

Republic of the Philippines  
**HOUSE OF REPRESENTATIVES**  
Quezon City

EIGHTEENTH CONGRESS  
First Regular Session

HOUSE BILL NO. 6411



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Introduced By Representative ALLAN BENEDICT S. REYES

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**EXPLANATORY NOTE**

It is a Constitutional guaranty that the State shall protect and promote the health of the people, instill among them a consciousness about healthy living, and ensure that every citizen lives in a balanced and healthful ecology in accord with the rhythm and harmony of nature.

Over the last decade, there is a growing concern about the possible health hazards from genetically modified crops and its derivatives that are directly and indirectly consumed by people as food and beverage. In the Philippines, local ordinances banning the planting of Genetically Modified Organisms (GMOs) like in Bohol, Mindoro, Benguet, Negros, have been enacted after the local government saw the possible negative effects of GMOs on the environment and on the health of the people.

Internationally, a number of nations, more particularly those from the European Union, has banned or restricted the planting and the entry of genetically modified organisms (GMOs) into their food system. To date, 64 countries around the world have imposed mandatory labeling of food and beverages that contain ingredients from GMO crops. GMO corn, soya, canola, and their derivatives, among other GMO crops, are used as feeds for livestock or ingredients for processed foods, and eventually become an integral component of processed and fresh food for human consumption.

It is thus imperative to address the dangers brought about by GMOs through appropriate regulation, awareness campaign, and mandatory labeling of food and beverages and from where and how they have been derived to assure safety in public's consumption as mandated by Section 12, Article XIII of the Constitution, to wit:

The State shall establish and maintain an effective food and drug regulatory system and undertake appropriate health, manpower development, and research, responsive to the country's health needs and problems.

Hence, the filing of this urgent health measure is earnestly sought.

*ABR*  
HON. ALLAN BENEDICT S. REYES  
Representative  
3rd District, Quezon City

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**AN ACT TO BE KNOWN AS “THE PHILIPPINE GENETICALLY MODIFIED ORGANISM (GMO) LABELING ACT, THE RIGHT-TO-KNOW-ACT” REQUIRING THE MANDATORY LABELING AND REGULATION OF FOOD WHICH ARE GMOs OR CONTAINING SUBSTANCES DERIVED FROM GMOs AND THOSE PRODUCED BY GENETIC ENGINEERING TECHNOLOGIES, PROVIDING FUNDS THEREFOR AND FOR OTHER PURPOSES**

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

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2       **SECTION 1. Short Title** – This Act shall be known as, “**THE PHILIPPINE**  
3       **GMO LABELING AND REGULATION ACT, THE RIGHT-TO-KNOW ACT**”  
4

5       **SECTION 2. Declaration of Policy** – It is hereby declared that the State  
6 shall protect and promote the right to health of the people and instill health  
7 consciousness among them. It shall also protect and advance the right of the  
8 people to a balanced and healthy ecology, as it is the duty of the State to inform  
9 and warn the people of the health and environmental risks of Genetically Modified  
10 Organisms or GMOs.

11       The State further guarantees the **establishment and maintenance of an**  
12 **effective food and drug regulatory system** and undertaking the appropriate  
13 health, manpower development, and research, responsive to the country’s health  
14 needs and problems.

15       To this end, the State recognizes the rights of the consumers and general  
16 public on their right to know whether the food, food products and their derivatives  
17 (processed or not) and agricultural products like rice, corn and vegetables, among  
18 others, contain genetically modified organisms (GMOs) or are produced by genetic  
19 engineering technologies (GET). Thus, the State shall endeavor to provide  
20 Filipinos with safe and healthy food and agricultural products that will not  
21 jeopardize their health and place the environment at risk as a result of  
22 developments in food manufacturing that utilizes the process and products of  
23 genetic engineering technologies through an efficient and effective regulation and  
24 labeling.

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### **SECTION 3. Definition of Terms.**

- 4     a) Genetically Modified Organism (GMO) – Living organisms whose genetic make-up  
5       were manipulated in a laboratory by non-natural means such as genetic engineering  
6       technologies in order to incorporate genes from different species that normally will not  
7       reproduce with each other. An example of this is a corn plant whose genes were  
8       inserted with a gene from a bacterium such as the Bacillus thuringiensis or Bt that acts  
9       as a pesticide to kill corn borers.
- 10    b) Genetic Engineering (GE) -- The process of inserting a foreign gene into a host  
11       organism to achieve a trait not normally held by that organism, such as the ability to  
12       produce its own pesticides, or to be immune to certain herbicides.
- 13    c) Genetically Modified Food (GM Food) – Processed food that contain GMOs, or  
14       products derived from them, fresh meat and edible plants that are modified through  
15       genetic engineering.
- 16    d) Genetically Modified Crops (GM Crops) – Refer to agricultural crops like, but not limited  
17       to Bt Corn, Round Up-Ready (RR) soya, RR canola, Bt cotton, etc., which contain  
18       genes of other organisms inserted in their own genes thru genetic engineering.
- 19    e) Genetically Modified Feed (GM Feed) – Any agricultural crop like corn, soya, etc. that  
20       are genetically modified and used as feed for the livestock industry
- 21    f) Livestock – Any animal, bird, fish, etc, that are raised as an industry for human  
22       consumption.
- 23    g) Derivative – Refers to products derived from processing GMO crops, such as but not  
24       limited to, extracts, oil, syrup, powder, flour, etc.
- 25    h) Food – In addition to Republic Act 9711 or the Food and Drug Administration Act of  
26       2009's definition of food, it is any natural and processed food intended for human  
27       consumption that includes agricultural crops like rice, corn, vegetables, among others;  
28       beverages, whether natural or processed; and any substance, whether natural or  
29       processed which has been used as an ingredient in the manufacture, preparation or  
30       treatment of food.

- 1      i) Processed Food – Any food that has undergone any procedure that has changed it  
2      and made it different from its natural state. Examples of these procedures are  
3      preserving, packaging, irradiation, pasteurization, fortifying, genetically modifying, etc.
- 4      j) Threshold Level -- the percentage limit of traceable GMO contained in all food, crops,  
5      feeds and derivatives

6  
7      **SECTION 4. Coverage and Scope** – The following persons, whether natural or  
8      juridical, are covered by this Act:

- 9            a. Any person who are involved in the development, production and/or sale  
10          of GMO agricultural crops, GMO marine products and other unnatural  
11          substances obtained by genetic engineering technologies, as defined by  
12          this Act, that are intended for propagation and commercialization;
- 13            b. Any person who are involved in the propagation, planting, marketing  
14          and/or sale of the items, in whole or in part, as stated in Section 4.a. and  
15          defined by this Act;
- 16            c. Any person who are involved in the manufacture and processing of  
17          derivatives, as defined by this Act, that are sourced from the items as  
18          stated in Section 4.a. and defined by this Act;
- 19            d. Any person who are involved in the marketing and sale of derivatives, as  
20          defined by this Act;
- 21            e. Any person who are involved in the production, marketing, distribution  
22          and/or sale of food and beverage that are sourced from the items as  
23          stated in Section 4.a. and defined by this Act; and
- 24  
25            f. Any person who are involved in importing or bringing in Genetically  
26          Modified food, crops, animals, feeds and derivatives, as defined by this  
27          Act, into the Philippines.

28  
29      **SECTION 5. Mandatory Labeling of Food, Food Products and their Derivatives**  
30      **and Agricultural Products like rice, corn and vegetables, among others, that contain**  
31      **GMOs.** – All food, food products that have been processed and their derivatives and  
32      agricultural products like rice, corn and vegetables, among others, containing genetically  
33      modified organisms or GMOs, as well as those produced by genetic engineering  
34      technologies shall be labeled in accordance with the provisions of this Act and the existing  
35      regulations issued by the Department of Health through the Food and Drug Administration  
36      (FDA).

- 1           a. For the regulation and mandatory labeling of processed food and  
2           beverages containing GMO as defined by this Act, the Department of  
3           Health, through the FDA, will be the implementing agency;
- 4           b. For the regulation and mandatory labeling of genetically modified crops,  
5           livestock and marine products, some of which have been specifically  
6           approved by Philippine government in the past for commercialization in  
7           the Philippines, and as defined by this Act, the Department of Agriculture,  
8           in collaboration with the FDA, will be the implementing agency;
- 9           c. **"THIS PRODUCT CONTAINS GENETICALLY MODIFIED  
10 ORGANISM"** or **"THIS PRODUCT CONTAINS GMO"** shall be  
11 permanently printed as a stand-alone, conspicuous label in the front and  
12 back sides of the general packaging materials like boxes, sacks, etc.,  
13 containing the bulk of the food, beverages and all other items, and  
14 individual packaging materials of the items as proportioned by the  
15 producer. The size of the printed label shall not be less than five (5)  
16 percent of the total volume area of the container;
- 17           d. In the case of GMO agricultural crops, livestock and marine products, as  
18           defined by this Act whose cultivation/propagation is ongoing, the same  
19           label as stated by Section 5, subsection c of this Act shall be printed on  
20           the front side of a tag, banner, or any other medium where the label can  
21           be printed. The size of the tag, or a corresponding combination of number  
22           and volume thereof, shall not be less than one-half (0.5) square meter  
23           per hectare of the total area of the cultivation/propagation. The size of  
24           the tag shall also be same even if the area is less than one (1) hectare.  
25           The size of the label on the tag shall not be less than forty percent of the  
26           size of the tag;
- 27           e. In the case of GMO agricultural crops, livestock and marine products, as  
28           defined by this Act, that are retailed and displayed as a mass -- like rice,  
29           corn, soya beans, vegetables, fish, meat, etc. – the same label as  
30           mandated by Section 5, subsection c of this Act shall be printed on a tag  
31           that will correspondingly identify the specific item being sold; and
- 32           f. The labeling line shall be as follows:
- 33               1) The producers of seeds for GMO crops shall label their products so  
34               that the farmers will be informed;  
35               2) Farmers who plant GMO crops shall tag their fields and label their  
36               containers so that their buyers will be informed;  
37               3) Buyers of GMO crops, who will sell the same as a mass product,  
38               should tag and label so that the consuming public will be informed;

- 1           4) Buyers and processors of GMO crops for the production of derivatives  
2           as defined by this Act shall label their products so that their food and  
3           beverage-processor buyers will be informed;  
4           5) Food and beverage processors who use GMO crop derivatives as  
5           defined by this Act shall label their products so that the consuming  
6           public will be informed; and  
7           6) Importers of GMO-containing processed or whole food into the  
8           Philippines, including derivatives, all defined by this Act, shall secure  
9           prior clearance from the Authority created by this Act for the effective  
10          administration of the labeling mandate of this Act.

11  
12         **SECTION 6. Threshold level** -- The tolerance level is nine-tenth (0.9) percent.  
13         Above the threshold level, a food is regarded as genetically modified. Hence, must be  
14         properly labeled. The threshold of nine-tenth (0.9) percent still allows for some unintended  
15         contamination of food with GM ingredients.

16  
17         **SECTION 7. Institutional Mechanism** – There is hereby created an office  
18         to be called the GMO CROP, FOOD AND BEVERAGE LABELING COUNCIL  
19         (COUNCIL) that will be attached to the Department of Health, thru the FDA, and  
20         chaired by the same agency. With the mandate to oversee and coordinate the  
21         implementation of this Act, supervise the Authority, the institution of a continuous  
22         information and education campaign on GMOs to be led and undertaken by its  
23         sectoral members and propose policies to Congress for the improvement of this  
24         Act, the Council shall be composed by the following:

- 25           a. Food and Drug Administration Head – Ex-Officio Chairman  
26           b. Department of Agriculture – Ex-Officio  
27           c. Department of Trade and Industry – Ex-Officio  
28           d. Department of Environment and Natural Resources – Ex-Officio  
29           e. Two (2) representatives Academic Sector – Members  
30           f. Two (2) representatives Consumer Sector – Members  
31           g. Two (2) representatives Peasant Sector – Members  
32           h. Two (2) representatives Health Advocacy Sector – Members  
33           i. Two (2) representatives Business Sector – Members

34         The sectoral members of the Council shall be selected by the FDA through  
35         a nomination process from the respective sector.

36  
37         **SECTION 8. ACCREDITATION OF SCIENTIFIC LABORATORIES** – All scientific  
38         laboratories operating in the Philippines, as well as those located abroad, shall secure  
39         prior accreditation to be recognized as lawful and competent entities that are able to  
40         conduct analyses of GMO presence in food, beverages and other items as defined by this  
41         Act.

42  
43         **SECTION 9. GMO Crop, Food and Beverage Labeling and Regulation Authority**  
44         (Authority) – There is hereby created an office to be called the GMO CROP, FOOD AND  
45         BEVERAGE LABELING AND REGULATION AUTHORITY that will be attached to the  
46         Department of Health, thru the FDA.

- 47           a) Functions:

- 1        1. Even without prior notice, to visit within the Philippines the offices,  
2           facilities, or farms, etc., of all the entities involved in all the articles  
3           defined and cited by this Act;
- 4        2. To enlist and secure the assistance of any government agency,  
5           including the police, in the performance of its functions;
- 6        3. After due process, to establish the evidence and culpability of the  
7           violators of this Act;
- 8        4. With warranted evidence, to file the case against the violators before  
9           the court of law;
- 10      5. To conduct *motu proprio* investigations against violators;
- 11      6. To act immediately on all verified reports of violations;
- 12      7. To issue subpoena duces tecum and ad testificandum to any parties  
13           in the complaint;
- 14      8. To conduct regular research on GMOs and track the uses of the same  
15           in the food chain;
- 16      9. To maintain a database of its research and other documents. It shall  
17           post in its website all the labeled articles in the Philippines by virtue  
18           of this Act. It shall also post all the articles and products all over the  
19           world that are equally labeled, including the identification and tracking  
20           of derivatives of GMO crops as defined by this Act;
- 21      10. After due process, to collect all the imposable penalties and fines from  
22           the violators;
- 23      11. To regulate and approve all the applications for the label as defined  
24           and mandated by this Act. The label becomes lawful when it is  
25           accompanied by the logo of this Authority;
- 26      12. To validate the label annually from the date of approval;
- 27      13. To collect a nominal fee annually for the application, processing,  
28           validation, approval and administration of the label. The fee shall be  
29           determined and approved by the Council;
- 30      14. Prior to importation, to verify and impose a clearing mechanism on  
31           all processed and whole food importations into the Philippines and  
32           collect a nominal fee therefor. The fee shall be determined and  
33           approved by the Council;
- 34      15. To accredit scientific laboratories and collect a nominal fee  
35           therefrom. The fee shall be determined and approved by the  
36           Council; and
- 37      16. To acquire the necessary competence by hiring expertise and  
38           purchasing scientific equipment to ascertain the presence of GMO  
39           in all the products and articles as defined by this Act. This shall be  
40           determined and approved by the Council.

41           b) Officers, staff and domicile:

- 42        1. The Authority will be headed by a Director. The Director reports  
43           directly to the head of the FDA, who is also the chair of the Council;
- 44        2. The Authority shall have an Assistant Director, who reports directly to  
45           the Director, to act as the officer-in-charge in the absence or  
46           incapacity of the Director;

3. The salaries of the Director and Assistant Director shall be commensurate to the promulgated rates for civil service;
4. The employment of the Director and Assistant Director shall be processed, scrutinized and recommended for approval by the Council. The Secretary of Health shall approve the said recommended employment and administer the oath of office;
5. The Authority shall have an initial three (3) administrative staff. Upon justification for any additional staff at any given time, the Council will process, scrutinize and approve the same; and
6. The office of the Authority shall be located in FDA.

**SECTION 10. Inapplicability of Regulation-Making Exemptions.** – Due to the urgent and extreme public necessity and the need to keep the public adequately protected and to preserve the right of the consumers to choose, the power of the Secretaries of Trade and Industry, and Health to make exemptions to the regulations as specified in the Consumer Act of the Philippines and the FDA Act is hereby rendered inapplicable unless it can be shown that public order, public health and public safety are not placed in jeopardy with its exemption, actual application and implementation.

**SECTION 11. Penalties.** –Any person who knowingly distribute GMO plants, food, food products and their derivatives and agricultural products such as rice, corn and vegetables, among others, containing or derived from GMOs without the proper label shall be subject to a fine per improper label of not less than five hundred thousand pesos (P500,000.00) and imprisonment ranging from six years and one day to twelve years and/or revocation of license to operate upon the discretion of the court.

**SECTION 12. Funding** – For the effective implementation of this Act, there is hereby appropriated an initial fund of ten million pesos (Php 10,000,000.00) for the administration and operation of the Council and the Authority, of which the Council shall be funded with two million pesos (Php 2,000,000.00) and the rest consists the fund for the Authority.

The total initial fund of ten million pesos (Php 10,000,000.00) shared by the Council and Authority, which shall be rated by future Congressional appropriations at twenty percent (20%) for the Council and eighty percent (80%) for the Authority, shall be sourced initially, during the first fiscal year of the effectivity of this Act, from the funds of the Department of Health and Department of Agriculture at five million pesos (Php 5,000,000.00) each.

The succeeding, combined funding requirement of the Council and Authority, rated at 20:80 percent, respectively, shall be determined and approved by the Council. The two bodies shall submit to Congress their budget proposals for final approval and appropriation.

To augment the funds of the Council and Authority for the purpose of implementing this Act, the monetary penalties, imposable fees and all other monies and resources collected by the Authority shall always remain in the coffers of the respective bodies for the sole purpose of the effective implementation of this Act. The accruing funds, including

1 the annual appropriated budget therefor, shall build the competency, capability and  
2 efficiency of the Council and Authority in the implementation of this Act.

3 And where the two bodies have sufficiently generated their own funds, determined  
4 through annual audited reports to Congress, they shall momentarily cease from proposing  
5 for appropriated funds until their own funds can last.

6  
7       **SECTION 13. Implementing Rules and Regulations.** – The Council and the  
8 Authority, with the participation of the appropriate government agency and in consultation  
9 with the concerned business and industry representatives, government agencies, civil  
10 society organizations and people's organizations, shall promulgate the necessary  
11 regulations for the effective implementation of this Act within six (6) months from the date  
12 of the passage of this Act.  
13

14       **SECTION 14. Repealing Clause.** – All laws, decrees, orders, rules and regulations,  
15 policies, programs or parts thereof, that are inconsistent with any of the provisions of this  
16 Act, are hereby repealed, amended or modified accordingly.

17       **SECTION 15. Separability Clause.** – If for any reason, any section or provisions  
18 of this Act is declared to be unconstitutional or invalid, the other sections or provisions  
19 hereof, which are not affected hereby, shall continue in full force and effect.  
20

21       **SECTION 16. Effectivity Clause.** – This Act shall take effect fifteen (15) days after  
22 its publication the *Official Gazette* or in the two (2) national newspaper of general  
23 circulation.  
24  
25  
26       Approved,