

CAMFIL SERVICE IN PHARMA INDUSTRIES

Your challenge:

Protection of employees, protection against contamination of the manufacturing processes

Our solution:

- We work according to DIN EN ISO 14644-3, VDI 2083-3, VDI 6022
- Changing Hepa/Ulpa, compact & bag filters
- Qualification / requalification of the filters and rooms according to GMP requirements
- GMP-compliant documentation
- Volume flow, temperature, humidity measurements, flow visualization
- Air exchange rate determinations, room pressure measurements
- Recovery time measurements, channel filter measurements
- Air germ measurements, microbiological investigations (Evaluation by an independent institute)



You would like to take advantage of our full service?

We would be happy to present our extensive services to you in more detail.

We look forward to your inquiry - contact us now!

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