Cleaning and Disinfection – Environmental Monitoring

Your	Ohi	iectives:
1001	\sim	LCCLIVCS.

At the end of the lesson, you should be able to recognise the environmental monitoring equipment.

While microorganisms in the manufacturing environment are common, it is important to keep
them under control, especially during critical steps. In order to ensure that the environment
where occurs is clean, and that microorganisms in that
environment remain within acceptable , Biogen
regularly environmental and
utility
The 21 CFR states that drug manufacturing premises must be of a suitable size and design in
such a way as to facilitate cleaning, and proper operations.
Additionally, there should be control systems in place to prevent
during the course of aseptic operations, which includes a
system for monitoring environmental conditions.

Routine monitoring Routine monitoring provides that are used to detect the possibility of a trend towards of the environment or utility over time. Conclusions based on the collected data assist in defining new procedures, refining existing procedures, determining equipment maintenance intervals and initiating further monitoring Static monitoring is performed either after initial construction or modification or after sanitisation of a . It is intended to establish the baseline of the facility and focuses on the air system performance in the absence of routine activity within the room. Dynamic monitoring is performed during periods of routine use, when personnel and materials are present and when equipment is in . The results of dynamic monitoring are the basis for the establishment of alert and action During routine monitoring, the environment or the being

monitored may occasionally fail to conform to expected results. Depending on the

course), two different responses are defined:

and duration of the excursion (deviation from an expected

Alert level values are slightly outs	side the range of				
cleanliness. No	is required, but because this level may				
indicate potential problems, further	is needed.				
Action level values, which include a series of alert level values, pertain to levels that are					
outside of the adequate	of cleanliness. Action level				
values quite simply require immediate investigation &/ intervention.					
Monitoring zones					
Clean areas at Biogen are classified according to (1) the extent of product to environment, (2) the extent of personnel gowning, and (3)					
the degree of asepsis required.					
Biogen also uses a multiple zone approach, whereby more than one zone in an area is					
monitored. The	frequency, as well as particulate and microbial				
limits, differ in each zone. Separate	and procedures are				
specified for each operational area in the SOPs for monitoring specifics for a given area.					

Types of monitoring:

Utility monitoring

Utilities typically include a production/generation element and a distribcution system. Utilities at Biogen include, but are not limited to, the following:

- Purified water system
- WFI system

• Clean	system
Glycol system	
Heating,	and air-conditioning system (HVAC)
These systems are monitored on a rota	ting schedule delineated for each system.
Water monitoring	
According to the USP (U.S. Pharmacop	peia), organic molecules and other contaminants can be
introduced into the water	either from source water or from
& distri	bution system materials.
	purified water at Biogen includes collection of
on whic	ch TOC testing is performed.
Total organic carbon (TOC) is a testing	that measures the total
water. TOC sampling is a sensitive pro	TOC provides a useful way of measuring the purity of ocess. So, for example, for TOC sampling, one does not national materials with 70% alcohol, since alcohol is an organic
; any a	lcohol residue, therefore, would impart a positive TOC
reading.	
• •	to collect 40-millilitre samples. These vials must be e is not compromised. One must follow proper TOC
sampling and documentation	described in the SOPs.

Viable air sampling					
Biogen uses several different devices for air	. Technicians				
typically use a hand-held battery-powered	sampler. That device				
is cleaned with a 70% alcohol	prior to sampling.				
The operator places	plates into the retaining slots in the air				
sampler, aspirates the specified volume of air, typ plates are used for routine sampling. Sabouraud De and mold sampling.					
The plates are removed and sent to QC (Quality incubator. After incubation, QC counts the number of					
Results are compared with alert	to determine if any level of				
contamination has been exceeded.					
Please follow air sampling and documentation proce	dures described in the SOPs.				
Particulate air sampling					
Particulates consist of mobile, extraneous	, other than gas, that				
cannot be analysed chemically because of their heterogeneous .					

Air particulate samplers collect particulates on a filter, which can then be analysed for its constituents. Results are compared with alert levels so as to determine if any level has been exceeded.

One will have to follow air sampling and documentation procedures described in the SOPs.

Settling plates
Settling plates contain trypticase soy agar (TSA). They are typically used to monitor
in horizontal laminar airflow workstations, bio-safety
, or other room surfaces while
operations are being performed.
The room temperature plates are placed face down on the area surface. Maximum exposure time is 1 hour per plate. At the conclusion of the operation or the end time, the plate is covered
and sent to QC, where it is . After the incubation period, QC
determines whether the number of bacterial colonies is within appropriate limits.
One must always follow proper sampling and documentation procedures described in the SOPs.
Viable surface-monitoring
Replicate Organism Detection and Counting (RODAC) plates are used to monitor both personnel and surfaces.
TSA plates are used for routine surface and personnel monitoring.
Sabouraud Dextrose Agar (SDA) plates are used for yeast and mold sampling.
The agar in the room temperature plates is against the test site
surface and removed. The plate is and sent to QC where it is
incubated. After the incubation period, QC determines if the number of bacterial, mold, or
yeast is within appropriate limits.

One need follow the proper sampling and documentation procedures described in the SOPs.

Personnel monitoring

horizontal laminar airflow worl	•	• •	s performed in		
The	gloved fingertips ar	gloved fingertips are placed on the room-temperature TSA			
plate. The plate is covered and determines if the number of ba			ition period, QC		
One is required to follow pro SOPs.	oper sampling and docu	mentation procedures d	escribed in the		
video: <u>https://youtu.be/F9K8D</u>	AupOFg				
Toxic gas monitoring At Biogen, toxic gas monitoring	z typically occurs in:				
•	rooms, where		gases are		
used and where there i	s a potential for oxygen				
•	areas where cryo	ogenic liquids are stored,			
Areas where chillers a	are located and where .	there is a potential fo	r a refrigerant		

Personnel that work in areas where continuous gas monitors are installed must be trained, by their supervisor, to recognise the fault, to caution, warn, and alarm conditions associated with the gas-monitoring equipment.

Aufgabe Lückentext:

Folgende Wörter bitte in den Lückentext einfüllen. Jedes Wort kommt einmal vor. Bitte Gross- und Kleinbuchstaben beachten.

air, asphyxiant, contact, Cold, conditions, cabinets, contamination, compound, clean, covered, colonies, composition, data, deterioration, deficiency, Enclosed, exposure, intervention, incubated, limits, levels, levels, leak, manufacturing, monitoring, monitoring, maintenance, operation, observation, operator's, protocols, processes, performs, pressed, procedure, procedures, range, room, solution, substances, surfaces, scheduled, system, severity, steam, purification, samples, sampling, tolerable, utility, ventilation