

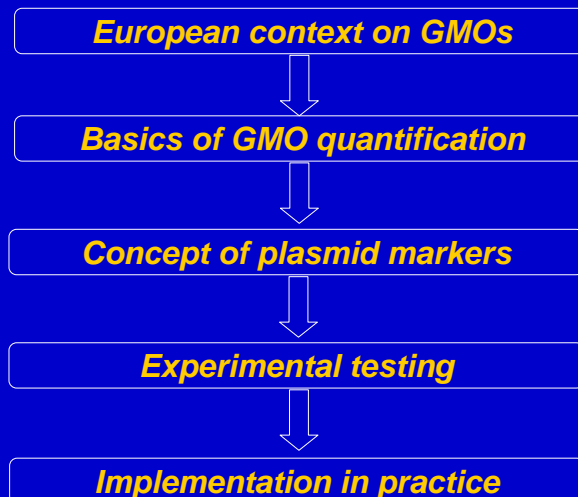


Cloned plasmid DNA molecules as a tool for GMO analysis

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Overview



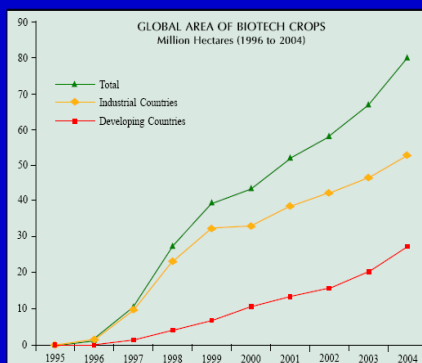
Overview

European context on GMOs



European context on GMOs

Worldwide
increasing area of
transgenic crops



James, 2004



Europe

opposition from consumer groups
and environmentalists



1998-2004
moratorium for cultivation
and marketing approvals

> April 2004
new, strict and comprehensive
regulatory framework



European context on GMOs

Contained use Deliberate release into the environment Labelling and traceability

Field trials Marketing

← Pre-market Post-market →



Introduction

Taverniers-qPCR-2005



European context on GMOs

Contained use Deliberate release into the environment Labelling and traceability

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Introduction

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European context on GMOs

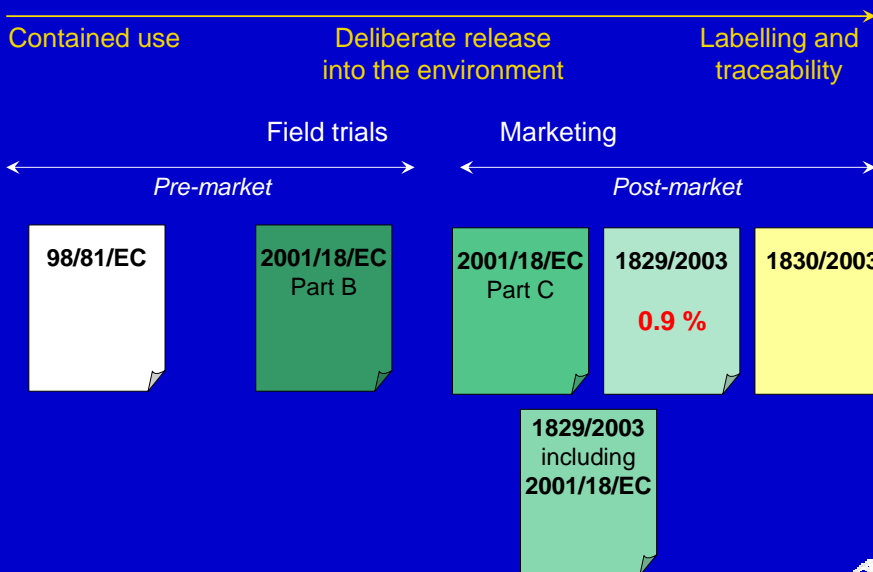


Introduction

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European context on GMOs

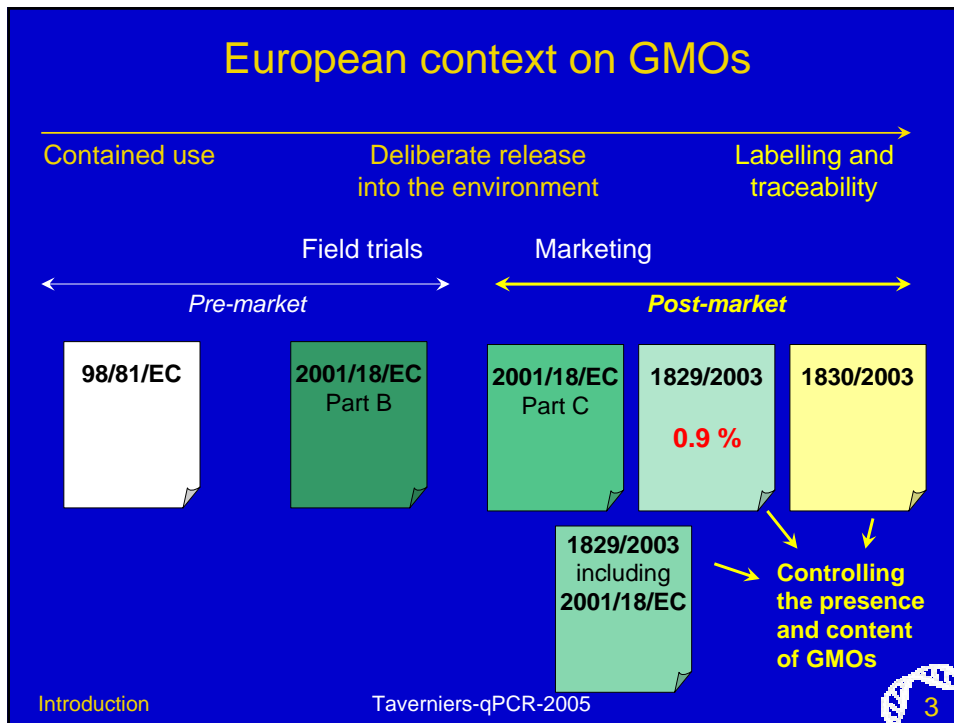


Introduction

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European context on GMOs



European context on GMOs

- ❑ **Reg. (EC) 1829/2003** on food and feed: Labelling is mandatory above a certain threshold value
- ❑ Quantification on ingredient level (single species)
- ❑ **Rec. (EC) 2004/787:** Haploid genome basis for measuring and expressing GMO contents
 $100\% \times (\# \text{ copies } \text{GMO target} / \# \text{ copies } \text{plant target})$
- ❑ Authorized GM food & feed products: **0.9 %**
- ❑ Non-authorized GMOs:
 - with positive risk evaluation: **0.5 %**
 - no evaluation: **0.1 %**
- ❑ GM seeds:
 - cross-pollinators (*rapeseed*): **0.3 %**
 - self-pollinators + exception cross-pollinator (*maize*): **0.5 %**
 - exception self-pollinator (*soybean*): **0.7 %**

Introduction

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Overview

European context on GMOs



Basics of GMO quantification



Overview

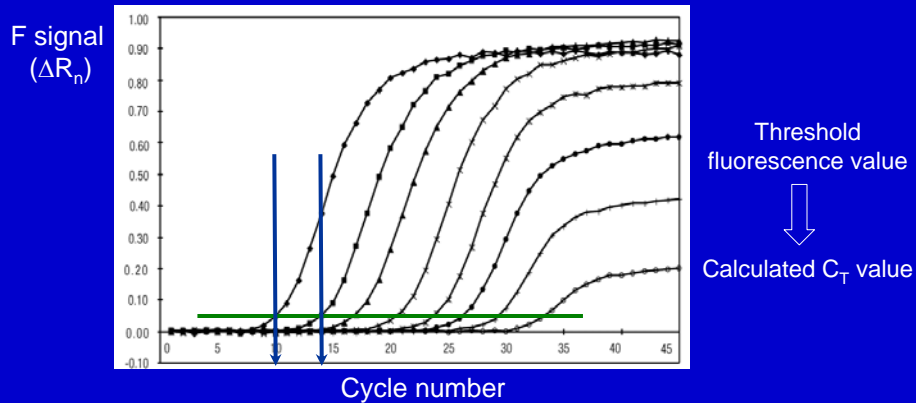
Basics of GMO quantification

- ❑ Real-time PCR technique
- ❑ Units of measurement and expression
- ❑ Calibrators for GMO quantification



DNA quantification with real-time PCR

- Exponential amplification of a DNA target molecule gives a **measured fluorescent signal**



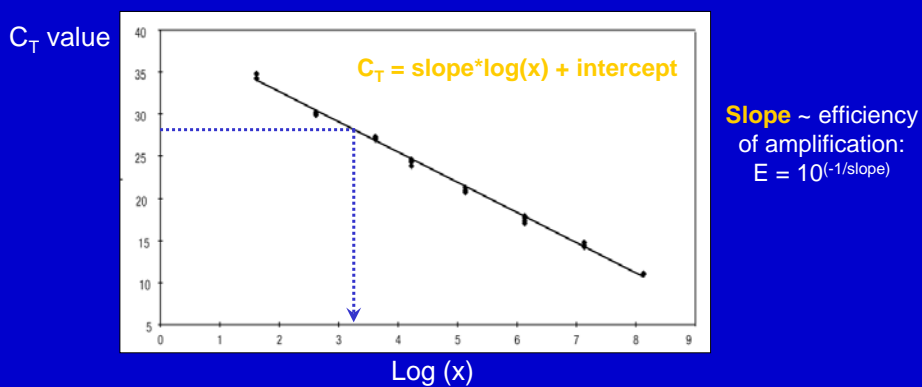
Basics of GMO quantification

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DNA quantification with real-time PCR

- Relationship between **calculated C_T value** and initial target DNA concentration is basis for quantification



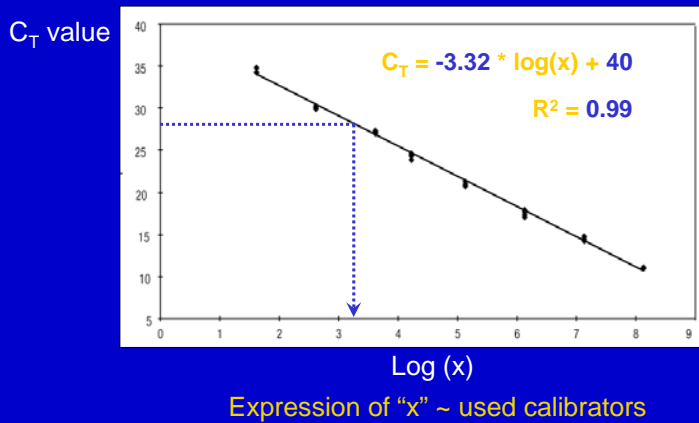
Basics of GMO quantification

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DNA quantification with real-time PCR

- Relationship between calculated C_T value and initial target DNA concentration is basis for quantification



Slope of -3.32 ~
 Maximum efficiency
 of amplification:
 $E = 2$

Units for measurement and expression

0.9 % of what ?



% GM maize

$$100 \times \frac{\# \text{ GM seeds}}{\# \text{ total seeds}}$$

% GM maize

$$100 \times \frac{\text{GM maize mass}}{\text{total maize mass}}$$

% GM maize

$$100 \times \frac{\# \text{ GMO-specific DNA copies}}{\# \text{ plant-specific DNA copies}}$$

Calibrators for GMO quantification



matrix RM

mass %



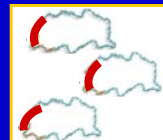
pure DNA RM

DNA copy n° %

genomic DNA



plasmid DNA



Calibrators for GMO quantification

Matrix RMs:

CRMs (EC-IRMM, Fluka); 0-5 % GMO w/w

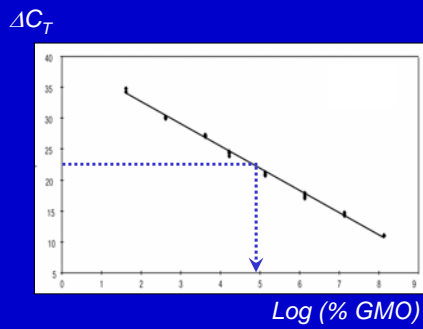
Pure DNA RMs:

- genomic DNA (EC-IRMM, Fluka); 0-5 % GMO w/w
- plasmid DNA fragments
 - single target plasmids - STP (*Taverniers et al., 2001; 2004; others; BNL & ENGL plasmid database*)
 - duplo target plasmids – DTP or 'tandem-marker plasmids' (*Mattarucchi et al., 2005; others; BNL & ENGL database*)
 - multiple target plasmids - MTP (*Kuribara et al., 2002; others; Nippon Gene Japan, Diagenode Belgium*)
- PCR amplicon fragments
 - single PCR amplicons (*Holck et al., 2002; others*)
 - hybrid amplicons (*Pardigol et al., 2003; others*)

Approaches for GMO quantification

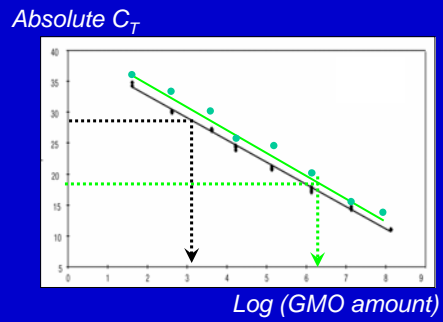
- Two means of deriving a relative % result

Relative quantification
Single calibration curve



Calibrators in %

Absolute quantification
Double calibration curves



Calibrators in amounts

Basics of GMO quantification

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Overview

European context on GMOs

Basics of GMO quantification

Concept of plasmid markers

Overview

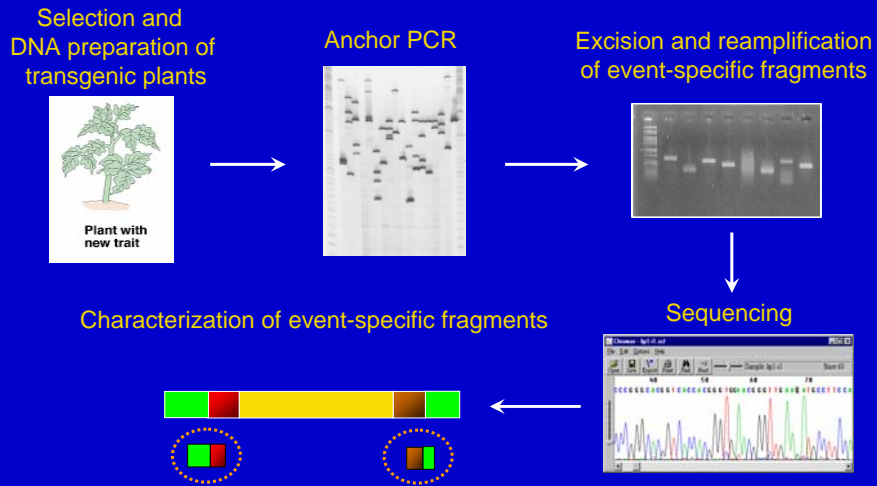
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Plasmid DNA calibrators

1. Generation of the target DNA sequences



Concept of plasmid markers

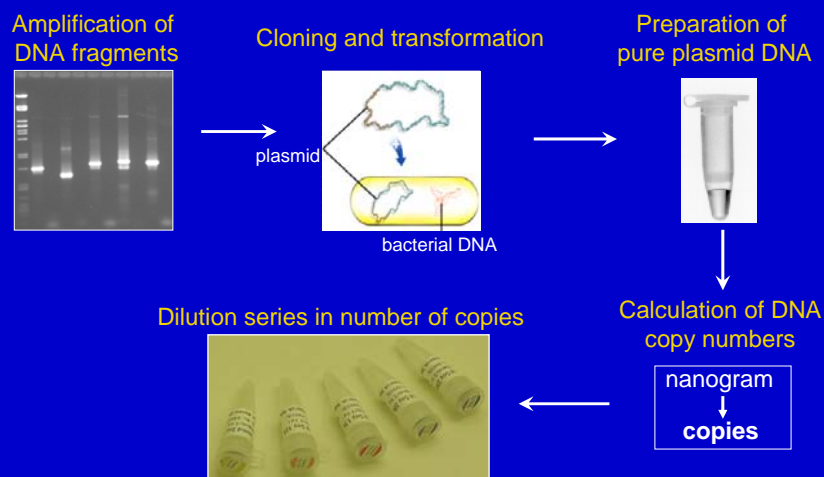
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Plasmid DNA calibrators

2. Cloning of DNA fragments into plasmid vectors



Concept of plasmid markers

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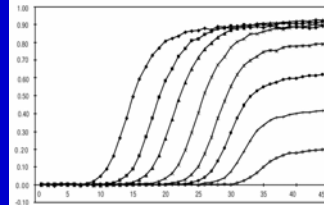
Plasmid DNA calibrators

3. Use of plasmid DNA series in real-time PCR

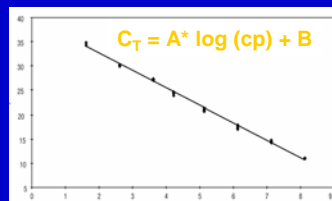
Real-time PCR



Setting up of calibration curves



C_T



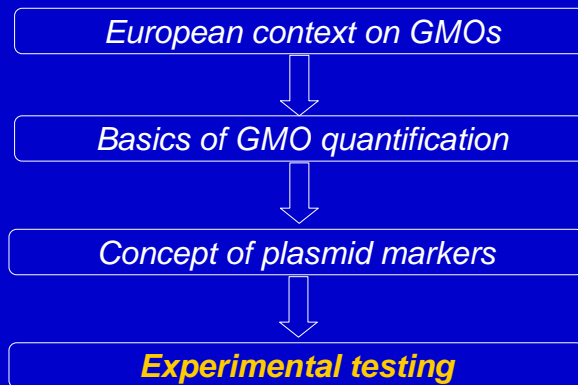
Log (# copies)

Concept of plasmid markers

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Overview



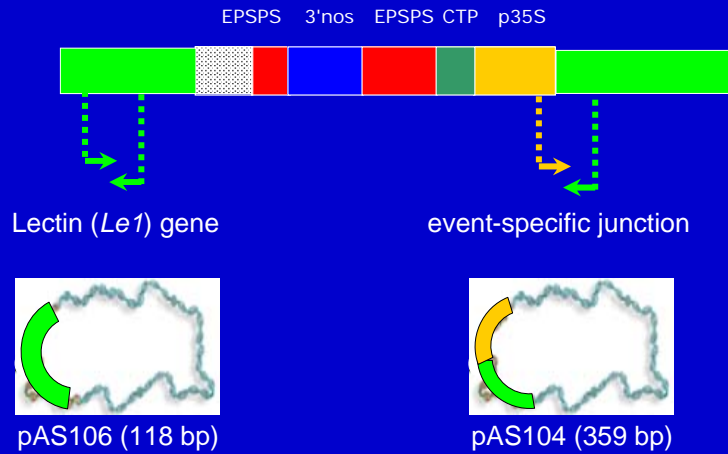
Overview

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Case study: RR soybean quantification

□ Construction of plasmids



Experimental testing

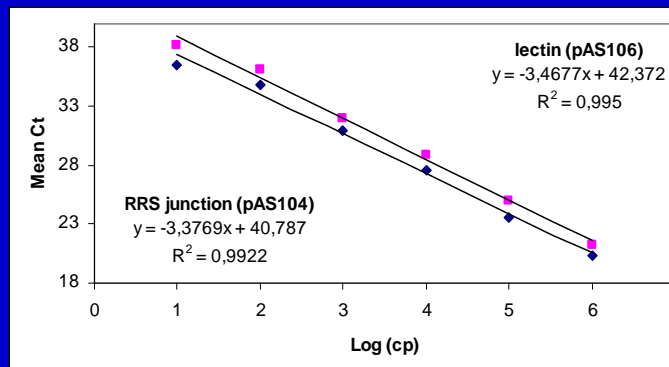
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Case study: RR soybean quantification

□ Absolute simplex PCR quantification

2 STP series (10^6 - 10 cp) - 2 standard curves (C_T)



Experimental testing

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Case study: RR soybean quantification



Sample	Cp junction	Mean cp junction	Cp lectin	Mean cp lectin	Conc (%)	Mean conc (%)	SD	%RSD	%error
NTC		NTC		NTC					
10		10		10					
100		100		100					
1000		1000		1000					
10000		10000		10000					
100000		100000		100000					
1E+06		1000000		1000000					
0	19,76 8,54	14,15	31595 19594	25594,1	0,063 0,044	0,054	0,013	24,07	5,4
0,1	1194,23 1317,9	1256,07	1E+06 1E+06	1185553	0,099 0,11	0,105	0,008	7,62	5
0,5	2163,36 1949,07	2056,22	521604 472012	496808	0,41 0,41	0,41	0	0	18
1	5863,98 7074,68	6469,33	683686 511536	597611	0,86 1,38	1,12	0,37	33,04	12
2	7743,99 11515,2	9629,59	306760 330960	318860	2,52 3,48	3	0,68	22,67	50

Experimental testing

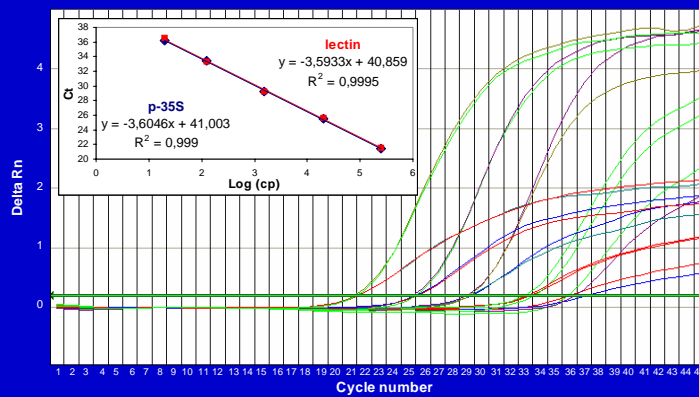
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Case study: RR soybean quantification

- Absolute simplex PCR quantification
- Absolute duplex PCR quantification

1 MTP series (10⁶-10 cp) - 2 standard curves (C_T)



Experimental testing

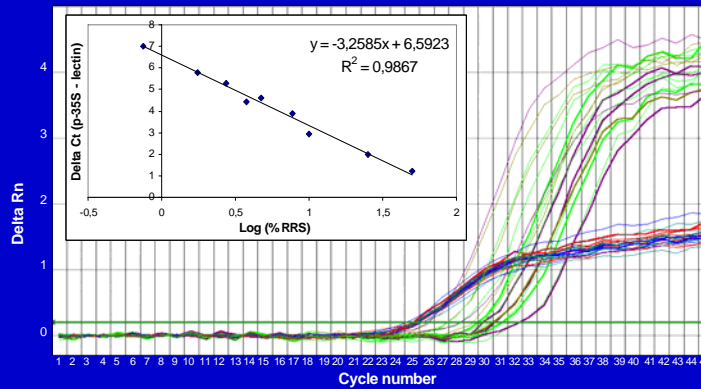
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Case study: RR soybean quantification

- Absolute simplex PCR quantification
- Absolute duplex PCR quantification
- Relative duplex PCR quantification

1 STP series in gDNA (%) - 1 standard curve (ΔC_T)



Experimental testing

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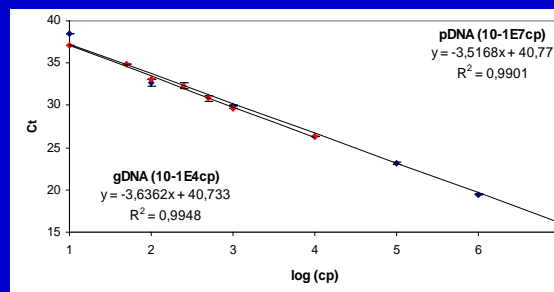
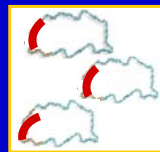


Comparability between gDNA and pDNA

genomic DNA



plasmid DNA



*Bt176 maize,
event-specific
method*

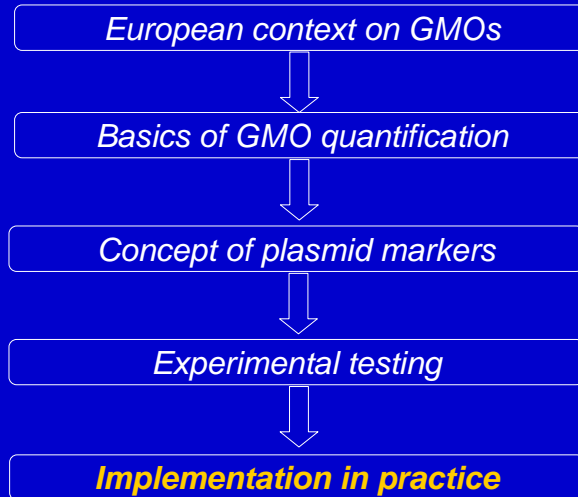
Comparable behaviour in the PCR

Experimental testing

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Overview



Implementation of plasmid system



In-house construction and testing of plasmids

pAS104, pAS106, pCM1, pCM2, ...

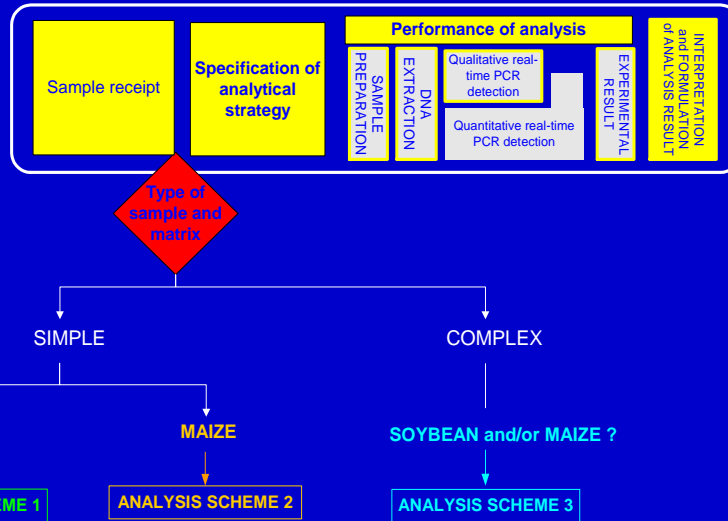


Subcloning to pUC18

pENGL1, pENGL2, pENGL3, ...

Officialization and standardization: ENGL database

Use of plasmids in routine practice

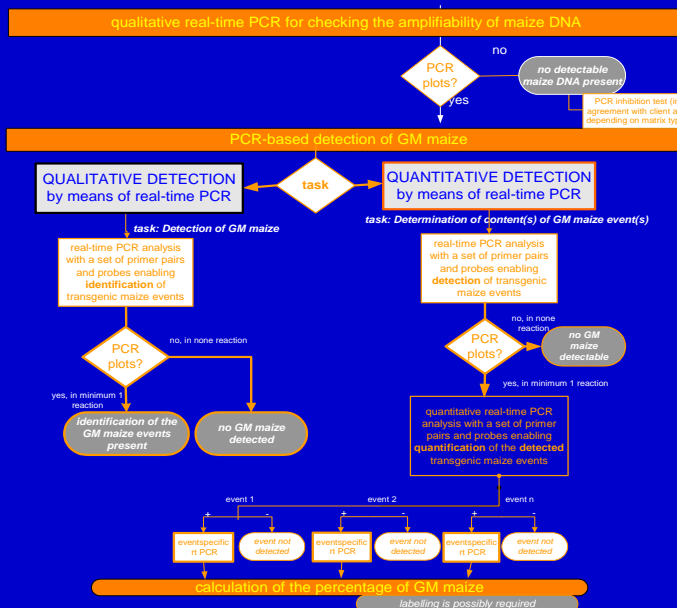


Implementation in practice

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Use of plasmids in routine practice



Implementation in practice

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Implementation of plasmid system

- ❑ Copy number-based DNA calibrators are in compliance with Comm. Rec. **2004/787/EC**, Reg. (EC) **1829/2003** and Reg. (EC) **1830/2003**
- ❑ Plasmid DNA markers are easy in production, storage, distribution and wide in applicability
- ❑ Plasmid DNA and genomic DNA calibrators show perfectly comparable behaviour (commutability)
- ❑ Reduction of analytical errors and improvement of efficiency with:
 - duplo-target plasmids (DTP) or multiple-target plasmids (MTP)
 - duplexing GM-specific and plant-specific target amplification
- ❑ Future challenges:
 - certification of copy number-based reference materials?
 - multiplexing?

Conclusions

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Cloned plasmid DNA molecules as a tool for GMO analysis



Questions ?