Where GMOs Stand Today

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What I want to communicate

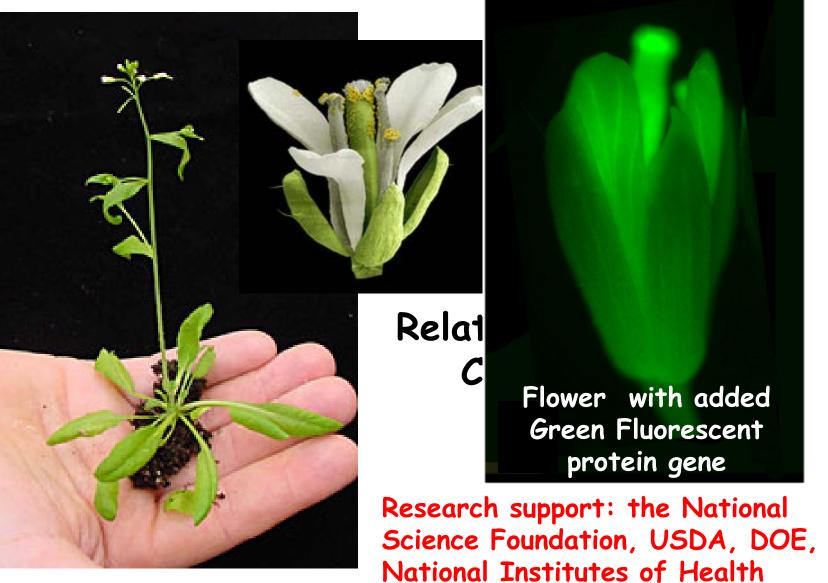
- What is my bias?
- How do we put genes into plants?
- What are the major "transgenic crops" today?
- What about fruit crops?

Will new technologies "replace" GMO plants? What do we need to do next?

What is my bias?

- Use <u>knowledge-based</u> processes to understand potential risks and rewards of new (and old) technologies
- Protect the right for farmers to farm in <u>different sustainable</u> ways (and make a living), and for consumers to choose foods of their preference
- Concern: Over-heated rhetoric is obscuring the risks and rewards of GMOs.
 - GMOs will double yields and solve all agricultural problems!!!
 - GMOs will kill you, or at least make you sick, and besides...it's MONSANTO (buy organic) !!!

I use GMO "technology" to put genes into plants for basic research and discovery.



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coli,

Major Scientific Societies Worldwide Agree that Genetic Engineering Technology is Safe

NETIC LITERACY PRO.

WHERE SCIENCE TRUMPS IDEOLOGY www.geneticliteracyproject.org



THE AMERICAN MEDICAL ASSOCIATION

THE AMERICAN ASSOCIATION FOR THE ADVANCEMENT OF SCIENCE

THE NATIONAL ACADEMY OF SCIENCES

FOOD STANDARDS AUSTRALIA NEW ZEALAND

THE FRENCH ACADEMY OF SCIENCE

THE ROYAL SOCIETY OF MEDICINE

THE EUROPEAN COMMISSION

THE UNION OF GERMAN ACADEMICS OF SCIENCES AND HUMANITIES

SEVEN OF THE WORLD'S ACADEMIES OF SCIENCES

WORLD HEALTH ORGANIZATION

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Putting a GENE of choice into plants using Agrobacterium was first achieved in 1983

NATURE VOL. 303 19 MAY 1983

NATURE VOL. 303 19 MAY 1983

ARTICLES

Expression of chimaeric genes transferred into plant cells using a Ti-plasmid-derived vector

Luis Herrera-Estrella*, Ann Depicker*, Marc Van Montagu* & Jeff Schell**

* Laboratorium voor Genetica, Rijksuniversiteit Gent, B-9000 Gent, Belgium † Max-Planck-Institut f
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üchtungsforschung, D-5000 K
öln 30, FRG

Foreign genes introduced into plant cells with Ti-plasmid vectors are not expressed. We have constructed an expression vector derived from the promoter sequence of nopaline synthase, and have inserted the coding sequences of the octopine synthase gene and a chloramphenicol acetyltransferase gene into this vector. These chimaeric genes are functionally expressed in plant cells after their transfer via a Ti-plasmid of Agrobacterium tumefaciens.

CROWN gall formation on dicotyledonous plants by Agrobacterium tumefaciens is the result of the transfer and covalent integration of a segment (called T-region) of the Ti-plasmid into the chromosomal DNA of plant cells (for reviews see refs 1-4). Insertion of foreign DNA sequences within the T-region of Ti-plasmids leads to their co-transfer and integration into the plant genome⁵ To date inserts of up to 50 kilobases (kb) from pTiT37 (refs 15, 16). Although both genes are encoded by plasmids of bacterial origin, they share more characteristics with eukaryotic genes than with prokaryotic genes. Both octopine and nopaline synthase genes, designated ocs and nos respectively, have a sequence similar to the so-called 'TATA' or 'Goldberg-Hogness' box¹⁷ in the 5' region upstream of the start of transcription and a sequence 'AATAA' similar to the

209

Proc. Natl. Acad. Sci. USA Vol. 80, pp. 4803-4807, August 1983 Genetics Proc. Natl. Acad. Sci. USA Vol. 80, pp. 4803–4807, August 1983 Genetics

Expression of bacterial genes in plant cells

(plant protoplasts/transformation/foreign DNA/antibiotic resistance/selectable markers)

ROBERT T. FRALEY, STEPHEN G. ROGERS, ROBERT B. HORSCH, PATRICIA R. SANDERS, JEFFERY S. FLICK, STEVEN P. ADAMS, MICHAEL L. BITTNER, LESLIE A. BRAND, CYNTHIA L. FINK, JOYCE S. FRY, GERALD R. GALLUPPI, SARAH B. GOLDBERG, NANCY L. HOFFMANN, AND SHERRY C. WOO

Monsanto Company, 800 North Lindbergh Boulevard, St. Louis, Missouri 63167

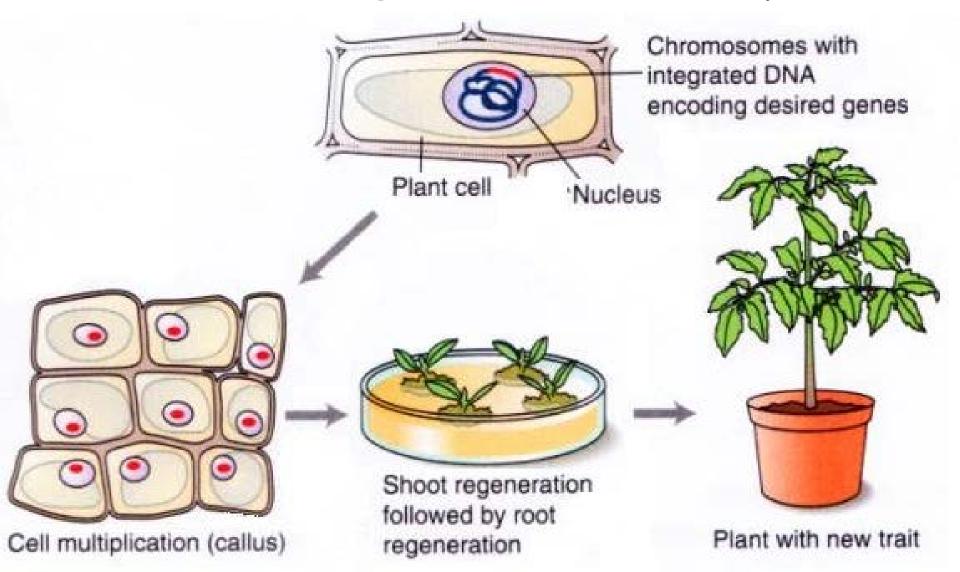
Communicated by Howard A. Schneiderman, April 25, 1983

Monsanto Company, 800 North Lindbergh Boulevard, St. Louis, Missouri 63167

Using Agrobacterium to make "Transgenic" Plants

We insert DNA with The bacterium puts it Our Favorite Gene into the plant for us! into the bacterium Ti plasmid Agrobacterium Plant Cell

We can introduce a gene into a plant cell and the "regenerate" a whole plant



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INTERNATIONAL SERVICE FOR THE ACQUISITION OF AGRI-BIOTECH APPLICATIONS

http://www.isaaa.org/default.asp



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http://cera-gmc.org/

The Center for **Environmental Risk** Assessment (CERA) is dedicated to developing and applying sound science to the environmental risk assessment of agricultural biotechnologies so their contributions to the sustainable production of food, fuel and fiber may be safely realized.

"Biotech" crops currently grown Countries listed in order of number of acres

Country	Note: There is NO "GMO" rice, wheat, peanut
USA*	Maize, soybean, cotton, canola, sugar beet, alfalfa, papaya, squash
Brazil*	Soybean, maize, cotton
Argentina*	Soybean, maize, cotton
India*	Cotton
Canada*	Canola, maize, soybean, sugar beet
China*	Cotton, papaya, poplar, tomato, sweet pepper

Major genes currently in transgenic crops

Herbicide tolerance (HT): Corn, soybean, canola, cotton, sugar beet, alfalfa "Roundup Ready" Gene: EPSP synthase

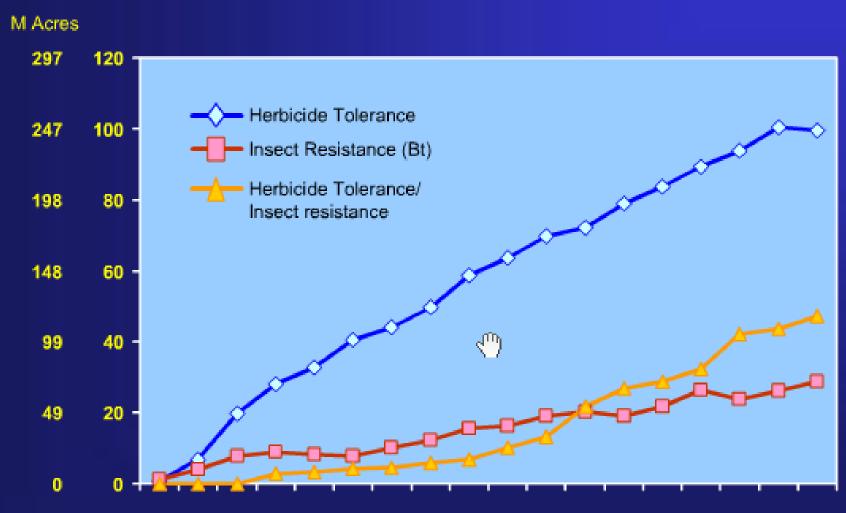
Insect Resistance (Bt): Corn, Cotton

 "<u>Bacillus thuringensis</u> toxin"
 Gene: Bt toxin

 Papaya Ring Spot Virus resistance: Papaya
 Gene: RSV protein

Global Area of Biotech Crops, 1996 to 2013: By Trait (Million Hectares, Million Acres)

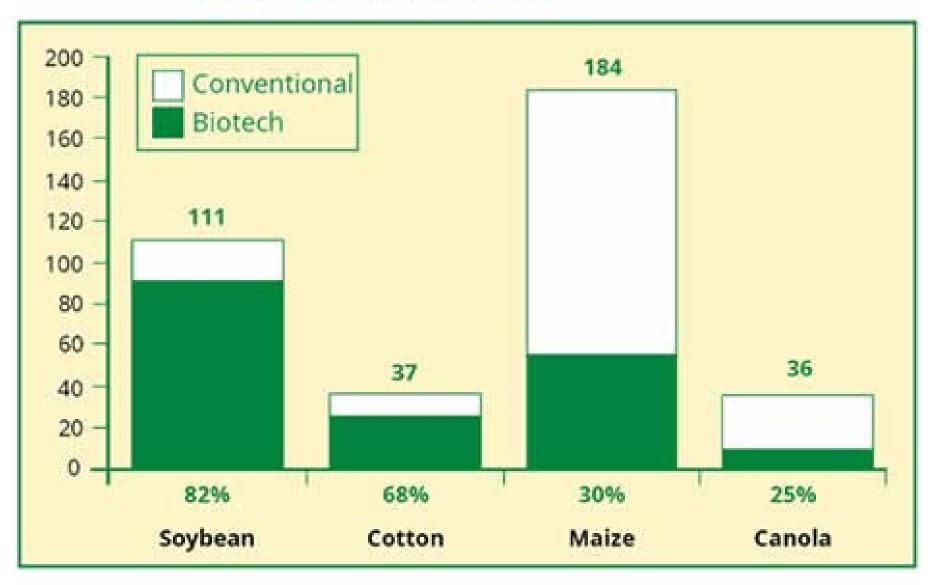




1996 1997 1998 1999 2000 2001 2002 2003 2004 2005 2006 2007 2008 2009 2010 2011 2012 2013

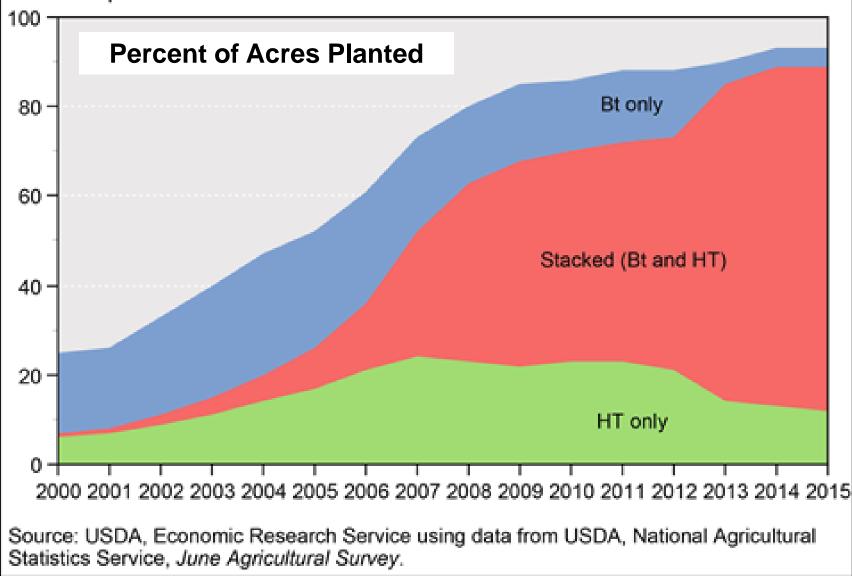
Source: Clive James, 2013

Figure 3. Biotech Crop Area as % of Global Area of Principal Crops, 2014 (Million Hectares)



Source: Clive James, 2014.

Adoption of Genetically Engineered Corn in the US



Why I am perfectly happy eating <u>herbicide tolerant</u> plants!

The gene used specifies a protein found in all plants. We eat it all the time!

The change that makes the protein "herbicide tolerant" involves a few amino acids – less difference than there is between two different plants.

Why I am perfectly happy eating insect resistant plants!

Bt toxin is very INSECT SPECIFIC, specific types affect only certain insects

Crystals and spores are ingested by insect larvae. Toxins are activated to active form by gut enzymes.

Midgut membrane damage leads to starvation or septicemia.

> Activated toxin binds to the receptor, subsequently inserts into the membrane and causes leakage of ions and small molecules.

Bt toxin is very INSECT SPECIFIC, specific types affect only certain insects

In order to kill the insect, the Bt protein must be attached to a specific receptor found only in insects not in humans

Midgut membrane damage leads to starvation or septicemia.

> Activated toxin binds to the receptor, subsequently inserts into the membrane and causes leakage of ions and small molecules.

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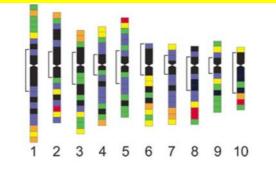
Will new technologies "replace" GMO plants?

What do we need to do next?

Why are GM methods used sometimes and molecular breeding others? Molecular breeding



1. Desired trait must be present in population



2. Genetic resources must be available



3. Plant should be propagated sexually



1. Gene can come from any source







3. Plant can be propagated vegetatively

GM Disease Resistant Papaya has replaced 80% of the Hawaiian Papaya crop

Show abstract

GM Crop Database

Database Product Description

UFL-X17CP-6 (X17-2)

Host Organism Trait

Trait Introduction

Company Information

Proposed Use

Carica papaya L. (Papaya) Resistance to viral infection, papaya ringspot virus (PRSV). Agrobacterium tumefaciens-mediated plant transformation. Production of papaya for human consumption, either fresh or processed. University of Florida

<u>http://cera-gmc.org/index.php</u>? action=gm_crop_database



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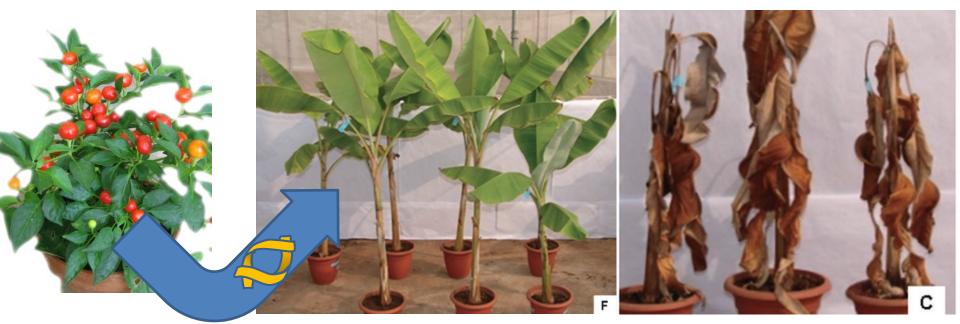
Transgenics have been successful with other diseases of fruit crops



Bananas Bacterial wilt disease

Apples Fire blight **Bacterial disease** Viral disease

GM Example: Disease resistant banana by introduction of a gene from pepper



Resistant Susceptible Banana bacterial wilt is destroying plants in eastern Africa. Transgenic plants carrying a resistance gene from pepper are resistant to the disease

Tripathi, L., Mwaka, H., Tripathi, J.N., and Tushemereirwe, W.K. (2010). Expression of sweet pepper Hrap gene in banana enhances resistance to Xanthomonas campestris pv. musacearum. Molecular Plant Pathology 11: 721-731.

Erwinia amylovora (Apple Fireblight) Resistance from "Crab apple" Engineered into Commercial Cultivars

Engineering fire blight resistance into the apple cultivar 'Gala' using the *FB_MR5* CC-NBS-LRR resistance gene of *Malus* × *robusta* 5. Giovanni A. L. Broggini et al.

Plant Biotechnology Journal Volume 12, Issue 6, pages 728–733, August 2014

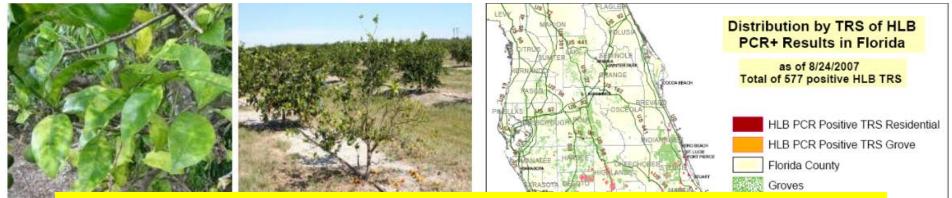
Plum Pox (viral disease) Resistance Engineered with Gene Silencing of Viral Gene

> Current treatment - Destruction of infected trees.

- Field tests of transgenics were conducted in Poland from 1996 to 2006, Spain from 1996 to 2012, Romania 1996-2006 and Czech Republic 2002-2013. Clearly demonstrated resistance to PPV infection through aphid vectors and by graft inoculation.
- > Not yet in commercial production.

http://www.ars.usda.gov/is/br/plumpox/

The spread of Citrus Greening disease



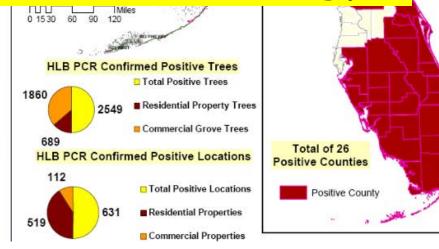
We have no way to breed citrus for resistance to this disease, but it could be done with GMO technology



/24/ 2007 rican 1983 Harn







"Artic" Apple - Non-browning

Okanagan Specialty Fruits Inc. PO Box 1533 Summerland, BC VOH 1ZO Canada

Engineered to inhibit expression of the apple polyphenol oxidase gene.

Inhibits browning but not softening.

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<u>New Technologies for</u> <u>Crop Improvement</u>

> Marker Assisted Selection/Breeding

> CRISPR/Cas gene editing

Marker Assisted Selection

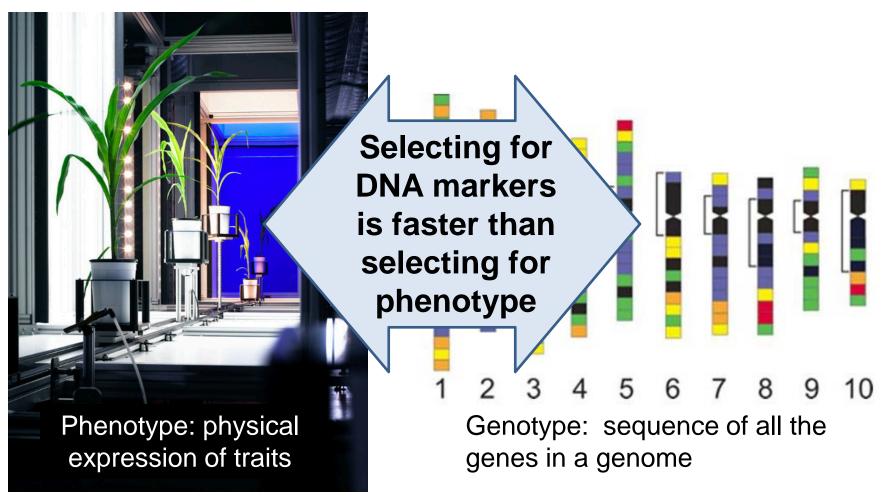
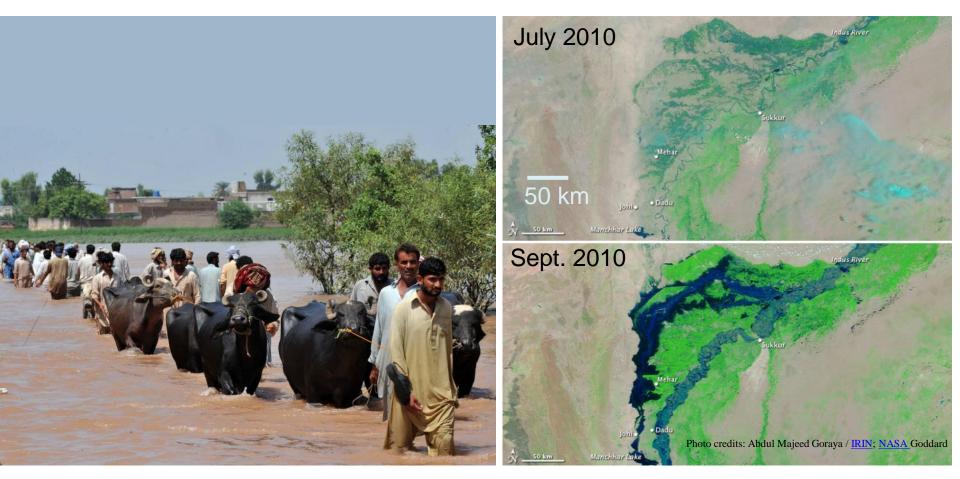


Photo credit LemnaTec; Anderson, L.K., Lai, A., Stack, S.M., Rizzon, C. and Gaut, B.S. (2006). Uneven distribution of expressed sequence tag loci on maize pachytene chromosomes. Genome Research. 16: <u>115-122</u>.

MAS was used to produce of submergence tolerant rice (*Sub1*)



Many rice-growing regions are prone to flooding. In Pakistan a 2010 a huge flood submerged 17 million acres (69,000 km²) and destroyed much of the harvest

<u>CRISP/Cas</u>

A technique for "Editing" a genome Remove genes, change existing genes

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Worldwide, <u>preharvest</u> crop loss estimates:

We need to use every tool at our
 disposal, including GMO technology, to
 solve agricultural and environmental
 problems to create food security and
 sustainability

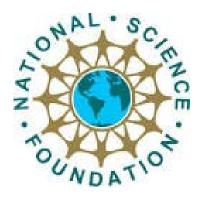
Other losses due to stress: drought, cold heat, salinization

Where to go from here?

- Each crop, each "trait" (modification) needs to be evaluated separately.
- > We need more agricultural research
 - To understand how to combat pests and disease
 - To move away from monoculture towards sustainable practices

> Let's be passionate about the science!

Thanks to many members of my research lab over the last 30 years



Thank you for your attention!