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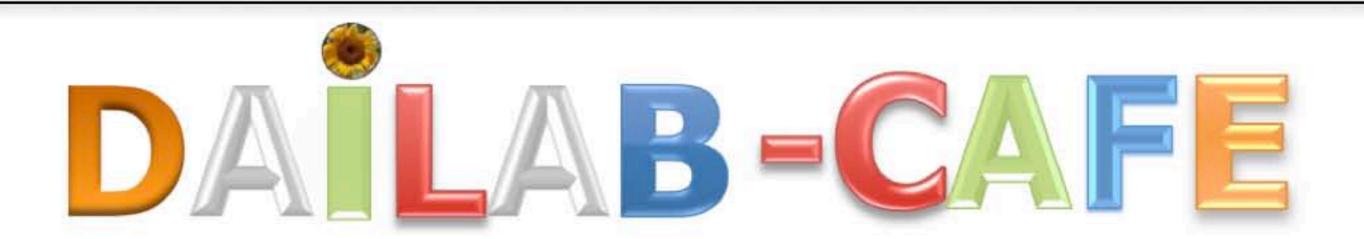
DBT -AIST International Laboratory for Advanced Biomedicine



## Classroom for Advanced & Frontier Education







## Series - 12

E-mail: t-tamura@aist.go.jp

Date and Time – Oct. 7, 2015 (12:30~13:30)

Venue - Central 4 (2F) Meeting Room 1

Speaker – Tomohiro TAMURA

Affiliation –Bioproduction Research Institute, AIST, Japan



## Title – Efficient production of active form of vitamin D<sub>3</sub> by microbial conversion

Vitamin  $D_3$  (VD<sub>3</sub>) is a fat-soluble prohormone that plays a crucial role in bone metabolism, immunity, and the control of cell proliferation and differentiation. The most active form,  $1\alpha,25(OH)_2VD_3$ , is used to treat osteoporosis, hyperparathyroidism, psoriasis, and VD<sub>3</sub> metabolic abnormality. The industrial production of  $1\alpha,25(OH)_2VD_3$  is performed chemically or microbiologically, but the processes for the microbiological production of the active form of VD<sub>3</sub> are simpler than those for chemical synthesis.

The actinomycete *Pseudonocardia autotrophica* is capable of bioconversion of  $VD_3$  into its physiologically active forms,  $25(OH)VD_3$  or  $1\alpha,25(OH)_2VD_3$ . We identified vitamin  $D_3$  hydroxylase (vdh) from *P. autotrophica* and characterized it structurally and enzymatically. Biotransformation of  $VD_3$  into  $25(OH)VD_3$  was then accomplished with a Vdh-expressed recombinant strain of actinomycete *Rhodococcus erythropolis*. We have recently succeeded in significant improvement of cellular permeability of vitamin D3 by using nisin-treated cells, and have developed a new platform for vitamin  $D_3$  hydroxylation process.

In this seminar, I would like to introduce how to improve the efficiency of production of hydroxylated form of vitamin D3 by using *Rhodococcus erythropolis* as a host cell.







Dear Tamura san,
Thank you!
for
An extremely
interesting
CAFÉ-TALK
It was thoroughly
enjoyed!