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Eurofins PHAST offers over 9,000 recognised Reference Standards from USP, Ph. Eur./EDQM, and MHRA/BP – ideal for pharmaceutical analytics. As a USP Authorised Distributor, we are your reliable partner for high-quality standards.

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USP Reference Standards for Your Analytics

USP provides high-quality Reference Standards and Reference Materials for:

- **Chemical reference substances** – drug substances, excipients, degradation products, food additives and supplements, compendial reagents, Dissolution Performance Verification Standards



Reference Standards

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- **Biologics** – Monoclonal antibodies (mAbs), cell, gene and tissue standards, vaccine standards (mRNA), oligonucleotides, proteins, peptides, antibiotics, carbohydrates, heparins as Reference Preparations
- **Specialty Reference Materials** – Analytical Reference Materials (ARM), Pharmaceutical Analytical Impurities (PAI), Nitrosamine impurities, Excipients as Documentation Standards

USP Product Lines – The Highest Quality

Just like chemically derived pharmaceuticals, biologic developers and manufacturers must ensure product purity and quality. For biologics and biosimilars, USP focuses on product families and classes, emphasising analytical methods, Reference Materials, and standards to support testing procedures.

Ensuring that original drugs and generics/biosimilars achieve their expected therapeutic effect requires high-quality standards. USP products from Eurofins PHAST, your authorised distributor, provide the support you need.

Contact Us:

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