Monitoring microbiological cleanliness of surfaces

Juha Koivisto, CTO CleanDet Oy

www.cleandet.com juha.koivisto@cleandet.com

Automatic detection of cleanliness

Automatic

- Software
- Internet of Things

Detection

- Sensors
- Illumination

Cleanliness

- Microbiology
- Material characterization



Global Examples of Emerging Infection Diseases



200 nm

- Harmless and normally live in the intestines of healthy
- Pathogenic strains cause relatively brief diarrhea, abdominal cramps and 18%

200 nm

5.11.2020

2013 -2015

Modified from Morens et al. 2004 Nature 430:242

Currently 2 weeks from infection to diagnosis

How we can prevent spreading of bacteria on touch surfaces in airports, ships, trains, hospitals, etc?

What we have?

- Sampling & sample preparation •
 - Chemicals
 - Trained personnel
 - Expensive equipment/laboratory
- Detection time ٠
 - from 1 hours to several days

What we need?

- Easy ٠
- **Real-Time**
- Rapid



Person is exposed to E.Coli Time to illness 1-5 days

Person becomes to carrier of infection 1-5 davs

Bacteria detection: Cultural techniques Biochemical tests Molecular methods Time to diagnosis 1-7 days



Microbial cultivation

Photometry

Analysis with microscopes















Earth surface is possible, Why not are touch surfaces?

Crop deseases can be detected from space



Could we detect

Microbial contaminants by the same method ?







It works because light interacts differently between different objects





This is a leaf.

Compare with optical coherence tomography or soap film





5.11.2020 6

Simple proof-of-concept: oils on steel

Separation of oils from metal

Uses fluorescence

- Input: UVA (black light, disco light)
- Output: red, green and blue

Off-the-shelf technology

- Sita Process Solutions Ltd
- Detection of storage oil residues





5.11.2020 7

Realtime classification



Video also at: http://youtu.be/Op_mQyjUuTIs



Materials & Method

- Matrix The solvent where bacteria is grown is Gibco LB Broth liquid
- Concentration of bacteria were from 0 to 6,5 x 10⁷ cells/50 μL
- Metal & plastic plates surfaces
- 3 illumination (red, green, blue) &
 16 imaging wavelengts were used

I_{bacteria} = I_{bacteria spot} - (I_{background} + I_{matrix})



Actual measurements with bacteria

Now you see it, now you don't

- Same gray
- bacteria either visible or hidden
- Depends on wavelengths





Bacteria identification

Plastic plate



Metal plate

Separation of signals of background & matrix from bacteria Separation of signals from *E.coli* & *B.Subtilis*

Bacteria identification

Plastic plate



Separation of signals of background & matrix from bacteria Separation of signals from *E.coli* & *B.Subtilis*

Metal plate

Decreasing consentration



Average location of the cluster 10^4 to 10^7 cells/50 µL

Adenosine Triphosphate testing (ATP) & Photometry for developed method validation

- Lower Limit of Detection is 10-20 times less
- The Relative Standard Deviation is ±3%
- The OD_{AutoDet} is higher than for validation methods and 2 times more intense for 250 filter than for 324 nm



Real life example: hospital waiting room

Take images of various surfaces and classify regions of images to dirty and clean



Hospital waiting room

Real data, real environment Visualize locations Reduce all physics to "hygiene index"







Clear surface – blood on steel

- Stain is clearly visible to human eyes
- Now you see it, now you don't principle can be reproduced
- Automation is possible



Oxidized surface

- Stain is very hard to see with human eyes
- Now you see it, now you don't principle can be reproduced
- Automation is possible



Remote detection contaminants possible

- More sensitive than ATP swab
- Skip the sample preparation
- Eliminate the usage of chemicals
- Detection in minutes
- Automated with infinitely long memory



- Now: detect biofilms
- Next: identify bacteria, then viruses

CleanDet Oy





