

ATMP Workshop

Let's talk ATMP – GMP for ATMP and their regulatory requirements

Advanced Therapy Medicinal Products (ATMPs) promise groundbreaking therapies – yet the regulatory requirements are dynamic and complex. From the proper classification of individualized therapies and the requirements set out in the GMP guidelines to expectations for starting and ancillary materials, supplier qualification, and collaboration with CMOs: a lack of regulatory clarity quickly leads to delays, inspection findings, and additional costs.

Our workshop “Let's talk ATMP – GMP for ATMPs and their regulatory requirements” cuts through the complexity. We provide a concise update on the current regulatory landscape and the GMP framework, translate requirements into practical, actionable steps, and work hands-on together – from the QTPP to a robust specification for a starting material. You'll also receive best practices for supplier qualification, including audit approaches, as well as an open exchange on selecting, qualifying, and managing CMOs.

Outcome: clear orientation, immediately applicable approaches, and confident decision-making in your day-to-day ATMP operations.



Our Competence

- Deep ATMP regulatory and documentation expertise
- End-to-end pharmaceutical process know-how
- Interdisciplinary team of scientists and engineers

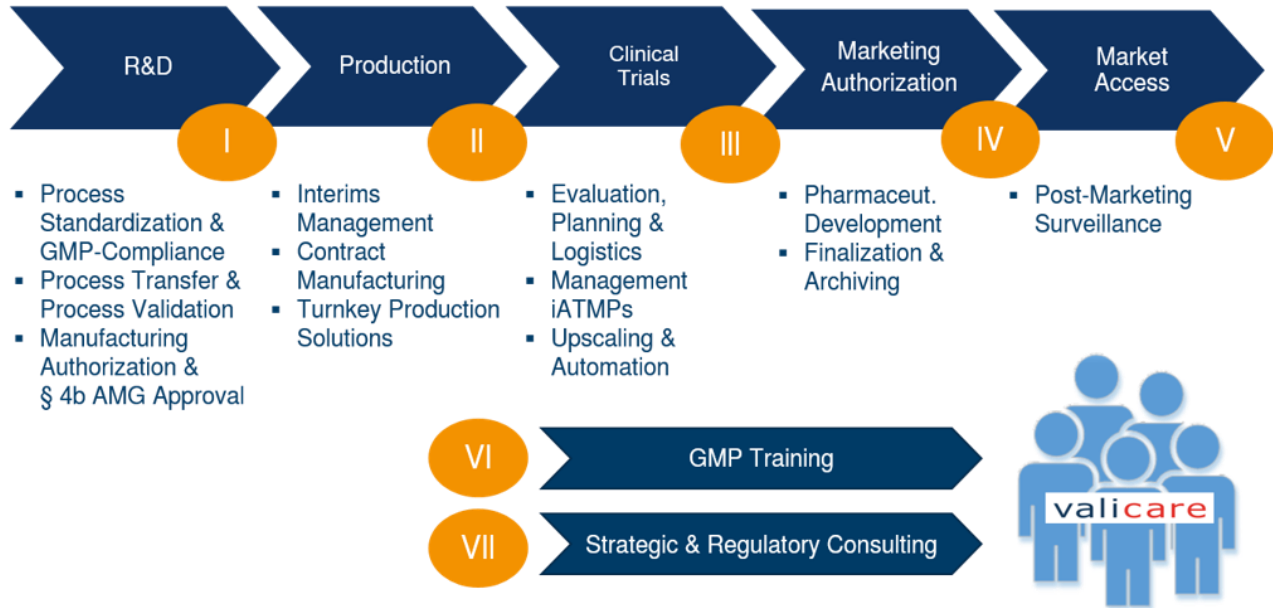
Our Services and Support

- Interactive, action-oriented workshop with current regulatory and GMP update
- Hands-on: QTPP to robust specifications for starting/ancillary materials
- Expert Q&A and real-world case studies tailored to your needs

Your Benefits

- Real-world case studies
- Hands-on exercise for deeper understanding
- Expert live Q&A tailored to your questions
- Ready-to-use checklists, templates, and examples

The Valicare-Team supports you along the entire value-added chain!



Contact us if you need clear, practical answers for your ATMP project on regulatory, GMP, and supplier/CMO management. We will support you!



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