

T47D/S5 Cell Line

Catalogue number: 152111

Sub-type: Continuous

Images:

Contributor

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Images:

Tool details

***FOR RESEARCH USE ONLY**

Name: T47D/S5 Cell Line

Alternate name:

Class:

Conjugate:

Description: The T47D/S5 cell line is a control cell line for the fulvestrant (Faslodex) resistant T47D/182R-1 and T47D/182R-2 cell lines. T47D/S5 is adherent and the morphology is polygonal epithelial. The cell passage number is 155 (AL3043, AL3044). T47D/S5 cells express oestrogen receptor alpha and progesterone receptor. This cell line allows the study of the mechanisms involved in fulvestrant resistant breast cancer cell growth.

Purpose:

Parental cell: T47D

Organism: Human

Tissue: Breast

Model: Tumour line

Gender:

Isotype:

Reactivity:

Selectivity:

Host:

Immunogen:

Immunogen UNIPROT ID:

Sequence:

Growth properties:

Production details: T47D/S5 cells grows continuously in presence of 5% fetal calf serum.

Formulation:

Recommended controls:

Bacterial resistance:

Selectable markers:

Additional notes:

Target details

Target: Oestrogen receptor

Target alternate names:

Target background:

Molecular weight:

Ic50:

Applications

Application: Determining molecular mechanisms around fulvestrant resistance

Application notes:

Handling

Format: Frozen

Concentration:

Passage number: Passage 155 (AL3043, AL3044)

Growth medium: Phenol red free RPMI 1640 + 5% FCS + glutamax + 8ug Insulin/ml

Temperature: 37° C

Atmosphere: 5% CO2

Volume:

Storage medium:

Storage buffer:

Storage conditions:

Shipping conditions: Dry ice

Related tools

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References

References: Larsen et al. 2015. PLoS One. 10(2):e0118346. PMID: 25706943. ; Larsen et al. 2015.

BMC Cancer. 15:239. PMID: 25885472. ; Aurora kinase B is important for antiestrogen resistant cell growth and a potential biomarker for tamoxifen resistant breast cancer. ; SRC drives growth of antiestrogen resistant breast cancer cell lines and is a marker for reduced benefit of tamoxifen treatment. ; Thrane et al. 2014. Oncogene. :. PMID: 25362855. ; A kinase inhibitor screen identifies Mcl-1 and Aurora kinase A as novel treatment targets in antiestrogen-resistant breast cancer cells.

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