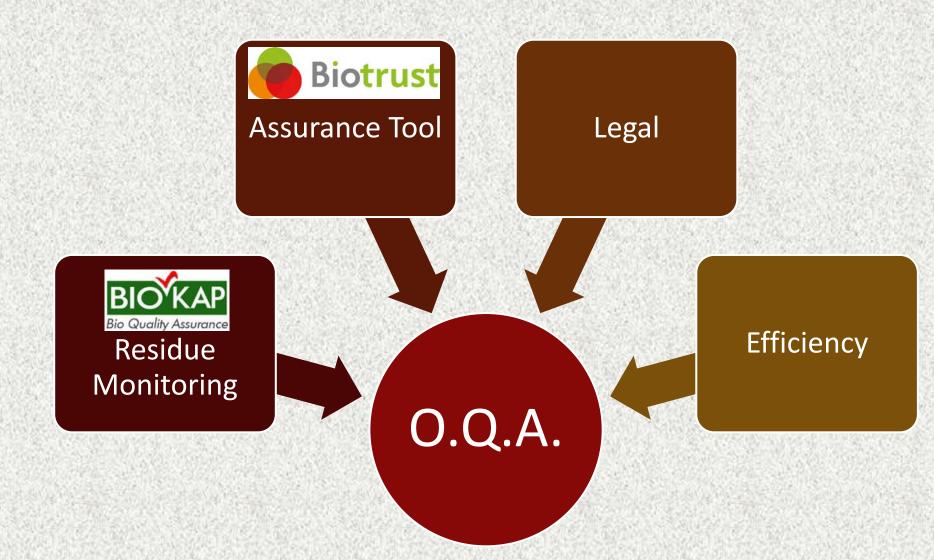


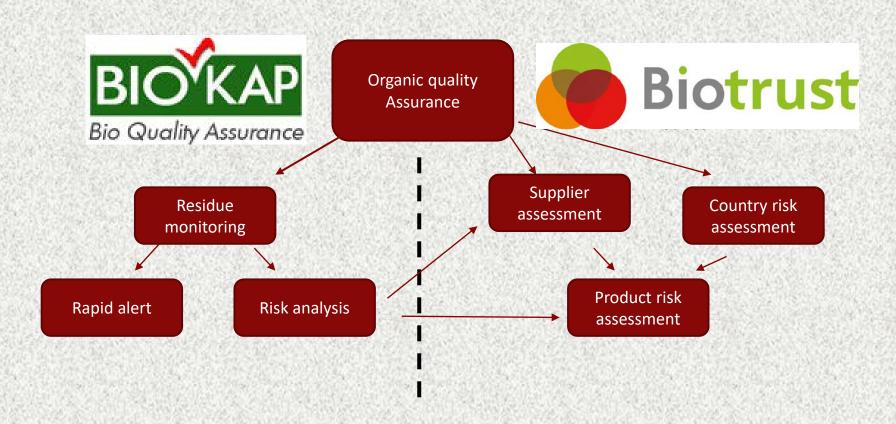
Operators perspective for residue analyses and sampling

Bavo van den Idsert

BioKap and BioTrust



BioKap and BioTrust





Aims and objectives

- Risk based priority setting for verification of raw material risks
- Risk based process control based on open source
- Practical tool that can be used in existing QA systems
- Risks comparable and discussable among active members
- Demonstrable
- Meets the requirements of the organic law
- Cooperation and exchange between companies possible

Approach to Food Fraud Prevention

Food Safety Nanagement System

Food Safety

Food Defence

Food Fraud

HACCP

Hazards

TACCP

Threats

VACCP

Vulnerabilities

Prevention of unintentional / accidental adulteration

- Science based
- · Food borne illness

Prevention of intentional adulteration

Behaviourally or ideologically motivated Prevention of intentional adulteration

Economically motivated

6



PDCA cycle

Commitment and Mandate Communicate and Train Plan **Policy Statement** Communications and Risk Management Plan reporting plan Define & Analyse a Assurance plan Training strategy Problem and Identify the RM Network Standards **Root Cause** Procedures/Guidelines Do Act Common Approach Devise a Solution Standardise Solution Used in ISO Develop Detailed Action Review and Define Plan & Implement It Management Next Issues Systematically System Standards Organise and Allocate Check **Board RM Committee** Measure and review Exec RM Committee Confirm Outcomes Control assurance Manager, RM RM Plan progress **Against Plan** RM Champions Governance reporting Risk, Control, Risk owners Identify Deviations and Benchmarking Assurance providers Performance criteria Issues

BioTrust process: 6 steps

STEP 1: Define Risk Matrix

Likelihood Impact

STEP 2: Define General Product –Process Hazard

f.e. Contamination with conventional



STEP 3: Modify Risk

Country Risk

Relationship risk

STEP 4: Validate the Risk

Low

Medium

High

STEP 5: Reduce the risks

Design an action plan

STEP 6: Implement and evaluate acc. PDCA cycle

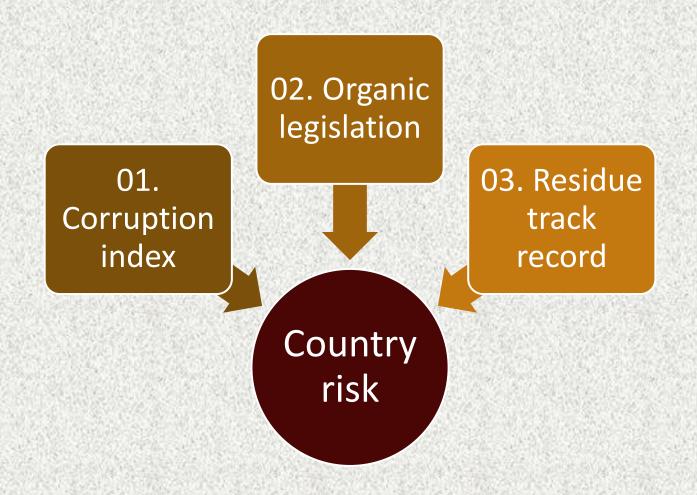
Step 1: define the risk matrix

Description	Extra clarification	Impact				
consequences and violation of the organic principles. Or a	Impact on more products or lots over a longer period, likely to cause media attention, e.g.: high residue levels, large scale mixture with regular, systematically violation of the legislation	3	Low	Medium	High	High
 CPM(#)*CF(**B)**********************************	Impact on one product or lot, likely no media attention, e.g.: low residue levels around limits, GGO below 0,9	2	Low	Low	Medium	High
A non-conformity with no (direct) implication on the product	No impact on the product, e.g.: administration	1	Low	Low	Low	Medium
		Likelihood	1	2	3	4
		Description	Very small	Small	Medium	High
		% of non- conformities	<1	<5	<10	>10

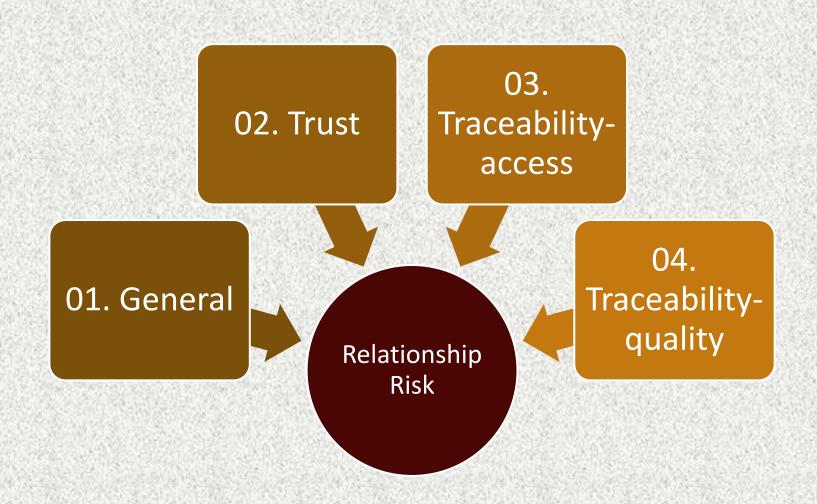
Step 2: define the general hazards

Proces Proces section		Proces section	Hazards			
primairy sector	crop and feedcrop	cultivation	inconversion product sold as organic	3		
primairy sector	crop and feedcrop	cultivation	use or misuse of not permitted basic materials (seeds)	3		
primairy sector	crop and feedcrop	cultivation	use or misuse of not permitted fertilizers	2		
primairy sector	crop and feedcrop	cultivation	incorrect application of crop rotation	2		
primairy sector	crop and feedcrop	cultivation	use of not permitted ground covers	2		
primairy sector	crop and feedcrop	cultivation	use of not permitted pesticides			
primairy sector	crop and feedcrop	cultivation	use of not permitted crop enhancers	2		
primairy sector	crop and feedcrop	cultivation	use of not permitted products for cleaning and sanitazion/decontamination of installations or buildings	2		
primairy sector	crop and feedcrop	cultivation	not permitted crops cultivated on substratum	2		
primairy sector	crop and feedcrop	cultivation	use of sanitazion/decontamination products during substratum cultivation	2		
primairy sector	crop and feedcrop	cultivation	use of not permitted "dekaarde"	2		
primairy sector	crop and feedcrop	cultivation	use of not permitted raw materials in substratum	2		
primairy sector	crop and feedcrop	cultivation	presence of GGO in raw materials used in substratum	2		
primairy sector	crop and feedcrop	cultivation	insufficient seperation between organic and regular plots	3		
primairy sector	crop and feedcrop	cultivation	cultivation of the same crop as organic and regular on the same farm	3		
primairy sector	crop and feedcrop	cultivation	exceding the duration of use regular livestock on a plot	2		
primairy sector	crop and feedcrop	cultivation	exceding the nitrogen/hectare using livestock	2		
primairy sector	crop and feedcrop	administration crop and feed	insufficient or no cultivationplan for past en present year	1		
primairy sector	crop and feedcrop	administration crop and feed	insufficient or no fertilize administration	1		

O What do we know of the country?



O What do we know of our relation?



Final Risk

03.Country 02. Impact Risk 04. 01.Likelihood Relationship Final risk Risk

STEP 5: Final Risk output

Raw material		Risk Analysis								
	Hazard	Impa ct		Likelihood motivation	Country Risk Factor	Relationship Risk factor	(Country + 2*Relations hip)/3	Likelihood with Country and Relationship	Total Raw material Risk	Which verification measurements are necessary?
apricot	contaminati on conventiona	3	2	difficult crop	1,2	1,4	1,3	2,7	MEDIUM	audit supplier
apricot	contaminati on conventiona	3	2	difficult crop	0,5	_1	8,0	1,7	LOW	

Actual practices at trading operators

- O I. Apply risk-based matrix with the following parameters:
 - Relationship & knowledge about the supplier: + 5 years = low risk
 - Risks related to the specific product / ingredient
 - Risk of the country of origin
 - Risk in the chain of transport, storage and transparacy till source
 - History of the supplier
 - Minimal control frequency per supplier / ingredient-product
- II. Follow new developments with substances based on labs and market-info
- III. Aim to reduce the pressure on increasing number of analyses

Something about sample taking

- O In general only systematic approach towards ingredients (and not the end product).
- Experiences that residues can pop-up after processing steps (very disturbing and confusing when the ingredient was free from substances).
- Operators in general still working with concentration factors.
- O In general the way of sampling is very much depending on the type of product.
- In most cases operators choose for representative samples: take different samples from the whole batch and put the together as one representative sample.
- Operators keep a second sample in case a quality issue appears (brought forward by a buyer abroad).
- There are no real sampling requirements from CB's towards operators.
- Challenges ingredients for processed food are much higher than for fresh products