

# **THE FUTURE OF MICROBIOLOGY**

## **The ongoing revolution**

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# Agenda

- **Primary missions**
- **Background**
- **Reduction of time for microbial detection and identification**
- **Non cultural approach & POCT**
- **Automation of culture process**
- **Medical microbiologist: the profession**
- **Key messages**

# Primary missions

**TO IMPROVE THE MANAGEMENT OF INFECTIOUS DISEASE**

## **CONTRIBUTION TO DIAGNOSTIC**

**Presence /absence of pathogens  
Identification +/- quantification**

## **CONTRIBUTION TO CHOICE OF ANTIBIOTHERAPY**

**Antimicrobial susceptibility testing, identification of resistance  
mechanisms and resistance genes**

## **SUPPORT TO INFECTION CONTROL**

**To provide useful, accurate and relevant results**



# Useful results

## POSITIVE IMPACT ON

- **Therapeutic decision?**
- **Optimized management of patients?**
- **Morbidity, mortality?**
- **Length of hospitalization?**



*Reduction of Turn-Around-Time  
for result and its notification to clinician*

- **Control of nosocomial infections?**
- **Antibiotic consumption?**
- **Control of antimicrobial resistance?**

# Background

*To understand priorities for the future*

# Medical evolutionary background

## Impact factors on development and daily practice of microbiology

- **Economic environment**
  - Cost-effective use of available resources
- **Trained human resources**
  - Population pyramid and labour shortage
- **Medical environment**
  - Increasing emphasis on evidence-based medicine and adherence to guidelines
- **Technological background**
  - Exponential progress: molecular biology and robots
- **Quality assurance, traceability**
- **Global increase of antimicrobial resistance**

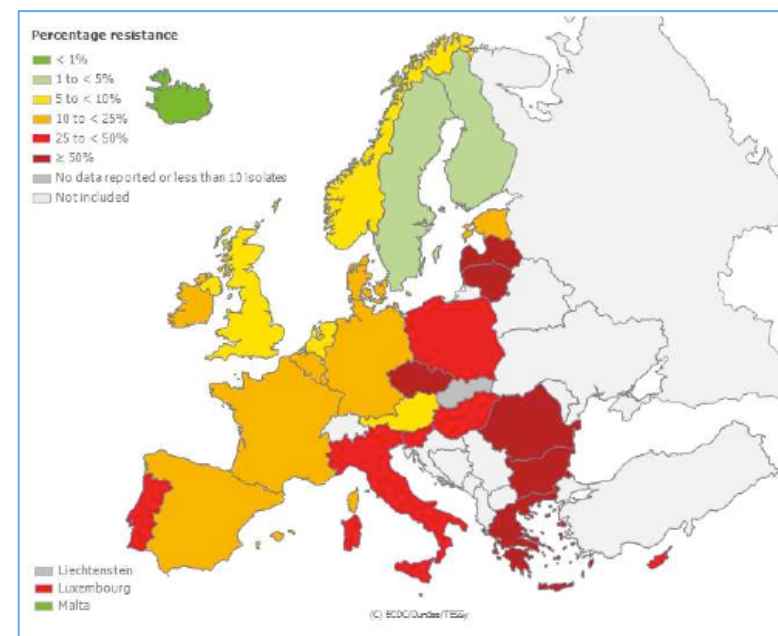
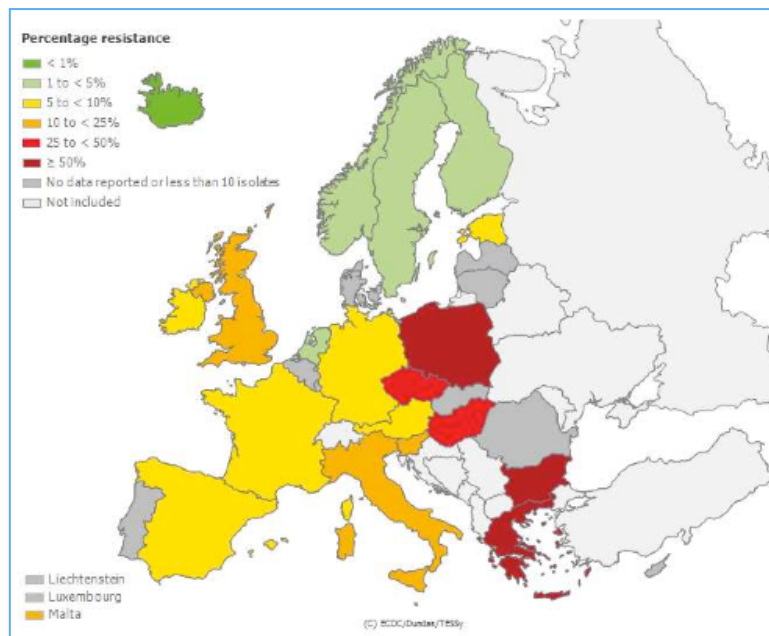


# Global increase of antimicrobial resistance

## Emerging superbug

Invasive isolates in Europe : 2005 vs 2009 (EARSS-ECDC)

Proportion of 3<sup>rd</sup> generation cephalosporin resistant *Klebsiella pneumoniae* (R+I)



Antimicrobial selective pressure  
Abusive use of broad-spectrum antibiotics  
Delay to identifying MDR bacteria carrier = bad control of transmission

# Challenges greater than ever in all parts of the world

- **Numbers of emerging infections**
- **AND Global increase of antimicrobial resistance**
- **AND Drug development pipeline for new antibiotics virtually empty !!**

**THREAT within the next 10 years**

**Shortage crisis of effective drugs for treating  
bacterial infection**

# European research priority

## To combat growing problem of antimicrobial R

- **Multidisciplinary research**
- **EU Framework Programmes**
  - **Studies on molecular mechanisms leading to emergence and spread of R genes**
  - **Epidemiological studies with aim of developing control and preventive measures**
  - **Translational, clinical and public health research**
    - **For developing guidelines for evidence-based clinical practice and patient management: for prudent use of antibiotics**



# European research priority

## To combat growing problem of antimicrobial R

### ■ EU Framework Programmes

- To promote the rational use of antibiotics
  - Mobilization of European biotechnology industry to develop **POC diagnostic tests**
    - *For early identification and differentiation of infectious disease-causing agents*
- To seek for novel molecular targets in pathogens and candidate drugs against them

### ■ Strengthened Transatlantic cooperation

- Transatlantic Taskforce on Antimicrobial Resistance (TATFAR since 2009)

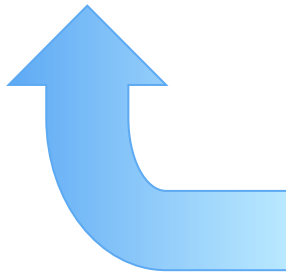
# Reduction of time for microbial detection and identification

***“Need for speed”***

# Medical microbiology laboratory Challenges in 2011

## *Theranostic approach*

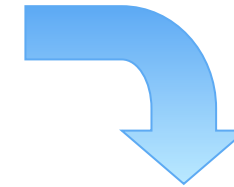
**Optimized  
management  
of patient and  
Infectious  
diseases**



**Turnaround time  
collection of specimen**



**Identification  
AST**



**Specimen  
Analysis:  
Relevant  
pathogens**



***Delayed results are unhelpful for clinicians !***



# How could rapid bacterial identification improve the management of septic patients?

*“... every hour without adequate treatment in septic shock is clearly associated with a 7.6% decrease in survival ...”*

## ■ Severe sepsis and septic shock

- Major motivation to identify bacteria present in blood quickly
  - Tools exists for positive blood cultures (MALDI-TOF MS, PCR, microarray, ...) but far more rapid needed

## ■ Clinical impact of rapid ID even if not AST

- Clue towards primary site of infection and targeted investigation
- Reduction of empirical broad-spectrum antibiotics to a more narrow pre-emptive therapy treatment
  - Based on usual susceptibility (or natural R) of pathogens
  - If *E.coli*: stopping vancomycin and macrolides whenever prescribed

# Rapid identification of a pathogen

**Prime importance for effective provision of care to patients with infections**

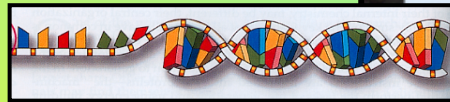
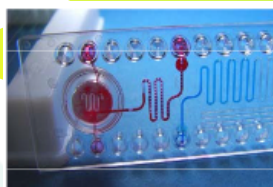
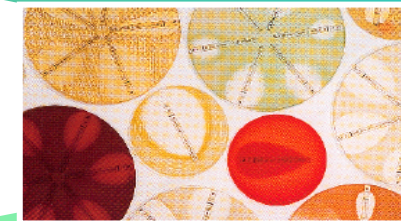
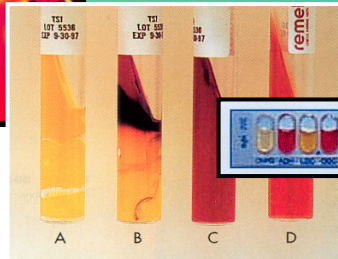
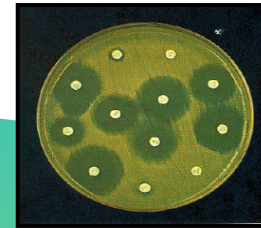
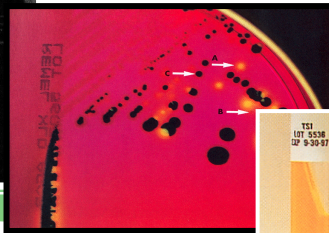
## ■ Includes

- Use of rapid bedside diagnostic tests
  - Need to be backed up by conventional culture techniques (for AST)
- Rapid reporting of results
- Pre-emptive culturing !!
- Molecular diagnostics
  - Bacteremia is different from DNAemia
  - Sometimes challenge to interpret

# Non cultural approach & POCT



# Technological evolution



# Identification and MALDI-TOF MS (>2009)



- 20-years old technology
  - Becomes the **standard for ID**
    - < Isolated colonies
    - < Positive blood cultures
- C.Meex's poster*
- Bacteria, yeasts, fungi
  - Results within a few minutes
  - Results 24hours prior to conventional methods
  - Interface LIS
  - Cost-effective
  - Clinical and therapeutic impacts

# Various mass spectrometry methods & application in clinical microbiology

- **MALDI-TOF MS** (Bruker; Shimadzu/bioMerieux)
  - Bacterial ID
  - More specific databases in development
    - Virulence and resistance markers
- **PCR-ESI-MS** (*Electro Spray Ionization*), **Plex-ID**
  - Directly from sample : DNA extraction
  - 4-6 hours
  - Semi-quantitative ID of all organisms present including virus
    - Syndromic panels of infection
  - Coupled with amplification by PCR, broad range of primers
  - High sensitivity
  - Detection of virulence or resistance genes
  - Results // to microarrays but faster
  - Still very expensive

# Various emerging applications of sequence-specific DNA amplification

- **PCR and Real-time PCR**
  - **Single or multiplex**
  - **Screening, detection and quantitative assays**
- **Loop-mediated isothermal AMPLification (LAMP)**
  - **Direct detection**
  - **Fast, sensitive, robust**
    - *Salmonella, C.difficile, group B streptococci, ...*
- **Broad-range 16S rDNA PCR followed by sequencing**
  - **Major milestone in diagnostic of infectious disease**
    - **Negative cultured bone and joint infections, endocarditis,**  
**Al Masalma et al., 2009; Fenollar and Raoult, 2007**

# Point of-care-test platforms for early diagnosis of infection

To provide an integrated solution addressing technological challenges

- For rapid detection of bacteria, mycobacteria, fungi, viruses, host markers and resistance to antimicrobial drugs
- To enhance clinical decision-making
- To improve quality of care and clinical outcomes
- To improve targeted therapy and reduce overuse
  - Specific probes
  - Novel methods of sample preparation
  - Ultra-high sensitive detection methods

Results availability

in less than 2 hours/30 min for IN/OUT patients

# Point of-care-test platforms for early diagnosis of infection

## Among EU funded projects

- **RAPP-ID** - Development of **RA**pid **P**oint-of-Care test **P**latforms for **I**nfectious **D**iseases
  - Bloodstream infections
  - Lower respiratory tract infections
  - Tuberculosis
  - *From Be: UIA, KUL, RUG and Imec*
- **C4L** - **C**hips **f**or **L**ife
  - Lower respiratory tract infections
  - *From Be: CHU-Ulg, UIA and Coris*

**Huge challenges and synergies:  
Biotechnologies, microtechnologies and clinical practice**



# Automation of culture process

# Automation in bacteriology

## Medical and economical plus values

- Labour shortages
- Increasing test volumes
- Reduction of TAT
- Reduction of laboratory errors associated with specimen handling
- Full traceability
- Improved workflow
- Enhanced quality

**The ongoing challenge**

### Availability of high-throughput robots

- Gradually being introduced into everyday microbiology diagnostics
- Most cost effective use of resources

# Automation in bacteriology

## Medical and economical plus values

### The ongoing challenge

#### Most cost effective use of resources

- Rational throughput if
  - Large volume of specimens
  - Often centralisation of services
- Danger of large-scale microbiology labs
  - Logistics rather than diagnostics
  - Risk of lost of clinical focus on the patient

# First fields of automation and improvement : bloodcultures and mycobacteria



**BactAlert**



**MGIT  
Bactec**

**2011: 10<sup>th</sup> generation !!**

- **Sensitivity**
- **Flexibility**
- **Ergonomics**
- **Continuous measures**
  - **Earliness of detection**
- **Interface with LIS**
- **Evolution of culture broths**

*Since 2009*

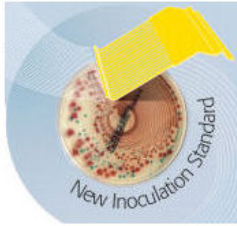
# Plating and inoculation of specimens

## ■ **Systems**

- **PREVI™ Isola, bioMérieux**
- **WASP, Copan**
- **INNOVA, BD**
- **Inocula, Kiestra**
  - **Semi-manual**
  - **Full-automated**

## ■ **Characteristics**

- **Media storage**
- **Types of specimen container**
- **Inoculation control**
- **Tips, swabs, volumes**
- **Plating method**
- **Streak patterns**
- **Traceability**
- **Biosecurity**
- **Sorting**
- **Interface with LIS**
- **Integration in a chain**
- **Slide module**
- **Output: plates/hour**



# PREVI™ Isola, bioMérieux



- **Liquid specimen**
- **Tips**
- **1 pattern : comb**
- **Max of 5 different media**
- **180 plates/h**



# WASP, Copan Slide module (option)

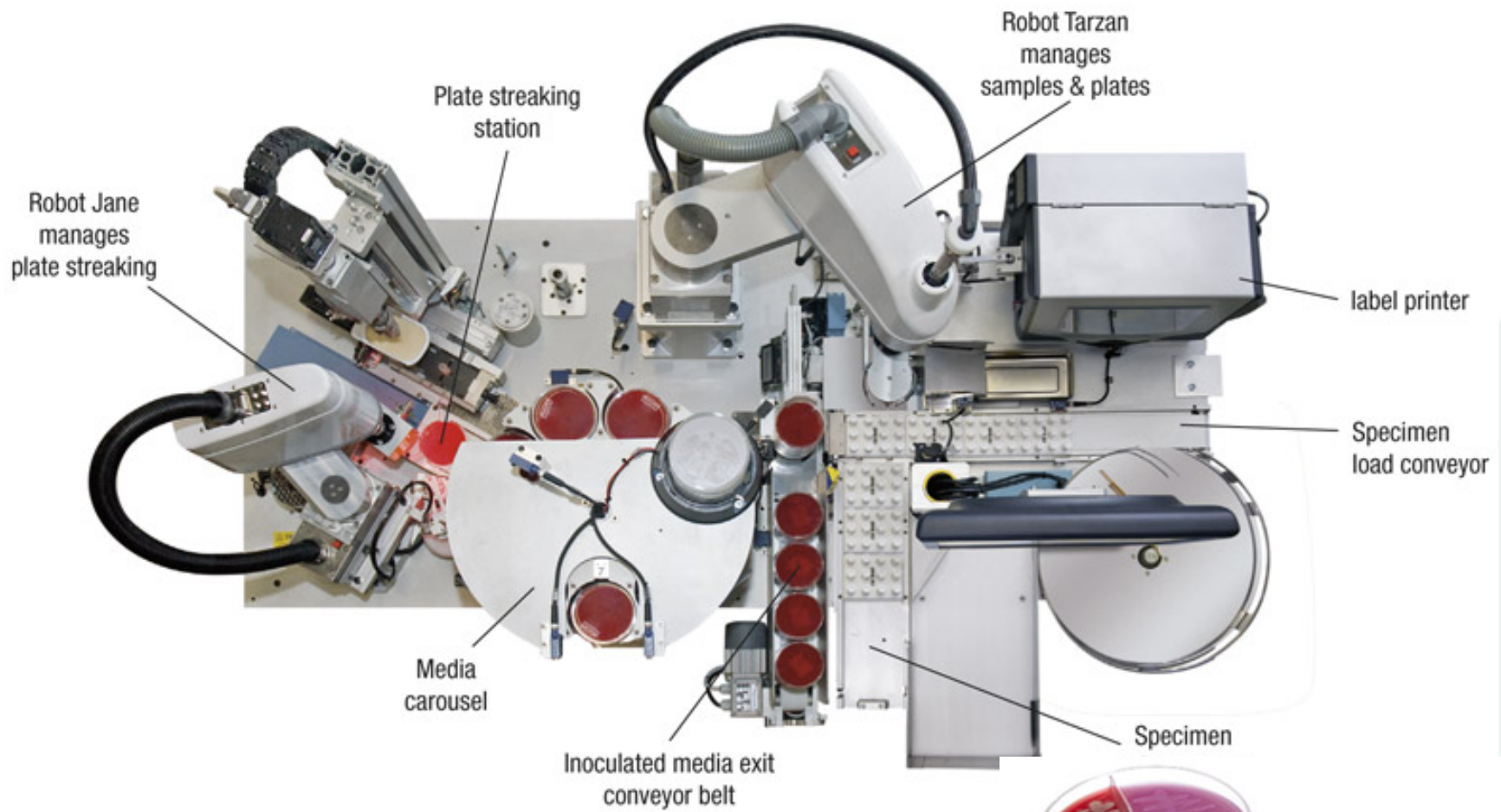


- Liquid specimens
- 1, 10 et 30  $\mu$ l
- Max 9 media



The WASP manages all specimen types & styles





# Inoqua, Kiestra Semi- and Full-automated inoculation

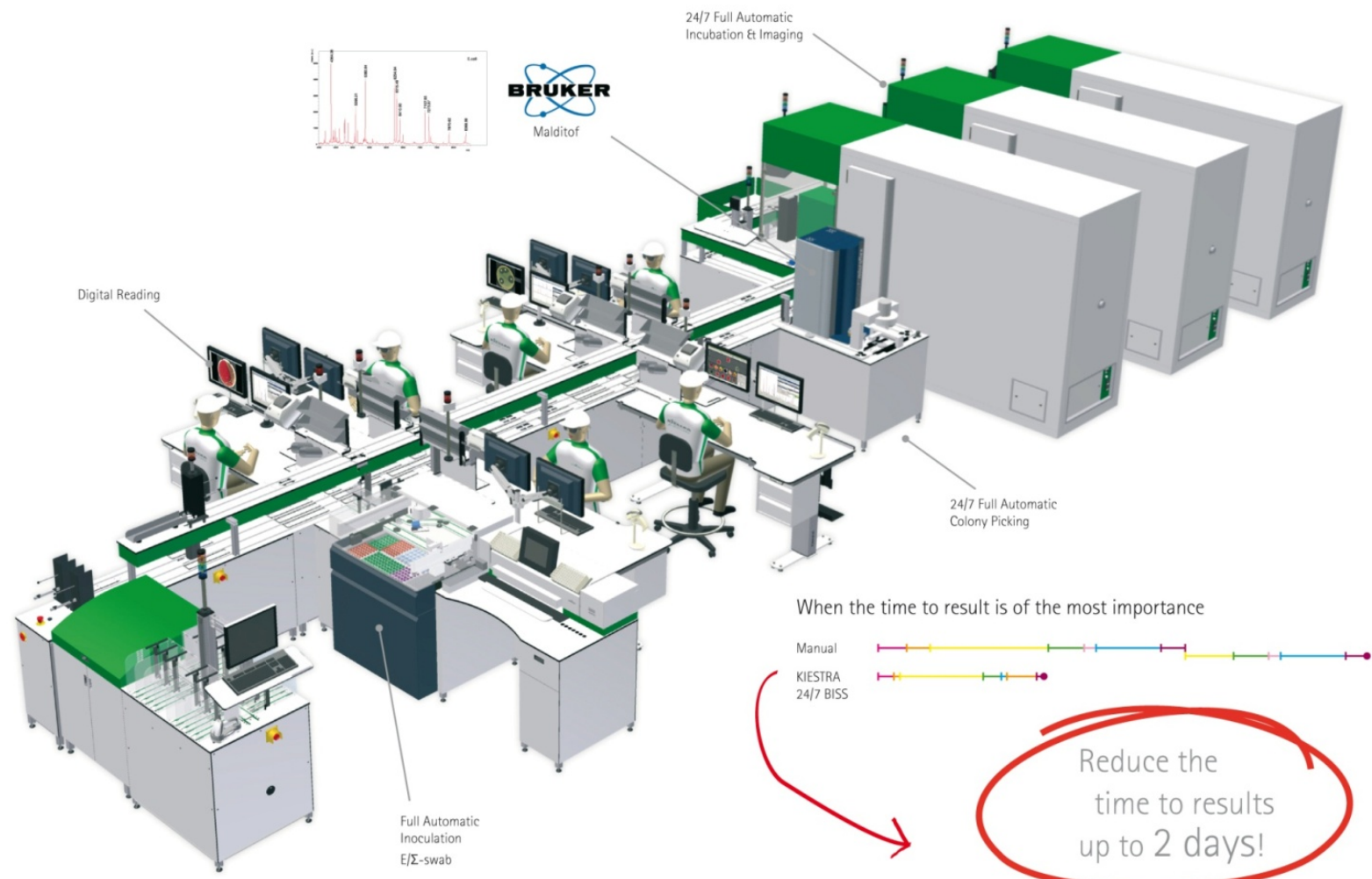


- 6 – 30 media
- Barcode labelling at side dish
- Full Automatic ( decap / cap, pipet ) in plates, slides and tubes
- Shaking / vortexing
- Manual Interactive inoculation
- All specimen types
- Sorting + Stacking for incubators
- 300 – 500 inoculations per hour

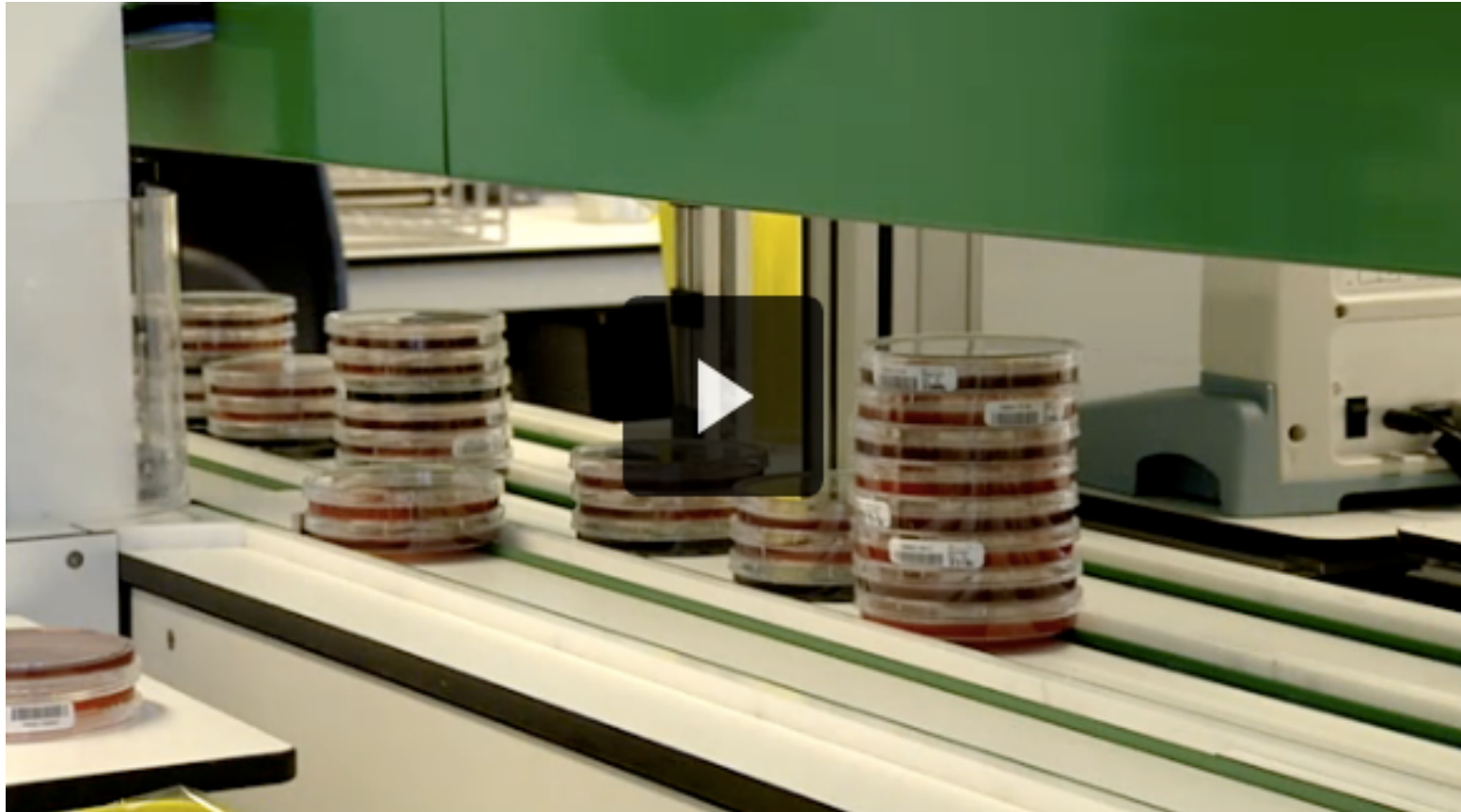




# «Future» : Total laboratory automation Kiestra, Wasplab (Copan), bioMerieux concept

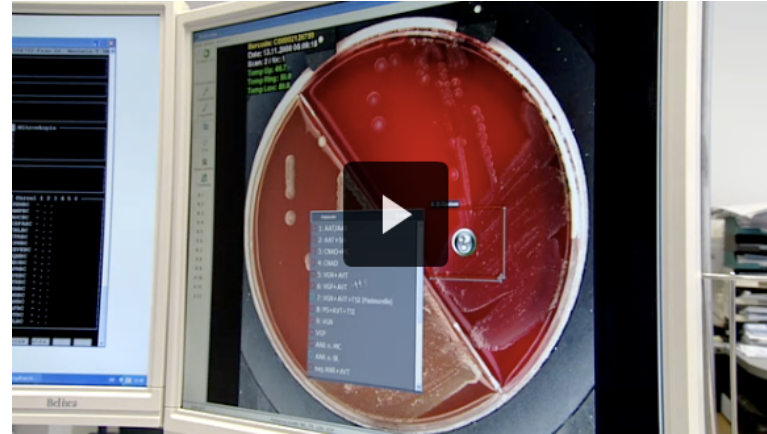


# Transport

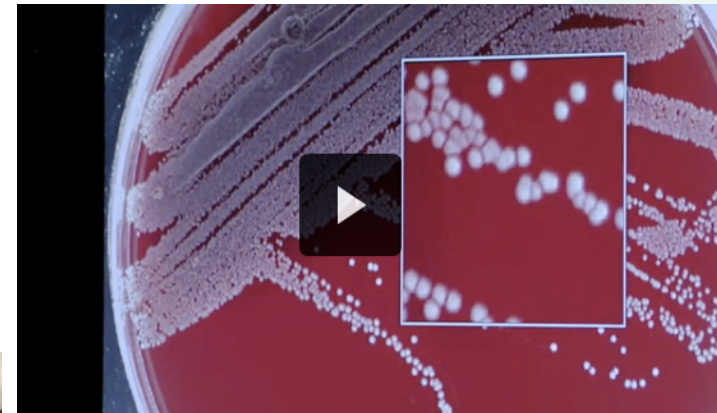


# Incubation, digitized images of cultures

## 24H/24, 7J/7



**Reading**  
-/patient  
-/type of specimen  
-/...



**Telebacteriology**  
**Training – teaching**  
**Verification, comparison**  
**Archives**



# Medical microbiologist

*To face the future*

# Medical microbiologist according to Union of European Medical Specialties (UEMS)

## Main tasks

- Advice on diagnosis, treatment and prevention of infections
- Implementation of antibiotic policy
- Generation of surveillance data

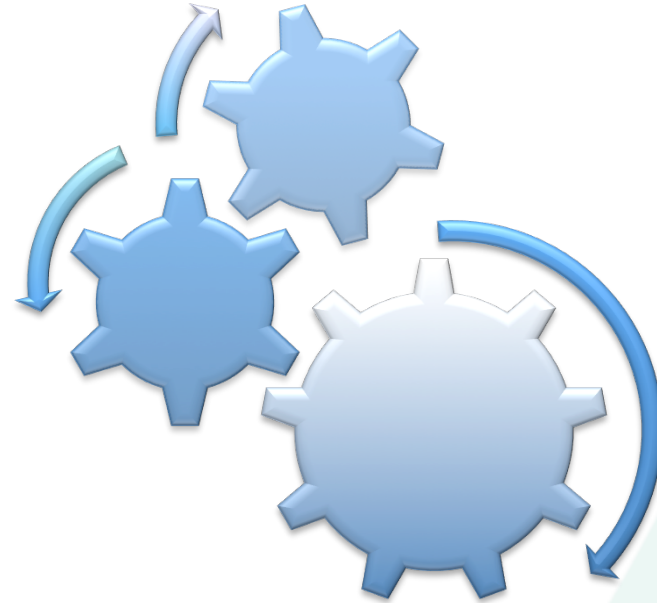
## To go to

- Integration of activities with Infectious Disease specialists to optimize the efficiency

# Medical microbiologist in Europe

**A specialty in many European countries**

- **Integrated training with Infectious disease**
- **Specialist's mobility through Europe**
- **In Belgium**
  - **> 10.2012 – A 2-year certificate in “Infectious disease and clinical microbiology”**
    - **25 modules**
    - **ULB, UCL and ULg**



***Mutations & a new culture are necessary to enjoy over the future of microbiology***

## **KEY MESSAGE**

- **Improving time to result**
  - **Clinical impact only if used without any further delay by clinicians**
- **Non-culture approach and back-up cultures**
  - **Cost-effectiveness & clinical relevance still to show**
- **To develop and master technological advances**
  - **Laboratory automation and improved workflow**
  - **Avalanche of new methods for rapid diagnostics**
- **Increasing pressure for**
  - **Concentration, amalgamation and outsourcing of lab service**

***Each laboratory should carefully think about how they can positively reorganize to speed up the diagnostics***

***A lot of opportunities to enjoy  
over the next few years !***

**Thank you**