

SHIPPED: The Model G device was transported during 1955 or 1956, by Dr. C. W. Harper of Bethany, Okla., from L. L. Roby Mfg. Co., Tiffin, Ohio, and the *Anapathic device* was shipped in May 1950, from Kansas City, Mo., by Radiation Laboratories, Inc.

LABEL IN PART: "Electronic Magnetic Model G" and "Anapathic * * * Radiation Laboratories, Inc."

ACCOMPANYING LABELING: Leaflets entitled "Electronic Magnetic Instrument Model G * * * Prices F.O.B. Tiffin Ohio," "Electronic Magnetic Instrument Model G * * * Instructions"; and sets of instructions entitled "Anapathic Instructions."

RESULTS OF INVESTIGATION: The *Electronic Magnetic Instrument* contained components to provide an electronic circuit of low voltage, low frequency faradic current, and a magnetic circuit of low voltage electromagnetic energy of 220 cycles per second.

The *Anapathic device* was a wooden cabinet approximately 24" x 20" x 12" with an instrument control panel. The panel face contained two wells labeled "specimen" and "anapathic," an anapathic switch labeled "Neut 1" "Neut 2" "off" "scan" and "therapy," a power on-off switch, a start switch, power and scan lights and test switch number 0, 1, 2, 3, 4, two electrode pads approximately 4" x 5", with twin leads to plug into the panel which were labeled "collector" and "therapy." In the upper center of the front panel was a large timing dial. The back panel had prongs to which the power cord and ground leads were connected. There was also a 115v. AC convenience outlet.

LIBELED: 12-26-57, W. Dist. Okla.

CHARGE: 502(a)—when shipped, the labeling which accompanied the articles contained false and misleading representations that the Model G device was capable of treating body areas of congestion, inflammation and irritation; and of supplying stimulating energy to the body; and that the *Anapathic device* was capable of amplifying body energy; of implanting body energy in a media which could be used in the treatment of acute and chronic conditions; and of treating undulant fever, Streptococcus and meningococcus infections, carcinoma, tuberculosis, all colon disturbances, gas and all metallic poisons.

DISPOSITION: On 3-17-58, a default decree of condemnation was entered against the *Anapathic device* and it was delivered to the Food and Drug Administration.

C. W. Harper appeared and filed a claim for the Electromagnetic device.

Thereafter, the Government filed written interrogatories. Following the filing of answers to the interrogatories the Government filed a motion for summary judgment which was sustained on 5-27-60.

On 6-21-60, the Electromagnetic device was ordered condemned and delivered to the Food and Drug Administration.

DRUG FOR VETERINARY USE*

6380. Worm control preparation (veterinary). (F.D.C. No. 44302. S. No. 35-342 P.)

INFORMATION FILED: 5-13-60, M. Dist. Tenn., against John W. Griffith, t/a Blue Ace, Nashville, Tenn.

SHIPPED: 4-14-58, from Tennessee to Pennsylvania.

*See also No. 6362.

LABEL IN PART: (Btl.) "Blue Ace 1 Qt. Active Ingredients—Combined Iodine 3% Nicotine 18%—Inert Ingredients 79% Blue Ace Nashville, Tennessee Poultry—Turkey—Broiler (Large Round) Worm Control Preparation."

ACCOMPANYING LABELING: Circular entitled "Blue Ace from Incubator to Maturity."

CHARGE: 502(a)—when shipped, the labeling contained false and misleading representations that the article would be adequate and effective in the treatment of worms in poultry; would provide larger, better quality eggs with harder shells; would contribute to greater egg production, more weight, and to lowered mortality rate in poultry; and that the use of the drug in controlling large roundworms would result in greater profits.

PLEA: Guilty.

DISPOSITION: 10-11-60. \$100 fine and probation for 1 year.

INDEX TO NOTICES OF JUDGMENT D.D.N.J. NOS. 6361 TO 6380

PRODUCTS

	N.J. No.		N.J. No.
Allergy capsules-----	6364	Lone Indian Herb Tonic-----	6368
Amphetamine, -dextro, sulfate tablets-----	¹ 6367	Lumbago, remedy for. <i>See</i> Rheumatism, remedy for.	
sulfate tablets-----	¹ 6367	Neuralgia, remedy for. <i>See</i> Rheumatism, remedy for.	
Anapathic device-----	6379	Neuritis, remedy for. <i>See</i> Rheu- matism, remedy for.	
Appetite depressant tablets-----	6374	Obesity, remedies for. <i>See</i> Re- ducing preparations.	
Arthritis, remedy for. <i>See</i> Rheu- matism, remedy for.		Penicillin G potassium tablets---	6365
Black Widow Smear (veteri- nary)-----	6362	Prophylactics, rubber-----	6372, 6373
Bursitis, remedy for. <i>See</i> Rheu- matism, remedy for.		Reducing preparations-----	6362, 6374
Contact lens wetting solution---	6371	Rheumatism, remedy for-----	6368
Delfetamine tablets-----	6363	S. F. vitamin tablets with phenyl- propanolamine-----	6374
Devices--- 6361, 6372, 6373, 6376,	6377	Scholl's, Dr., Foot Exercizer (sandals)-----	6377
Dexobarbital tablets-----	¹ 6367	Sciatica, remedy for. <i>See</i> Rheu- matism, remedy for.	
Dextro-amphetamine sulfate tab- lets-----	¹ 6367	Sopamycetin chloramphenicol capsules-----	6366
Electro-galvanic bracelets-----	6376	Thiede's head harness-----	6361
Electronic Magnetic Instrument.	6379	Trim-All capsules-----	6364
Filtronair Air Purifier device---	6378	Vegetable juices-----	² 6375
Folabin-----	6370	Veterinary preparations-----	6362, 6380
Gout, remedy for. <i>See</i> Rheu- matism, remedy for.		Vitamin preparations-----	6370, 6374
Herb Tonic, Lone Indian-----	6368	Worm control preparation (vet- erinary)-----	6380
Laxative without required warn- ing statement-----	6368		
Lobelia herb-----	6369		

¹ (6367) Seizure contested.

² (6375) Injunction issued.

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U.S. Department of Health, Education, and Welfare
FOOD AND DRUG ADMINISTRATION

**NOTICES OF JUDGMENT UNDER THE FEDERAL FOOD,
DRUG, AND COSMETIC ACT**

[Given pursuant to section 705 of the Food, Drug, and Cosmetic Act]

6401-6420

DRUGS AND DEVICES

The cases reported herewith were instituted in the United States district courts by United States attorneys, acting upon reports submitted by the Department of Health, Education, and Welfare. They involve drugs which are required at time of interstate shipment to bear a label containing the statement "Caution: Federal law prohibits dispensing without prescription," and which were dispensed after such shipment without a prescription or by refilling a prescription without authorization. This dispensing was contrary to Section 503(b)(1) and thereby resulted in the dispensed drugs being misbranded while held for sale.

Published by direction of the Secretary of Health, Education, and Welfare.

GEO. P. LARRICK, *Commissioner of Food and Drugs.*

WASHINGTON, D.C., *July 28, 1961.*

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