UNITED NATIONS

E/NL 1955/84
February 1956
ORIGINAL: English

LAWS AND REGULATIONS

PROMULGATED TO GIVE EFFECT TO THE PROVISIONS OF THE CONVENTION OF 13 JULY 1931 FOR LIMITING THE MANUFACTURE AND REGULATING THE DISTRIBUTION OF NARCOTIC DRUGS, AS AMENDED BY THE PROTOCOL OF 11 DECEMBER 1946

ISRAEL

Communicated by the Government of Israel

NOTE BY THE SECRETARY-GENERAL -- In accordance with Article 21 of the Convention of 13 July 1931 for Limiting the Manufacture and Regulating the Distribution of Narcotic Drugs, as amended by the Protocol of 11 December 1946, the Secretary-General has the honour to communicate the following legislative text.

SPECIAL CONDITIONS FOR THE MANUFACTURE OF SYNTHETIC DANGEROUS DRUGS*

1) The manufacture of dangerous drugs should be done solely within the factory premises and not without, under the strict supervision of a pharmacist holding an Israeli licence who is personally responsible.

2) Each dangerous drug should be named according to the official pharmacopeia recognized in Israel, and it is not to be indicated by any other name. In case there is no pharmacopeial name, the recognized scientific name should be used.

3) The manufacturer is required to inform the Director of the Ministry of Health in advance, through the District Pharmacist, on the estimated quantity of the dangerous drug to be synthesized.

4) The manufacturer is required to keep a Production Book, the pages of which should each be numerated and stamped with the Ministry of Health's seal. The first and last page should bear the signature of the District Pharmacist on behalf of the District Medical Officer.

5) The manufacturer is required to register in this book all the stages of the synthesis and should include the following items: date, number, batch, quantity of the raw material, residue, transitory substances produced in all stages (each transitory substance should be registered on a separate page), the final residue including absolute and relative figures and the percentage of the dangerous drug remaining in the residue. The residue should be preserved and not destroyed or used without the authorization of the District Pharmacist.

6) The manufacturer is required to register the final quantity of the produced drug in the Dangerous Drug Register (in addition to the Production Book mentioned above) and state besides this figure the respective batch number.

7) On the container or wrappings of each dangerous drug the official (pharmacopeial) name should be inserted in large characters and underlined with a double red line. Under the name of the drug the words "Narcotic Dangerous Drug" should be included.

8) The manufacturer is required to submit to the Director, Pharmacy Division of the Ministry of Health, through the District Pharmacist, annual reports of production figures of each synthetic dangerous drug and the quantities of dangerous drugs used for conversion in the production stages.

9) The manufacture of any preparation containing one or more dangerous drugs, in whatever form, requires an authorization from the Director of the Ministry of Health, according to the special condition set down for the manufacture of medicinal preparations.

Signed

Minister of Health

Note by the Secretary-General:
Transmitted by the Government of Israel by note of 29 September 1955.