



# **IMPORTANT RULES**

## **FOR SUCCESSFUL CELL CULTURING**

### **1, Do not expose the cryovials to ambient temperature**

- Keep the cryovials in liquid nitrogen for long term storage.
- Bury the cryovials in dry ice at all times when out of liquid nitrogen prior to use.

### **2. Do not over-thaw the cryovials**

Thaw cryovials in a 37°C water bath no longer than 90 seconds; there should still be ice crystals left in the vials.

### **3. Do use treated tissue culture dishes**

Corning or Greiner culture dishes are recommended for optimum performance.

### **4. Do not heat subculture reagents to 37°C**

Use the Trypsin/EDTA at or below room temperature. DO NOT heat them to 37°C in incubator or water bath at any time.

### **5. Do not over-trypsinize the cells**

- Use Trypsin/EDTA with the correct concentration.
- Knock the cells loose when the cells round up. Do not wait for the cells to detach by themselves.

### **6. Read the instructions closely for optimum cell growth**

Prevent mistakes and save time in the long run.