

PRÉLÈVEMENT DE L'AIR PAR TECHNOLOGIE CYCLONIQUE : OUTIL EXISTANT ET ÉTUDE DE CAS

M. Romain Verollet
Head of product management

www.bertin.fr

▷ Precellys® Homogenizers for biological sample preparation

- A full range of systems and consumables to prepare any type of sample in less than 1 min with high reproducibility



▷ Coriolis® bio air samplers for indoor and outdoor controls

- Bio air samplers based on airborne particles transfer into a liquid to go beyond traditional methods



▷ Sterilwave® medical waste management solution

- Based on microwave validated technology, on-site solution for bio-hazardous waste sterilization without effluent



10 YEARS OF WET CYCLONE TECHNOLOGY IMPLEMENTATION

▷ MonaLisa project – 2006-2009

- Validation of a new method for pollen and allergen detection



▷ French army project – since 2006

- Portable air sampler for airborne pathogens detection.

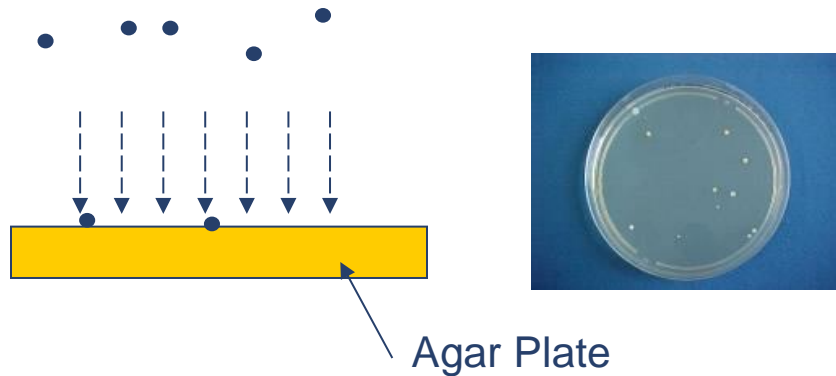


▷ Lab equipment – since 2009

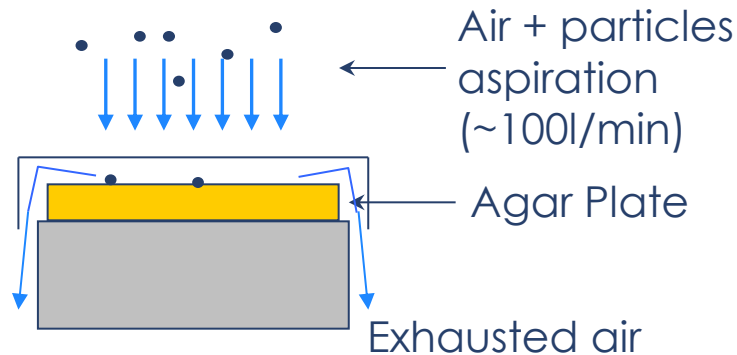
- Microbial air sampler for indoor and outdoor air bio-contamination.



▷ Passive: Settle plate exposed max 4h



▷ Active: Impaction is the reference

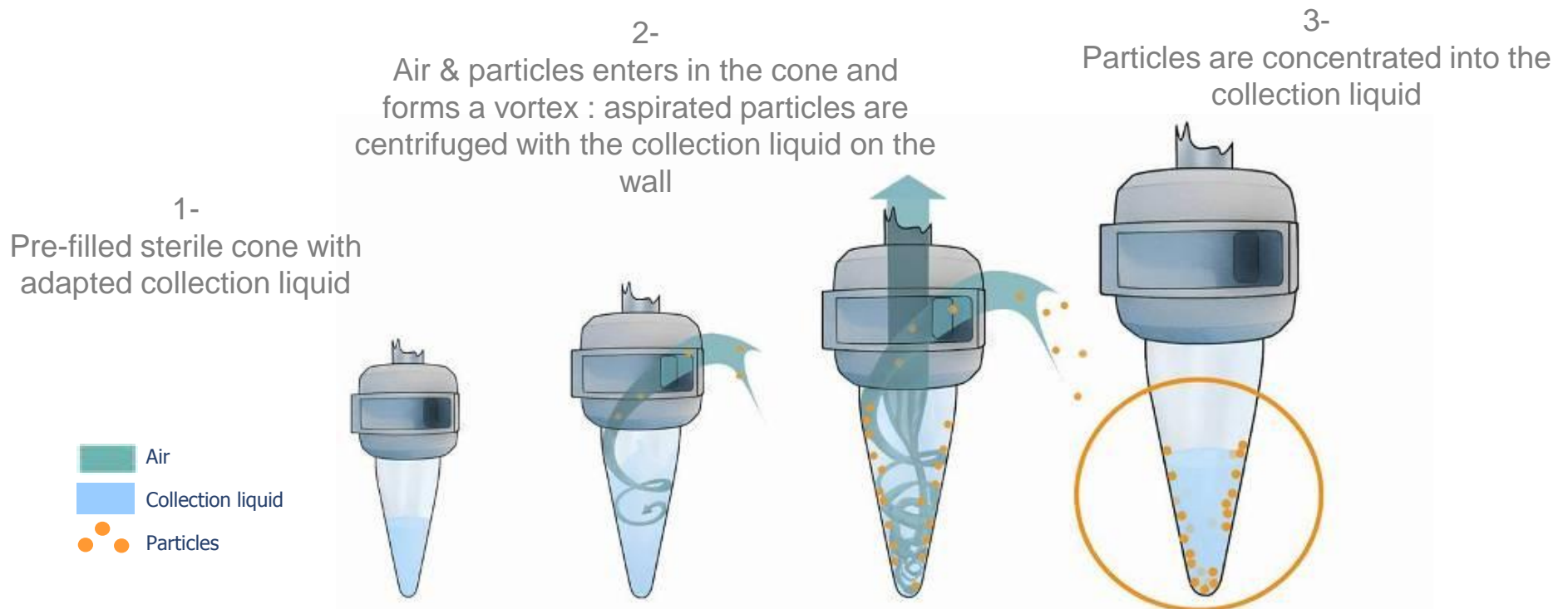


▷ Limitation of traditional technologies

- Information only on cultivable flora (what can grow on the nutritive agar)
- Long time to results (incubation step for growth) = several days (2 to 14)
- Limited volume of air collected (1m^3)
- Short collection time (10 min)
- Air flow rate limited (to avoid the stress of the microorganisms) (~100 L/min)
- Saturation of the collection media in case of charged environments

ALTERNATIVE SOLUTION: CORIOLIS® TECHNOLOGY

- ▷ Patented cyclonic technology concentrating particles from 0.5 to 20µm into a sterile liquid collection media
- ▷ Captures and concentrates all airborne particles (bacteria, fungi, spores, viruses, pollens, allergens, endotoxins...)



WET CYCLONE COLLECTION ADVANTAGES

- ✓ High flow rate
 - Representative air sampling (10 lpm vs 300 lpm)

- ✓ Liquid sample output (10 mL)
 - No saturation of the liquid sample
 - Compatible with **qPCR**
 - Compatible with **ELISA**
 - Compatible with **Titration**

- ✓ Long time monitoring (up to 6 hours)
 - Concentration of the target in the collection liquid (generally sampling of 30 min)
 - Unpredictable pollution event (for area monitoring)

The Coriolis® micro gives access to more information than traditional methods

CORIOLIS[®] MICRO PRODUCT

▷ Designed for clean rooms, hospital and indoor air control

- High air flow rate: 100 to 300 L/min
- Light: 3 kg
- Collect viable, non cultivable and total flora, pollens, viruses...
- Easy decontamination (single use consumable, H2O2 decontamination)
- Battery operated
- Long time monitoring option (up to 6 hours)



www.coriolis-air sampler.com





▷ Topic:

- ALBEDO Project 2012-2015
- Evaluation of sampling device with field biological analysis incorporated
- Target: aspergillus fumigatus

▷ Sampling site

- Outside composition site

▷ Analysis:

- magnetic beads magnisense

▷ Conclusion

- Translating manual to automatic analysis requires development
- Define the pathogen target

CASE STUDY: CHARACTERIZATION OF BIOLOGICAL FILTER EFFICIENCY



Contact : Vincent Moulès



▷ Topic:

- Efficiency evaluation of indoor air decontamination system
- Nebulization chamber in BSL3 laboratory to simulate different in situ conditions

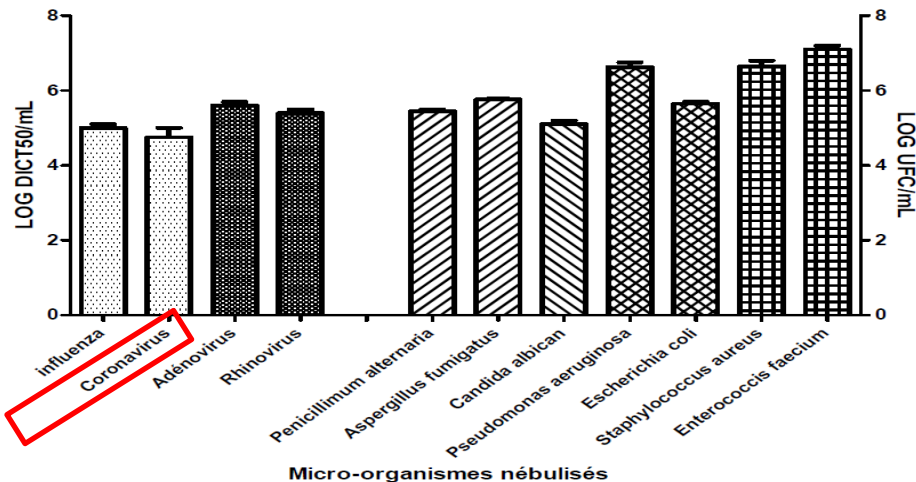
▷ Sampling site

- Air chamber : Suction of total volume of chamber with the Coriolis® micro with different conditions according to nature of micro-organisms (collection media and time/flow rate)

▷ Analysis: Titration, culture, RT-qPCR etc.

▷ Conclusion

- Collection of various pathogens
- Enable efficiency results of MERS
- Quantify respiratory viruses in air samples with specific RT-qPCR molecular procedures



CASE STUDY: GLUTEN IN THE AIR IN A FOOD PROCESSING

Sampling parameters	[Gluten] (mg of gluten/kg of liquid collection)	[Gluten] (mg of gluten/m ³ of air)	Results
Control (distilled water)	<5 ppm	-	OK
A1-300L/min-20 min Machines stopped	<5 ppm	< 0,075 mg de gluten/6000L of air < 12,5 µg de gluten/m ³ of air	Low gluten contamination
A1-300L/min-10 min Machines stopped	<5 ppm	< 0,075 mg de gluten/3000L of air < 25 µg de gluten/m ³ d'air	
A1-300L/min-20 min Production time	<5 ppm	< 0,075 mg de gluten/6000L of air < 12,5 µg de gluten/m ³ of air	Low gluten contamination
A1-300L/min-10 min Production time	8 ppm	0,12 mg de gluten/3000L of air 40 µg de gluten/m ³ of air	Increase of the level of gluten during the production

Sampling parameters	[Gluten] (mg of gluten/kg of liquid collection)	[Gluten] (mg of gluten/m ³ of air)	Results
A2-300L/min-20 min Production time	35 ppm	0,525 mg de gluten/6000L of air 87,5 µg de gluten/m ³ of air	Detection of gluten at a high concentration -> the production of gluten-free products close to this area is not possible
A2-300L/min-10 min Production time	54 ppm	0,81 mg de gluten/3000L of air 270 µg de gluten/m ³ of air	
	44 ppm	0,66 mg de gluten/3000L of air 220 µg de gluten/m ³ of air	
A2-200L/min-5 min Production time	18 ppm	270 µg de gluten/m ³ d'air	

▷ Topic:

- Threat of contamination of gluten-free food by air (FDA regulation: 20ppm in food)

▷ Sampling sites

- A1: This area has been emptied out of all products containing gluten
- A2: Raw materials containing gluten are continuously used in this area.

▷ Analysis: ELISA (R5-Mendez)

▷ Conclusion

- Thanks to its high flow rate, Coriolis Micro is a valuable tool for the collection of gluten even in a low contaminated environment.

CASE STUDY: STUDIES ABOUT THE DETECTION AND QUANTIFICATION OF BIOAEROSOLS WITHIN AND IN THE VICINITY OF PIG AND POULTRY BARNS



Contact : Jochen Schulz



Ahmed, Schulz, Hartung (2013): Air samplings in a *Campylobacter jejuni* positive laying hen flock. *Annals of Agricultural and Environmental Medicine* 20: 16-20

Topic:

- The air of laying hen houses can contain high concentrations of airborne bacteria including zoonotic pathogens.
- The numbers of these bacteria can be influenced by the efficiency of the chosen sampling method.
- sampling aerobic mesophilic bacteria in a *Campylobacter jejuni* (airborne *C. jejuni* is suggested to be a potential health risk when it is swallowed, Wilson 2004)

▷ Sampling methods

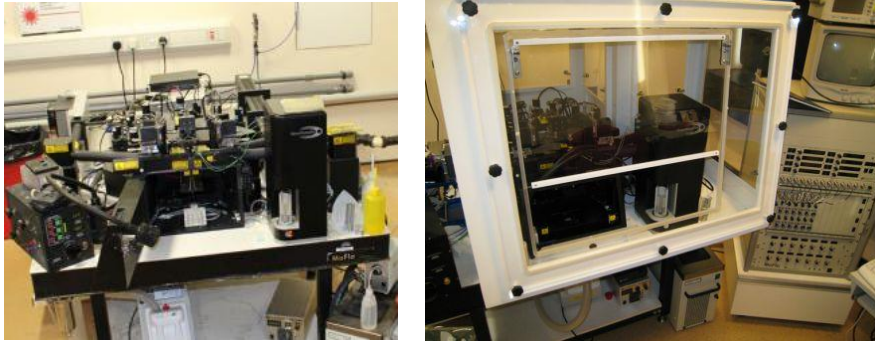
- AGI-30 Impinger and the Coriolis@µ

▷ Conclusions

- The Coriolis µ Air Sampler showed higher bacteria concentrations than the AGI-30 impinger. The differences were highly significant



Contact : Jan Baier



Cytometers without and with Bio Safety Cabinet (BSC)

▷ Topic:

- Operator protection from aerosols, potentially carrying bio hazardous sample constituents (e.g. HIV), generated during cell sorting experiments
- In failure mode, high amounts of aerosol are produced because of (partial) obstruction of the “nozzle” of the cytometers

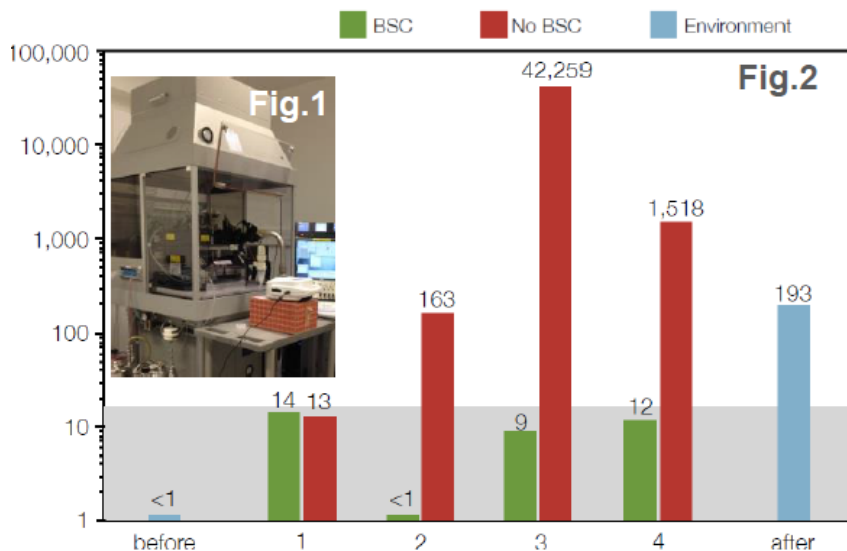
▷ Sampling site: in front of the cytometer

- Exp. 1: no operation (= background)
- Exp. 2: normal operation
- Exp. 3: failure mode
- Exp. 4: 30 seconds after failure mode
- Exp. 5: 2 meters away from cytometer

▷ Analysis: fluorescent micro beads.

▷ Conclusions

- The results of this preliminary study clearly indicate the efficiency of providing operator safety by running jet-in-air flow cytometric cell sorters inside a BSC.



Bead concentration (beads per m³) in “operator safety zone” under different operation conditions,



BERTIN TECHNOLOGIES

Romain Verollet
Head of Product Management Life Science

E.MAIL

Romain.verollet@bertin.fr

PHONE

+33 1 39 30 61 18

HEAD OFFICE

Parc d'Activités du Pas du Lac
10 bis avenue Ampère
78180 Montigny-le-Bretonneux
FRANCE

www.bertin.fr