Cleaning and Disinfection – Environmental Monitoring

Your Objectives:	
At the end of the lesson, you should be able to requipment.	ecognise the environmental monitoring
While microorganisms in the manufacturing environment them under control, especially during critical steps. I	•
where occurs is clo	ean, 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	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
and duration of the excursion (deviation from an expected
course), two different responses are defined:
Alert level values are slightly outside the range of
cleanliness. No six 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.

Monit	oring zo	nes										
Clean	areas	at	Biogen	are	classified	according	to	(1)	the	extent	of	produc
				to	o environr	nent, (2) the	exte	nt of	perso	nnel gov	vning	g, and (3
the de	gree of a	asep	sis requir	ed.								
Bioger	n also u	ses	a multip	le zor	ne approa	ch, whereby	, mor	e th	an on	e zone	in ar	n area is
monito	ored. Th	e				frequen	cy, as	well	as pa	rticulate	and	microbia
limits,	differ	in e	each zone	e. Sep	parate					and pr	oced	lures are
specifi	ed for ea	ach d	operation	al are	a in the SC	Ps for monit	oring	speci	fics fo	r a given	area	ı .
	<u>of moni</u> monito		<u>ıg:</u>									
		-	-		_	ration eleme following:	ent an	id a d	listribo	cution sy	stem	ı. Utilities
•	Purified	d wa	ter syster	n								
•	WFI sys	stem	I									
•	Clean					system						
•	Glycol	syste	em									
•	Heating	5,				and air-	condi	tionir	ıg syst	em (HVA	rC)	

These systems are monitored on a rotating schedule delineated for each system.

Water monitoring

According to the USP (U.S. Pl	harmacopeia), orgai	nic molecule	es and oth	ner contar	minants can	be
introduced into the water		€	either fro	m source	water or fro	om
	& distribution sys	stem materia	als.			
Routine monitoring of W	/FI and purified	water at	Biogen	includes	collection	of
	on which TOC tes	ting is perfo	rmed.			
Total organic carbon (TOC) is	a testing			that mea	sures the to	otal
organic material present in a water. TOC sampling is a ser sanitise either gloves or san	sitive process. So,	for example	, for TOC	sampling	, one does r	not
	; any alcohol res	idue, theref	ore, wou	ld impart	a positive T	OC
reading.						
Specially cleaned TOC vials carefully handled so that the			-			
sampling and documentation			describe	d in the SC	Ps.	
Viable air sampling						
Biogen uses several differer	nt devices for air				. Technicia	ans
typically use a hand-held l	pattery-powered				sampler. T	hat
device is cleaned with a 70% a	alcohol		рі	rior to sam	npling.	
The operator places		plates	into the	retaining	slots in the	air
sampler, aspirates the specif	ied volume of air, t	typically 320	litres. T	rypticase s	Soy Agar (TS	5A)

plates are used for routine sampling. Sabouraud Dextrose Agar (SDA) plates are used for yeast and mold sampling.
The plates are removed and sent to QC (Quality Control), where they are placed into an incubator. After incubation, QC counts the number of colonies that appear on the agar surface.
Results are compared with alert to determine if any level of
contamination has been exceeded.
Please follow air sampling and documentation procedures 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 pla	ates is	against the test site
surface and removed. The plate is		and sent to QC where it is
incubated. After the incubation peri	iod, QC determines if the nu	mber of bacterial, mold, or
yeast	is within appropriate limits.	

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

Personnel monitoring

Personnel monitoring is done at the asepsis of critical operations, which is performed in horizontal laminar airflow workstations or bio-safety cabinets.

The	gloved fingertips are placed on the room-temperature TSA

plate. The plate is covered and sent to QC where it is incubated. After the incubation period, QC determines if the number of bacterial colonies is within appropriate limits.

One is required to follow proper sampling and documentation procedures described in the SOPs.

video: https://youtu.be/F9K8DAupOFg

Toxic gas monitoring

At Biogen, toxic gas monitoring typically occurs in:	
• rooms, where	gases are
used and where there is a potential for oxygen	
areas where cryogenic liquids are	e stored,
Areas where chillers are located and where there is a pote	ential for a refrigerant
•	
Personnel that work in areas where continuous gas monitors are instable by their supervisor, to recognise the fault, to caution, warn, and alarm 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