



MOLECULAR PLANT PATHOLOGY

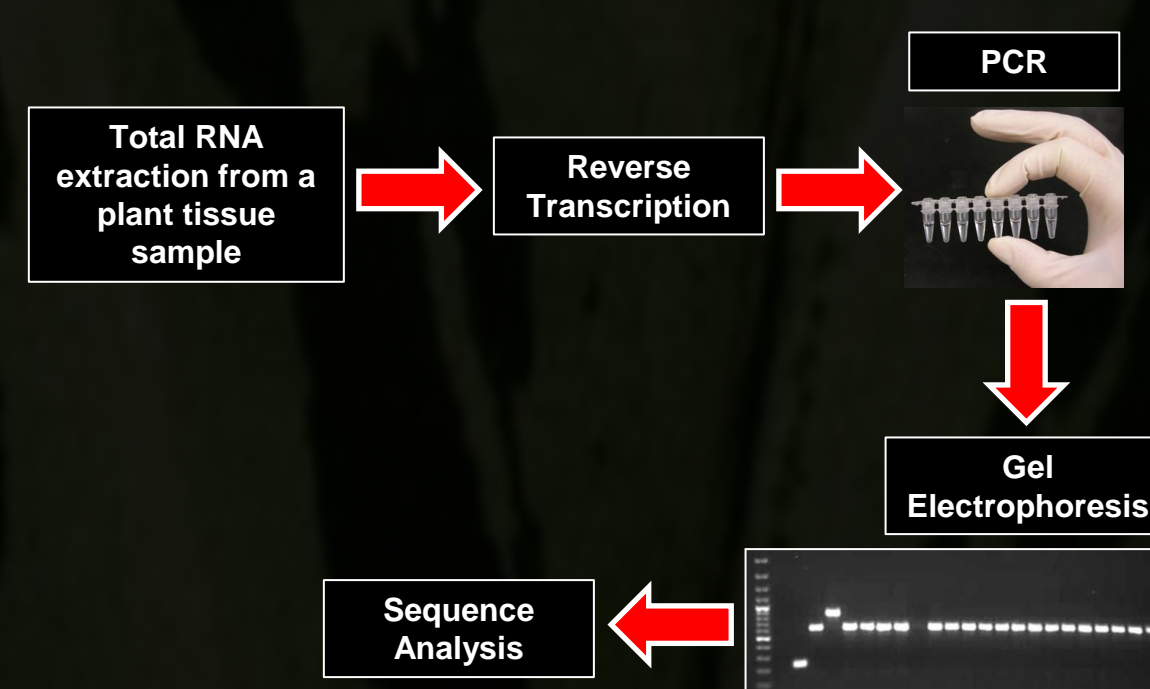
Objectives

Molecular techniques and technologies are used to accurately and quantitatively detect plant pathogens, to study their interaction with plants on a molecular and/or cellular level, and to investigate plant defense responses against them.

Pathogen identification and quantification

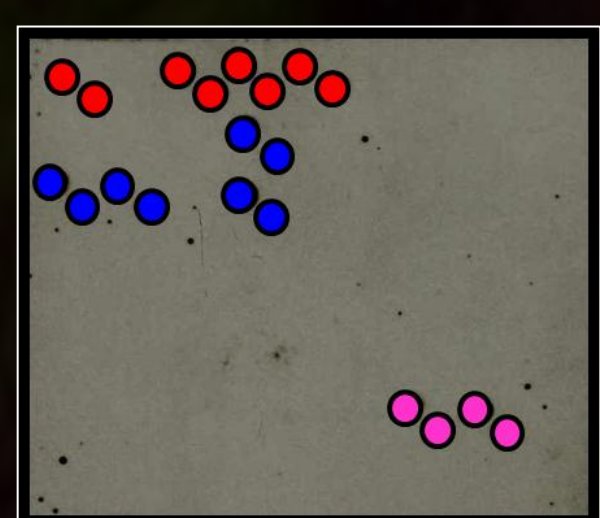
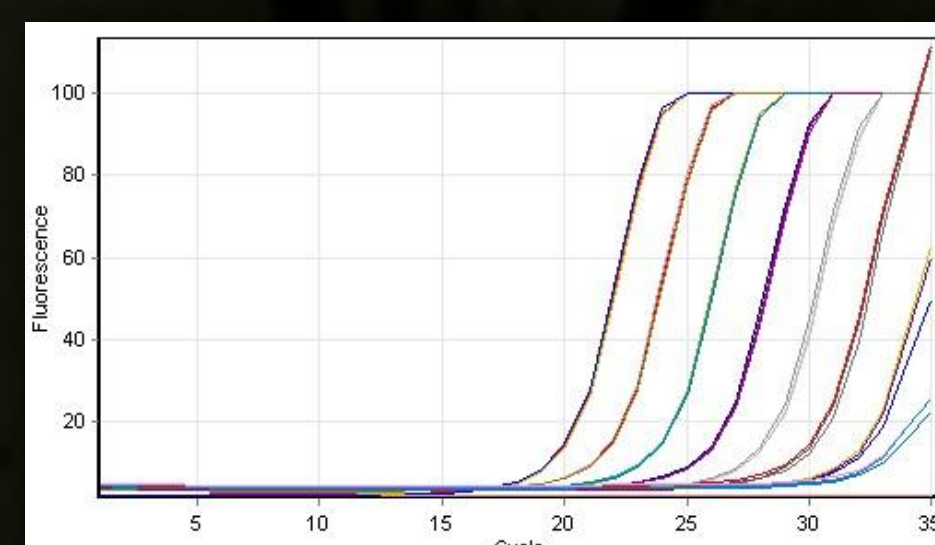
• PCR

The use of species-specific primers, molecular fingerprinting and sequence comparisons to identify important plant pathogens and distinguish between closely-related organisms.



• Quantitative RT-PCR

The quantitative detection of fungal pathogen DNA in plants, water, soil and the air – replacing spore counts. The infection co-efficient (IC) is determined by calculating the ration of pathogen DNA to host DNA.



• Macro-array analysis

Oligonucleotide arrays are developed and used for rapid detection and differentiation of oomycete and fungal species.



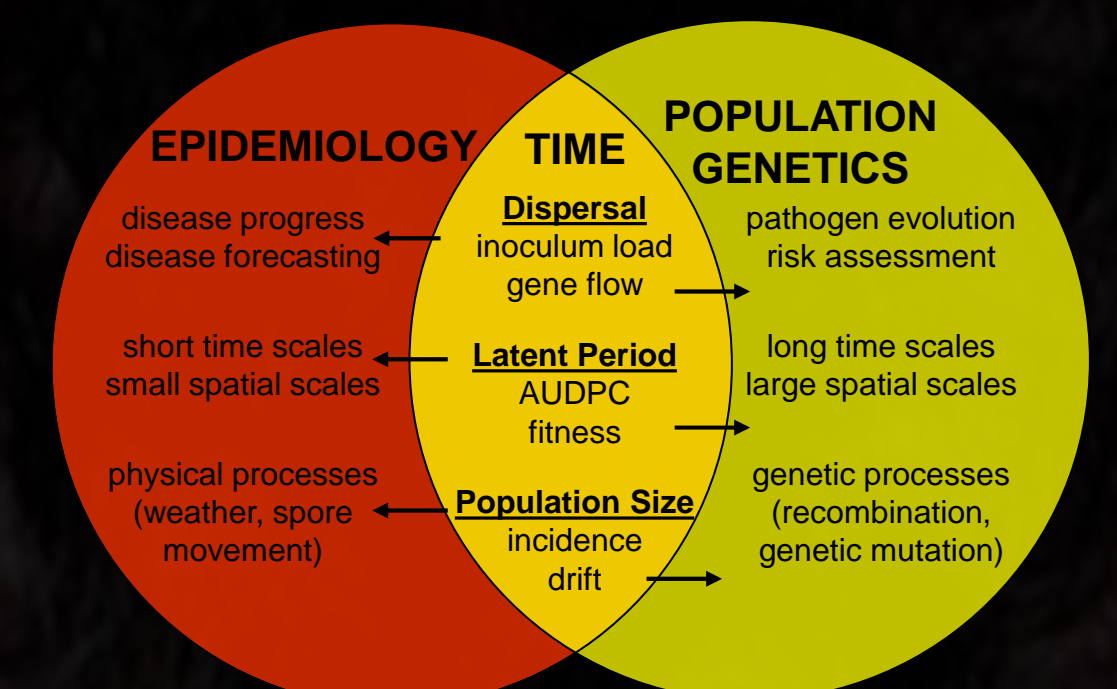
• Vegetative compatibility groups (VCGs)

Fusarium isolates that share common alleles at the same loci are grouped into vegetative compatibility groups following the generation of *nit* mutants that are paired with a known set of VCG testers.

Population genetics and phylogenetics

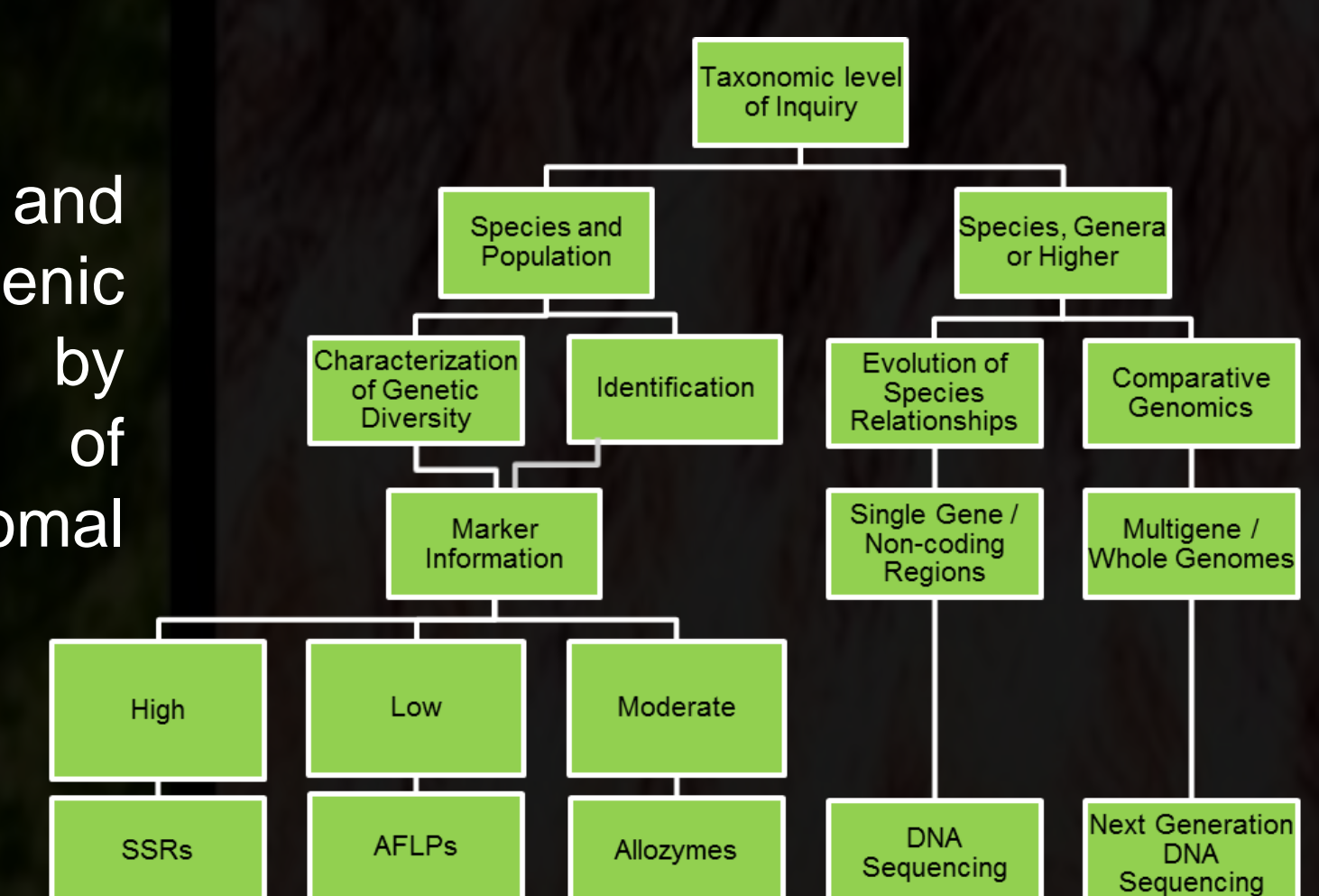
• Population genetics

Molecular markers, such as short-sequence repeats (SSRs, microsatellites) and AFLPs are used to study variation, migration, gene flow and recombination in fungal populations.



• Phylogenetics

The evolutionary history and taxonomy of plant pathogenic species are determined by phylogenetic analyses of relevant nuclear, ribosomal and mitochondrial genes.



• Comparative genomics

Genes and genomes of plant pathogens are compared by multi-gene and new generation sequencing to study their evolution and diversification.

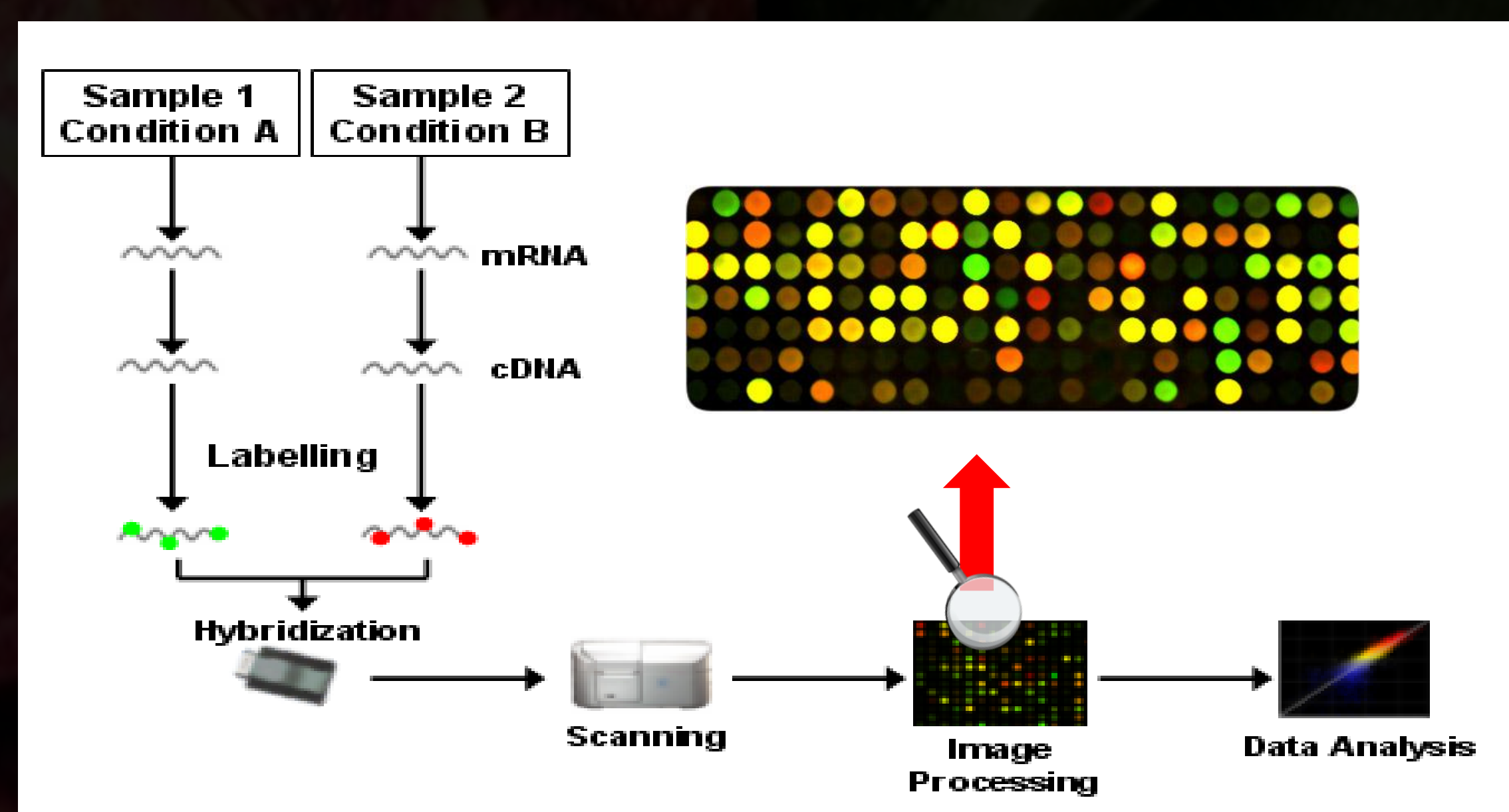
Gene expression analysis

• Differential gene expression

Technologies such as suppression subtractive hybridisation (SSH) are used to compare populations of mRNA in plants and pathogens and isolate differentially expressed genes.

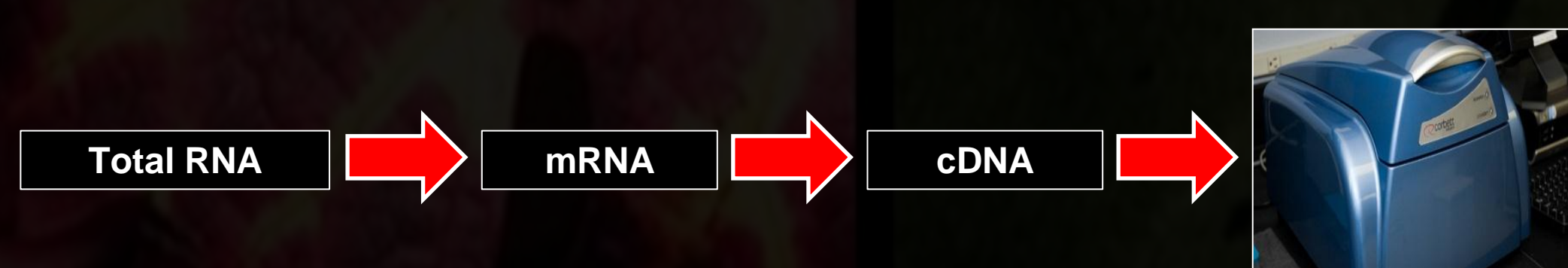
• Micro-array/Genechip technology

The simultaneous expression of large numbers of genes in plants and pathogens are determined by using technologies such as Affimetrix and Agilent.



• Reverse transcription quantitative PCR (RT-qPCR)

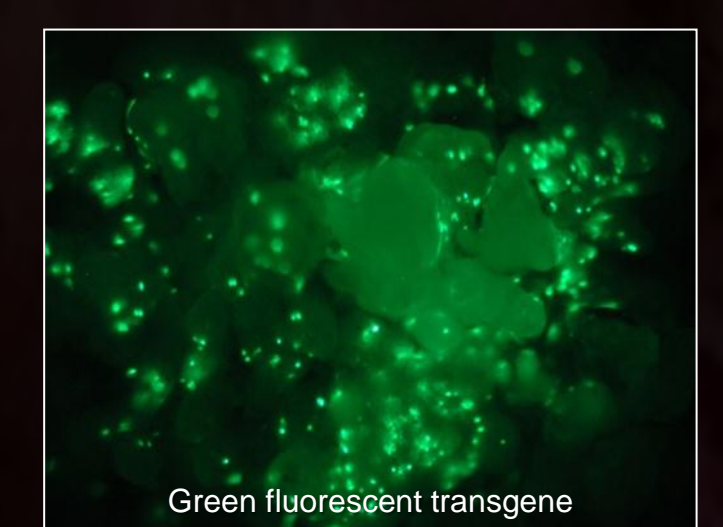
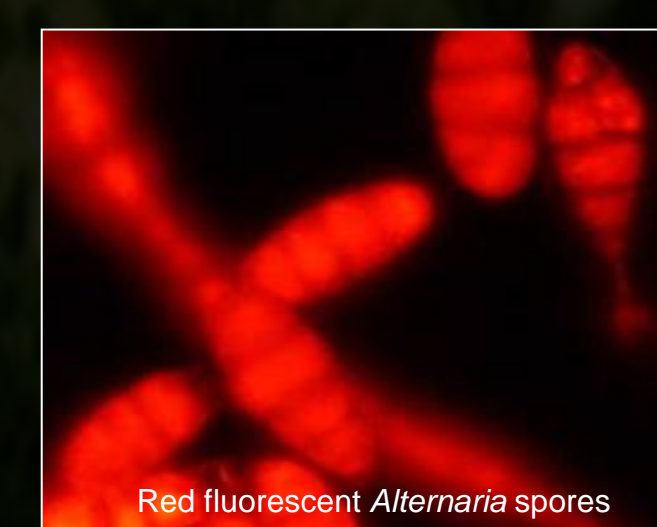
RT-qPCR is used to quantify differences in gene expression levels of a specific target (gene) between different samples.



Genetic modification

• Recombinant DNA technology

Transgenes are inserted in agricultural crops and plant pathogens for studies on host-pathogen interactions and plant resistance.



• Mutation breeding

Agricultural crops are modified by irradiation for improved resistance to fungal pathogens and their secondary metabolites.



Molecular plant breeding

• Marker-assisted selection (MAS)

Molecular markers are used to identify quantitative trait loci (QTLs) in resistance and susceptible plants that can in future be used for MAS.

