

## CATEGORY 3 CSP MEDIA-FILL TEST KIT LOG SHEET

Employee Name: \_\_\_\_\_

Employee Signature: \_\_\_\_\_

Date: \_\_\_\_\_

Lot Number: \_\_\_\_\_

Expiration Date: \_\_\_\_\_

Vial Number	Hood Number	Incubation Start (Date & Time)	Temperature One <sup>1</sup> (°C/°F)	Temperature Adjusted (Date & Time)	Temperature Two <sup>2</sup> (°C/°F)	Incubation End <sup>3</sup> (Date & Time)	Result <sup>4</sup>	Notes
<b>Control 1</b>							Growth / No Growth	
<b>Control 2</b>							Growth / No Growth	
<b>Control 3</b>							Growth / No Growth	
<b>1</b>							Pass / Fail	
<b>2</b>							Pass / Fail	
<b>3</b>							Pass / Fail	
<b>4</b>							Pass / Fail	
<b>5</b>							Pass / Fail	
<b>6</b>							Pass / Fail	

1: 20°– 25°C (68°– 77°F)

2: 30°– 35°C (86°– 95°F)

3: Per USP <797>, incubate for a minimum of 7 days at each temperature range. A total incubation period of at least 14 days is recommended to confirm the absence of microbial growth.

4: Any signs of microbial growth, such as turbidity or precipitation, at any point during the incubation period, indicates a failure in the sterility test. Microbial growth is expected for the vials in the control group.

Supervisor Name: \_\_\_\_\_

Supervisor Signature: \_\_\_\_\_

Date (results acknowledged): \_\_\_\_\_

INTRAVENOUS QUALITY ASSURANCE

6260 River Crest Dr., Suite E  
Riverside, CA 92507  
(626) 629-0418  
support@ivqa.com  
www.ivqa.com