Master Formulation Record

Name of compounded product:		Protocol number and version:	
Concentration:		Effective date:	
Pharmaceutical form: Route of administration:		Developed by: Verified by:	
Formula			
Ingredients:	Quantities:	Physical description:	Other information (i.e. DIN, lot number, manufacturer, expiry date, expected yield):
Additional information about the ingredients:			
Notes on calculations and measurements:			
Required equipment, instruments, and materials:			
Compounding method:			
Quality controls:			

Packaging:	
Stability and storage:	
Labelling:	Sample label:
	Patient label:
Training:	
References consulted:	
Preparation date sheet history #:	
Revised: Change made:	Revised by: Version number changed: ☐ Yes ☐ No
Revised: Change made:	Revised by: Version number changed: ☐ Yes ☐ No