Republic of the Philippines
HOUSE OF REPRESENTATIVES
Quezon City
EIGHTEENTH CONGRESS
Third Regular Session
HOUSE BILL NO. 9855

Introduced by HON. LUIS RAYMUND “LRAY” F. VILLAFUERTE, JR.

AN ACT
ESTABLISHING THE DRUG PRICE REGULATORY BOARD TO REGULATE THE
PRICES OF DRUGS AND MEDICINES IN THE PHILIPPINES AMENDING FOR
THE PURPOSE REPUBLIC ACT NO. 9502, OTHERWISE KNOWN AS THE
"UNIVERSALLY ACCESSIBLE CHEAPER AND QUALITY MEDICINES ACT OF
2008" AND FOR OTHER PURPOSES

The Philippine Statistics Authority reports that the average Filipino spent P2,024 on medicines alone in 2018. This equates to P10,120 spent by a family of five and translates to gross sales amounting to P206.7 billion.

The prohibitive price of health care in the Philippines results in self-treatment and self-prescription, and those pose their own problems. The affordability of medicines are especially difficult for those who suffer from debilitating illnesses requiring maintenance drugs. Restraints on costs lead to irregular treatment and complications.

This bill aims to introduce amendments to Republic Act 9502 or the Universally Accessible Cheaper and Quality Medicines Act of 2008 in order to simplify the procedure of lowering prices of essential medicines. It proposes to establish a Drug Price Regulatory Board, which will be tasked to, among others, regulate the prices of drugs and medicines in the country.

In view of the foregoing, passage of this bill is earnestly sought.

LUIS RAYMUND “LRAY” F. VILLAFUERTE, JR.
Republic of the Philippines
HOUSE OF REPRESENTATIVES
Quezon City

EIGHTEENTH CONGRESS
Third Regular Session

HOUSE BILL NO. 9855

Introduced by HON. LUIS RAYMUND “LRAY” F. VILLAFUERTE, JR.

AN ACT
ESTABLISHING THE DRUG PRICE REGULATORY BOARD TO REGULATE THE
PRICES OF DRUGS AND MEDICINES IN THE PHILIPPINES AMENDING FOR
THE PURPOSE REPUBLIC ACT NO. 9502, OTHERWISE KNOWN AS THE
"UNIVERSALLY ACCESSIBLE CHEAPER AND QUALITY MEDICINES ACT OF
2008" AND FOR OTHER PURPOSES

Be it enacted by the Senate and House of Representatives of the Philippines in Congress assembled:

SECTION 1. Chapter 3 of Republic Act No. 9502 will now be titled, "Drugs and Medicines Price Regulatory Board."

SEC. 2. Section 17 to Section 22 of Republic Act No. 9502 will be amended to read as follows:

SECTION 17. CREATION AND COMPOSITION OF THE DRUG PRICE REGULATORY BOARD. –

(A) THERE IS HEREBY CREATED THE DRUG PRICE REGULATORY BOARD, WHICH SHALL BE ATTACHED TO THE DEPARTMENT OF HEALTH AND COMPOSED OF SEVEN (7) MEMBERS AS FOLLOWS:

(1) SECRETARY OF HEALTH OR HIS DULY DESIGNATED REPRESENTATIVE AS CHAIRPERSON;
(2) SECRETARY OF TRADE AND INDUSTRY OR HIS DULY DESIGNATED REPRESENTATIVE AS VICE CHAIRPERSON;
(3) DIRECTOR, FOOD AND DRUGS ADMINISTRATION OR HIS DULY DESIGNATED REPRESENTATIVE AS MEMBER;
(4) CHAIRMAN, PHILIPPINE HEALTH INSURANCE CORPORATION AS MEMBER;
(5) ONE (1) ECONOMIST FROM THE ACADEME AS MEMBER; AND
(6) TWO (2) REPRESENTATIVES FROM THE CONSUMER SECTOR AS MEMBERS.

SECTION 18. POWERS OF THE BOARD. – THE BOARD SHALL HAVE THE FOLLOWING POWERS:

(A) POWER TO DETERMINE THE MAXIMUM RETAIL PRICE OF DRUGS OR MEDICINES SUBJECT TO PRICE REGULATION –

(1) UPON APPLICATION OR MOTU PROPIO WHEN THE PUBLIC INTEREST SO REQUIRES, THE BOARD SHALL HAVE THE POWER TO REGULATE THE RETAIL PRICES OF DRUGS AND MEDICINES LISTED UNDER SECTION 19 HEREOF, INCLUDING THEIR DOSAGE FORM AND PACKING, AND, IN ORDER THAT THEY SHALL BE MADE AVAILABLE TO THE PUBLIC AT AFFORDABLE RETAIL PRICE FROM THE DIFFERENT MANUFACTURERS, IMPORTERS, TRADERS, DISTRIBUTORS, WHOLESALERS OR RETAILERS AND AFTER A PROPER DETERMINATION AS THE BOARD MAY DEEM FIT, FIX FROM TIME TO TIME, BY PUBLICATION THE MAXIMUM RETAIL PRICE AT WHICH SUCH FORMULATIONS SHALL BE SOLD;

(2) NO RETAILER SHALL SELL DRUGS AND MEDICINES AT A RETAIL PRICE EXCEEDING THE MAXIMUM RETAIL PRICE FIXED BY THE BOARD: PROVIDED, THAT UNTIL THE MAXIMUM RETAIL PRICE OF DRUGS AND MEDICINES SUBJECT TO PRICE REGULATION IS FIXED BY THE BOARD, THE RETAIL PRICE THEREOF SHALL BE THE PRICE WHICH PREVAILED IMMEDIATELY BEFORE THE EFFECTIVITY OF THIS ACT AND NO MANUFACTURER, IMPORTER, TRADER, DISTRIBUTOR, WHOLESALER OR RETAILER OF SUCH DRUG OR MEDICINE SHALL SELL THE SAME AT A RETAIL PRICE EXCEEDING THE PRICE PREVAILING IMMEDIATELY BEFORE THE EFFECTIVITY OF THIS ACT.

FOR PURPOSES, HEREOF, DRUGS AND MEDICINES SHALL INCLUDE BUT IS NOT LIMITED TO SINGLE- AND MULTI-INGREDIENT MEDICINES INCLUDED IN THE PHILIPPINE NATIONAL DRUG FORMULARY (PNDF) ESSENTIAL DRUG LIST AND SOLD UNDER THEIR GENERIC AND BRAND NAMES.

(B) POWER TO INCLUDE OTHER DRUGS OR MEDICINES IN THE LIST SUBJECT TO PRICE REGULATION – UPON APPLICATION OR MOTU PROPIO WHEN THE PUBLIC INTEREST SO REQUIRES AND AFTER PROPER DETERMINATION, THE BOARD MAY ORDER THE INCLUSION OF DRUGS AND MEDICINES TO THE LIST SUBJECT TO PRICE REGULATION UNDER SECTION 19 THEREOF.

(C) POWER TO IMPLEMENT COST-CONTAINMENT AND OTHER MEASURES –
(1) THE BOARD SHALL HAVE THE POWER TO DETERMINE THE FAIR PRICE OF DRUGS OR MEDICINES FOR PURPOSES OF PUBLIC HEALTH INSURANCE AND GOVERNMENT PROCUREMENT; AND

(2) THE BOARD SHALL HAVE THE POWER TO IMPLEMENT ANY OTHER MEASURES THAT THE GOVERNMENT MAY AVAL OF TO EFFECTIVELY REDUCE THE COST OF DRUGS OR MEDICINES THAT SHALL INCLUDE, BUT IS NOT LIMITED TO, COMPETITIVE BIDDING, PRICE-VOLUME NEGOTIATIONS, AND OTHER APPROPRIATE MECHANISMS THAT INFLUENCE SUPPLY, DEMAND, AND EXPENDITURES ON DRUGS AND MEDICINES.

(D) POWER TO IMPOSE ADMINISTRATIVE FINES AND PENALTIES – AFTER DUE NOTICE AND HEARING, THE BOARD SHALL HAVE THE POWER TO SUSPEND OR REVOKE THE LICENSE TO OPERATE (LTO), PROFESSIONAL OR BUSINESS LICENSE, AS THE CASE MAY BE, OF ANY PERSON, MANUFACTURER, IMPORTER, TRADER, DISTRIBUTOR, WHOLESALER, RETAILER, OR ANY OTHER ENTITY, AND IMPOSE ADMINISTRATIVE FINES IN SUCH AMOUNT AS IT MAY DEEM REASONABLE WHICH SHALL IN NO CASE BE LESS THAN FIFTY THOUSAND PESOS (P50,000.00) NOR MORE THAN FIVE MILLION PESOS (P5,000,000.00) FOR VIOLATIONS OF THE MAXIMUM RETAIL PRICE FIXED PURSUANT TO THIS SECTION.

SEC. 3. Delete Section 19 to Section 22 and renumber the succeeding sections accordingly.

SEC. 4. The renumbered Section 22 will be amended to read as follows:

SECTION 22. DISPLAY OF PRICE FIXED BY THE BOARD FOR DRUGS OR MEDICINES SUBJECT TO PRICE REGULATION. –

(a) Within a reasonable period as may be determined by the [Secretary of the Department of Health] BOARD, and Provided, That it conforms to existing drug product labeling requirements, every manufacturer, importer, distributor, wholesaler, trader, or retailer of a drug and medicine intended for sale shall display the retail price which shall not exceed the maximum retail price [approved by order of the President of the Philippines] FIXED BY THE BOARD. The maximum retail price shall be printed on the label of the immediate container of the drug and medicine and the minimum pack thereof offered for retail sale with the words "RETAIL PRICE NOT TO EXCEED" preceding it, and "UNDER DRUG PRICE REGULATION" on a red strip: PROVIDED, THAT IN CASE OF A CONTAINER CONSISTING OF SMALLER SALEABLE PACKS, THE RETAIL PRICE OF SUCH SMALLER PACK SHALL ALSO BE DISPLAYED ON THE LABEL OF EACH SMALLER PACK AND SUCH PRICE SHALL NOT BE MORE THAN THE PRO-RATA RETAIL PRICE OF THE MAIN PACK ROUNDED OFF TO THE NEAREST CENTAVO.

(b) Within a period as may be determined by the [Secretary of the Department of Health] BOARD from time to time, every manufacturer, importer, or trader shall issue a price list to wholesalers, distributors, retailers and to the [Secretary of the
Department of Health] BOARD, indicating the retail price, the maximum retail price, and such other information as may be required by the [Secretary of the Department of Health] BOARD.

SEC. 5. Insert a new Section 23 and renumber the succeeding sections accordingly. The new Section 23 shall read as follows:

SECTION 23. DISPLAY OF PRICE LIST OF DRUGS OR MEDICINES EXCLUDED FROM THE LIST SUBJECT TO PRICE REGULATION. – EVERY MANUFACTURER, IMPORTER, TRADER, DISTRIBUTOR, WHOLESALER, OR RETAILER OF A DRUG OR MEDICINE EXCLUDED FROM THE LIST SUBJECT TO PRICE REGULATION UNDER SECTION 19 HEREOF SHALL DISPLAY IN INDELIBLE PRINT MARK ON THE LABEL OF THE IMMEDIATE CONTAINER OF THE DRUG OR MEDICINE AND THE MINIMUM PACK THEREOF OFFERED FOR RETAIL SALE, THE WORDS "NOT UNDER PRICE REGULATION" ON A GREEN STRIP.

SEC. 6. Chapter 8 on Miscellaneous Provisions will be amended to read as follows:

CHAPTER 8: AMENDMENTS TO REPUBLIC ACT 9994 OR THE EXPANDED SENIOR CITIZENS ACT OF 2010.

SECTION 42. EXEMPTION OF DRUGS AND MEDICINES UNDER PRICE REGULATION FROM THE "EXPANDED SENIOR CITIZENS ACT OF 2010." – DRUGS AND MEDICINES UNDER PRICE REGULATION AS FIXED BY THE BOARD WILL NOT BE INCLUDED IN THE GRANT OF TWENTY PERCENT (20%) DISCOUNT AND EXEMPTION FROM THE VALUE-ADDED TAX (VAT) TO SENIOR CITIZENS AVAILING THE PROVISIONS OF REPUBLIC ACT 9994 OR THE "EXPANDED SENIOR CITIZENS ACT OF 2010."

SEC. 7. Separability Clause. – Any portion or provision of this Act that may be declared unconstitutional or invalid shall not have the effect of nullifying other portions and provisions hereof as long as such remaining portions or provisions can still subsist and be given effect in its entirety.

SEC. 8. Repealing Clause. – All laws, decrees, executive orders, proclamations and administrative regulations or parts thereof inconsistent herewith are hereby repealed or modified accordingly.

SEC. 9. Effectivity Clause. – This Act shall take effect fifteen (15) days after its publication in at least two national newspapers of general circulation.

Approved,