

*If You Don't Understand It, You Can't Translate It!*

## At a glance

Accurate translation of pharmaceutical Master Batch Records (MBRs) is critical to ensuring manufacturing quality, regulatory compliance, and product consistency worldwide. Precise, compliant translations of MBRs support global manufacturing operations, safeguard patient safety, and streamline international regulatory submissions.

## Key metrics



**>30%**

Reduction in Manufacturing Downtime



**99.7%**

On Time Delivery



**100%**

Compliance with GMP regulations

## CHALLENGES



Master Batch Records contain precise manufacturing instructions and regulatory specifications that must be translated flawlessly. Any inaccuracies or inconsistencies can lead to manufacturing errors, regulatory rejections, production delays, or compromised product quality. Additionally, translating MBRs requires strict adherence to global GMP guidelines and precise terminology across multiple languages, posing significant challenges under tight production timelines.

## SOLUTIONS



AI-optimized translation combined with specialized human review ensures Master Batch Records meet global regulatory and manufacturing standards with precision, accuracy, and consistency.



**GMP Trained  
Linguists**



**12 Step Quality  
Control Process**



**Scalable To Meet  
Your Needs**

## BENEFITS



**1**

### Improved Manufacturing Compliance

Translations align perfectly with GMP standards, ensuring consistent manufacturing quality and reducing regulatory risk.

**2**

### Enhanced Global Production Efficiency

Accurate, timely translations streamline global manufacturing operations, eliminating production delays and reducing costly errors.

**3**

### Reduced Risk of Regulatory Delays

Precisely translated MBRs facilitate smoother audits, quicker approvals, and faster international product launches.