TERANGA GROUPE

CLEANING VALIDATION



Are you 100% confident in your protocols and their implementation?

Our multidisciplinary teams can help you.

Within TERANGA Group, ACM Pharma, Cebiphar and UPS Consultants teams combine their analytical & microbiological and regulatory expertise to assist manufacturers through each step of the process:

- **Overview of regulatory framework** and requirements (Europe and US)
- Consulting and assistance for **cleaning validation strategy** and its implementation
 - Organisation of the Steering Committee
 - Writing of documents (Validation plans, protocols, reports..)
 - Return on investment : optimisation of methods, grouping, worst-case, special products...
 - Which contaminants should be monitored?
 - How to calculate acceptance thresholds?

- Training of managers and operational teams and qualification of sample collecting staff
- Development and validation of **sampling techniques**
- Development, validation and transfer of analytical methods
 - Microbiological analysis
 - Physicochemical analysis : specific (HPLC, GC..) or non-specific methods (TOC..)
- **Sample analyses**

Our assets

Multidisciplinary teams Strategic and operational approach Tailored training plans







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