

About **Ecolab**

A trusted partner at nearly three million customer locations, Ecolab is the global leader in water, hygiene and infection prevention solutions and services. With annual sales of €12 billion and more than 50,000 associates, Ecolab delivers comprehensive solutions, data-driven insights and personalised service to advance food safety, maintain clean and safe environments, optimise water and energy use, and improve operational efficiencies and sustainability for customers in the food, healthcare, hospitality, Life Sciences and industrial markets in more than 170 countries around the world.

Our **Assurance**

Ecolab is committed to providing exceptional service, total plant solutions and unsurpassed industry expertise to help you achieve your business goals.

Our breadth of resources and integrated business model ensure consistency across your operations.

It All Begins with **Regulatory**& Compliance in Cell and Gene Therapy

With new regulatory standards being introduced as the field advances, the Cell and Gene Therapy industry provides a unique challenge to navigate. For that reason, compliance very often relies on a case by case risk assessment.

From early process development to commercial manufacturing, GMP and regulatory guidance become increasingly stringent.

However, in this complex regulatory landscape, some key highlights and trends should be considered a priority:

Protocols that demonstrate process control and prevent product contamination are required to ensure patient safety during clinical trials.¹ Closed systems with high levels of protection are more frequently recommended for Grade A areas to achieve a closed system with consistent Sterility Assurance.²





Ecolab has extensive expertise in GMP compliance related to cleaning and disinfection applicable to Cell & Gene Therapy manufacturing.

We will guide you through appropriate contamination risk assessments to achieve compliance thereby avoiding delays and expensive process redesigns.

Contamination control strategies should ensure appropriate contamination risk management and proactive procedures to prevent ATMP contamination.³ Cleaning and disinfection of regulated areas shall be performed according to a written and validated procedure using broad spectrum disinfectants.³

Isolators in GMP areas shall be contained in Grade C/D cleanrooms.³

FUNDAMENTAL RESEARCH

Research & Development

CLINICAL TRIALS

Preclinical Trial

Phase One Phase Two

Phase

MANUFACTURING TRANSFER & SCALING UP

First Commercial Batch Routine Production

INCREASE IN GMP REQUIREMENTS

RESEARCH & DEVELOPMENT PHASE

- A controlled environment can decrease process variability
- Saves time and money by preventing process redesign at later phases
- ▲ Ease readiness for clinical trial

CLINICAL TRIAL PHASES

- Increased contamination control improves patient safety, reducing delays to your study
- ▲ Helps to get prepared for significant increase of GMP requirements for Phase 2 & 3
- Facilitates drug dossier approval before first commercial batch

MANUFACTURING PHASE

- Helps ensure products' quality assurance
- Offers a scalable solution with regulatory compliance at the forefront



ISOLATORS WITH
INTEGRATED
BIO-DECONTAMINATION
TECHNOLOGY

Bioquell Qube

The Bioquell Qube is the perfect fit to secure your most critical operations in both GLP and GMP environments.

CLEANING AND DISINFECTION SOLUTIONS

Automated Bio-Decontamination Options

Discover Bioquell Hydrogen Peroxide Vapour generators for repeatable, automated and validated biodecontamination of rooms, equipment and large zones.

Manual Cleaning and Disinfection

In addition, Ecolab provides sterile and non-sterile biocides suitable for manual cleaning and disinfection in both classified and non-classified areas. A complete range of hand hygiene products and dispensers are also available.







From GLP to GMP, Mitigate Your Contamination Risk

The development of novel Cell & Gene therapies such as CAR
T-Cells, mRNA or stem cell therapy requires extended Research and
Development activity before starting clinical trials. For that reason, the
Bioquell Qube has been designed to fit both GMP and GLP environments.

At this stage it is important to find a closed containment system that offers the least invasive option while retaining an aseptic space for you to work within. This would mean no building work, no HVAC connections needed, and no additional electrical work to use the device. Discover the perfect fit for your working environments with the Bioquell Qube



Be prepared for the next development phase with closed systems in your laboratory and GLP environments Help protect
your product
from contamination
in production rooms
and other GMP
environments

The Ideal Closed Environment Solution **The Bioquell Qube**

Ecolab's Bioquell Qube is an isolator with numerous configurations that uses Bioquell 35% Hydrogen Peroxide Vapour technology to create aseptic operating areas that can be maintained for long periods of time.

It is a key tool to reduce the risk of operator and product exposure to viral and other biological contaminations with regulatory compliance and GMP in mind.



Creates validated Grade A / ISO 5 environment

Environmental control and monitoring





21 CFR Part 11 compliance

The Bioquell Qube is the perfect fit for your Cell and Gene Therapy manufacturing operations







Bioquell Qube Benefits are Aligned with Cell & Gene Therapy Manufacturing Main Challenges

The Bioquell Qube is tried and trusted in the most demanding of environments, including high grade pharmaceutical facilities, and can quickly create a Grade A / ISO 5 environment in any area.

Integrated Bioquell bio-decontamination technology



SPEED

- Quick implementation and short lead time
- ▲ Fast decontamination cycle



CONSISTENCY

 Create a Grade A / ISO 5 area with validated and automated cycles for repeatable results



COST

- Cost effective option for scalable and effective work environments
- Operate a Grade A / ISO 5 environment while hosting in Grade C/D location



negative pressure

Operate in positive or

Supports GMP compliance



FLEXIBILITY & SCALABILITY

▲ Meet future capacity to grow as you do



No need of for HVAC connection



COMPLIANCE

- Bio-decontamination with Bioquell 35% Hydrogen Peroxide Vapour technology
- Meets requirements to help provide product consistency for regulatory bodies around the world
- ▲ 21 CFR Part 11 compliant audit trail
- Uses EU Biocidal Product Regulation (BPR) compliant sporicidal disinfectant

The Bioquell Qube provides a controlled and secured environment for the most critical operations in Cell & Gene Therapy like:

- ▲ Cell lines purification
- Genetic engineering
- Cellular culture and viral propagation
- ▲ Aseptic filling
- Sterility testing
- Media handling and preparation





ROOM AND ZONE BIO-DECONTAMINATION

Bioquell ProteQ

A mobile room to room system for small and large open spaces. Link multiple systems for even larger areas.



EQUIPMENT & MULTIPURPOSE BIO-DECONTAMINATION

Bioquell L-4

A versatile system that can bio-decontaminate existing isolators, biosafety cabinets, material airlocks (MALs), pass-throughs and, with optional equipment, small rooms.





PASS-THROUGHS, MALS AND SMALL ROOMS

Bioquell SeQure

A wall-mounted biodecontamination offering is ideal for material transfer areas, providing validated, trusted and repeatable results.



CORRECTIVE ACTION AND EMERGENCY RESPONSE:

Keep your laboratories or GMP area under control

Whether environmental level thresholds have been reached or a contamination event has occurred, Bioquell RBDS offers a 6-log sporicidal reduction on exposed surfaces, including difficult to clean areas.

PREVENTIVE ACTION:

Mitigate safety concern of viral vectors

In-process viral contamination, crosscontamination and operator safety are at risk due to exposure to viruses and biological agents used in Cell & Gene Therapy development. Bioquell Hydrogen Peroxide Vapour technology kills microbial life on every exposed surface.

Effective Manual Cleaning and Disinfection for Your Cell & Gene Therapy Facility

(lercide"

In addition to the Bioquell Hydrogen Peroxide Vapour generator, the Ecolab Klercide range of cleaners and disinfectants provides effective products for manual cleaning and disinfection in critical areas.



LARGE SURFACES IN CRITICAL AREAS

A wide range of both pre-impregnated wipes and ready-to-use liquids

- Alcohols
- ▲ Broad spectrum biocides & sporicides
- Cleaning and maintenance solutions
- With supporting equipment for efficient application



Klercide Low Residue Peroxide and Klercide **Neutral Detergent sprays** and wipes are specifically recommended to clean the Bioquell Qube.



SMALL SURFACES IN CONTROLLED ENVIRONMENTS

Alcohol wipes are a perfect fit for quickly cleaning work areas in laboratories and lowgrade cleanrooms.





HAND HYGIENE PRODUCT RANGE

Clean and sanitise hands in both GMP and GLP areas.



Our Complete Contamination Control Package For Cell & Gene Therapy

Annex 1 regulations require isolators to be contained in a classified cleanroom, with significant validation resources required to achieve compliance.

In the pharmaceutical industry, cleaning and disinfection validation can take 6-12 months to implement and cost up to €100k per facility. Ecolab has experience with global regulatory criteria that can significantly reduce the time and cost of validation.



Save **Time**Save **Money**Gain **Control**

References

1. Section V.a.2.E., "Process Validation And/Or Evaluation (3.2.S.2.5)," EMA Guidelines And/Or Evaluation (3.2.S.2.5)," EMA Guidelines

2. Annex 1

3. Guidelines on Good Manufacturing Practice specific to Advanced Therapy Medicinal Products

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