including those surfaces on which non-lead-based paints have been used to cover surfaces to which lead-based paints were previously applied; and

(3) any other actions which will reduce or eliminate lead-based paint poisoning.

(b) Each such program shall--

(1) be consistent with the appropriate local program assisted under section 101, and

(2) afford, to the maximum extent feasible, opportunities for employing the residents of communities or neighborhoods affected by lead-based paint poisoning, and for providing appropriate training, education, and any information which may be necessary to inform such residents of opportunities for employment in lead-based paint elimination programs.

TITLE III -- FEDERAL DEMONSTRATION AND RESEARCH PROGRAM

FEDERAL DEMONSTRATION AND RESEARCH PROGRAM

SEC. 302. The Secretary of Housing and Urban Development, in consultation with the Secretary of Health, Education, and Welfare, shall develop and carry out a demonstration and research program to determine the nature and extent of the problem of lead-based paint

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poisoning in the United States, particularly in urban areas, and the methods by which lead-based paint can most effectively be removed from interior surfaces, porches, and exterior surfaces to which children may be commonly exposed of residential housing. Within one year after the date of the enactment of this Act the Secretary shall submit to the Congress a full and complete report of his findings and recommendations as developed pursuant to such program, together with a statement of any legislation which should be enacted and any changes in existing law which should be made, in order to carry out such recommendations.

TITLE IV - PROHIBITION AGAINST FUTURE USE OF LEAD-BASED PAINT

PROHIBITION AGAINST USE OF LEAD-BASED PAINT IN FUTURE CONSTRUCTION AND REHABILITATION

SEC. 401. The Secretary of Health, Education, and Welfare shall take such steps and impose such conditions as may be necessary or appropriate to prohibit the use of lead-based paint in residential structures constructed or rehabilitated after the date of enactment of this Act by the Federal government, or with Federal assistance in any form.

TITLE V - GENERAL DEFINITIONS

SEC. 501. As used in this Act--

(1) The term "State" means the several States, the District of Columbia, the Commonwealth of Puerto Rico, and the territories and possessions of the United States;

(2) the term "units of general local government" means: (A) any city, county, township, town, borough, parish, village, or other general purpose political subdivision of a State; (B) any combination of units of general local government in one or more States; (C) an Indian tribe; or (D) with respect to lead-based paint poisoning elimination activities in their urban areas, the territories and possessions of the United States; and

(3) the term "lead-based paint" means any paint containing more than 1 per centum lead by weight (calculated as lead metal) in the total non-volatile content of liquid paints or in the dried film of paint already applied.

CONSULTATION WITH OTHER DEPARTMENTS AND AGENCIES

SEC. 502. In carrying out the authority under this Act, the Secretary of Health, Education, and Welfare shall cooperate with and seek the advice of the heads of any other departments or agencies regarding any programs under their respective responsibilities which are related to, or would be affected by, such authority.

APPROPRIATIONS

SEC. 503. (a) There is hereby authorized to be appropriated to carry out the provisions of title I of this Act not to exceed \$2,330,000 for the fiscal year 1971 and \$4,000,000 for the fiscal year 1972.

(b) There is hereby authorized to be appropriated to carry out the

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provisions of title II of this Act not to exceed \$1,000,000 for the fiscal year 1971 and \$10,000,000 for the fiscal year 1972.

(c) There is hereby authorized to be appropriated to carry out the provisions of title III of this Act not to exceed \$1,875,000 for the fiscal year 1971 and \$3,340,000 for the fiscal year 1972.

(d) Any amounts appropriated under this section shall remain available until expended when so provided in appropriation Acts and any amounts authorized for the fiscal year 1971 but not appropriated may be appropriated for the fiscal year 1972.

Approved January 13, 1971.

LEGISLATIVE HISTORY

HOUSE REPORTS No. 81-1021 (Comm. on Banking and Currency, and No. 81-1022 (Comm. on Conference).

SENATE REPORT No. 81-1021 (Comm. on Labor and Public Welfare).

CONGRESSIONAL RECORD Vol. 116 (1970).

Oct. 1, considered and passed House.
Dec. 17, considered and passed Senate amendment.
Dec. 22, House disagreed to Senate amendment.
Dec. 23, Senate agreed to conference report.
Dec. 31, House agreed to conference report.

It will be noted that Title II of the act authorizes the secretary of Health, Education and Welfare to make grants to units of "general local government in any State" to eliminate existing lead-based paints from "all interior surfaces, porches, and exterior surfaces to which children may be commonly exposed, of residential housing on which lead based paints have been used as a surface covering. . . This is paint already applied.

Title IV deals with future use of lead-based paint. In short and terse form it directs HEW to "take such steps and impose such conditions as may be necessary or appropriate to prohibit the use of lead-based paint in residential structures constructed or rehabilitated after the date of enactment of this Act by the Federal government, or with Federal assistance in any form."

John Montgomery, general counsel for the National Paint, Varnish and Lacquer Association, who followed the legislation closely from the time it was introduced until the time the act was approved, said that the paint and coatings industry and the raw material suppliers need to be vitally concerned with

the regulations which HEW sets up to implement carrying out of the Title IV directive. The national association already has been in close contact with HEW and has offered to assist in drawing up these regulations, Mr. Montgomery said.

These regulations will be drafted over a period of time, Mr. Montgomery said, "and we will be constantly and alertly taking such actions as are possible to protect the industry's interests."

Mr. Montgomery said however, that Title III is of extreme immediate importance, because Congress directs the secretary of Housing and Urban Development to report back within one year after the date of the enactment of the act with certain findings and recommendations regarding lead-based paint poisoning to assist in implementing provisions in Title I and II.

The NPVLA general counsel said that there is much to be done regarding the act. He said that while NPVLA will be watching developments constantly, everyone in the industry should familiarize himself with the provisions of the act and be prepared to give assistance in any way possible to insure that

be held confidential and a determination made that a proper showing in support of the request has been made on the grounds that its disclosure could adversely affect such person by disclosing information in the nature of trade secrets or commercial or financial information obtained from any person and privileged or confidential. If it is determined that a proper showing has been made in support of the request, the material will be held confidential; otherwise, notice will be given of denial of such a request and an opportunity afforded for withdrawal of the submission. Requests for confidential treatment will be held confidential (1 CFR 1.27 (c)).

Comments on the proposal should bear a reference to the date and page number of this issue of the Federal Register.

Date at Washington, D.C., on October 27, 1971.

Kenneth M. McEwen,
Deputy Administrator, Meat
and Poultry Inspection Program.
[FR Doc. 71-18067 Filed 11-1-71; 9:47 am]

DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

Food and Drug Administration

[21 CFR Parts 27, 31]

CERTAIN ORANGE JUICE PRODUCTS

Extension of Time for Filing Comments
on Proposed Regulations Stayed
Identity Standards

The notice published in the Federal Register of September 9, 1971 (36 F.R. 18068), proposing to revise the stayed standards for the diluted orange juice beverages (27.120, 27.121, 27.122) and proposing establishment of additional identity standards for certain related products, provided for the filing of comments within 60 days after said date.

The Commissioner of Food and Drugs has received a request for a 30-day extension of such time and, good reason therefor appearing, the time for filing comments on the subject proposals is hereby extended to December 8, 1971. Requested comments may be sent in the office of the Hearing Clerk, Department of Health, Education, and Welfare, Room 6-01, 1400 Fishers Lane, Rockville, Md. 20851, during working hours, Monday through Friday.

This action is taken pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (am. 481, 761, 68 Stat. 1944-1945), as amended, 76 Stat. 912, 78 Stat. 941; 21 U.S.C. 361, 371 and under authority delegated to the Commissioner (21 CFR 2.120).

Dated: October 15, 1971.

Sam D. Fann,
Associate Commissioner
for Compliance.

[FR Doc. 71-18068 Filed 11-1-71; 8:46 am]

[21 CFR Part 1481]

COMBINATION DRUG CONTAINING NISONICIN SULFATE AND AMPHO- TEBICIN FOR ORAL USE

Proposed Revocation of Provisions
for Certification

In the Federal Register of July 2, 1970 (35 F.R. 10795, DEB 11313), the Commissioner of Food and Drug announced the provisions of the Food and Drug Administration following evaluation of reports received from the National Academy of Sciences-National Research Council Drug Efficacy Study Group on a certain oral preparation containing nisonicic sulfates and amphotericin. The drug was regarded as lacking substantial evidence, as defined in the Federal Food, Drug, and Cosmetic Act, that it is effective as a fixed combination for its claimed clinical effect and that each component of the drug contributes to the total effects claimed for such drug.

In addition to the section of the antibiotic drug regulations describing conditions for certification of such preparations, another section describes conditions for certification of a related drug, Fungicide Tablets containing nisonicic sulfates and amphotericin B. An approved Form 1 antibiotic drug application (FDA 68-514) is held for this drug by E. R. Squibb & Sons, Inc., 5 Greenwich Road, New Brunswick, N.J. 08901.

This product was not reviewed by the Agency; however, the Food and Drug Administration, having evaluated data originally filed in support of efficacy of this preparation, concludes that there is a lack of substantial evidence, as defined in the Federal Food, Drug, and Cosmetic Act, that this drug is effective as a fixed combination for its claimed clinical effect, and that each component of the drug contributes to the total effects claimed for such drug.

Accordingly, the Commissioner concludes that the antibiotic drug regulations providing for certification of such drugs should be revoked.

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (am. 502, 507, 68 Stat. 1944-51 as amended, 68 Stat. 655 as amended; 21 U.S.C. 361, 367) and under authority delegated to him (21 CFR 2.120), the Commissioner proposes that Part 1481 be amended by revoking § 1481.65. Nisonicic sulfates/amphotericin B tablets.

Interested persons may, within 30 days after publication hereof in the Federal Register, file with the Hearing Clerk, Department of Health, Education, and Welfare, Room 6-01, 1400 Fishers Lane, Rockville, Md. 20851, written comments (preferably in triplicate) regarding this proposal. Comments may be accompanied by a memorandum or brief in support thereof. Requested comments may be sent in the above office during working hours, Monday through Friday.

Dated: October 15, 1971.

Sam D. Fann,
Associate Commissioner
for Compliance.

[FR Doc. 71-18069 Filed 11-1-71; 8:45 am]

[21 CFR Part 191]

HAZARDOUS SUBSTANCES

Proposal To Declare Certain Heavy
Metal-Containing Paints and Other
Surface-Coatings To Require Special
Labeling for Child Protection

The heavy metals antimony, arsenic, cadmium, lead, mercury, selenium and soluble barium may be toxic when ingested, even in small amounts. Because children may show on interior and certain exterior residential surfaces and on toys and other children's articles, the Commissioner of Food and Drug concludes that paints and other surface-coating materials are hazardous substances requiring special labeling warning against their use on such surfaces if they contain more of these heavy metals than is essential under good manufacturing practices, or in any event if they contain more than 0.5 percent lead; or a total of 0.50 percent of the other heavy metals listed above on a dried weight basis, except for soluble barium which shall constitute no more than 1 percent of the total barium present. The special labeling to be required shall include on the main panel a cautionary signal word "Warning", a statement of the hazard "Contains _____" (the blank being filled in with the name(s) of the applicable heavy metal(s)) and instructions to read carefully the summary information placed on other panel(s) where located. Existing technology will permit industry to produce household paints containing no more of these heavy metals than is specified above. As new technology becomes available, the acceptable levels may be further reduced in accordance with good manufacturing practices.

Although paints containing small amounts of lead in excess of the proposed levels may not be toxic in themselves, when considered in conjunction with other sources of lead in the environment they constitute a substantial addition to the body burden that can reasonably be avoided through the application of available technology. The Commissioner has concluded that the scope of the Federal Hazardous Substances Act fully encompasses the question of cumulative toxicity.

The American National Standards Institute, in its voluntary standard Z96.1, has specified a maximum limit since 1955 of 1 percent lead in paint intended for use on children's toys, furniture, or interior surfaces. A 1-percent lead level was also adopted by Congress in the Lead-Based Paint Poisoning Prevention Act (Public Law 91-603; 84 Stat. 3078), which is administered by the Bureau of Community Environmental Management, Health Services and Mental Health Administration. That act prohibits the use of paint with a lead content above 1 percent in residential structures constructed or rehabilitated by the Federal Government or with Federal assistance. As an additional margin of safety, the

this regulation, containing any of the heavy metals specified below at a level exceeding that which is essential under good manufacturing practices or in any event containing amounts of such heavy metals as follows:

(2) Lead compounds of which the lead content (calculated as the metal) is in excess of 0.6 percent of the total weight of the contained solids or dried paint film; or

(11) Antimony, arsenic, cadmium, mercury, and selenium of which the metal content individually or in total (calculated as the metal) amounts 1.56 percent of the total weight of the contained solids or dried paint film; or

(2) Barium compounds of which the water soluble barium (calculated as the metal) exceeds 1 percent of the total barium present.

• 191.7 Products requiring special labeling under section 3(b) of the act.

(b) The Commissioner finds that these substances present special hazards and that the labeling required by section 2(p)(1) of the act is not adequate for the protection of the public health. Under section 2(b) of the act the following specific label statements are deemed necessary to supplement the labeling required by section 2(p)(1) of the act:

• • • • •

(7) Paint and other surface-coating materials declared to be hazardous under § 191.9(a)(2) shall bear on the main panel of their label, in addition to the requirements of § 191.103(a), the statement "contains....." the blank

being filed in with the name of each heavy metal present in the amount specified in § 191.6a) E7. Such paint and other surface-coating materials shall also bear on their labeling the signal word "Warning" and the following additional statement or its equivalent:

Print this on your page map to illustrate
 your own design.

Do not apply on maps and other children's
 books, newspapers, or similar articles of any
 kind, or on any other article, that is intended
 to be used by children.

Do not apply on those various articles of
 clothing, such as undershirts, pajamas,
 coats, or mittens, in which children may be
 commonly exposed.

Do not use on the inside of children's shoes.

the blank being filled in with the name of each heavy metal present in the amount specified in a list of 17.

Submitted comments may, within 30 days after publication, be used in the Personnel Review. Be with the Hearing Clerk, Department of Health, Education and Welfare, Room 4-05, 5600 Fishers Lane, Bethesda, MD 20895. Written comments (preferably in triplicate) regarding this proposal. Comments may be accompanied by a memorandum or brief in support thereof. Received comments may be seen in the above office during working hours. Member Through Print.

CHARLES C. INWALD,
Commissioner of Food and Drugs
[FDR Doc. 71-18061 Filed 11-1-71; 8:01 am]

BANNED HAZARDOUS SUBSTANCES

The Commissioner of Food and Drugs has received a petition, submitted pursuant to section 715(a)(1)(B) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 371(a)(1)(2)), proposing the issuance of a regulation under section 715(a)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 371(a)(2)) identifying as hazardous certain substances meant for household use containing more than minute trace of lead.

The petitioner proposes that § 101.5(a) of the hazardous substances regulations be amended by adding thereto a new subparagraph (§ 101.5(a)(6)), as follows.

§ 101.9 Banned hardware (software).

(a) Under the authority of section 2(a)(1) (C) of the act, the Commissioner declared as having hazardous substances the following articles because they possess such a degree of hazard that adequate cautionary labeling cannot be written and the public health and safety can be served only by keeping such articles out of interstate commerce:

(d) Faint containing lead, except for minute traces which no reasonable manufacturer could preclude from his product, as measured by the atomic absorption method, intended for interior or exterior use, and packaged in a form suitable for use in the household as defined by 18.1.1(e).

The following is the statement of grounds as presented in the petition:

SUBJECT: IN THE MATTER OF THE ESTATE OF WILLIAM FREDERICK HUNT FOR THE ESTATE OF THE SAME.

Human experience has established that the human brain is a very plastic organ (plastic meaning that it can change its shape) (through eating of and using chemicals). Human experience has also established that the human brain is a very plastic organ (plastic meaning that it can change its shape) (through eating of and using chemicals). Human experience has also established that the human brain is a very plastic organ (plastic meaning that it can change its shape) (through eating of and using chemicals).

children a child with lead poisoning 1 gram of lead based paint (1 percent) and by ASTM Standard 581-1; only with 10,000 micrograms of lead of which 1,000 micrograms will be absorbed. In addition to current legislation by setting safety paint a child will be exposed to from 14 to 500 micrograms of lead from other conventional sources. "Health Hazards of Environmental Lead," at Table 3 (April 1971).

It is known that blood levels of above 40 micrograms per 100 milliliters of blood represent a serious exposure to lead absorption of lead. "Research on R.E. 1700, R.E. 1500, R.E. 1470, before the Subcommittee on Housing of the House Committee on Banking and Currency," 91st Cong second sess 70 (1970). At levels of 80 micrograms per 100 milliliters a child is treated as a medical emergency. Dr. Whitehouse in his paper has extrapolated the effect of lead in children with lead at A-0, the same must be done with 1 percent lead paint to determine the result in children who ingest paint lead.

In his study, Dr. Hahn found that an adult man had 1,000 micrograms of lead daily. In addition to the usual amount in his diet, he had a blood lead level after 4 months of 50 micrograms lead per 100 grams whole blood. It was estimated that he would have absorbed a "total" level of 20 micrograms lead per 100 grams whole blood if he had had consumed for 4 additional months. "Hahn A. A. 30 Jour. Roy Soc. Med. 519, 521-527, 1951, 120-124, 177-180 (1951). As Hahn has suggested, this would mean 40 micrograms per 100 grams whole blood in a 10 microgram lead paint child with lead weighing 10 micrograms ingested lead to the same degree of Hahn's subject. As Hahn has suggested, then a proper assumption would be that a daily replacement of 40 micrograms lead would produce toxicity within 4 months. A child ingesting a 1 gram chip of 1 percent lead based paint would have a daily intake of almost 10 times the amount of Hahn's subject. Concomitantly, intake of this rate would produce an unusual medical emergency much more quickly than Hahn foresaw for his subject.

Medical experiments cannot be performed on young children and infants to determine whether their absorption rate for lead is greater than that of adults. The child with lead must be protected from lead in his environment. The only adequate way to protect him is to eliminate the lead hazards that confront him. It is within the existing state of the art for the paint industry to eliminate lead (except for matrix traces) from paint. "Research on R.E. 1700," report of Mr. American have been failed into a feeling of security with regard to today's paint. See EPA, "Environmental Lead and Public Health," at 25 (March 1971). Many believe that lead in paint has been banned. On the day of New York Times, July 24, 1971, at 1, August 4, 1971, at 15. This date's industry's ANSI standard 20.1 and a law banning the sale of paint for interior use with more than a 1 percent lead content without an explicit warning label.

Warning labels are, however, absolutely inadequate to protect injury to children from lead based paint. In today's mobile society a family has no idea when their landlord or predecessor accepted the need to coat the walls of their living quarters. The government that is suffering from lead based paint poisoning today is taking the bulk of the burden and the cost. The government that this proposed regulation aims to protect is not unborn. This parents are unborn. It would be impossible to surmise that warning in-

less would have any significant impact on the future children of America. Moreover, there is no evidence that warning labels are effective for their intended purpose. The parents of the children and the children will have no opportunity to read the warning labels of the paint.

It is not necessary to weigh the advantages of lead based paint against the health and welfare of the Nation's children and the future children. The paint industry can produce paint without lead (except for matrix traces) as measured by the atomic absorption method. Furthermore, therefore, urge the Commission of Food and Drugs to proceed with all due speed to publish this proposal in the Federal Register in order that all interested persons may present their views thereon.

Interested persons may, within 90 days after publication hereof in the Federal Register, file with the Hearing Clerk, Department of Health, Education, and Welfare, Room 4-01, 5600 Fishers Lane, Rockville, Md. 20851, written comments (preferably in quadruplicate) regarding this proposal. Comments may be accompanied by a memorandum or brief in support thereof. Received comments may be seen in the above office during working hours, Monday through Friday.

Dated: October 22, 1971.

CHARLES C. EDWARDS,
Commissioner of Food and Drugs.
[Fm Doc. 71-1000 Filed 11-1-71; 9:58 am]

DEPARTMENT OF TRANSPORTATION

National Highway Traffic Safety
Administration

[49 CFR Part 566]

(Docket No. 71-11; Notice 1)

MANUFACTURER IDENTIFICATION

Proposed Motor Vehicle Safety Regulations

The purpose of this proposed amendment to Part 566 in Title 49, Code of Federal Regulations, is to provide for the coverage of "incomplete vehicles," as defined in Part 569, Vehicles Manufactured in Two or More Stages.

A previous notice of proposed rule making published on April 23, 1971 (36 F.R. 7979), proposed a new Part 566 that would require manufacturers of motor vehicles and of motor vehicle equipment to which a motor vehicle safety standard applies to submit identifying information and a description of the items which they produce. A notice of issuance of that part is published in this issue of the Federal Register, 34 F.R. 5677.

In responding to a comment from an incomplete vehicle manufacturer, it was noted that while the proposed regulation clearly covers intermediate and final-stage manufacturers (as defined in Part 569) it makes no reference to incomplete vehicle manufacturers. The

proposed amendment is intended to clarify this ambiguity by specifically providing for coverage of incomplete vehicles.

The incomplete vehicle manufacturer stated that he was unaware of the final use of his light truck vehicles and requested that he be permitted to submit a brief description of the incomplete vehicle expressed in the terminology of the industry as an alternative to the description in terms of final use. This proposed alternative for incomplete vehicle manufacturers has been found acceptable, and the NHTSA accordingly proposes to grant this request.

In consideration of the foregoing the NHTSA proposes that the following amendments be made to Part 566 of Title 49, Code of Federal Regulations:

1. § 566.4 would be amended to read as follows:

§ 566.4 Definitions.

All terms defined in the Act and the rules and standards issued under its authority are used as defined therein. Specifically, "incomplete vehicle", "intermediate manufacturer", and "final-stage manufacturer" are used as defined in Part 569, Vehicles Manufactured in Two or More Stages.

2. Paragraph (c) (3) of § 566.5 would be amended to read as follows:

§ 566.5 Requirements.

Each manufacturer of motor vehicles, and each manufacturer of covered equipment, shall furnish the information specified in paragraphs (a) through (c) of this section to: Administrator, National Highway Traffic Safety Administration, 400 Seventh Street SW, Washington, DC 20590.

(a) . . .

(3) In the case of motor vehicles produced in two or more stages, if the manufacturer is an incomplete vehicle manufacturer, the description shall so state and include a description indicating the stage of completion of the vehicle and, where known, the type of use for which the vehicle is intended.

EXAMPLE: "Chassis-cab", "van-type truck".

If the manufacturer is an intermediate manufacturer, or a final-stage manufacturer, the description shall so state and include a brief description of the work performed.

EXAMPLE: "Manufactured passenger vehicle: Motor home with GVWR from 5,000 to 10,000 pounds. Final-stage manufacturer—add body to bare chassis".

Proposed effective date: February 1, 1972.

Interested persons are invited to submit comments on the proposed amendment. Comments should identify the docket number and be submitted to: Docket Section, National Highway Traffic Safety Administration, Room 3231, 400 Seventh Street SW, Washington, DC 20590. It is requested but not required that 10 copies be submitted.