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Report Highlights:

Sri Lanka does not produce genetically engineered (GE) crops or animals. Some GE research is carried out at the laboratory level, but no development exists at a commercial level. The country does not allow the import of GE food, crops, or animals. Sri Lanka signed and ratified the Cartagena Protocol on Biosafety in 2004. Sri Lanka is in the process of developing policies to regulate and promote biotechnology. Most of the policies are incomplete, or not fully implemented.

Executive Summary

The United States and Sri Lanka enjoy a healthy agricultural trade relationship. Although the country does not allow imports of food, crops, animals, or agricultural products derived from genetic engineering, the United States exported 146 million U.S. dollars' worth of agricultural commodities to Sri Lanka in 2018.

Sri Lankan trade regulations require mandatory labeling of GE ingredients for imported goods, and approval of imports for food products containing more than 0.5 percent (by volume) of GE-derived ingredients. GE-free certification is required for crops with "Genetically Modified (GM)" varieties. However, the absence of a functioning approval mechanism in effect leads to a ban on the sale of agricultural products derived from genetic engineering.

No GE crops are produced in Sri Lanka. Some GE research is carried out at the laboratory level, but the research does not reach commercialization. The unavailability of a legal framework and proper biosafety procedures are a major setback.

Sri Lanka is developing multiple policies to regulate and promote biotechnology. These include the National Biotechnology Policy, the National Biosafety Framework (which includes the National Biosafety Policy and the National Biosafety Act), and the Control of Import, Labeling and Sale of Genetically Modified Foods regulation of 2006. Sri Lanka's National Biosafety Framework has been developed in conformity with the country's commitments to the Cartagena Protocol, which it has signed and ratified. However, these policies are still at various stages of development or implementation.

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CHAPTER 1: PLANT BIOTECHNOLOGY

PART A: PRODUCTION AND TRADE

a) PRODUCT DEVELOPMENT:

Biotechnology applications in Sri Lanka are quite limited. Of those, the most common are Polymerase Chain Reaction (PCR) based detection of pathogens and genes, and Marker Assisted Selection. Molecular biotechnologies, such as recombinant DNA and RNA technologies, are used to a limited extent in Sri Lanka. The country does not have GE plants or crops under development that can be commercialized in the near future; however, some GE crop research is underway at the laboratory level. In addition, tissue culture with biotech applications is common.

Biotechnology is mainly used for DNA finger printing, molecular detection of plant pathogens, molecular characterization, disease diagnosis, marker-aided selection, gene identification, and RNA-level expression studies on select crops such as rice, vegetable crops, root and tuber crops, cucumber, field crops, oil seeds, and fruit crops. Biotech research is mostly conducted for genome studies and to improve yields through development of varieties resistant to biotic and abiotic stress (bacteria, parasites, and pests, or drought, salinity and floods). The applications of genetically engineered plants are limited to laboratory research, although green-house production has been tested in a handful of instances.

Current biotech research on crops includes:

- Diagnosing resistance to: anthracnose in chilies, yellow vein virus in mung beans, and bruchid in cow peas.
- Transgenic development of chilies. Other field crops undergoing biotech research are finger millet, onion, and maize.
- Marker-aided selection on rice, mainly for development of rice varieties with tolerance to salinity, drought, and bacterial leaf blight disease. The new varieties are not yet released.
- Developing varieties resistant to brown plant hopper and stem borer.
- Disease diagnosis of cowpea weevil, and viruses on chilies and tomatoes.

b) COMMERCIAL PRODUCTION:

Genetically engineered crops are not being produced at a commercial level; however, tissue culture production with biotechnological applications is widely used at a commercial level for several crops.

c) EXPORTS:

Sri Lanka neither produces nor exports GE products.

d) IMPORTS:

Genetically engineered products are effectively banned for import into Sri Lanka. The country does not import GE products except for a few pharmaceutical products that contain GE microbes and drugs produced from excretions of GE organisms. Food which contains GE ingredients in amounts less than 0.5 percent can be imported for human consumption, if the presence of such “genetically modified organisms (GMOs)” are considered technically unavoidable and the organisms have been subjected to a scientific risk assessment.

Imports of animal feed are governed by the Animal Feed Act No. 15 of 1986. The Act does not restrict the import of animal feeds containing GE content; however, the Department of Animal Production and Health (DAPH) prevents the imports of GE animal feed by provisions in the existing regulations. If there is a request to import GE animal feed, the DAPH will make the decision in concurrence with the Department of Agriculture and the Ministry of Environment.

e) FOOD AID:

Sri Lanka has been a food aid recipient from the United States and other countries. Nevertheless, regulations prohibit importing of GE food items even as food aid.

f) TRADE BARRIERS:

Sri Lanka has not yet passed any laws to specifically deal with the issue of genetic engineering, except for the Control of Import, Labeling and Sale of Genetically Modified Foods regulation of 2006 under the [Food Act, No. 26 of 1980](#) (GM Food regulation). However, some provisions in the existing laws are used to control, check, and even ban the introduction of certain genetically engineered (GE) products. As a result, the importation or sale of GE products, including ingredients for human consumption, is banned in Sri Lanka. Products intended for human consumption that contain GE ingredients must receive the approval of Sri Lanka’s Chief Food Authority. The general quarantine procedure for import of plant and plant products states that “genetically modified organisms (GMOs)” and “living modified organisms (LMOs)” are not allowed to be imported into Sri Lanka. However, the absence of a functioning approval mechanism in effect leads to bans on the sale of seeds and other agricultural products derived from genetic engineering.

Under the “GM” Food regulation of Sri Lanka, food products for human consumption containing GE ingredients must be labeled, but Sri Lanka has yet to approve any food product containing GE-derived ingredients, creating a trade barrier. Importers lament the burden and complexity of the labeling regulations.

As a result, uninformed consumers are needlessly being made skeptical of GE products, causing no pressure to improve the process.

PART B: POLICY

a) REGULATORY FRAMEWORK:

Except for the Control of Import, Labeling and Sale of Genetically Modified Foods regulation of 2006 under the Food Act, No. 26 of 1980 (“GM” Food regulation), Sri Lanka has not yet passed any laws to specifically deal with genetically engineered products. Sri Lanka is developing multiple policies to regulate and promote biotechnology. These include the National Biotechnology Policy, the National Biosafety Framework of 2005 (which includes the National Biosafety Policy and the National Biosafety Act), and the “GM” Food regulation. The country is expecting the new legal framework to take effect with the enactment of the National Biosafety Act. The Act is under legal review and expected to be in effect soon. Regulations under the Act are being prepared. The Act supposedly will provide guidelines for contained and confined laboratory and field trials.

National Biotechnology Policy:

Sri Lanka’s [National Biotechnology Policy \(NBP\)](#) was promulgated in July 2010, although its enforcement remains inconsistent. The NBP acknowledges the importance of biotechnology in the economic development of Sri Lanka.

Biotechnology is defined by the NBP as “all technologies involving the use of organisms, cells and bio-molecules leading to industrial, agricultural, medical, energy and environmental applications”. The policy has an extensive scope, covering biotechnologies in the following areas.

- (a) All areas of agriculture, livestock, fisheries, forestry, human and animal health, food production, energy and environment;
- (b) All research and development in biotechnology
- (c) All promotional and regulatory activities for product development and commercialization
- (d) All measures to ensure public health and environmental safety with regard to biotechnological application in Sri Lanka

The policy intends to use biotechnology to:

- (a) Support economic development;
- (b) Provide an economic and legal framework to facilitate development of research and commercialization of biotechnology;
- (c) Provide an institutional framework; the policy proposes to establish a National Biotechnology Council to plan, coordinate, monitor and evaluate all activities related to biotechnology, including facilitating and supporting bio-industries while ensuring safe and ethical practices.
- (d) Promote ethical and biosafety considerations of biotechnology; support research and human resources development related to biotechnology;
- (e) Ensure biodiversity innovations have sustainable use of environment and biodiversity;

- (f) Safeguard intellectual property rights and traditional knowledge;
- (g) Promote public-private –partnership in biotechnology.

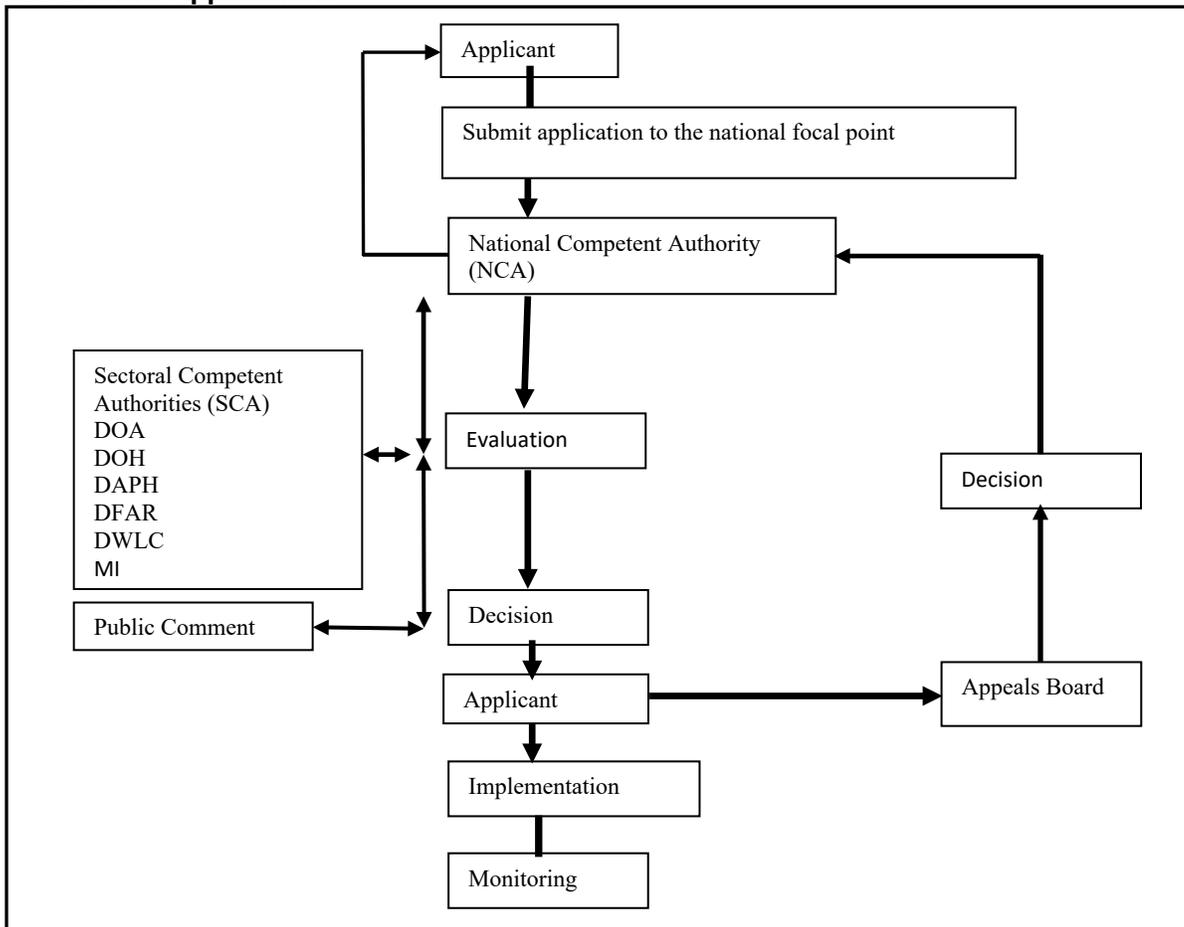
A National Biotechnology Strategy is described in the policy document for development of biotechnology and related industry, while safeguarding human health, environment and biodiversity. The key themes of the strategy are: government commitment for research, development and commercialization of biotechnology; promote public awareness and position biotechnology in society; human resources development to build capacity in biotechnology; sustainable use of biodiversity, enhance opportunities for biotechnology related industries and establishment of centers of excellence and biotechnology parks.

Implementation of the policy is inconsistent; neither the national Biotechnology Council nor the National Biotechnology Strategy suggested under the Act is yet finalized, and the work on Biotechnology Park is progressing at a slow place.

The National Biosafety Framework:

The National Biosafety Framework of Sri Lanka (NBFSL) was developed in 2005, in conformity with the country’s commitments to the Cartagena Protocol (see part B: Policy, paragraph I). The NBFSL was created to ensure an adequate level of protection for the safe transfer, handling, and use of LMOs. Specifically, the NBFSL aims to minimize risks caused by modern biotechnology to biodiversity, human health, and the environment by regulating transboundary movements through use of relevant policies, regulations, technical guidelines and establishment of management bodies and supervisory mechanisms. The NBFSL is a first step towards a more permanent legislative framework for biosafety.

Figure 1: The proposed Administrative System of Circulation and Assessment of the Notifications for the use of GMO Applications



Source: National Biosafety Framework, 2005

The National Biosafety Policy and The National Biosafety Act:

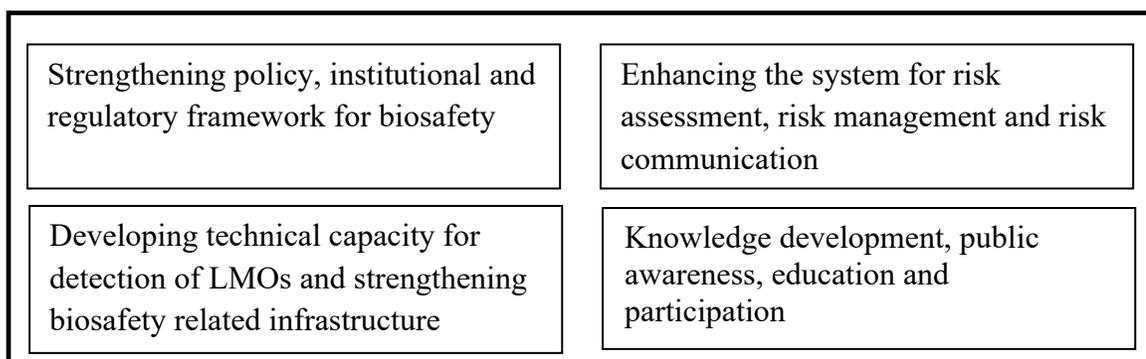
The Government of Sri Lanka has created a National Biosafety Policy as part of the National Biosafety Framework of Sri Lanka (NBFSL). The National Biosafety Policy is based on a precautionary approach, in compliance with the Cartagena Protocol on biosafety. It defines biotechnology in accordance with Cartagena Protocol as “any technological application that uses biological systems, living organisms, or derivatives thereof, to make or modify products or processes for a specific use”. Both the NBFSL and the National Biosafety Policy identifies “modern biotechnology” as application of (a) *in vitro* nucleic acid techniques, including recombinant deoxyribonucleic acid (DNA) and direct injection of nucleic acid into cells or organelles, or (b) fusion of cells beyond the taxonomic family, that overcome natural physiological, reproductive or recombination barriers and that are not techniques used in traditional breeding and selection, which have the potential advances in many fields but with potential adverse effects not yet known. The Biosafety Policy in this view ensures the safe application of modern biotechnology; no adverse effects on conservation, and sustainable use of biological diversity.

The National Biosafety Policy differs from the NBFSL; the NBFSL is the Cartagena Protocol-mandated framework of legal, technical and administrative mechanisms for the regulation of biosafety, whereas the National Biosafety Policy implements specific aspects of the NBFSL. The draft National Biosafety Act also intends to implement the NBFSL, which is expected in effect soon. No current draft of the National Biosafety Act is publicly available.

The new Act will regulate and monitor the applications of modern biotechnologies, including all “GMOs”, “LMOs”, and products that affect food consumption, research, commercial production, and imports and exports. The new Act will detail procedures for approval, monitoring, and enforcement of penalties for violations.

The Global Environmental Facility (GEF) funds a project (National Biosafety Project) in Sri Lanka partnered with the Food and Agriculture Organization (FAO) from 2017 to 2020. The project aims to strengthen institutional, regulatory and technical capacities, for effective implementation of the National Biosafety Framework, in conformity with the Cartagena Protocol on Biosafety. The Ministry of Environment is the government counterpart. The project has identified four components (I) Strengthening policy, institutional and regulatory framework for biosafety; (II) Enhancing the system for risk assessment, risk management and risk communication; (III) Developing technical capacity for detection of “LMOs” and strengthening biosafety related infrastructure; (IV) Knowledge development, public awareness, education and participation.

Figure 2: Components of the National Biosafety Project



Source: Based on the Workshop on Draft Guidelines for Risk Assessment of Living Modified Organisms, FAO, Ministry of Mahaweli Development and Environment, GEF, 2019

Regulations for Import, Labeling, and Sale of Genetically Modified Food:

The Control of Import, Labeling and Sale of Genetically Modified Foods regulation of 2006 comes under the Food Act of Sri Lanka, No. 26, 1980. This is the only regulation that applies to import of products and is binding only for products imported for human consumption. The regulation requires that biotech products for human consumption in Sri Lanka receive rigorous testing and risk assessments.

The regulation prohibits import, storing, transportation, distribution, selling or offering for sale any form of “GMOs” in food for human consumption, without the permission of the Chief Food Authority. This includes any food produced from or containing ingredients produced from genetic engineering.

The regulation requires importers to declare food products with more than 0.5 percent GE content for prior approval by the Ministry of Health. According to the regulations, a risk assessment should be conducted by a technical evaluation committee as defined in the Act, but Sri Lanka lacks capacity to conduct monitoring or risk assessment. Although no GE foods have been approved to date, lack of capacity to monitor could mean Sri Lanka has GE food products in retail stores which are not labeled as such.

Plant Protection Act 1999 No. 35

[The Plant Protection Act No. 35 of 1999](#) replaced the Plant Protection Ordinance. The existing Act does not contain restrictions on the import of GE plants, but based on the powers vested by the Act, the Director General of Agriculture can impose regulations. As a result, the general quarantine procedure for importing plants and plant products states that “GMO” and “LMO” are not allowed to be imported into Sri Lanka. In practice, at the time of applying for import permits, GE-free certification is requested for products derived from crops having GE varieties. The importer is requested to declare if the products have GE components when applying for import permits. Such requests are forwarded for review to the Director General of Agriculture or, in the case of animal feed, to the Department of Animal Production and Health. The regulations under the Act are being considered for revision to regulate import of GE plants and plant products.

Sri Lankan Ministries and Their Policy Roles (Table 1): There is no single regulatory authority to oversee biotechnology products. The NBFSL recommended the formation of a national competent authority on biotechnology, to be known as the National Biosafety Council. The recommended council, which comprises representatives of various concerned Ministries and civil society, will be required to:

- a. screen applications and forward them to the relevant Sectoral Competent Authorities (SCA) and;
- b. make the applications available for public comment. These authorities are required to carry out risk assessments and report back to the council. SCAs may involve the following agencies:

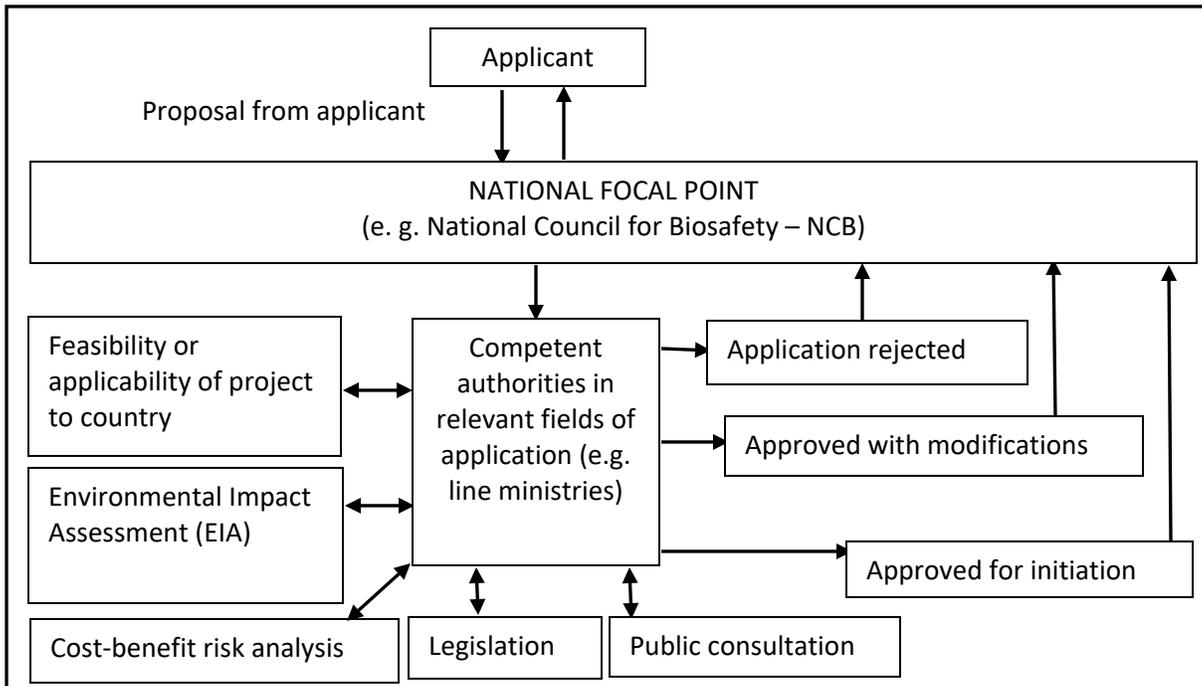
Table 1. Sri Lanka: National Council for Biosafety Sectoral Competent Authorities as identified in National Biosafety Framework

TITLE AGENCY	AREA OF OVERSIGHT
Department of Agriculture	Agricultural and non-agricultural (e.g. forest species, ornamentals) plants and planting materials, microorganisms and animals.
Department of Health	“GM” food and pharmaceuticals.
Veterinary Drug Control Authority (Department of Animal Production and Health)	Domestic animals, including fish, birds, bees, and any other domesticated or wild animals kept in captivity. “GM” fish and/or veterinary pharmaceuticals. Animal feed including “GM” feed ingredients.
Department of Wildlife Conservation	All animals except listed tropical aquarium fish and domestic animals (“GM” fish not in the excluded list).
Department of Fisheries & Aquatic Resources	All aquatic animals and aquatic plants.

Source: National Biosafety Framework, 2005

The following is the proposed flow chart for the evaluation of biotech applications:

Figure 3. Sri Lanka: Proposed Regulatory Mechanism – Application and Review Procedures for Biotech Application



Source: National Policy on Biosafety

Local funding agencies for biotechnology:

Only a handful of agencies fund biotechnology research in Sri Lanka. The main institutions are the National Science Foundation (NSF), Sri Lanka Council for Agricultural Research Policy (SLCARP), and National Research Council (NRC). SLCARP has a National Agricultural Research Plan (NARP), which identifies the biotechnology research priorities for Sri Lanka. The research grants are provided based on the identified priority areas for the country.

b) APPROVALS:

No GE crop is approved for cultivation or import in Sri Lanka. There are no regulations that mandate prior approval for GE research. Nonetheless, the National Science and Technology Commission was vested with power by the Science and Technology Development Act, No. 11 of 1994 to review the science and technology activities in the country, carried out by both public and private institutions.

c) STACKED or PYRAMIDED EVENT APPROVALS:

Existing regulations do not address how stacked or pyramided events would be approved.

d) FIELD TESTING:

Existing regulatory framework does not allow field-testing of GE crops in Sri Lanka

e) INNOVATIVE BIOTECHNOLOGIES:

Sri Lanka has not discussed or determined their position on the research, development, application, or regulation of innovative biotechnologies.

f) COEXISTENCE:

As there is no cultivation of GE crops, there is no coexistence guidelines.

g) LABELING AND TRACABILITY:

According to the Control of Import, Labeling and Sale of Genetically Modified Foods regulation of 2006, if the application has been approved and permission is granted in accordance with the regulation, the product is permitted to be placed in the market subject to appropriate labeling. The label of a package of genetically engineered (GE) food, or food ingredients used in the preparation of food, must include the statement 'genetically modified' in conjunction with the name of that food or ingredients, or processing aid irrespective of the size of the label or package. If the product is displayed for retail sale without packaging, the same information must be attached as a label on the food. Food that has GE content of less than 0.5 percent is exempted from these regulations, provided that the presence of such GE content is considered technically unavoidable and the organisms have been subject to a scientific risk assessment.

In Sri Lanka, the acronyms genetically modified (GM), "GMO," and "LMO" are widely used.

h) MONITORING AND TESTING:

Sri Lanka lacks testing facilities at the ports of entry/exit to test for GE. The laboratories equipped for GE content testing at present have limited capacities and are not accredited. Three laboratories are earmarked for upgrading for GE content testing; National Plant Quarantine Service, Industrial Training Institute (ITI) and Biotechnology Centre of the University of Peradeniya. No interceptions have been reported of import consignments containing unapproved GE events. There is no routine monitoring of products in the market place for GE content. Similarly, authorities do not regularly monitor field crops for unapproved GE events, as the regulations prohibit entry of GE seeds or plants.

i) LOW LEVEL PRESENCE POLICY:

Sri Lanka has a Low-Level Presence (LLP) policy only for food products imported for human consumption. Foods that have GE content of less than 0.5 percent are exempted from these regulations, if the presence of such content is considered technically unavoidable and the organisms have been subjected to a scientific risk assessment. Sri Lanka has zero tolerance for unapproved GE events, although the LLP policy or other regulations do not specify a penalty for undeclared imports of GE products.

j) ADDITIONAL REGULATORY REQUIREMENTS:

Nothing to report.

k) INTELLECTUAL PROPERTY RIGHTS (IPR):

The Intellectual Property Act of Sri Lanka makes it possible to patent GE microbes. However, provisions in the Act allow regulators to deny patents upon recommendation of other relevant authorities if they are considered detrimental.

The draft Plant Breeders Rights Act attempts to comply with obligations under the Trade Related-aspects of the Intellectual Property Rights Agreement and international legal agreement between all member nations of the World Trade Organization. While the Act gives Plant Breeders Rights on new plant varieties, if a variety is a GE plant, it first needs approval under the provisions of the new law before it can be given the Plant Breeders Rights.

l) CARTAGENA PROTOCOL RATIFICATION:

Sri Lanka signed the Cartagena Protocol on Biosafety on 24 May 2000, in Nairobi, Kenya, when it was first open for signatories. Sri Lanka ratified the Cartagena Protocol on 28 April 2004, which took effect on 28 July 2004. The Ministry of Environment is identified as the National Focal Point for Cartagena Protocol on Biosafety and has responsibility for developing the National Biosafety Framework.

m) INTERNATIONAL TREATIES and FORUMS:

Sri Lanka is a member of the International Plant Protection Convention. It is also a member country of the Codex Alimentarius since 1972. It has been a WTO member since 1995 and a member of GATT since 1948. Sri Lanka has not stated in international forums its positions -- either positive or negative -- on genetic engineering of plants.

n) RELATED ISSUES:

Nothing to report.

PART C: MARKETING

a) PUBLIC/PRIVATE OPINIONS:

In general, the public and regulators in Sri Lanka have negative perceptions or attitudes toward genetically engineered (GE) products and research. Although the research community certainly recognizes the benefits of GE products, they are constrained both by the lack of commercial marketing opportunities, clarity in regulations for GE research and by the lack of basic research funding.

A survey carried out in Sri Lanka on perceptions on “GM” food and organisms reveals the significant knowledge gap on biotechnology and biosafety that has led to misconceptions. A majority of the participants of the survey (68%) believe, Sri Lanka can benefit from “GMO” if proven safe and, 60% believe “GMO” products are available in the country. Out of the sample, the government officials and research community were more knowledgeable about “GMO”s. Out of the rest of the sample, more than half had a poor understanding of “GMO”s. Growers, importers, biotech-related organizations, media, and some fractions of the general public were uncertain about the difference between genetic engineering and conventional breeding techniques.¹

b) MARKET ACCEPTANCE/STUDIES:

Nothing to report.

CHAPTER 2: ANIMAL BIOTECHNOLOGY

PART D: PRODUCTION AND TRADE

a) PRODUCT DEVELOPMENT:

Genetic engineering research for animals is not taking place in Sri Lanka. Some field trial research exists on nutritional biotechnology such as rumen bypass feed development, digestibility, and rumen microflora quality improvement. Also, other ongoing research includes optimization for synchronization protocols for reproductive efficiency, disease diagnosis, early pregnancy detection, and vaccine development. Still other research includes that for molecular characterization, especially genetic conservation. There is no research and development happening on animal cloning.

b) COMMERCIAL PRODUCTION:

There is no commercial production of GE animals, insects, birds, or fish in Sri Lanka, nor is there commercial production of cloned animals.

c) EXPORTS:

Sri Lanka does not export any GE animals, animal clones, or products from these animals.

d) IMPORTS:

There is no legal framework governing the controls for importing GE animals or animal products to Sri Lanka. However, some provisions in the existing regulations are used to control, check, and even ban the introduction of certain GE products. The importers are instructed to declare such imports to the Department of Animal Production and Health which will approve or deny such imports.

¹ Kandanaarachchi, M. (2019), “A Preliminary Survey on Sri Lankans’ Knowledge and Understanding of Biosafety and GMOs”, 7th Annual South Asia Biosafety Conference, Dhaka, Bangladesh.

e) TRADE BARRIERS:

The trade barriers applicable to plant products are also applicable for GE animal products.

PART E: POLICY

f) REGULATORY FRAMEWORK:

Imports of animals are governed under the Animal Disease Act No. 59 1992. The Act does not restrict the import of GE animals, however, in practice the Department of Animal Production and Health prevents imports of GE animals based on the provisions in the existing regulations.

g) APPROVALS

No regulations detail requirements on labeling or traceability of GE animals and products, including cloned animals.

h) INNOVATIVE BIOTECHNOLOGIES:

Nothing to report.

i) LABELING AND TRACEABILITY:

No regulations detail requirements on labeling or traceability of GE animals and products, including cloned animals.

j) INTELLECTUAL PROPERTY RIGHTS (IPR):

No specific regulations exist on IPR for animal biotechnology.

k) INTERNATIONAL TREATIES and FORUMS:

Sri Lanka is a member of World Organization for Animal Health (OIE). The Director General of the Department of Animal Production and Health is a permanent delegate of the OIE. Sri Lanka is also a member of the Codex Alimentarius since 1972. Sri Lanka is not known to have stated positions on GE animals or cloning in international forums.

l) RELATED ISSUES:

Nothing significant to report.

PART F: MARKETING

m) PUBLIC/PRIVATE OPINIONS:

Similar to plant biotechnology.

n) MARKET ACCEPTANCE/STUDIES:

Nothing to report.

Attachments:

No Attachments