

## ПРЕЗЕНТАЦИЯ

### PRESENTATION

International conference on  
Food Safety Risk Analysis and  
Antimicrobial Resistance

Moscow, Russian  
Federation, 17- 18 12/2019

Международная  
конференция по вопросам  
анализа риска  
безопасности пищевой  
продукции и устойчивости  
к противомикробным  
препаратам

Москва, Российская  
Федерация, 17 -18 12/2019

## Risk Assessment of Novel Technologies and Novel Foods

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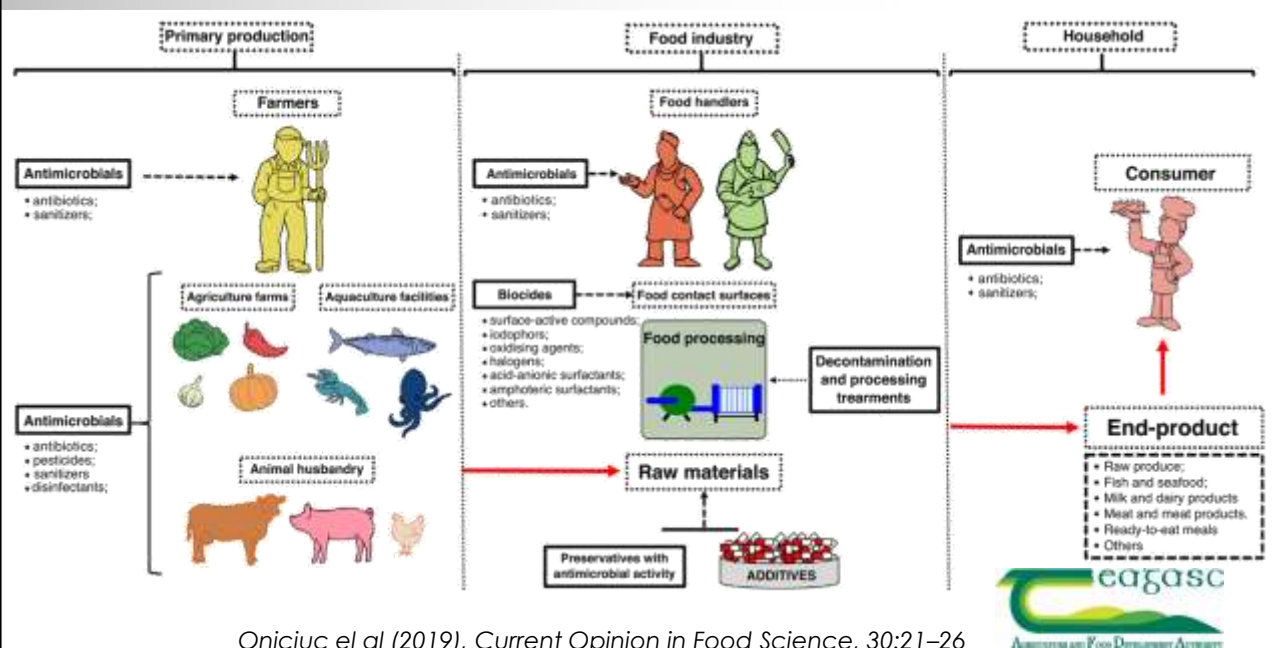


## Teagasc –Agriculture and Food Development Authority

- Animal and Grassland Research and Innovation
- Crops, Environment and Land Use
- **Food**
- Rural Economy and Development



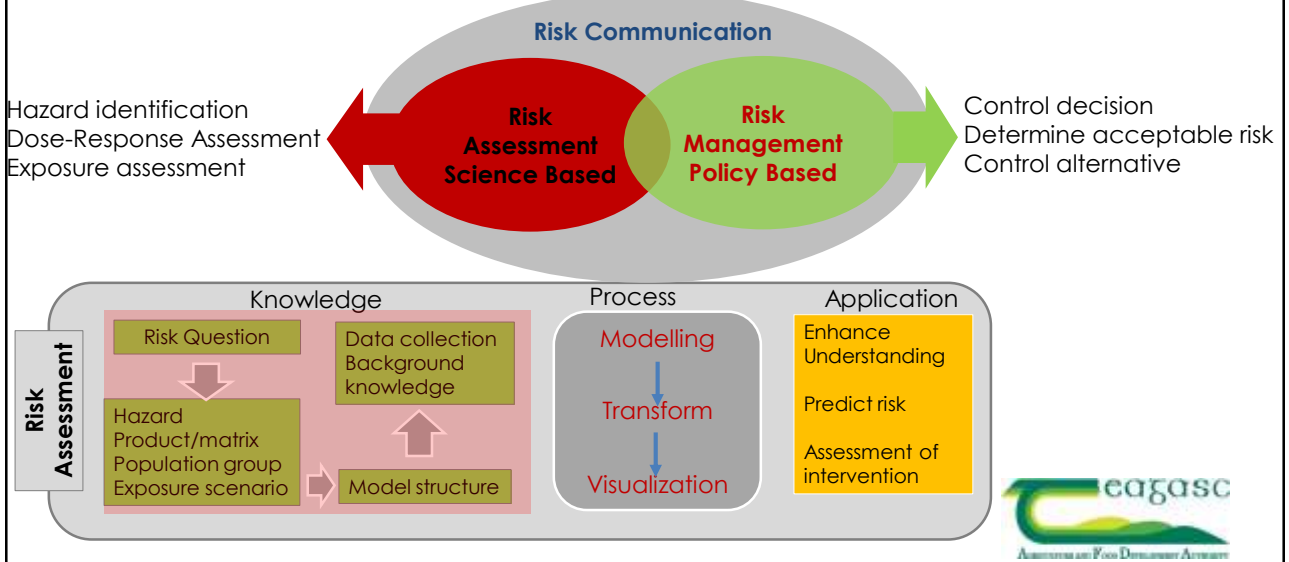
## Causes of AMR in Food Chain



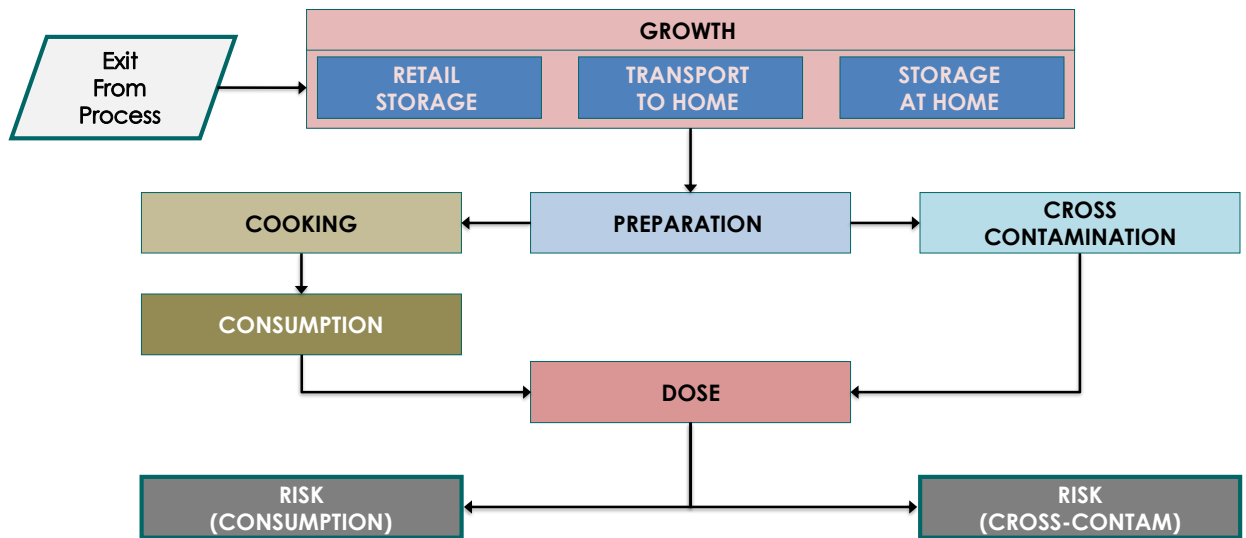
Oniciuc et al (2019), *Current Opinion in Food Science*, 30:21–26

## Risk assessment of novel technologies

RISK ASSESSMENT is a process to scientifically evaluate the probability of occurrence and severity of known or potential adverse health effect resulting from human exposure to foodborne hazards



## Exposure assessment *Salmonella* spp in Chicken



# Risk assessment of novel technologies

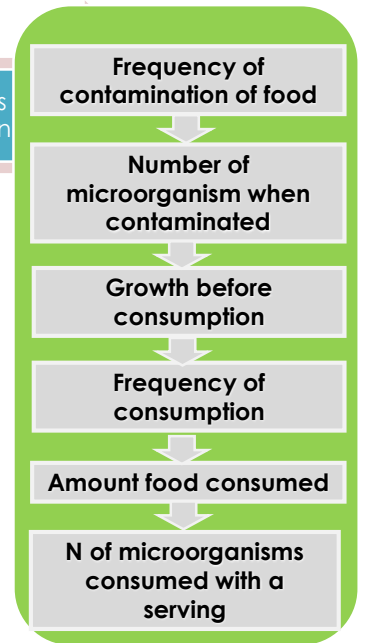


## Key Elements of Microbial Risk Assessment

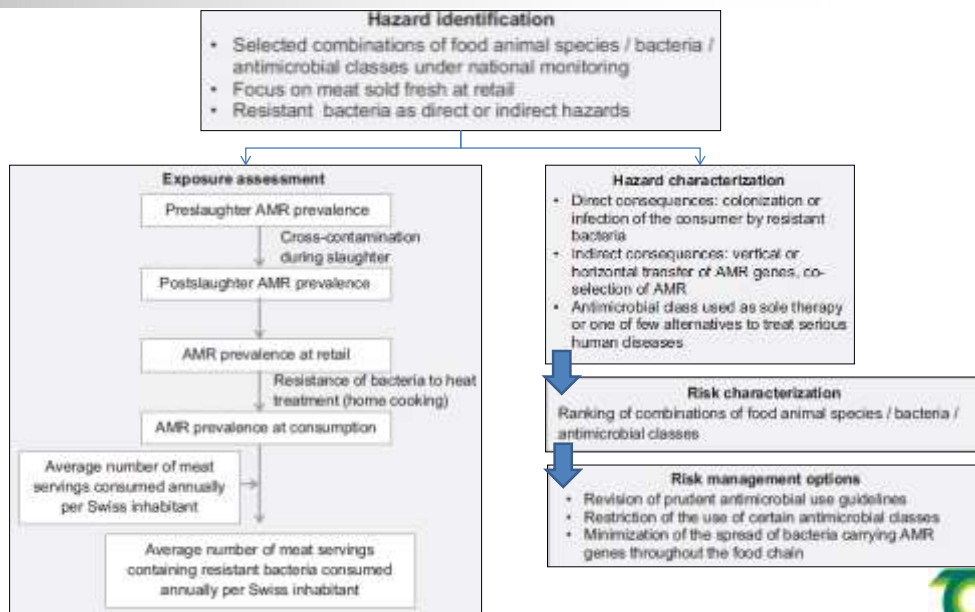
Hazard Identification  
Exposure Assessment  
Hazard Characterization (Dose-Response Relation + Severity Assessment)  
Risk Characterization

## FAO/WHO Microbial Risk Assessments

*Listeria monocytogenes* – ready-to-eat foods  
*Salmonella* in broilers  
*Salmonella Enteritidis* in eggs  
*Campylobacter* in broilers  
*Vibrio parahaemolyticus* in fish and shellfish  
*Enterohemorrhagic Escherichia coli* in ground beef and produce



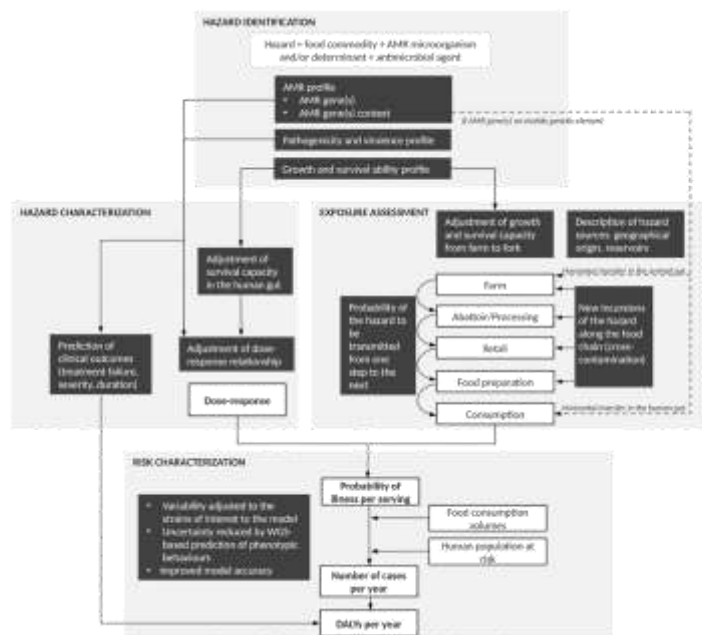
# Overview of the risk assessment framework



Collineau et al (2018) Risk Analysis, Vol. 38, No. 5, 2018



## Overview of the risk assessment framework

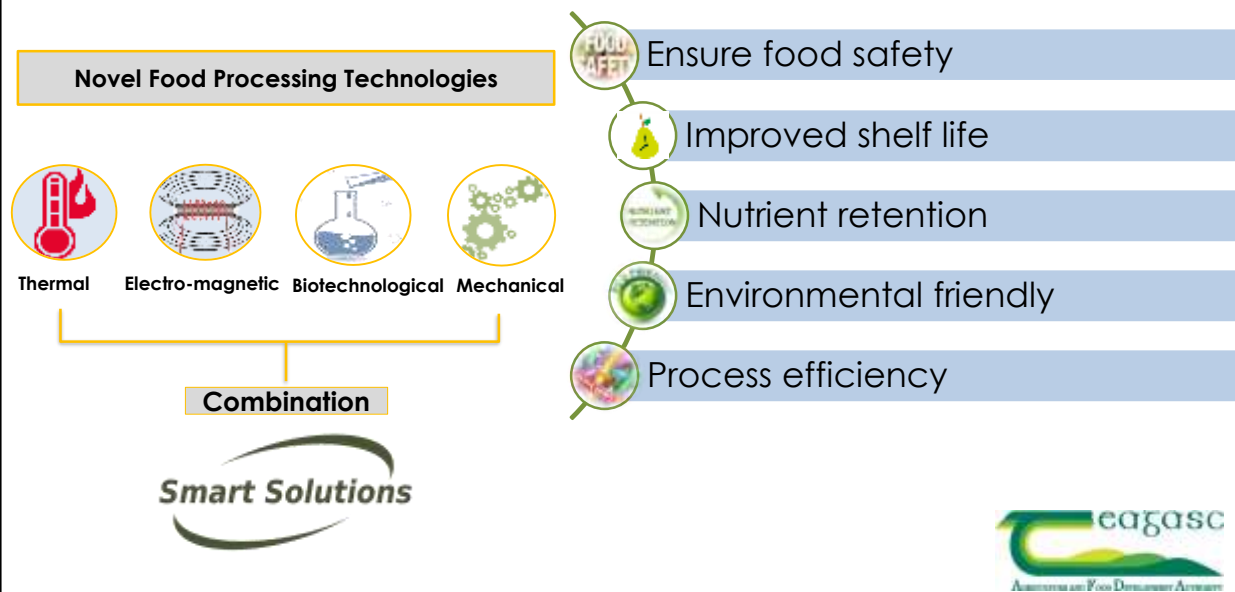


## Evidence of cross-resistance between biocides and clinically relevant antibiotics

Microbial species	Biocides used	Antibiotics to which resistance was developed
E. coli Pseudomonas spp.	Triclosan benzalkonium chloride, Didecylidimonium chloride and sodium hypochlorite	Levofloxacin, amoxicillin, tetracycline and chloramphenicol Colistin, ceftazidime, amikacin, meropenem, gentamicin, piperacillin-tazobactam, ciprofloxacin, Polymyxin B, tetracycline, ciprofloxacin
Bacillus spp., Enterococcus spp. and Staphylococcus spp.	Cetrimide, chlorhexidine, benzalkonium chloride, hexadecylpyridinium chloride	Ampicillin, sulfamethoxazole, cefotaxime, Ceftazidime and cefotaxime, among others
Klebsiella pneumoniae E. coli, C. coli, S. enterica, L. monocytogenes C. jejuni	Chlorhexidine Didecyl dimethyl ammonium chloride	Colistin Ampicillin, cefotaxime, ceftazidime, chloramphenicol and ciprofloxacin Kanamycin, streptomycin
S. enterica serovar Typhimurium	Trisodium phosphate, sodium hypochlorite, acetic acid and a commercial alkaline biocide which contains sodium lauryl ether sulfate, linear alkyl benzene sulfonic acid, sodium salt, propylene glycol monomethyl ether, and dipropylene glycol n-propyl ether A mixture of aldehydes and QAC; a QAC; an oxidative compound; a halogenated tertiary amine compound, A blend of oxidizing compounds; a QAC containing formaldehyde and glutaraldehyde; a biocide composed of organic acids and surfactants	Nalidixic acid, ciprofloxacin, chloramphenicol, tetracycline, Nalidixic acid, chloramphenicol, tetracycline, ciprofloxacin, Ciprofloxacin, chloramphenicol, tetracycline, and ampicillin

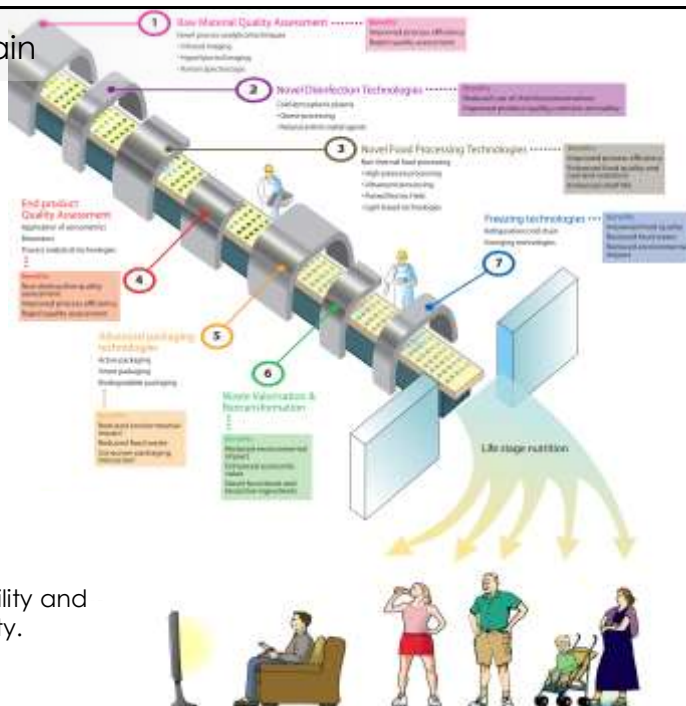


# Novel food processing technologies



## Novel technologies across food chain

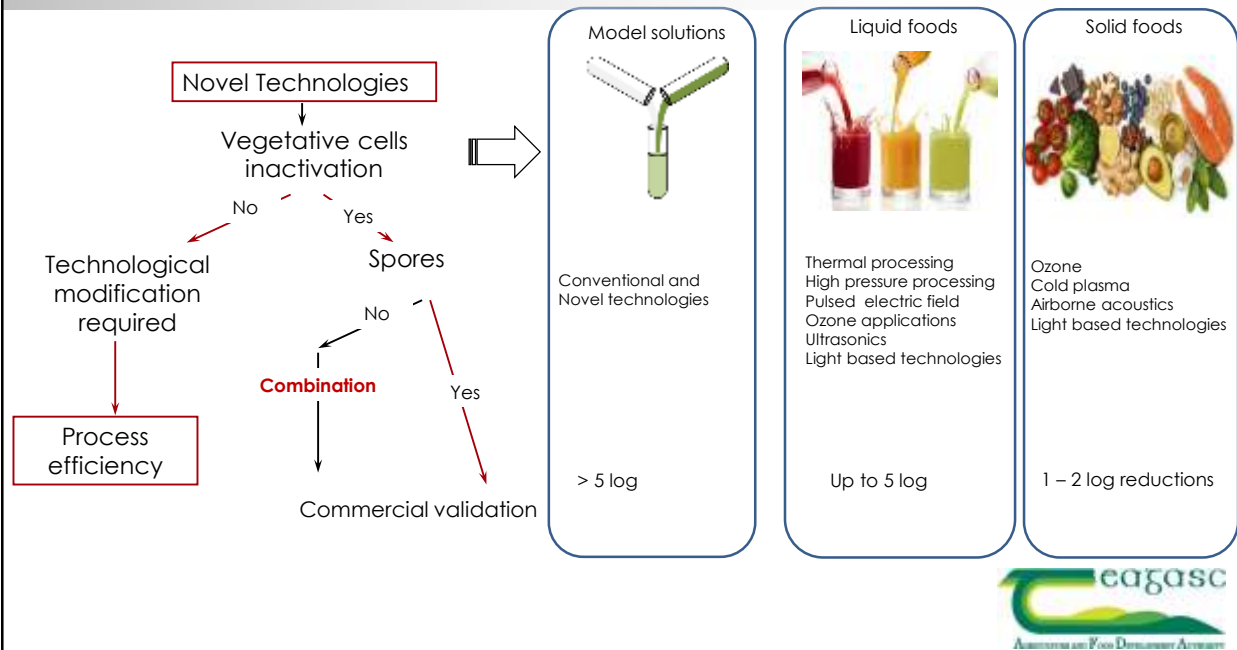
- ✓ Improved food quality
- ✓ Reduced **energy** and **water** consumption (clean and green solutions to key challenges faced by the food industry)
- ✓ Employ new interventions for developing new food products underpinning key health, nutrition and wellness challenges



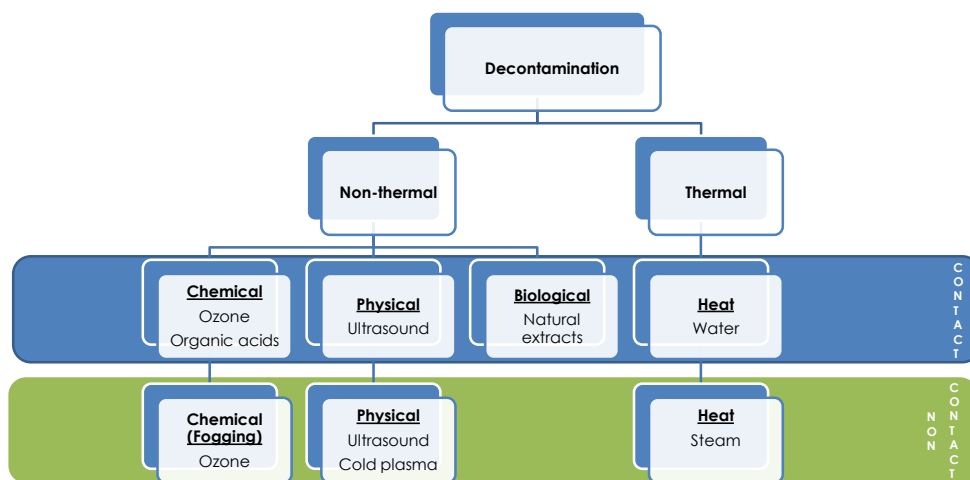
### Mechanism of AMR

- ✓ Enzymatic degradation of antibiotics,
- ✓ Antibiotic target modification,
- ✓ Changing the bacterial cell wall permeability and
- ✓ Alternative pathways to escape the activity.

## Selection of technologies



## Surface Decontamination Technologies





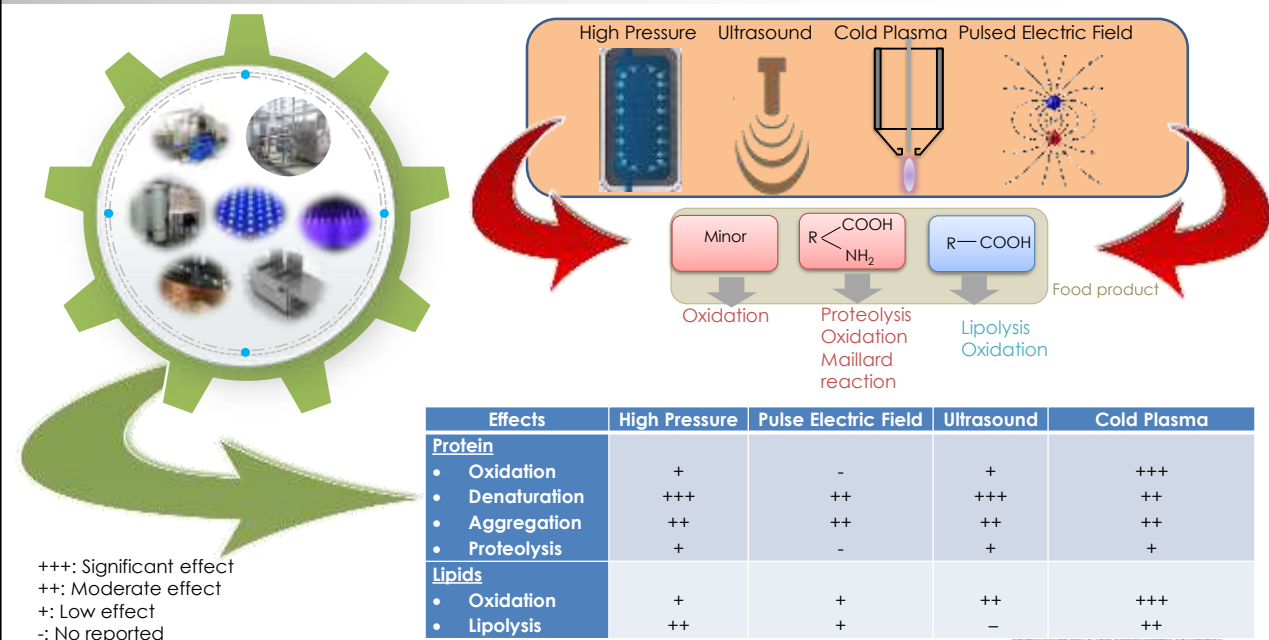
## Microbial aspects of novel technologies

Key mechanisms		PEF	US	Cold plasma	UV light	Ozone	Thermal
	Pressure	Electroporation	Sonoporation	Oxidation		Heating	
<b>Vegetative cells</b>							
Damages to cell membrane	+++	+++	+++	+++	+++	++	+++
Inactivation of key enzymes modulating growth of cells	++	++	++	+	+		+++
Oxidative damages to cell membrane constituents (peptidoglycan layers)	++	++	+	+++	+++	+++	+++
Damage to DNA/Nucleic acid	-	-	-	++	+++	++	+++
<b>Spores</b>							
Damage to spore coat	++	++	+++	++	++	++	+++
Chemical modification in spore core and cortex	+	++	+	++	+	+	+++

+++: key mechanism of action  
 ++: some synergistic effects  
 +: based on limited scientific information  
 -: Not reported

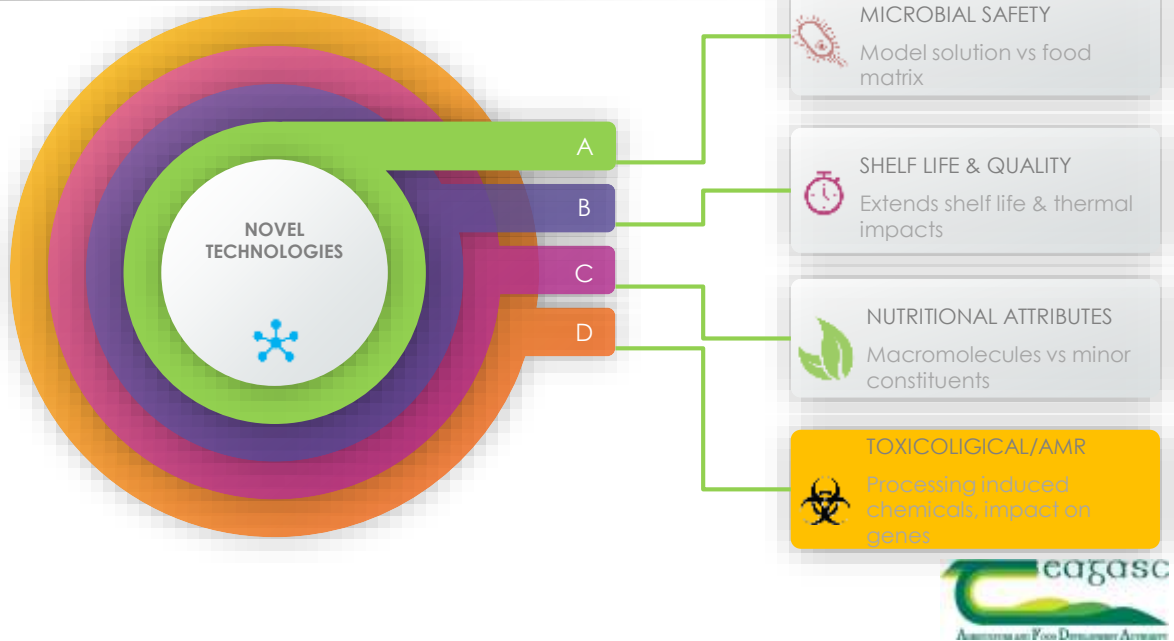


## Chemicals aspects of novel technologies



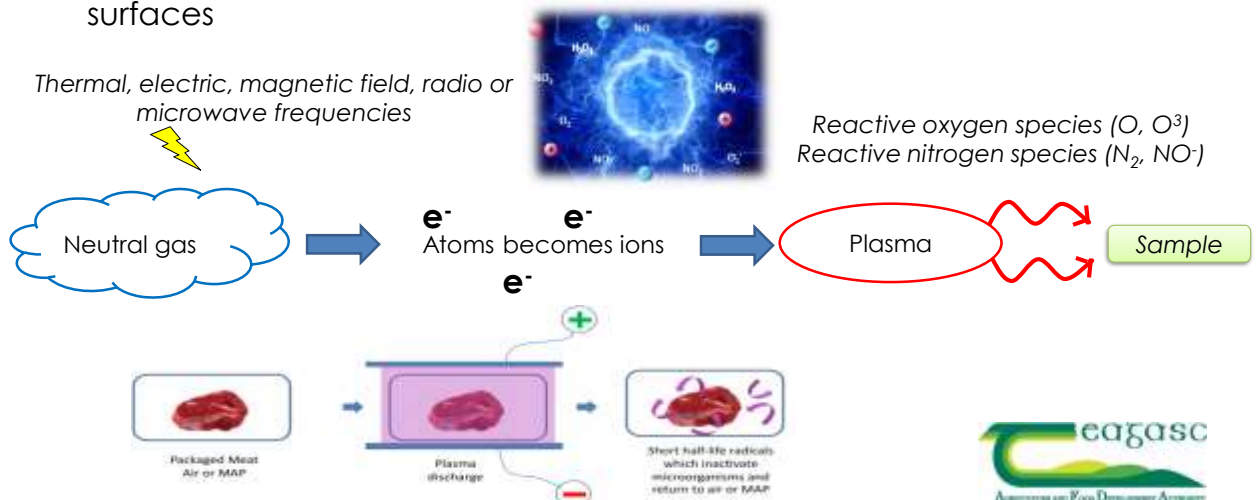


## Known and unknowns of novel technologies

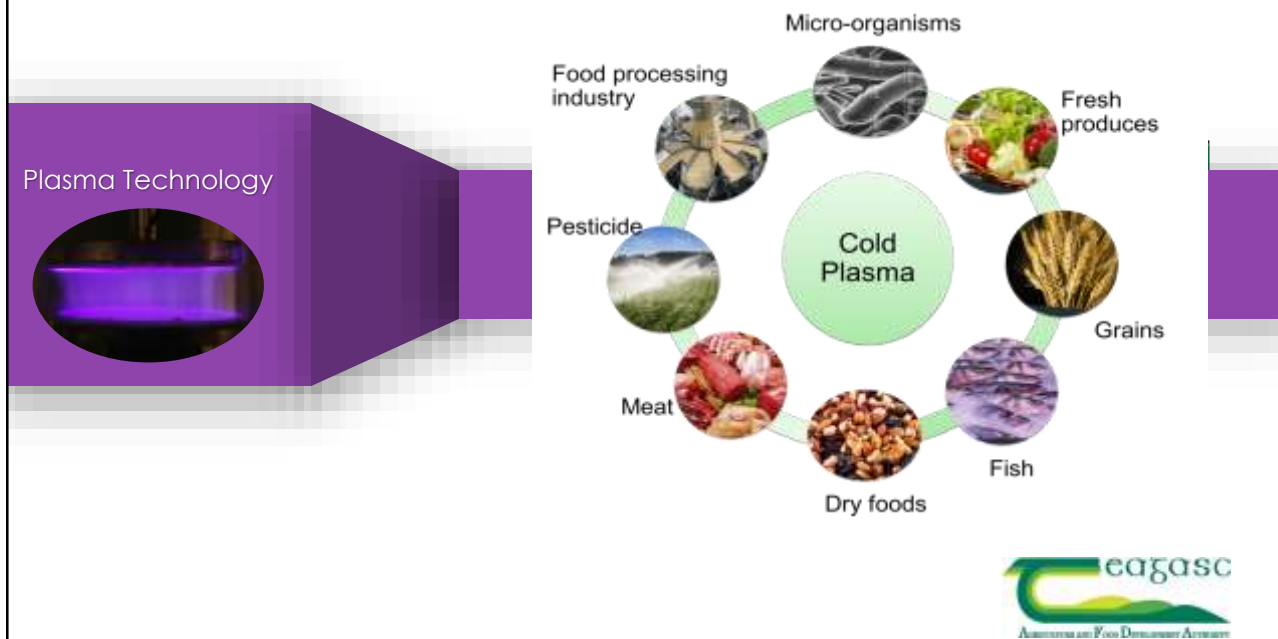


## Cold Plasma Technology

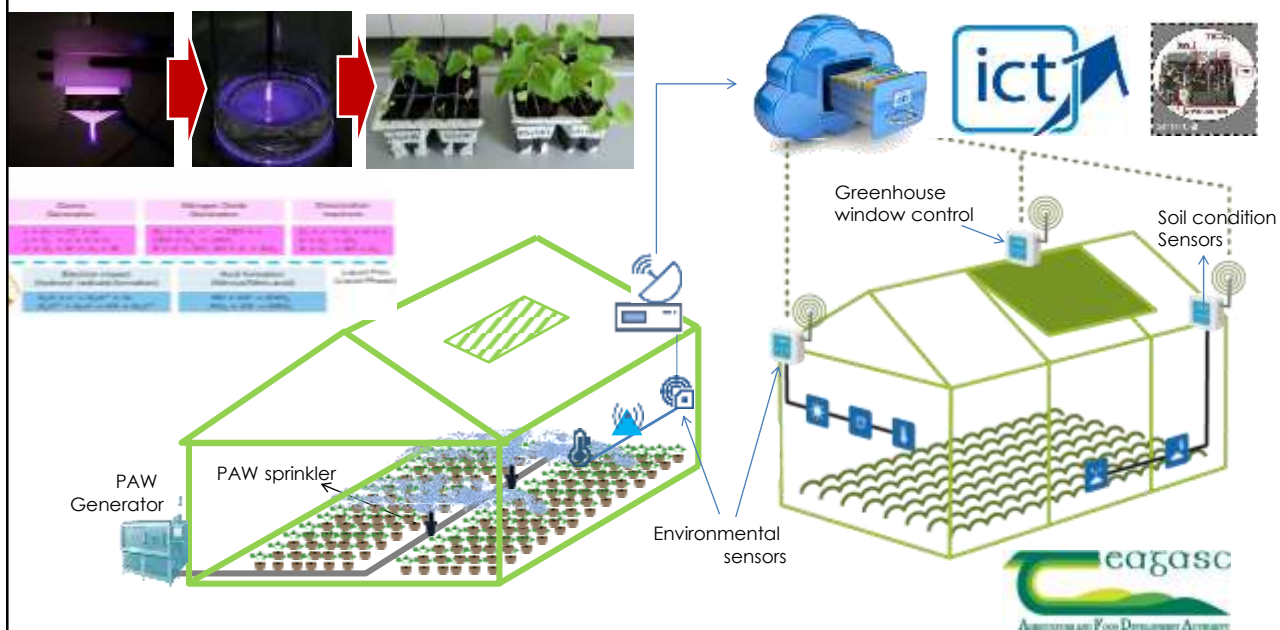
- Plasma (an energetic ionised gas) is widely used for industrial material processing.
- A novel nonthermal approach for reducing the microbial populations on surfaces



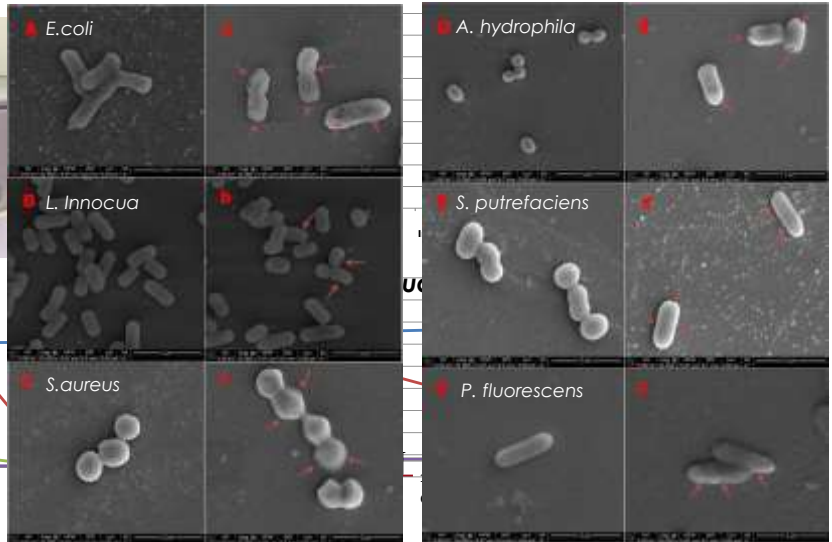
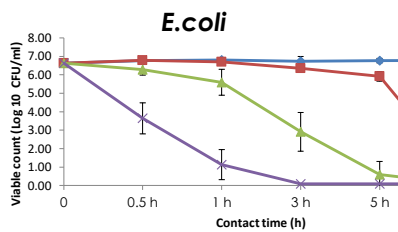
## Application of Cold Plasma Technologies



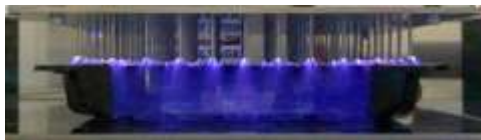
## Crop production



## Effect of PAW on microbial load



## Effect of Plasma on biofilms



Direct treatment of bacterial biofilms

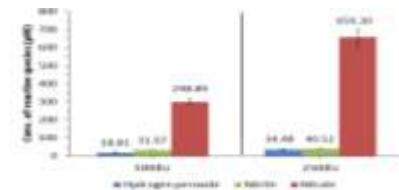


Control

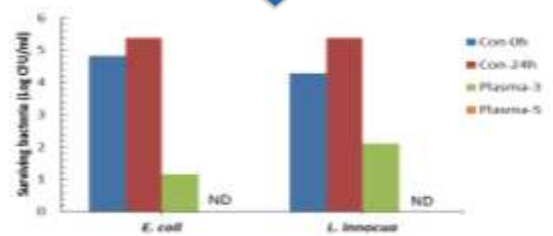


3min ACP treated

Confocal analysis: green (live cells), red (dead cells)



Reactive species generated





## Risk assessment – associated with Cold Plasma Technology

Potential risk factors using cold plasma technology:

- electrical charge system
- thermal damage
- formation of UV radiation on the product surface during treatment



Although no evidence on undesired side effects of a plasma treatment has yet been presented, research currently are in process to detect

- Short term → Cytotoxic effects
- Long term → Mutagenic effects



Electric currents, UV radiations displays certain impact on cell performance, however, so far no adverse effects have been attributed

No mutagenic action of the kinpen plasma in mammalian cell line



Complex bio-fluids showed increased mutation frequency over time

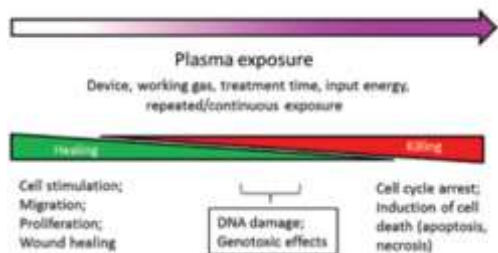


Treatment reduced the toxicity of the food sector effluents, however, Cyto-eco-toxicity varied depending on treatment parameters.



Endotoxin responsible for pathogenicity in bacteria is reduced with cold plasma and non-toxic to Galleria mellonella





The mix of these parameters has to be adapted individually for each application

Optimum treatment parameters should be set to minimise risk factors

Further investigations regarding the nature and safety of plasma-treated food products



## Conclusions

- Available risk assessment framework for QMRA should modify to focus on resistant micro-organisms
- Scientific information available on technologies is mainly associated with food safety/quality
- Limited studies are available on AMR/WGS of novel technologies
- Approval of Novel technologies for food application for food safety purposes is not yet approved in many countries.
- Majority of novel technologies are for niche application
- Replacement of antibiotics and chemical preservative with natural alternative would provide solution to AMR.