

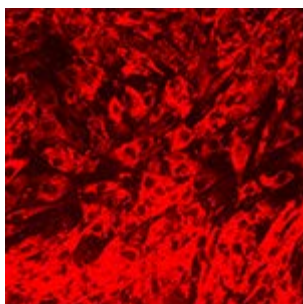


October 2017

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cell passages

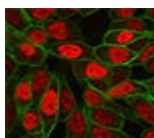


New Tumor Microenvironment Tools

Tumor progression is not solely determined by the mutated cell, but also by the tumor's microenvironment. ATCC now offers normal-associated fibroblasts (hTERT SMC PM151T; [ATCC® CRL-3291™](#)) paired with prostate cancer-associated fibroblasts (hTERT PF179T CAF; [ATCC® CRL-3290™](#)) to provide you with the ideal platform to investigate the transformation of prostate cells in the context of the tumor microenvironment.

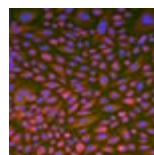
Explore hTERT-immortalized Primary Prostate Cells>>

- Media collected from cancer-associated fibroblasts promotes increased LNCaP growth as compared to media collected from normal-associated fibroblasts and control growth medium
- Increased α -SMA upon TGF- β stimulation
- Cells undergo smooth muscle-dependent differentiation



Tumor Xenograft Prostate Cancer Cells

ATCC now offers cell line-derived explant models of prostate cancer. C4 ([ATCC® CRL-3313™](#)) and subline C4-2 ([ATCC® CRL-3314™](#)) are clones of LNCaP ([ATCC® CRL-1740™](#)) that were introduced subcutaneously into an athymic male nude mouse and isolated from the resulting tumor. These cell lines offer an advanced model of prostate cancer tumorigenicity, androgen-independent



Primary Vaginal Epithelial Cells

ATCC Human Primary Vaginal Epithelial Cells ([ATCC® PCS-480-010™](#)) may be used to study the cellular physiology of the reproductive tract, cellular response to infectious agents, and female reproductive tract cancer development. ATCC also offers a complete solution for culturing these cells, including a Vaginal Epithelial Cell Basal Medium ([ATCC® PCS-480-030™](#)) and a Vaginal Epithelial Cell

®

progression, and bone metastasis.

Growth Kit ([ATCC PCS-480-040™](#)).

[Order cell line C4 \(ATCC® CRL-3313™\) >](#)

[Order cell line C4-2 \(ATCC® CRL-3314™\) >](#)

[Primary Vaginal Epithelial Cells >](#)

[Vaginal Epithelial Cell Basal Medium >](#)

[Vaginal Epithelial Cell Growth Kit >](#)



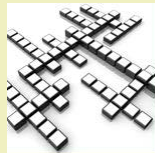
Human Renal Proximal Tubule Epithelial Cells (RPTEC/TERT1) Modified to Express Drug Transporters - A New Model for Toxicity Studies

Presenter: Chaozhong Zou, Ph.D., *Senior Scientist*, ATCC

October 19, 12:00 PM ET

This presentation will introduce hTERT-immortalized RPTEC that stably overexpress the OAT1 or OCT2 gene. These modified cell lines provide kidney tissue-relevant results, improved consistency over time, and more predictability for clinical trials versus current models.

[Register today>](#)



ATCC Puzzle

Try this [month's crossword puzzle](#). The solution will appear in next month's issue.

For the solution to last month's puzzle [click here](#).

Resources

- [Poster: hTERT-immortalized Prostate-derived Stromal Cells](#)
- [Reproductive Cancer](#)
- [Gynecologic Cancer Cell Panel](#)
- [Prostate Cell Lines](#)



Frequently Asked Questions

Q: What is cholera toxin and why is it used in cell culture media?

A: Cholera toxin is an adenylate cyclase activator. It stimulates the growth of epithelioid cells from normal breast *in vitro*. Biochemically, it is involved with the transportation of calcium. This supplement has also been found to be growth stimulating for cells derived from colon, lung, prostate, and skin.

[Have more questions?](#)

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