

# Certificate of Compliance

No.:166W20200901

<b>Product Code</b>	82141 12X75mm culture tube	<b>Specification</b>	Rimed,Dual-position cap,Graduated,PS,EO Sterile		
<b>Lot No.</b>	09/02/20	<b>Expire Date</b>	09/01/25		
<b>Packaging</b>	Individual peel bag pack,50PCS/bag,10bags/C ase	<b>Samples Tested</b>	32pcs	<b>Lot Quantity</b>	1000pcs
SPECIFICATIONS				Findings	
<b>Material</b>	PS, Non-cytotoxic.			yes	
<b>Appearance</b>	There is no Impurity, Oiliness, incompleteness, scratch onto tube/cap.			yes	
	No contamination from dust or something else outside.			yes	
	Color of body: transparency-the pure color of PS material			yes	
	Color of Cap: transparency-the pure color of PE material			yes	
	Printed figures are correct, clear and complete. No blur.			yes	
	Printed lines are in correct position, no blur and segmentation.			yes	
	The printed figures&lines will not disappear or lose color after scratch for at least 20 times.			yes	
<b>Performance</b>	When pushing on the cap onto tube, the hand-feeling should be better.			yes	
	External Diameter of Cap: 16.1mm ± 0.3mm			yes	
	Overall Height of Cap: 13.7mm ± 0.3mm			yes	
	External Diameter of Tube: 12.3mm ± 0.3mm			yes	
	Overall Length of Tube: 75.0mm ± 0.3mm			yes	
	Total height(w/cap): 76.2mm ± 0.3mm			yes	
<b>Weight (EA)</b>	Cap: 0.63±0.50(g); Tube: 2.86±0.50(g)			yes	
<b>Sterilization</b>	Ethylene Oxide sterile			ETO	

**Quality System Compliance** –Runlab Technology Co.,Ltd. products are manufactured under the ISO 13485 system. Management system 13485 with SOPs and clean control room guarantee quality, and Stringent quality control protocols are followed at every process of production ranging from qualitative raw materials and process of production till final products.

**Quality Control Testing** –Products are Inspected and controlled through whole production processing in accordance with current applicable product specifications and QC SOP. Inspection records are reviewed and signed off by qualified personnel for product release.

**Sterilization** – Runlab Products have been sterilized with Ethylene Oxide and found to conform to the sterility requirements of EN ISO11135:2014 .

The details and process of sterilization are as follows:

1. Type of sterilization cycle: overkill cycle
2. Gas mixture: EO density of 1Kg/m3
3. Maximum level of gas residue: 10ppm
4. Sterilization process:
  - a. Load the products in the sterilization chamber.
  - b. Load and locate 20 bio-indicators in the sterilization chamber for proving and check the sterility.
  - c. Precondition at 45±5°C temperature and 50-70%humidity for 30min.
  - d. Initial vacuum: 50±1Kpa
  - e. Inject the 20% ETO into the sterilization chamber and operate sterilization cycle at temperature 50±1°C and 40-70%humidity with exposure pressure 135Kpa for 8 hours.
  - f. Evacuate the gas to 70±2Kpa, then turn off the vacuum valve.
  - g. Inject the fresh air into the sterilization chamber, when the pressure of chamber is atmosphere, turn off the air valve.
  - h. Repeat the procedure of f and g for 3 times.

QA Dept. Director: Wang Guoli

Date: September 01, 2020

