

Coating systems for cleanroom and clean manufacturing areas- Floor, Wall, Ceiling

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Essential of Cleanrooms and clean manufacturing areas

Essential for a growing number of companies, in order that they can safeguard their process operations and the quality of their components.

The requirements of the various industrial sectors differ significantly. Cleanrooms and clean manufacturing areas are used primarily in the following industrial sectors:

Clean rooms

Biotechnology

Semiconductor industry

Photovoltaic

Microsystem technology

Food industry

Pharmaceutical industry

Aviation and space industry

Clean manufacturing areas

Automobile and supplier industries Machine engineering

Cleanliness areas

Cleanliness areas are set up to protect sensitive surfaces and items. The purpose of a cleanliness area is to maintain as far as possible the specified cleanliness quality of components, ancillary materials and assemblies during processing. The cleanliness level should not be reduced by the influence of environ- mental factors. Bringing contamination into a cleanliness area must be avoided. Any contamination, which is incurred there, is to be kept in check and then eliminated. The design, required measures and method of use for cleanliness areas are based on cleanliness requirements, which are related to the particular product. The critical particle sizes are generally between 5 µm and 1,000 µm.

The classification of the cleanliness levels in accordance with VDA 19

Part 2 is in 4 levels:

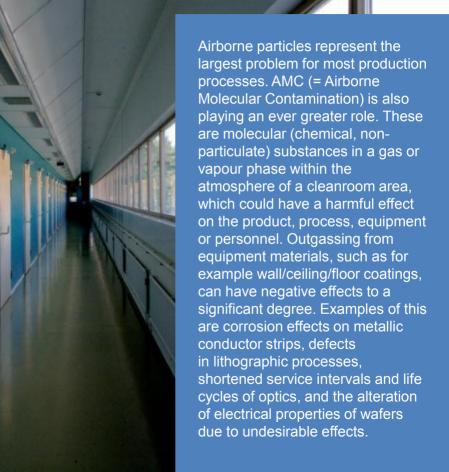
- Cleanliness level 0 (SaS0):
- Non-controlled area
- Cleanliness level 1 (SaS1):
- Clean zone
- Cleanliness level 2 (SaS2):
- Clean manufacturing area
- Cleanliness level 3 (SaS3):
- Cleanroom



Classifications bring safety - Regulations & standards

The requirements of different industrial sectors vary considerably. While a low level of outgassing from equipment is mandatory for the manufacture of semiconductors, this has mostly played no role as yet in the manufacture of pharmaceutical products. this has mostly played no role as yet in the manufacture of pharmaceutical products.

There are similarly differences in the particle cleanliness classes between DIN EN ISO 14644-1 and GMP (Good Manufacturing Practice) or cGMP, which applies to the manufacture of human and veterinary medical products. The particle cleanliness classes for air in accordance with DIN EN 14644-1 are divided into Classes 1 to 9, whereby the highest permitted particle count is the lowest in Class 1. In GMP the division is made into Classes A to D, whereby Class A corresponds approximately to ISO Class 5.



In accordance with DIN EN ISO 14644-1 and VDI 2083 Page 1: "A room, in which the concentration of airborne particles is controlled, and which is designed and used in such a way, that the number of particles, which have been brought into the room or which have arisen and been deposited in the room, is as small as possible, and where other parameters relevant to cleanliness, such as temperature, humidity and pressure are controlled as required." In contrast to the particle sizes in cleanliness areas, particle sizes of 0.1 µm to 5 µm are relevant here. In order to keep the share of more or less severely contaminated outside air as small as possible, normally rooms or storeys outside the actual cleanroom are used for air circulation. This means that it is also necessary there, that floors, walls and ceilings, which mostly consist of reinforced concrete, have a cleanroom-compatible surface. These measures have a significant influence on the lifespan of the filter elements for the cleanroom.

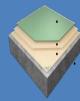
In addition to the quality of the supply air and method of introducing it as well as to the surfaces and personnel, the other critical influencing factor in the cleanliness of a clean-room is the equipment in the room. The equipment includes internal fixtures and fittings such as walls, doors, ceilings and floors. Significant factors in the cleanroom compatibility of equipment are:

- Cleanability
- Emission of airborne particles
- · Outgassing behaviour
- Electrostatic discharge properties
- Resistance to chemicals and disinfectants
- Metabolising potential and microbicidity
- Smooth and crack-free surface



Depending on the area of application, the following requirements are placed on wall & floor coating systems:

- Dissipative
- Crack-bridging
- Good abrasion resistance (low particle formation)
- Good mechanical resistance
- Good chemical resistance
- Low outgassing
- Smooth, easily cleanable surface
- Resistant to disinfectants
- · Biostatic or microbicidal





Tobslak Tobage Floor Coating System

Concentra

of chemical concentration (ACC) Concentration µg/m3

Classification of air purity on the basis

10⁻⁶ (1 000 000)

10⁻⁵ (0.000 01)

10-6 (0.000 001)

10⁻⁵ (100 000)

Concentration ng/m3

10⁹ (1 000 000 000)

-3

-5

ISO ACC

class

10-1 10^{-2}

tion

g/m3

10⁰

10-3 10-4 10-5 10-6 10^{-7} 10-8 10-9

10⁻¹⁰

10-11

10-12

10-4 (10 000) 10⁻³ (1 000) 10^{-2} (100) $10^{-1}(10)$ $10^{-0}(1)$ $10^{-1}(0.1)$ $10^{-2}(0.01)$ $10^{-3} (0.001)$ 10-4 (0.000 1) 108 (100 000 000) $10^7 (10\ 000\ 000)$ 106 (1 000 000) 10⁵ (100 000) 10⁴ (10 000)

 $10^3 (1\ 000)$ $10^2 (100)$ $10^{1}(10)$ $10^{0}(1)$ $10^{1}(0.1)$ $10^2 (0.01)$ $10^3 (0.001)$

-6 -7 -8 -9 -10 -11 -12

Sources of contamination in a clean room

In order to control contamination operators and those in charge of a cleanroom need to be cognizant of sources of contamination. These include:

- 1 .Facilities : Walls, Floors and ceiling, paints and coatings, spills and leaks.
- 2. People: Skin flakes and oil, cosmetics and perfume, spittle, clothing debris (lint, fibers etc.)hair
- 3. Tool-generated :Friction and wear particle, lubricants and emissions, vibrations, brooms, mops, and dusters .
- Fluids: Particulate floating in air, bacteria, organics and moisture, floor finishes or coatings, cleaning chemicals, plasticizers, (outgasses), water
- Product- generated :glass flakes, cleanroom debris, aluminum particles from vial caps.

Tobslak Cleanroom Coating systems -

Coating systems generally consist of the following work stages:

- Substrate preparation
- Primer
- Levelling layer
- Finishing coat

Depending on the area of application, the following advantages are with Tobslak cleanroom coating systems:

- Good abrasion resistance (low particle formation)
- Good mechanical resistance
- Good chemical resistance
- Low outgassing
- Smooth, easily cleanable surface
- Resistant to disinfectants
- · Biostatic or microbicidal
- Dissipative
- Crack-bridging

Four safety levels, S 1 to S 4, in accordance with DIN EN 12128,

System structure Cleanroom Cleanroom

Tob Sauber Primer

and quartz sand

Tob Sauber Paint

2050

Scratch coat

Conductive

intermediate

Coating

Sealing

	Floor System 1 & 2	Floor System 3 & 4	Floor System 5 & 6
Prime coating	Tob Sauber Primer	Tob Sauberpoxy 2000	Tob Sauburethane 5000

Tob Sauberpoxy 2000

Tob Sauburethane Active

Tob Sauberpoxy 4000

Tob Aqua 1K PU

and quartz sand

Cleanroom coating systems for floors - An overview of the floor systems

Cleanroom

Tob Sauburethane 5000

Tob Sauburethane 4

Tob Aqua 1K PU

and quartz sand

the wall & ceiling systems System structure
Cleanroom Cleanroom Cleanroom Floor System 1 & 2 Floor System 3 & 4 Floor System 5 & 6

Cleanroom coating systems for Wall and ceiling - An overview of

Prime coating Tob Sauber Primer Tob Sauberpoxy 2000 Tob Sauburethane 5000

Scratch coat Tob Sauber Primer Tob Sauberpoxy 2000 Tob Sauburethane 5000 and quartz sand and quartz sand and quartz sand

Tob Aqua 1K PU

Tob Aqua 1K PU

Conductive Tob Sauburethane Active intermediate Coating Tob Sauburethane 4 Tob Clean Paint Tob Sauber Paint 2050

Sealing



Technical Info Center

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