“Man has an intense desire for assured knowledge.”[1]

Albert Einstein

Information is perhaps the most important resource of the 21st century. At the same time, we are literally overwhelmed with a flood of it.
How do you solve this dilemma?

For more than 20 years thousands of customers trust GMP Publishing and use the good manufacturing practice knowledge of industry and regulatory experts. Our founders have been pioneers and formulated a vision:

“With us our customers always find a GMP-safe answer:
up-to-date, practical & inspection-proven!”

Do you need the most current and comprehensive GMP information? Do you need to be on the safe side in your business? Do you need to be certain that you have made the right decisions by the end of the day?

Take advantage of this expert community’s knowledge and make your GMP life easier - just like thousands of other colleagues in the pharmaceutical industry, consulting and authorities.

Increase safety and reduce time and money

Why live with uncertainty when long-term and experienced professionals can provide security in daily decisions? It's up to you - just give it a try and test the GMP Compliance Adviser with a 3-month-license.

Benefit for your career

Use the most comprehensive GMP knowledge compilation for the commercial medicinal drug manufacturing. Make use of this knowledge and benefit from it to advance professionally. Consultants take between 100-200 €/hour. In comparison the GMP Compliance Adviser costs as little as 85 € / 102 US$ per month!

It's up to you - just give it a try and test the GMP Compliance Adviser with a 3-month-license.

[1] *in P. A. Schilpp (ed.), The Philosophy of Bertrand Russell (1971), Vol. 1, 285.*

From the practice for the practice

Theory and practice are steadily changing in the regulatory environment of the 21th century. Benefit from 24/7 expert support for successful implementation of GMP compliance. Use the advantages of quality management systems up to documentation and personnel requirements. Get information to premises and facilities as well as equipment, pharmaceutical water and equipment qualification. Learn about process, cleaning and computer system validation. Know about hygiene, production, sterile manufacturing, packaging and also about laboratory controls, storage, transportation. Have insights into contractors and suppliers' challenges, GMP inspections and quality risk methodologies. And finally improve your GMP compliance continually in pharma and API manufacturing.

It's up to you - just give it a try and test the GMP Compliance Adviser with a 3-month-license.

Regulations, regulations, regulations – always up-to-date

Use all relevant GMP regulations in one source – with full text search:

EU Directives and Guidelines – U.S. CFR and FDA Guidelines – ICH-Guidelines – PIC/S Guidelines – Canadian Regulations – Japanese Regulations – Chinese Regulations – Indian Regulations – Australian Regulations – Brazilian Regulations – WHO Guidelines

A reliable source that saves you search time and for current regulations and the effort to identify the relevant sections. It's up to you - just give it a try and test the GMP Compliance Adviser with a 3-month-license.

Simplify your GMP business

- Stay informed with the latest news:
  We check and observe 50+ websites of regulatory bodies, associations, etc. for you on a weekly basis
- Save time when you are looking for adequate regulations and interpretations
- Enjoy the safety of the global leading GMP knowledge portal
- Have current GMP information & regulations at your fingertips

It's up to you - just give it a try and test the GMP Compliance Adviser with a 3-month-license.

… and b*y the way:*
The GMP Compliance Adviser has received a SIPAward already in 2016 in the category "Best New Success Story" in Washington DC. The jury was impressed with the application "Safe Drugs through Better GMP Understanding".

Your advantages	GMP Compliance Adviser
GMP in Practice	
Details to proven GMP processes, procedures, documents and technical solutions	✔
Instructions for GMP compliant solutions, easy how-to-do descriptions	✔
5,000 pages exclusive GMP knowledge	✔
More than 700 checklists, tables, figures, templates, SOPs, etc.	✔
Further literature tips	✔
GMP Regulations	
Regulations of EU, FDA, ICH, PIC/S, WHO	✔
Canadian, Japanese, Chinese, Indian, Australian, Brazilian Regulations	✔
Features	
Advanced full text search function	✔
„Editorial Questions"-Button	✔
Mark favourites (for single user licence)	✔
Edited index	✔
No LOGIN necessary for corporate licenses (automatic IP address identification)	✔
Exclusive member extras: regularly changing specials	✔
Latest GMP news on the homepage	✔