Training organization registered
n° 1175 54659 75

Accreditation COFRAC Tests n°1-0019.
Scope available on www.cofrac.fr

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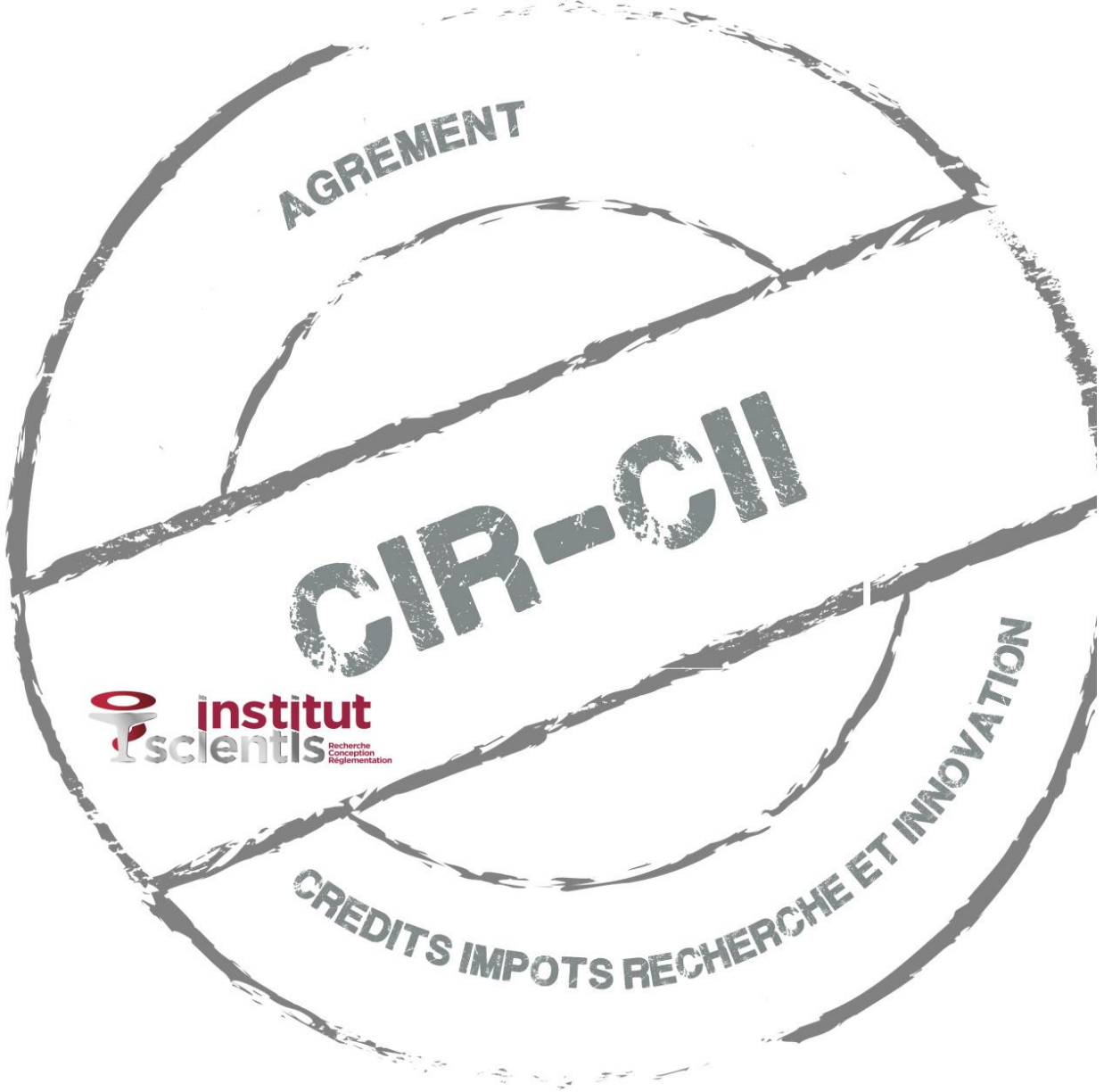
COSMETICS
BIOCIDES
DETERGENTS
DISINFECTANTS
MEDICAL DEVICES

TRAINING
MICROBIOLOGY
FORMULATION
REGULATIONS
SCIENTIFIC MARKETING

The addition of two companies skills for the evaluation of the safety of your products.



Authorized by the Ministry of National Education, Higher Education and Research
for your R&D work likely to give you the benefit of a Research Tax Credit (CIR) and Innovation (CII)

2019

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Safety assessment of the cosmetic products – *Toxicological expertise (European Registered Toxicologist)*

THE KEY POINTS

INDEPENDANCE

GLOBAL SCIENTIFIC EXPERTISE

SPECIFIC SUPPORT

INNOVATION

Registered training organization

Research tax credit agreement

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AGREMENT CIR-CII CREDITS IMPOTS RECHERCHE ET INNOVATION

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FORMULATION

A complete projects management

- Help with the marketing strategy / Setting up of yours development specifications
- Strict selection of the raw materials
- Development of INNOVATIVE and EXCLUSIVE formulas.
- Improvement of your existing formulas / preservative systems
- Anticipation of the related regulations / Toxicological pre-expertise
- Efficacy and safety tests / Microbiological screenings
- Stability test / Content-container compatibility / Flash point
- Scientific marketing / Writing of the communication media
- Redaction of the regulatory files
- Help with the industrial transfer

Research tax credit agreement and innovation

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REGULATIONS

Safety assessment

REGULATIONS

- Cosmetic products – European regulation
  PIF / safety assessment / Notification / Label check / PAO
- Cosmetic products – International
  Regulatory compliance of formulas
- Biocidal products – European regulation
- Material Safety Data Sheets
- Scientific and regulatory support

Safety assessment of the cosmetic products – *Europeen toxicologist registered ERT*

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Microbiological tests & scientific expertise

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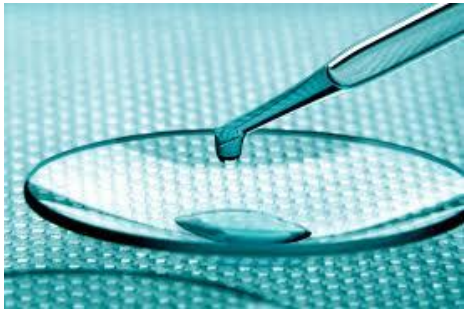
THE KEY POINTS

INDEPENDANCE

MICROBIOLOGICAL EXPERTISE

SPECIFIC SUPPORT

INNOVATION



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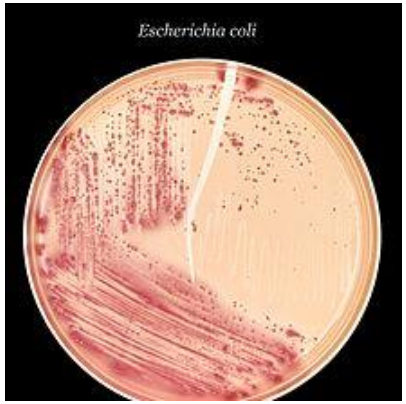
GENERAL SERVICES

BACTERIA-YEASTS-MOULDS-VIRUSES



- ✦ Evaluation of antimicrobial efficacy of biocidal products and medical devices (AFNOR, EN, ISO standards).
- ✦ Evaluation of the preservative system effectiveness of cosmetics-type risk products, preserved detergents, medical devices, drugs, (Preservative Efficacy Test).
- ✦ Microbiological assays of pharmaceutical products and medical devices.
- ✦ Determination of the antimicrobial activity of carriers such as plastics and textiles.
- ✦ Microbiological cleanliness tests of products not necessarily sterile.

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SPECIFIC SERVICES

VIROLOGY

EVALUATION OF THE VIRUCIDAL EFFICACY

- In compliance with the European (EN) standards methods.
- According to the method related to the plaque assay on cell cultures.
- Tested virus: Bovine enterovirus E (type 1), Bovine parvovirus (type 1), Bovine virus diarrhea disease (type 1) (Hepatitis C model), Canine adenovirus (type 1), Canine distemper virus (Morbillivirus), Canine parainfluenza virus, Canine parvovirus, Herpes simplex virus (type 1), Human Adenovirus, Influenza A virus, Murine norovirus, Simian rotavirus, Murine parvovirus, Pseudorabies virus (Hepatitis B model), Vaccinia virus.

BIOFILM

EVALUATION OF THE ANTIBIOFILM EFFICACY

- Method which has been developed and validated in a static mode.
- Installation of monobacterial biofilms on representative carriers to test the products for medical, food areas.
- Tests enabling to determine the activity of products with an antibiofim efficacy claim (mock-up of the real conditions).

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Associated experts – The addition of skills

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SCIENTIFIC EXPERTISE

DEFINITION OF THE TESTS STRATEGY

- To support the antimicrobial claims (bactericidal, fungicidal, virucidal…).
- For the placing on the market of biocidal products in compliance with the European Regulation n° 528/2012: definition of the tests strategy, to demonstrate the antimicrobial efficacy in compliance with the ECHA requirements: « Guidance on the BPR: Volume II Efficacy, Assessment + Evaluation »
- Definition of tests and related experimental conditions

PROOFREADING OF THE TEST REPORTS

The standardization evolves, the efficacy standards change for new versions.

- What do you have to do if your test was conducted many years ago?
- What version is required by the competent authorities for submitting a file to get a biocidal marketing authorization ?

A compliance upgrade with the last version of the standard is required if the changes can impact the results previously obtained.

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