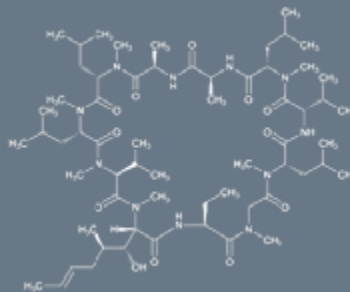# Cyclosporine
Quality Solutions
Category: Immunosuppressant

Authorized Distributor

USP can support your development and manufacturing activities on Cyclosporine-based medicines with these existing and upcoming standards.

Official and Proposed Documentary Standards and Associated Physical Materials

MONOGRAPH 1

CYCLOSPORINE Official as of 01-Dec-2021	
USP Cyclosporine RS	USP-1158504
USP Cyclosporine Resolution Mixture RS	USP-1158650

MONOGRAPH 2

CYCLOSPORINE INJECTION Official as of 01-May-2018	
USP Cyclosporine RS	USP-1158504

MONOGRAPH 3

CYCLOSPORINE COMPOUNDED OPHTHALMIC SOLUTION, VETERINARY Official as of 01-May-2020	
USP Cyclosporine RS	USP-1158504

MONOGRAPH 4

CYCLOSPORINE ORAL SOLUTION Official as of 31-Dec-2012	
USP Cyclosporine RS	USP-1158504

MONOGRAPH 5

CYCLOSPORINE CAPSULES Official as of 31-Dec-2012	
USP Cyclosporine RS	USP-1158504

PHARMACEUTICAL ANALYTICAL IMPURITY(IES)*	
USP Isocyclosporin A PAI I *New*	USP-1A03320
USP Cyclosporin D PAI I *New*	USP-1A04270
USP Cyclosporin H PAI I *New*	USP-1A04280
USP Sodium 4-Hydroxy-N-Methylleucinate PAI I *New*	USP-1A04290

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* Documents in PF Online are not official and not suitable to demonstrate compliance. They may never become official.

01-Dec-2023. For publications related to **Cyclosporine** in *PF* online after this date, please visit the integrated *USP-NF/PF* online.

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