

Product sheet

XXXXXXXX B95-8 | 601102

XXXXXX XXXXXX

Description XXX XXXXX B95-8 XXX XX XXXXXXXXXXXXXXX B XX XXXXXXXX, XXXXXX XXXX XXX XXXXXXX XXXXXXXXXXXX XX XXXXXXX XXX XXX (Saguinus oedipus). XXX
XXXX B95-8 XXXXXXXX XXXXXXX XXXXXXX XXXXXXX XXXXXXX XXXXXXX-XX XXXXXXX XXXXXXXXXX. XX XXXXXXX XXXXXXX XXXX XXXXXXX, XX XXXXXXX XXXX XXXX XXXX

Organism XXX XXXXX XXX-XXXX

Tissue XX

Synonyms B95.8, B 95.8, B 95-8, B-95-8, B958, GM07404, GM07404A, GM07404D

XXXXXXXXXXXX

Gender XXXXX

Morphology XXXXXXXXXX

Growth properties XXXXXXX

XXXXXXXXXX XXXXXXXXXXXXXXX

Citation B95-8 (XXXXX XXXXXXXX Cytion 601102)

Biosafety level 2

NCBI_TaxID 9490

CellosaurusAccession CVCL_1953

XXXXXXXXXX XXXX-XXXXXXXXXXXX

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Culture Medium RPMI 1640, w: 2.0 mM XXXXXXXX XXXXX, w: 2.0 g/L NaHCO3 (XXXXX XXXXXXXX XX Cytion 820700a)

Supplements XXXXX XXXXXXX 10% FBS

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Subculturing 1. Remove the medium from the flask and add 5 ml of fresh medium.

Split ratio 1:2 to 1:4

Fluid renewal 2 to 3 times per week

Freeze medium DMEM, 10% FBS + 10% DMSO

- Thawing and Culturing Cells**
1. Thaw the vial in a 37°C water bath.
 2. Add 1 ml of DMEM to the vial and centrifuge at 300 x g for 3 minutes.
 3. Remove the supernatant and resuspend the cells in 10 ml of DMEM.
 4. Seed the cells into a 150 cm² flask containing 70% medium.
 5. Incubate the cells at 37°C in 5% CO₂.
 6. Monitor the cells for confluency.
 7. Harvest the cells when they reach 80-90% confluency.
 8. Perform subculturing as described above.

Incubation Atmosphere 37°C, 5% CO₂

Flask Coating Not required

Freezing Procedure Freeze cells in DMEM + 10% FBS + 10% DMSO

Shipping Conditions Ship cells at -78°C

