



Mourmeche



*A **European** expertise for the quality of your IT projects
in the domain of pharmaceutical industry and compliance*

OUR COMMITMENT

We are committed to the **compliance** of our clients with the current regulations in all **pharmaceutical** activities from the research to the production of medicinal products.

OUR APPROACH

Our approach is based on the Standards, Norms, Guidelines and Best Practices used by the FDA and Europe Union :

- **FDA CFR 11, GMP, GCP, GLP**
- **FDA Guidelines**
- **GAMP**
- **ISO 13485:2003**
- **ICH**

OVERVIEW

Mourmeche is a Russian company created in 2008 by two frenchmen : Jérôme Pillard and Fabrice Jansen, consultants in IT Quality Assurance.

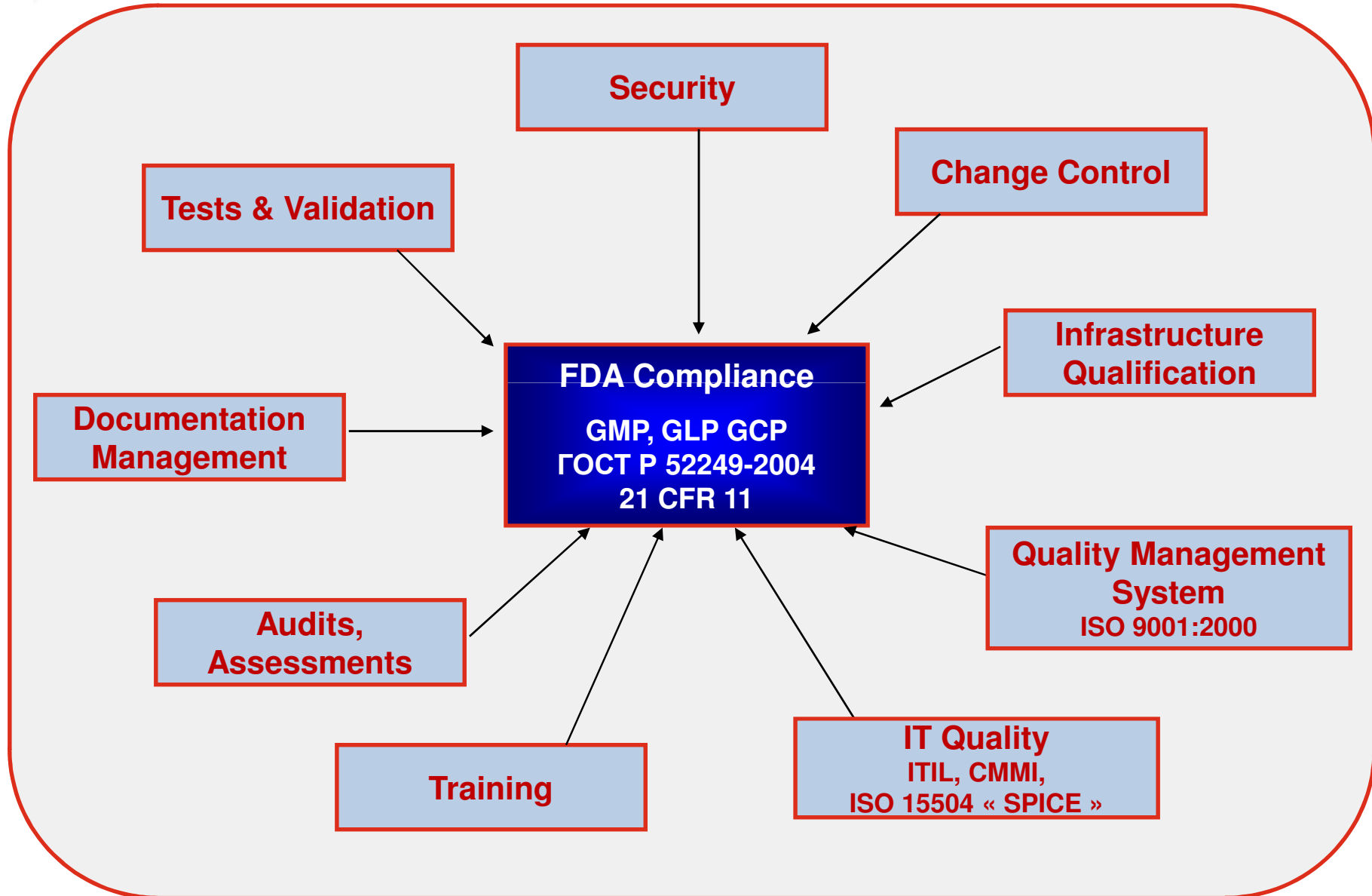
10 years experience in the pharmaceutical industry in France, Belgium, Switzerland, USA and Hungary.

Our major clients were **UCB, Glaxo SmithKline, Sanofi Aventis, and Pfizer.**

Independent of all software editors.



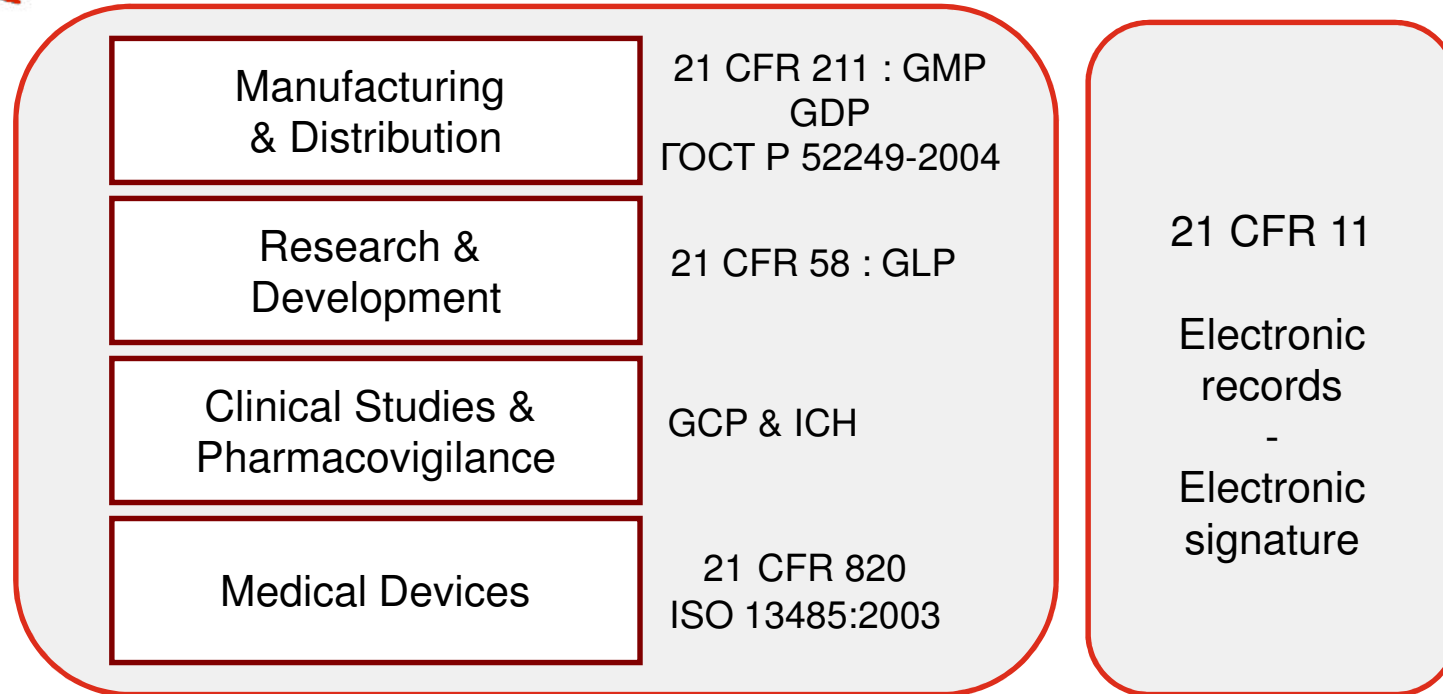
OUR SERVICES : PHARMA IT CONSULTING





Mourmeche

OUR SCOPE OF ACTIONS



OUR EXPERTISE

ERP, **Production** management & Distribution Systems (SAP...)

Electronic Documents Management Systems (Documentum...)

LIMS (HPLC Millenium Waters, Watson PSSI...)

Weighing & **Labelling** Systems (XFP Elan...)

IT Infrastructure (computer rooms, virtual servers, SAN, networks LAN WAN...)

Who we are



Jérôme Pillard, 35 years-old, is graduated from Staffordshire University, England (speciality : Computing Science and programming). From 1997, he has developed an expertise as an IT QA consultant.

He has performed many **assessments and audits of**

IT Quality Systems, for UCB, Merial, Stryker-Spine, Pfizer, Géodis, Aster-Cephac, Bouchara-Recordati.

He has performed **Training** in all his fields of expertise, for Pierre Fabre, UCB, Laboratoires Boiron, Geodis, and as an IT QA expert within consulting companies.

He has participated in numerous projects of **Software Testing** and **System Validation** (ERP, LIMS, Laboratory Systems...).

From 2001 to 2002, he worked as a **Quality Manager** in GSK (GlaxoSmithKline) IT Department and participated in the implementation and follow-up of the IT Quality System.

In 2004, he created Validation department in TESCO.

Since 2006, he lives in Russia.

Fabrice Jansen, 40 years-old, is graduated from La Sorbonne (Modern Literature) and from Innovaco Institute: Information Management.

From 2000, he has developed an expertise in **Validation and Engineering for Computerized**

Systems in the Health Industry:

- ▶ IT Infrastructure Qualification
- ▶ Validation of Computerized Systems
- ▶ Implementation of Electronic Signature
- ▶ Implementation of Quality Systems.

As **Quality Manager**, he has working in USA , Belgium and Switzerland for Novartis, Aventis, UCB, Schering SA, Galderma.

He was **Systems Analyst** during five years. He also intervened as Microprocessing and **Data Base Manager** during two years and as **IT Manager**.

Since 07-07-2007, he lives in Russia where he has married.



Mourmeche

THEY HAVE WORKED WITH US

Aster-Cephac

AstraZeneca

Aventis Pharma R&D

Bayer Pharma

Laboratoires Biocodex

Laboratoires Boiron

Laboratoires Bouchara-Recordati

Cetonia

Federa (Belgium)

Géodis

GlaxoSmithKline Biologicals
(Belgium)

GlaxoSmithKline France

Ibésis (groupe Beaufour Ipsen)

Imprimerie Tonnellier

Janssen-Cilag

Lipha (groupe Merck)

Merial

Novartis (in Switzerland)

Nufarm/SEAC

Pfizer (Pharmacia)

Pierre Fabre Dermo-Cosmétique

Porges

Sanofi Pasteur

Sanofi-Synthelabo Chinoin (Hungary)

Schering

Stryker-Spine

UCB Pharma (in Belgium, Switzerland
and USA)

Laboratoires Vétoquinol

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