

M-Clarity[™] Program

Your Guide to Quality and Portfolio Transparency



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The development and manufacture of products in the Life Science industry has become progressively more challenging in recent years. Because of the increasing complexity of the processes, regulatory requirements and local standards, it is crucial to understand, assess and manage risks while ensuring business continuity.

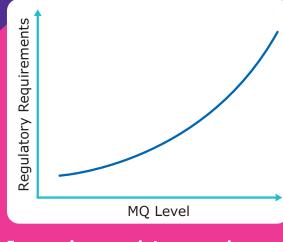
In this dynamic context, we present the **M-Clarity[™] Program** which defines product quality levels and improves product and service transparency throughout our broad Life Science portfolio.*





*Instruments, custom products, and contract manufacturing products are not part of the M-Clarity™ Program.

	Quality Segments		
	MQ100	MQ200	
Application scope	For non-regulated applications with no change notification requirements	For non-regulated applications with limited change control requirements	
Discriminating features	Standard control	Increased control	
Quality systems	ISO 9001	ISO 9001	
Quality attributes (e.g. specifications, Certificate of Quality)	•	•	
Basic change control		•	
Enhanced level of control			
Verified process			
Certified/validated process			
Highly regulated application			



Increasing regulatory needs require higher MQ levels Industry-driven regulations require that products of higher criticality or those used in highly-regulated industries, such as pharma or *in vitro* diagnostics manufacturing, need enhanced supplier quality support.

The MQ⁺ levels of the M-Clarity[™] Program provide transparency so that you can choose, with confidence, suitable products that meet your needs with respect to:

- Compliance with the appropriate quality and regulatory standards
- Portfolio transparency
- Change control notification support
- Documentation support

[†]MQ = MilliporeSigma Quality

MQ300	MQ400	MQ500	MQ600
For products used in applications requiring enhanced change control and quality agreement	For critical products and applications driven by high expectations and requiring verified process control or manufacturing control	For regulated applications	For highly-regulated applications under authority surveillance
Enhanced control	Driven by customer expectation	Driven by authority regulations	Driven by authority regulations and surveillance
ISO 9001	ISO 9001	IPEC or HACCP; FSSC 22000 or ISO 13485	ICH Q7 or 21 CFR medical device
•	•	•	•
•	•	•	•
•	•	•	•
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For more information go to SigmaAldrich.com/M-Clarity

select the right product to meet your needs

The M-Clarity[™] Program includes the majority of our Life Science products, classified into 6 MQ levels, MQ100 to MQ600:

- Each level provides specific documentation and services
- The levels have increasing attributes to meet your application and regulatory requirements
- Transparency allows you to select the right product for your needs

MOVE smoothly through product development phases

Developing and manufacturing products is a complex process involving multiple suppliers and raw materials. Minimizing disruptions when moving from development to manufacturing requires a clear risk assessment.

The M-Clarity[™] Program provides the perfect tool to guide the process of choosing components and raw materials, allowing for comparison of quality support and documentation, and ultimately minimizing costs and delays.



ENSUPE compliance by informed product selection

MQ levels provide transparency in the attributes of materials to support your regulatory requirements. The decision regarding the most relevant quality profile is driven by your specific needs for controlled and verified or validated processes as appropriate.

Leverage the M-Clarity[™] Program to choose the appropriate products to develop your own risk assessment.



To place an order or receive technical assistance

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MS_BR3925E 34617 01/2021 MilliporeSigma 400 Summit Drive Burlington, MA 01803

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