**Designation:**

**Zajdela-Hepatoma ZH-CLS**

**CLS order number:**

Cryovial: 500306  
Vital: 550306

### Origin and General Characteristics

| Depositor: | CLS |
| Organism: | Rat |
| Age/Stage: | 11 months |
| Strain: | Sprague-Dawley |
| Tissue: | Liver |
| Morphology: | Hepatoma |
| Growth Properties: | Monolayer, adherent |

**Description:**

This cell line was established as in vitro cell line from the Zajdela-Ascites-Hepatoma by CLS in 1998.

### Culture Conditions and Handling

**Culture Medium:**

McCoy's 5a medium supplemented with 2 mM L-glutamine and 10% fetal bovine serum (MG-200, CLS order number 821200).

**Subculturing:**

Remove medium and rinse the adherent cells using PBS without calcium and magnesium (3-5 ml PBS for T25, 5-10ml for T75 cell culture flasks). Add Accutase (1-2ml per T25, 2.5ml per T75 cell culture flask), the cell sheet must be covered completely. Incubate at ambient temperature for 8-10 minutes. Carefully resuspend the cells with medium (10 ml), centrifuge for 5 min at 300xg, resuspend cells in fresh medium and dispense into new flasks which contain fresh medium.

**Split Ratio:**

A ratio of 1:4 to 1:8 is recommended

**Fluid Renewal:**

Every 3 to 5 days

**Freeze Medium:**

CM-2 (CLS order number: 800225, 25ml, 800250, 50ml)

**Sterility:**

Fluorescence (DAPI) test: negative; Mycoplasma specific PCR: negative; Bacteria specific PCR: negative

**Biosafety Level:**

1

### Special Features of the Cell Line

**Tumorigenic:**

Yes, in Sprague-Dawley rat

**Viruses:**

SMRV: Negative, as confirmed by Real-Time PCR  
RAP-Test negative.

**Authentication:**

The rat origin was verified by Real-Time PCR.

**References:**

Wintzerith M et al. Comparative study of free uridylic nucleotides in the normal liver, the regenerating liver and in the Zajdela hepatoma. CR Seances Soc Biol Fil 156: 2114-8, 1962.
Recommendations for handling of adherent cell cultures following delivery

**Cryopreserved cells**
If immediate culturing is not intended, the cryovial(s) must be stored in liquid nitrogen (-196°C) or at least at -80°C after arrival.

If immediate culturing is intended, please follow these instructions:
Quickly thaw by rapid agitation in a 37°C water bath within 40-60 seconds. The water bath should have clean water containing an antimicrobial agent. As soon as the sample has thawed, remove the cryovial from the water bath. Note: A small ice clump should still remain and the vial should still be cold.

From now on, all operations should be carried out under aseptic conditions.
Transfer the cryovial to a sterile flow cabinet and wipe with 70% alcohol. Carefully open the vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of culture medium (room temperature). Resuspend the cells carefully. Centrifuge at 300xg for 3 min and discard the supernatant. The centrifugation step may be omitted, but in this case the remains of the freeze medium have to be removed 24 hours later.
Resuspend the cells carefully in 10 ml fresh cell culture medium and transfer them into two T25 cell culture flasks. All further steps are described in the Subculture section.

**Proliferating Cultures**
The cell culture flasks are completely filled with cell culture medium to prevent loss of cells during transit. Remove the entire medium except for a sufficient volume to cover the floor of the flask. Incubate at 37°C for 24 hrs.

Sometimes the cultures are handled roughly during transit, and most of the cells detach and float in the culture medium. If this has occurred remove the entire content of the flask and centrifuge at 300xg for 3 minutes. Take off the supernatant, resuspend the cells in 10 ml of culture medium and transfer the entire cell suspension into cell culture flasks of suitable size (do not seed in more than 1T75 flask).

Safety precautions for frozen cell lines
If the cryovial is planned to be stored in liquid nitrogen and to be thawed in the future, special safety precautions should be followed:

- Protective gloves and clothing should be used and a facemask or safety goggles must be worn when storing and/or thawing the cryovial.
- The removal of a cryovial from liquid nitrogen can result in the explosion of the cryovial creating flying fragments.