

Certificate US24/00000423

The management system of

# Sigma-Aldrich Corporation



3506 South Broadway, St. Louis, MO, 63118, United States Of America

has been assessed and certified as meeting the requirements of

## **EXCiPACT Certification Standard for Pharmaceutical Excipient Suppliers, 2021 Good Manufacturing Practices**

For the following activities

Manufacturing, quality control, quality assurance, and packaging, of dry powder and liquid cell culture media buffer solutions as pharmaceutical Auxiliary Materials.

This certificate is valid from 17 October 2024 until 16 October 2027 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 17 October 2024.

This certificate is not valid without a valid ISO 9001 certificate.

Certified activities performed by additional sites are listed on subsequent pages..



Authorised by  
Pieter Weterings  
Certification Manager

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Certificate US24/00000423, continued

# Sigma-Aldrich Corporation



## EXCiPACT Certification Standard for Pharmaceutical Excipient Suppliers, 2021

Issue 1

### Sites

Sigma-Aldrich Corporation  
3506 South Broadway, St. Louis, MO, 63118, United States Of America

Manufacturing, quality control, quality assurance, and packaging , of dry powder and liquid cell culture media buffer solutions as Pharmaceutical Auxiliary Materials.

Sigma-Aldrich Corporation  
3300 S. 2<sup>nd</sup> Street, St. Louis, MO, 63118, United States Of America

Quality control testing of dry powder and liquid cell culture media buffer solutions as Pharmaceutical Auxiliary Materials.



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To whom it may concern

**Scope of EXCiPACT® certificate for Sigma-Aldrich Corporation, Saint-Louis, MO, USA.**

Attached EXCiPACT® certificate US24/00000423 released by SGS and valid from 17 October 2024 until 16 October 2027, is applicable to “*Manufacturing, quality control, quality assurance, and packaging, of dry powder and liquid cell culture media buffer solutions as Pharmaceutical Auxiliary Materials*”, as mentioned in the scope.

We would like to share further details on the product portfolios covered by the certification:

Pharmaceutical Auxiliary Materials certified according to EXCiPACT®:2021 - PAM guide are exclusively dry powder or liquid cell culture media and buffer solutions, allocated to M-Clarity™ segment MQ500.

As stated on the label, our certified pharmaceutical auxiliary materials are not for direct animal or human use.

Sincerely,

Life Science - Quality Services

*This document has been produced electronically and is valid without a signature.*

*Date: November 13, 2024*