



















Design, Construction and Cost Estimate of Cleanroom for Pharmaceutical Industry

m+w zander

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What is Cleanroom ? : Sources of Contamination

External Impuritities	Internal Impurities
Introduction of contaminated outside air or circulating air *	Staff
Staff	Process
impure process media or raw materials	Production equipment, machines, tools etc.
Inadequately cleaned materials, tools etc.	Unsuitable building materials, work materials
	Mechanical abrasion in the cleanroom
* poor filter quality, none airtight filter seal surfaces, I recirculation equipment and in the ducting system	eakage in the ducting system, abrasion in air

gn, Construction and Cost Estimate	e of Cleanroom for F	Pharmaceutical Indus
What is Cleanroom ? : Sources of	f Contamination	
Emission of Particles by People	Particle Emission per Minute and Person	Type of Activity
making various Movements without Cleanroom Clothing	100 000	Standing and sitting without moving
	500 000	Sitting with gentle movement of head, hand or lower arm
	1 000 000	Sitting with moderate body and foot movement
	2 500 000	Standing up with full body movement
	5 000 000	Slow walking - approx. 3,5 km/h
	7 500 000	Walking at about 6 km/h
Germ emission per minute (according to Botzenhart) 000 - 13 000 CFU depending on activity	10 000 000	Walking at about 9 km/h
	15 - 30 000 000	Gymnastics and sports

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ew Clean	room Tecl	hnoloav fa	or Pharmac	eutical Ind	ustrv : Cla	assification of clean
1 Fed Table 1: I	leral Stan Federal Sta	dard 209 Indard 209 (E class limits			
Class	Particles /	′ ft³				
	\geq 0.1 μm	$\geq~0.2~\mu m$	$\geq~0.3~\mu m$	≥ 0.5 µm	\geq 5.0 μm	
1	35	7.5	3	1	NA	
10	350	75	30	10	NA	
	NA	750	300	100	NA	
100				1 000	7	
100 1,000	NA	NA	NA	1,000	1	E STOR
100 1,000 10,000	NA NA	NA NA	NA NA	10,000	70	LEEP

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v Cleanroom	Technolo	gy for Ph	armaceut	ical Industr	y : Classif	ication of cle
2 ISO St	tandards					
l able 2: Selected	ISO 14644-1 a	airborne parti	culate cleanli	ness classes for	cleanrooms a	nd clean zones
ISO Classification number	Maximum of larger than	concentratio the conside	n limits (par red sizes sho	ticles/m ³ of air) own below	for particles	equal to and
	\geq 0.1 μ m	\geq 0.2 μ m	\geq 0.3µm	\geq 0.5 μ m	$\geq 1 \mu m$	≥ 5.0µm
ISO Class 1	10	2				
ISO Class 2	100	24	10	4		
ISO Class 3	1 000	237	102	35	8	
ISO Class 4	10 000	2 370	1 020	352	83	
ISO Class 5	100 000	23 700	10 200	3 520	832	29
ISO Class 6	1 000 000	237 000	102 000	35 200	8 320	293
ISO Class 7				352 000	83 200	2 930
ISO Class 8				3 520 000	832 000	29 300
	1			35 200 000	8 320 000	293.000

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Review Cleanroom Teo	chnoloa	y for Pharmaceutical Industry :
Guidelines and standards f	or the des	sign of cleanroom and facilities
A. Guidelines for d	lesign of	cleanroom and pharmaceutical
GMP (ISPE I	Good Manufa nternational S	cturing Practice Society of Pharmaceutical Engineering
The most used G	MP guides	for cleanroom
PIC : GMP and Gu FDA cGMP EU GGMP	lidelines	Valid in European countries outside the EU and Australia Valid for the United States Valid for the EU area



















Desi	gn of Cleanroom and Facilities	for Ph	narmaceutical Industry :
Clear	nroom and Facilities Design		
4. E	Basis of Design		
Key	Point		
• •	To outline the following concept to deve	elop the	project
	 The site (Area Size and Location) 	٠	Production requirements
	 Room layouts and operation 	٠	Process & plant utilities
• •	To establish budget to develop the proj	ect (with	a contingency ± 20 %)







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Design of Cleanroom and Facilities for Pharmace	eutical Industry :
Cleanroom and Facilities Design	
5. Conceptual Design	
Construction Material and Surface Finishes for C	leanroom
Material design consideration	
 Non-dust generation from surface 	
Easy to clean	
• Air tight	
Conductivity	
• Out-gas	
Chemical resistance	
• Fire safe	











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Des Clea	sign of Cleanroom and Facilities for Pharmaceutical Industry :
5.	Conceptual Design Example of conceptual design of HVAC system
	System description for HVAC system
	The HVAC system provided for medicinal product buildings shall be the centralized chilled water system served by water-cooled chillers.
	HVAC system for Clean Room Class 100,000 will conform to the ISO standard 14644 and Federal Standard No. 290E of U.S.A. Air Conditioning System shall be the conventional clean room system (Turbulent Flow).
	Air handling units shall be double skin module type.
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