

# USP mAb Reference Standards

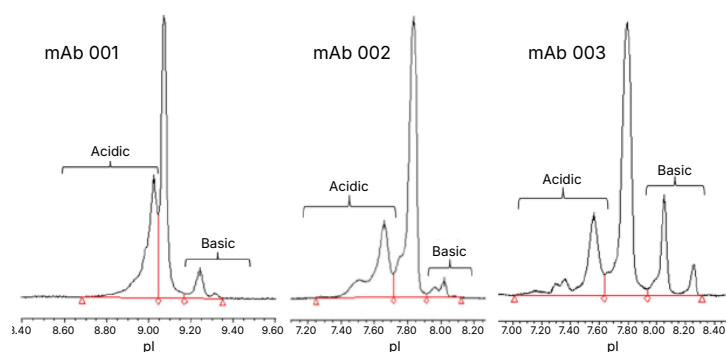
Bio-Techne now offers four monoclonal antibody U.S. Pharmacopeia (USP) Reference Standards (IgG System Suitability, mAb 001, mAb 002, and mAb 003) to overcome the limited availability of consistent, highly characterized mAb standards and to provide a range of reference products with different physicochemical properties. These USP mAb Reference Standards (RSs) aid in mAb development and manufacturing and:

- Provide a means to evaluate and monitor the performance of assays which measure physicochemical critical quality attributes (CQAs) of monoclonal antibody therapeutics
- Facilitate adoption of leading-edge analytical technologies by providing well-characterized standards
- Assist in reducing assay variability by providing materials for assay controls
- Are supported by USP quality and lifecycle management, including ongoing suitability for use studies for the lifetime of the RSs

The USP mAb Reference Standards are recombinant humanized IgG1s expressed in Chinese hamster ovary (CHO) cell culture and manufactured using industry standard upstream production and downstream purification. The Reference Standards have been rigorously tested and the results were evaluated during multi-laboratory studies using the methods described in USP General Chapter **<129> Analytical Procedures for Recombinant Therapeutic Monoclonal Antibodies**.

FIGURE // 01

Charge profile of USP mAb Reference Standards determined by icIEF on the Maurice system



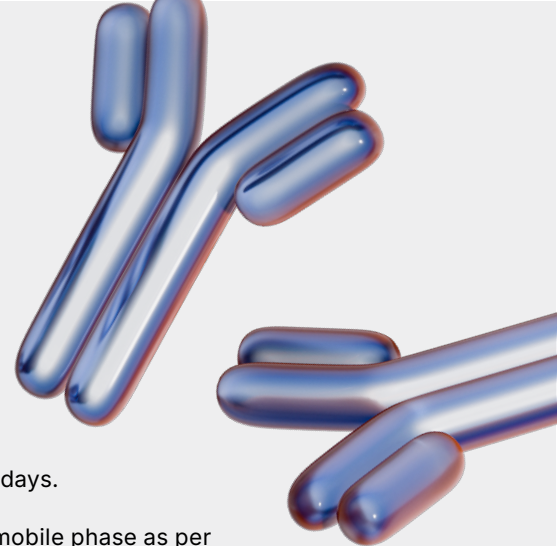
Electropherograms of USP mAb Reference Standards are representative of two instrument models, multiple preparations, and multiple injections from three independent laboratories.

## Characterization Data

- Capillary isoelectric focusing (cIEF)
- Imaged capillary isoelectric focusing (icIEF)
- Capillary electrophoresis sodium dodecyl sulfate (CE-SDS) (reduced and non-reduced)
- Intact (Protein) Mass by Spectrometry
- SE-HPLC (Size exclusion – High-performance liquid chromatography)
- N-Glycan analysis using Capillary electrophoresis – Laser Induced Fluorescence detection (CE-LIF) and Hydrophilic interaction chromatography – Fluorescence – Mass spectrometry (HILIC-FLR-MS)

## Applications

- Internal assay control
- Independent control material for method development
- Standardization of physicochemical testing
- Training activities
- Method transfer activities
- Optimization of platform methods (e.g., glycan analysis)
- Higher order structure



## IgG System Suitability Reference Standard

The USP IgG System Suitability Reference Standard enables analysts to efficiently evaluate system suitability criteria for aggregation and purity, which are critical quality attributes of mAbs, using methods outlined in USP General Chapter <129>. This standard is applicable during process development and routine method execution, ensuring consistent assay performance across different runs and days.

This USP Reference Standard is used to prepare a system suitability solution for the mobile phase as per USP Chapter <129> for the measurement of product-related impurities such as non-glycosylated molecules, half antibodies, and fragments by capillary sodium dodecyl sulfate (CE-SDS) electrophoresis.

FIGURE // 02

Conducting the USP <129> protocol on Maurice using the IgG System Suitability Reference Standard

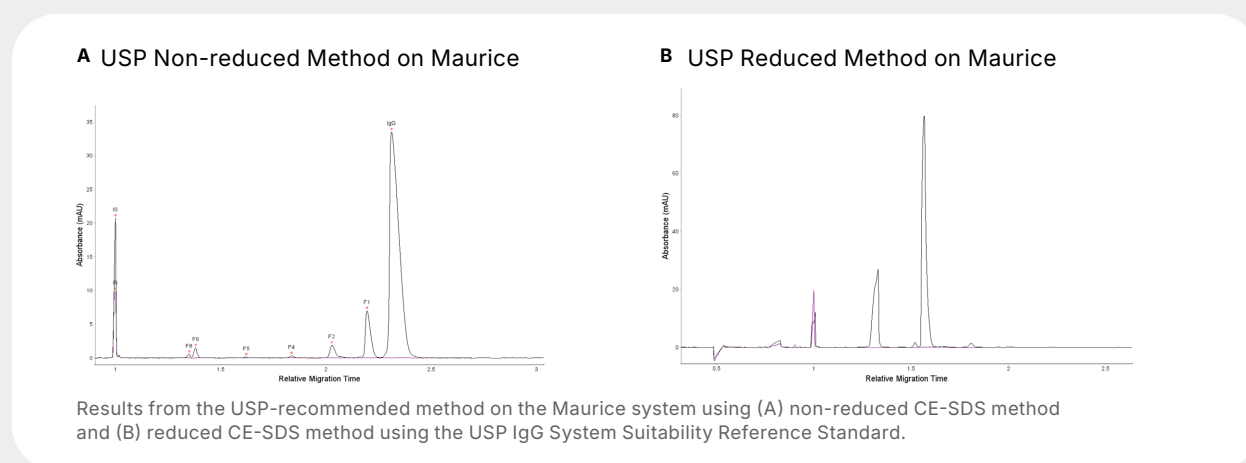


TABLE // 01

Ordering Information

Part No.	Description	Package Size
1445539	USP mAb 001, monoclonal IgG1	200 µl solution (2 mg protein content)
1445547	USP mAb 002, monoclonal IgG1	200 µl solution (2 mg protein content)
1445595	USP mAb 003, monoclonal IgG1	200 µl solution (2 mg protein content)
1445550	USP IgG System Suitability Standard	2 mg (lyophilized)



Learn more

[bio-techne.com/mab-standards](https://bio-techne.com/mab-standards)



Learn more about USP mAb Standards and Materials:

<https://www.usp.org/biologics/mabs>

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