Senator NELSON. I think the system has been a tragic and colossal failure. The scientists attacked Rachael Carson because she had not accumulated scientific evidence, which to the scientist means if the patient dies the hypothesis was correct. Well, it turned out that her whole thesis was correct, but we are late now in dealing with the problem of DDT in many States. The community is saying, for heavens sake, let's not destroy the insects, fish, animals, and birds, but it is awfully late in the game. That is what concerns me.

Dr. GODDARD. I think Rachael Carson actually did a very fine service in the very forceful manner in which she brought before the public and the Congress some of the abuses that are going on, just as Ralph Nader and others did.

Senator NELSON. Go ahead.

Dr. GODDARD. I would like to submit for the committee's information, Senator Nelson, this copy of our Current Good Manufacturing Practices regulations.

Senator NELSON. Is that a report to be printed in the record?

Dr. GODDARD. If it is your desire, sir.

Senator NELSON. We will print it in the record.

(The information referred to follows:)

PART 133—DRUGS; CURRENT GOOD MANUFACTURING PRACTICE IN MANUFACTURE, PROCESSING, PACKING, OR HOLDING

DEFINITIONS

Sec.
133.1 Definitions.

FINISHED PHARMACEUTICALS; MANUFACTURING PRACTICE

133.2 Current good manufacturing practice.
133.3 Buildings.
133.4 Equipment.
133.5 Personnel.
133.6 Components.
133.7 Master formula and batch-production records.
133.8 Production and control procedures.
133.9 Product containers.
133.10 Packaging and labeling.
133.11 Laboratory controls.
133.12 Distribution records.
133.13 Stability.
133.14 Complaint files.

MEDICATED FEEDS; MANUFACTURING PRACTICE

133.100 Current good manufacturing practice.
133.101 Buildings.
133.102 Equipment.
133.103 Personnel.
133.104 Components.
133.105 Formula and production records.
133.106 Production and control procedures.
133.107 Packaging and labeling.
133.108 Laboratory controls.
133.109 Distribution records.
133.110 Complaint files.

AUTHORITY: §§ 133.1 to 133.110 issued under secs. 501, 701; 52 Stat. 1050 as amended 76 Stat. 780, 781; 1055; 21 U.S.C.A. 351, 371.

DEFINITIONS

§ 133.1 Definitions.

(a) As used in this Part 133, "act" means the Federal Food, Drug, and Cosmetic Act, sections 201–902, 52 Stat. 1052 (21 U.S.C. 321–392), with all amendments thereto.

(b) The definitions and interpretations contained in section 201 of the Federal Food, Drug, and Cosmetic Act shall be applicable to such terms when used in the regulations in this Part 133.

(c) As used in this Part 133, the term "medicated feed" means any "complete feed," "feed additive supplement," or "feed additive concentrate," as defined in