

UV-C AIR DISINFECTION SCIENTIFICALLY VERIFIED

- UV-C AIR DISINFECTION **WHITE PAPER** BY DR. THIERRY K.S. JANSSENS
- **LAB REPORT** BY PROF. WACLAW DABROWSKI (INSTITUTE OF AGRICULTURE AND FOOD BIOTECHNOLOGY)
- **COMMENTS ON LAB REPORT** BY DR. THIERRY K.S. JANSSENS

UV-C Air Disinfection verified by Prof. Waclaw Dabrowski (Institute of Agriculture and Food Biotechnology)

The sudden Covid-19 pandemic has woken up the globalized society and economy about its susceptibility to emerging pathogens. The novel coronavirus that is spreading over the human population exhibited unexpected properties that put a challenge on health care systems worldwide. Individuals that are present as dense crowds are especially prone to the spread of respiratory infections, as has been shown in the last year how venues of recreation, sports and celebration were reported as hotspots for the spread of the SARS-CoV-2 virus. Therefore, all public indoor places with regular presence or turnover of crowds are potentially high-risk for transmission of the coronavirus. Consequently, the economic fallout of the Covid-19 pandemic most hit the arts, entertainment and recreation sector and accommodation and food services sector (McKinsey, 2020), as they are under strict guidelines from the authorities, suffer from consumer FUD and their recovery period to pre-pandemic levels would take up to five years and the high proportion of small businesses involved.

Measures such as hand washing, face masks, surface disinfection, social distancing, crowd management, testing and contact tracing have been instrumental to slowly open up the society after lock down measures during the first global wave of SARS-CoV-2 infections. Still, pre-pandemic numbers in crowds remain a challenge for e.g. recreational, business or religious events, educational organizations or the thriving of food services in restaurants and bars.

The scientific debate on the transmission of SARS-CoV-2 in the air is still going on (Lewis, 2020), and has caused public agitation about the formulation of guidelines, such as the distance to be kept between individuals and the contrast of the indoor versus outdoor risks. The distinction between droplets and aerosols in the spread of the virus remains in the fact that it could be spread respectively by coughing/sneezing only or also by talking or singing. There has been ample evidence of the detection of SARS-CoV-2 genetic material in aerosols, but the infectiousness is also highly determined by the magnitude of the viral load and duration of the exposure. Also, influenza virus (which is comparable in structural sensitivity to the novel coronavirus) has been shown to be most infective in aerosols at lower humidity levels (Noti *et al.*, 2013), such as the situation in heated indoor air. Following the concurrent growing consensus on the significant role of aerosols in the airborne transmission and the subsequent risk of super-spreading events in large crowds (The Lancet COVID-19 Commission, 2020), ventilation and disinfection of recirculated indoor air have become indispensable measures to assure a safer indoor ambient with minimized risk for infection by SARS-CoV-2 and other respiratory pathogens. In addition to the behavioral and maintenance guidelines, the proper treatment of indoor air will become an additional layer of biological safety in Covid-19 safe venues, as seen worldwide in initiatives to formulate guidelines for safety at venues; for example by REHVA (Federation of European Heating, Ventilation and Air Conditioning Associations (Kurnitski *et al.*, 2020)).

Ultraviolet germicidal inactivation (UVGCI) is an effective technique to eliminate micro-organisms. This high energy and low wavelength irradiation destroys the genetic material without any remaining residues and with limited chance of emergence of resistance compared to the use of biocides/antimicrobials. It can be applied on suitable surfaces and fomites in the absence of non-target organisms, such as humans, but has also shown its value in the topical treatment of infections, disinfection of culture water in aquaculture and indoor air in ventilation systems. It is a non-selective mode of eliminating micro-organisms, but not all cells and viruses are equally susceptible (Malayeri *et al.*, 2016). Cell walls, spore structures and dense viral capsids can require higher dose of UV-C radiation to reach the genetic material. Viruses - and especially the groups with single stranded genetic material - are more susceptible to UVGCI than cellular micro-organisms (Tseng *et al.*, 2005). Therefore the SARS-CoV-2 is quickly eliminated by this treatment, even at high densities (Heilingloh *et al.*, 2020).

The Luxibel type B air 2x55W TUV disinfection lamp makes use of this same principle to treat indoor air-and because of its aerodynamic design and vertical structure it samples in a laminar mode of flow in the middle of the air column above the crowds. As such, the MADS (Mid-Air Disinfection System)

avoids the turbulent distribution of aerosols and minimizes the spread of airborne pathogens by aerosols.

In a study executed by the IBPR (Prof. Waław Dąbrowski Institute of Agriculture and Food Biotechnology) in Poland, our apparatus exhibits an almost immediate effect on the microbial air quality, by a reduction of 71 and 49 % of respectively total microbial and fungal count after 2 hours of operation in a setup of 1,3 fold treatment of indoor air volume per hour (See Figure 1). After 20 hours the reduction in viable count were reduced with 98% and 93% of the initial burden respectively.

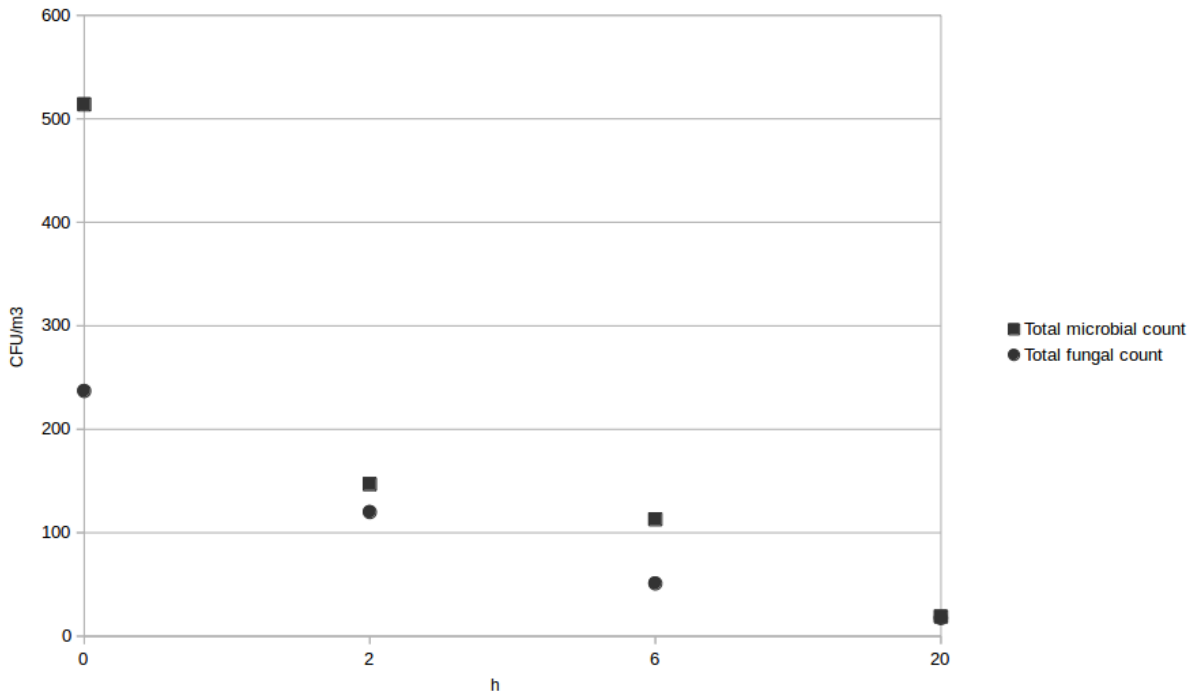


Figure 1: Total microbial and fungal viable counts upon sampling of 1 m³ of indoor air (MAS-100 ECO™ air sampler, MBV), after growth on media.

Upon 20 hours of treatment of indoor air in a closed room, the burden of viable of microbial cells in the sampled air was reduced to levels that are observed in operation theaters (Shaw *et al.*, 2018) or in occasional positive samples in production clean rooms (Tršan *et al.*, 2019).

The positioning of the air disinfection units along the principles of the patent-pending MADS principle can reduce the most dangerous aerosols in a more efficient way versus classic wall mounted systems, or clean room techniques.

In addition to the recommend measures to combat the spread of Covid1, as by the measures and guidelines issued by authorities, integrating the Luxibel type B air 2X55W UV disinfection unit into a well-dimensioned ventilation system will make any indoor public space or venue prepared to reduce the risk of infection by SARS-CoV-2 or any other airborne pathogens, both during the concurrent crisis and post-pandemic. This will results in a more hassle free use of public indoor spaces and a quicker than predicted economic recovery for the collapsed economic sectors.

Dr. Thierry K.S. Janssens
Molecular Biologist

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European Translation Agency – Certified Translation Department
PL - 00-336 Warsaw, Kopernika 30 Street, fax: 022 244 22 07
tel: +48 693 333 333 e-mail: info@e-ling.eu
Website: www.e-ling.eu – 24h service



Barbara Jarczyńska
Certified Translator of English

CERTIFIED TRANSLATION FROM POLISH INTO ENGLISH

INSTITUTE OF BIOTECHNOLOGY OF FOOD AND AGRICULTURAL INDUSTRY
named after prof Waław Dąbrowski

FOOD QUALITY DEPARTMENT

92-202 Lodz, ul. Marszałka J. Piłsudskiego no 84, tel (+48 42) 636 92 11, (+48 42) 636 55
72, (+48 42) 674 64 14 ext. 320, fax (+48 42) 674 81 24
zj@ibprs.pl
NIP [tax identification number]: 525-000-82-64 REGON number: 000053835-00026

(longitudinal seal)

Institute of Biotechnology of Food and Agricultural Industry
named after Professor Waław Dąbrowski
02- 532 Warsaw, ul. Rakowiecka 36

TAX ID: 525-000-82-81 Company's ID: 000053835-00026

FOOD QUALITY DEPARTMENT

92- 202 Lodz, ul. Marszałka J. Piłsudskiego no 84
Phone: (42) 674 64 14, (42) 636 92 11, tel./fax. (42) 674 81 24

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Lódź, 26-08-2020

Test report: No. K/313/01/2020

Research object: Flow disinfection luminaire UV type B Air 2 x 55W

Client: AED Distribution

Luxibel

Bedrijvenpark de Veert 13/004
2830 Willebroek, Belgium

The test object was collected and delivered by the client: 13-08-2020

Research started on: 19-08-2020



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 PL - 00-336 Warsaw, Kopernika 30 Street, fax: 022 244 22 07
 tel: +48 693 333 333 e-mail: info@e-ling.eu
 Website: www.e-ling.eu – 24h service



The research has been completed on: 25-08-2020

Type of marking / feature	Analytical method	Results	
Microbiological parameters			
Testing of the level of air pollution during the operation of the lamp in a room of 30 m ² and 2.9 m height	Own methodology using the MAS- 100 ECO™ microbial air sampler - MAS-100 Eco™ instruction manual	*[cfu/l m ³]	Reduction of microbes
- total number of microbes at time 0		514	-
- total number of microbes after 2 hours		147	R _{2h} = 71,40%
- total number of microbes after 6 hours		113	R _{6h} = 78,02 %
- total number of microbes after 20 hours		19	R _{20h} = 96,31%
- number of molds and yeasts at time 0		237	-
- number of molds and yeasts after 2 hours		120	R _{2h} = 49,37%
- number of molds and yeasts after 6 hours		51	R _{6h} = 78,48 %
- number of molds and yeasts after 20 hours		17.5	R _{20h} = 92,62 %

* The results are the average number of microbes from two measurements
 Authorised by:

[illegible handwritten signature]
 (longitudinal seal)
 LABORATORY OF MICROBIOLOGY
 Dr. Beata Paziak-Domańska
 Adjunct

Approved by:
 (longitudinal seal)
 [illegible handwritten signature]
 HEAD OF FOOD QUALITY DEPARTMENT



European Translation Agency – Certified Translation Department
PL - 00-336 Warsaw, Kopernika 30 Street, fax: 022 244 22 07
tel: +48 693 333 333 e-mail: info@e-ling.eu
Website: www.e-ling.eu – 24h service



Dr. Beasta Bartodziejska

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INSTITUTE OF BIOTECHNOLOGY OF FOOD AND AGRICULTURAL INDUSTRY
named after prof Waław Dąbrowski

FOOD QUALITY DEPARTMENT

92-202 Lodz, ul. Marszałka J. Piłsudskiego no 84, tel (+48 42) 636 92 11, (+48 42) 636 55
72, (+48 42) 674 64 14 ext. 320, fax (+48 42) 674 81 24
zj@ibprs.pl

NIP [tax identification number]: 525-000-82-64 REGON number: 000053835-00026

(longitudinal seal)

Institute of Biotechnology of Food and Agricultural Industry
named after Professor Waław Dąbrowski

02- 532 Warsaw, ul. Rakowiecka 36

TAX ID: 525-000-82-81 Company's ID: 000053835-00026

FOOD QUALITY DEPARTMENT

92- 202 Lodz, ul. Marszałka J. Piłsudskiego no 84

Phone: (42) 674 64 14, (42) 636 92 11, tel./fax. (42) 674 81 24

**Assessment of the effectiveness of air disinfection using the B Air 2 x 55W UV flow
disinfection luminaire**

Scope and purpose of the test

The aim of the test was to determine the effectiveness of air disinfection by means of the UV disinfection flow-through luminaire type B Air 2 x 55W (Test report K/313/01/2020) on the basis of testing the total number of microbes and the number of moulds and yeasts by the aspiration method after 2, 6 and 20 hours of lamp operation in a room with an area of 30 m² and height of 2.90 m.



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tel: +48 693 333 333 e-mail: info@e-ling.eu
Website: www.e-ling.eu – 24h service



Method of testing

The test was carried out in accordance with our own methodology and the MAS-100 ECO™ instruction manual (Microbiological Air Sampler) in a room with an area of 30 m². Before switching on the lamp, the total number of microbes and the number of moulds and yeasts in the air filling the room was tested. The degree of air pollution was measured at a distance of approx. 2 meters from the lamp after 2, 6 and 20 hours of operation. The test was carried out with the aspiration method using the MAS-100 ECO™ microbial air sampler, which draws 1000 litres of air through a perforated plate. The air stream containing the particles was directed to the surface of PCA or YGC agar in a standard petri dish. Upon completion of the air sampling cycle, the panes were incubated at 30°C for 72 hours or at 25°C for 5 days, then the colonies grown were counted and the number of microbes in 1 m² of the air was determined, taking into account the correction of the Feller statistical conversion table.

[illegible handwritten signature]
(longitudinal seal)
LABORATORY OF MICROBIOLOGY
Dr. Beata Paziak-Domańska
Adjunct

I, Barbara Jurczyńska, certified translator of English, entered in the list of certified translators under TP/2061/05 number, conducted by the Minister of Justice, hereby certify that this English text was translated from the Polish language.

Repertory number: 1924/2020
Date: 07/09/2020





INSTYTUT BIOTECHNOLOGII
PRZEMYSŁU ROLNO-SPOŻYWCZEGO
im. prof. Wacława Dąbrowskiego

ZAKŁAD JAKOŚCI ŻYWNOSCI

92-202 Łódź, Al. Marszałka J. Piłsudskiego 84
tel. (+48 42) 636 92 11, (+48 42) 636 55 72, (+48 42) 674 64 14 wew. 320, fax (+48 42) 674 81 24
zj@ibprs.pl
NIP: 525-000-82-64 REGON: 000053835-00026

Instytut Biotechnologii Przemysłu Rolno-Spożywczego
im. prof. Wacława Dąbrowskiego
02 - 532 Warszawa, ul. Rakowiecka 36
NIP 525-000-82-64 REGON 000053835
ZAKŁAD JAKOŚCI ŻYWNOSCI
92 - 202 Łódź, Al. Marszałka J. Piłsudskiego 84
tel. (42) 674 64 14, (42) 636 92 11, tel./fax. (42) 674 81 24

1/1



Łódź, 26-08-2020

Sprawozdanie z badań Nr K/313/01/2020

Obiekt badania: Przepływowa oprawa dezynfekcyjna UV typu B Air 2 x 55W

Klient: AED Distribution
Luxibel
Bedrijvenpark de Veert 13/004
2830 Willebroek, Belgium

Obiekt do badania pobral i dostarczył Klient: 13-08-2020
Badania rozpoczęto: 19-08-2020
Badania zakończono: 25-08-2020

Rodzaj oznaczenia / cecha	Metoda analityczna	Wyniki	
Parametry mikrobiologiczne			
Badanie poziomu zanieczyszczenia powietrza podczas działania lampy w pomieszczeniu o powierzchni 30 m ² i wysokości 2,9 m	Metodyka własna przy użyciu mikrobiologicznego próbnika powietrza MAS-100 ECO™ Instrukcja MAS-100 Eco™	*[jtk/1 m ³]	Redukcja drobnoustrojów
- ogólna liczba drobnoustrojów w czasie 0		514	-
- ogólna liczba drobnoustrojów po 2 godz.		147	R _{2h} = 71,40%
- ogólna liczba drobnoustrojów po 6 godz.		113	R _{6h} = 78,02 %
- ogólna liczba drobnoustrojów po 20 godz.		19	R _{20h} = 96,31%
- liczba pleśni i drożdży w czasie 0		237	-
- liczba pleśni i drożdży po 2 godz.		120	R _{2h} = 49,37%
- liczba pleśni i drożdży po 6 godz.		51	R _{6h} = 78,48 %
- liczba pleśni i drożdży po 20 godz.		17,5	R _{20h} = 92,62 %

*Wyniki stanowią średnią liczbę drobnoustrojów z dwóch pomiarów

Autoryzował:
B. Paziak-Domańska
PRACOWNIA MIKROBIOLOGII
dr Beata Paziak-Domańska
Adiunkt



Zatwierdził:

KIEROWNIK ZAKŁADU
JAKOŚCI ŻYWNOSCI
Beata Bartodziejska
dr Beata Bartodziejska

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INSTYTUT BIOTECHNOLOGII
PRZEMYSŁU ROLNO-SPOŻYWCZEGO
im. prof. Wacława Dąbrowskiego



ZAKŁAD JAKOŚCI ŻYWNOSCI

92-202 Łódź, Al. Marszałka J. Piłsudskiego 84

tel. (+48 42) 636 92 11, (+48 42) 636 55 72, (+48 42) 674 64 14 wew. 320, fax (+48 42) 674 81 24

Instytut Biotechnologii Przemysłu Rolno-Spożywczego
im. prof. Wacława Dąbrowskiego
02 - 532 Warszawa, ul. Rakowiecka 36

zj@ibprs.pl

NIP: 525-000-82-64 REGON: 000053835-00026

NIP 525-000-82-64 REGON 000053835
ZAKŁAD JAKOŚCI ŻYWNOSCI
ul. (42) 674 64 14, (42) 636 55 72, (+48 42) 674 64 14

Ocena skuteczności dezynfekcji powietrza przy użyciu Przepływowej oprawy dezynfekcyjnej

UV typu B Air 2 x 55W

Cel i zakres badania

Celem badania było określenie skuteczności dezynfekcji powietrza za pomocą **Przepływowej oprawy dezynfekcyjnej UV typu B Air 2 x 55W** (Sprawozdanie z badań K/313/01/2020) na podstawie badania ogólnej liczby drobnoustrojów oraz liczby pleśni i drożdży metodą aspiracyjną po 2, 6 i 20 godzinach pracy lampy w pomieszczeniu o powierzchni 30 m² i wysokości 2,90 m.

Sposób wykonania badania

Badania przeprowadzono zgodnie z własną metodyką oraz instrukcją MAS-100 ECO™ (Mikrobiologiczny Próbnik Powietrza) w pomieszczeniu o powierzchni 30 m². Przed włączeniem lampy wykonano badanie ogólnej liczby drobnoustrojów oraz liczby pleśni i drożdży w powietrzu wypełniającym pomieszczenie. Pomiaru stopnia zanieczyszczenia powietrza dokonywano w odległości ok. 2 metrów od lampy po 2, 6 i 20 godzinach pracy urządzenia. Badania wykonano metodą aspiracyjną przy użyciu mikrobiologicznego próbника powietrza MAS-100 ECO™, pobierającego 1000 litrów powietrza przez perforowaną płytkę. Strumień powietrza zawierający cząstki, kierowany był na powierzchnię agaru PCA lub YGC w standardowej szalce Petriego. Po ukończeniu cyklu pobierania próbki powietrza, szalki inkubowano w temperaturze 30°C przez 72h lub w temperaturze 25°C przez 5 dni, a następnie zliczano wyrosłe kolonie i określano liczbę drobnoustrojów w 1 m³ powietrza, uwzględniając korektę statystycznej tablicy przeliczeniowej Feller'a.

B. Paziak-Domańska

PRACOWNIA MIKROBIOLOGII
dr Beata Paziak-Domańska
Adiunkt



Comments on the test report (K/313/01/2020)

Luxibel Type B Air V2 (2 x 55W UV-C air disinfection lamp) by the IBPR (Prof. Waław Dąbrowski - Institute of Agriculture and Food Biotechnology)

In this report the effect of the UV-C disinfection lamp on the viability of cultivable microorganisms in indoor ambient air has been described. In order to assess the effect on the environmental microorganisms present in the room, an active microbial sampling method was applied, which better approaches the number of inhaled micro-organisms, as opposed to passive sampling methods, such as settling of microbial cells on exposed culture plates.

The applied microbial air sampler, makes use of the Andersen sampling method, by directing the sampled air through pores in a perforated plate to solid culture media, i.e. sterile agar plates. By applying the maximum sampling volume of 1m³, the limit of detection was most optimal to detect a decrease in microbial burden.

The growth on the culture media after incubation is reported as counts of (colony forming) units, recalculated according to the statistical corrections for the design of the sampling device. These counts of colony upon growth on the two culture media PCA and YGC respectively indicate the total number of viable of aerobic and cultivable micro-organisms (including moulds and yeast) and the number of viable yeast/mould cells (excluding bacteria). In this method no distinction is made between pathogenic and harmless micro-organisms and the presence of viruses is not taken into account. (The detection of infectious viral burden from environmental sources is very laborious and semi-quantitative, and would require the application of nebulized viral pathogens.)

The performance of the Type Luxibel B Air V2 (2x 55W UV-C air disinfection lamp) after 20 hours in reducing the microbial load in indoor air is effective, as it reduces the values of actively sampled total microbial counts to values below the average reported value in operation rooms (Shaw *et al*, 2018) and it approaches contamination levels that are observed in occasional positive samples in clean room situations (Tršan *et al.*, 2019). The relative reduction in counts (92,6%) is lower for the fungi (yeasts and moulds), as compared to the total microbial count (98,1%), as the former are known to be more UV-C resistant given their thick cell walls. Airborne viruses, and especially those with a single stranded nucleic acid genome (like SARS-CoV-2), are more susceptible than fungi and endospore-forming bacteria (Tseng *et al.*, 2005) (which are included in the total microbial count as well). Therefore, it is expected that the relative reduction of airborne infectious viral particles will be higher than 98,1%.

In order to assess the energy of UV-C radiation transmitted to the airflow, and how it relates to the susceptibility of the different groups of airborne micro-organisms, additional measurements and technical information on the disinfection device are required.

The methodology used in this study is appropriate and straightforward. However, I have some minor comments and questions.

- Given the midair sampling design of the Type B Air V2 (2 x 55W UV-C air disinfection lamp), it would be informative to indicate how the sampling device was located according to the generated airflow.
- The manufacturer recommends an application time of 24 h, therefore it would have been informative to extend the duration of the experiment and observe the long term effect of the disinfection lamp, by adding additional time points.
- What was the relative humidity (RH) and temperature during the test? The water film on microbial cells can scatter the UV light and make the UV treatment less effective. By providing the RH the results in the performance of the disinfection lamp could be better assessed in other conditions.

- How many microbial air samplers were used or how were the agar plates replaced sterilely? Has there been an influx of ambient air from outside?
- Replicate values/variances are lacking.

References

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About Dr. Thierry K.S. Janssens

Thierry Janssens is an experienced scientist with expertise in molecular biology, microbial ecology, virology and bioinformatics. He graduated with a MSc in Biology and an MSc in Biotechnology from Ghent University. Subsequently he attained his PhD in Biology from VU University Amsterdam, where he studied evolutionary processes in the framework of toxicity of heavy metals and oxidative stress. At later stage in his career he has conducted genomics and bioinformatics research on diverse applied toxicology, microbiology and bio prospection projects, but he also developed bioassays for studying the effects of novel antimicrobial candidate compounds. In the last few years he has conducted research at the Dutch National Institute of Public Health and the Environment (RIVM) on the use of metagenomics for an improved surveillance of respiratory viral pathogens.

About Prof. Waclaw Dąbrowski - Institute of Agriculture and Food Biotechnology

Prof. Waclaw Dąbrowski Institute of Agricultural and Food Biotechnology works on research, development and application of new biotechnological and technical methods in the food industry. These applications cover several areas, like: technical microbiology, food microbiology, cell engineering, process engineering, chemistry and biochemistry, food technology and human diet. We are working also in agricultural food areas, like: yeast production, alcohol production, winery, vinegary, brewery, fruits and vegetables processing, cereals processing and storage, starch and potatoes processing, bakery industry, food concentrates, cooled and frozen food, sugar industry, meat and fat industry as well as production of microbiologically active confection (enzymatic, probiotic, initiation cultures etc.).

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