

ANNEX E

Republic of the Philippines
Department of Agriculture
BUREAU OF PLANT INDUSTRY

PLANT QUARANTINE SERVICE
Manila

**Application for Biosafety
Permit for Direct Use as
Food and Feed, or for
Processing**

(Date)

The Director
Bureau of Plant Industry

Sir:

We -

Information	Applicant	Responsible Officer (RO)	Representative of RO (if applicable)
Name	MONSANTO PHILIPPINES, INC.	MR. IINAS IVAN LAO Country Commercial Lead	MS. MARIA LUISA C. PAHUYO Regulatory Science Lead
Address	2/F Bayer House, Canlubang Industrial Estate, Calamba, Laguna 4028	2/F Bayer House, Canlubang Industrial Estate, Calamba, Laguna 4028	2/F Bayer House, Canlubang Industrial Estate, Calamba, Laguna 4028
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hereby request for issuance of Biosafety Permit for Direct Use for the genetically modified (GM) plant/plant product described below.

Designation	Regulated Article	Donor Organism	Host Organism	Vector or Vector Agent	Constituents of the Regulated Article
Common Name	MON 95379 and all progenies derived from crosses of the product with any conventionally-bred maize, and/or maize	Common soil bacterium	Maize	MON 95379 was developed through <i>Agrobacterium</i> -mediated transformation utilizing plasmid vector PV-ZMIR52222 3.	Molecular characterization demonstrated that MON 95379 contains a single copy of the intended T-DNA at a single locus of the

	containing registered biotech events.				maize genome.
Scientific Name	<i>Zea mays</i> L.	<i>Bacillus thuringiensis</i>	<i>Zea mays</i> L.	-	-
Trade Name	-	-	-	-	-
Other Designations	-	-	-	-	-

Country of Origin	Importers of commodities/grains will have to provide their own sources.
Quantity	Importers of commodities/grains will have to provide the quantities they need to import.
Flight/Voyage No	Importers of commodities/grains will have to provide the flight/voyage no.
Port of Entry	Importers of commodities/grains will have to provide the port of entry.
Proposed Schedule	Importers of commodities/grains will have to provide the schedule of importation.

***The following supporting documents are attached**

1. In the case of imported regulated article, notification from the exporter or country of origin in accordance with existing international agreements on the transboundary movement of genetically modified organisms;
2. Information on socio-economic, cultural and ethical considerations;
3. Copy of the proposed PIS for Direct Use, (Annex "F") including information required for evaluation of environmental and health impacts;
4. In cases of renewal, certified true copy of expired Biosafety Permit for Direct Use issued by BPI; and
5. Proof of payment of fees
6. To facilitate review of the application, the applicant may submit documents to show that the regulated article is allowed for commercial distribution as food and feed by the regulatory authorities in the country of origin; and poses no greater risks to biodiversity, and human and animal health than its conventional counterpart. If the regulated article is intended for use as feed or for processing into feeds, the applicant may submit documents to show that the regulatory authorities in the country of origin have likewise determined that the regulated article poses no greater risks to biodiversity, and human and animal health than its conventional counterpart.

