



International Journal of Advanced Research in Biology, Engineering, Science and Technology (IJARBEST)

Vol. 2, Special Issue 8, February 2016 in association with

KAMARAJ COLLEGE OF ENGINEERING AND TECHNOLOGY, VIRUDHUNAGAR
DEPARTMENT OF BIOTECHNOLOGY

ORGANIZES

DBT, NEW DELHI SPONSORED NATIONAL LEVEL CONFERENCE ON CONTEMPORARY TRENDS IN
BIOENERGY AND GREEN TECHNOLOGY: CHALLENGES AND OPPORTUNITIES [ORA-2016]

(25-26TH FEBRUARY 2016)

IDENTIFICATION AND MOLECULAR CHARACTERIZATION OF OSTEOCLASTS IN CHRONIC OBSTRUCTIVE PULMONARY DISEASE

Vasanthakumar R¹, Radhakrishnan S², Tholcopiyan L², Shila S^{2*}

¹Rajalakshmi Engineering College, Chennai.

²VRR Institute of Biomedical Science (Affiliated to University of Madras), Chennai.

*Corresponding Author: E-mail: vrribms@gmail.com; Ph: +914426791700; Fax: +914424360219

Abstract

Background: Chronic Obstructive Pulmonary Disease (COPD) is one of the chronic diseases that facilitate the development of secondary osteoporosis. Prevalence of osteoporosis is 2 to 5 fold high in COPD patients compared to normal subjects. Molecular characterization of osteoclasts is crucial to understand the in-depth mechanism of osteoporosis development in COPD patients, since osteoclasts are the bone-resorbing cells.

Methods: A study was designed with 13 COPD and 7 normal healthy subjects. Monocytes were isolated from the whole blood of all the study subjects and cultured in the presence osteoclast differentiation factors (RANKL and M-CSF) for 14 days. Matured osteoclasts (3+) were identified by TRAP staining. NFATc1, c-Fos, TRAF6, and Fra-1 levels were determined by immunoblot analysis while cathepsin K levels were identified by RT-PCR.

Results: COPD subjects showed significant increase ($p < 0.001$) in the number of TRAP positive multinucleated osteoclast cells when compared to normal subjects. Expression of NFATc1, TRAF6, c-Fos, Fra-1 were significantly increased ($p < 0.001$) in the COPD subjects as compared to normal subjects and similarly, cathepsin K were found to be significantly increased ($p < 0.001$) in the COPD subjects as compared to normal subjects.

Conclusion: This clearly evidences that the COPD condition facilitates the development of osteoporosis by inducing RANKL mediated osteoclastogenesis.

Keywords: COPD; Osteoporosis; Osteoclasts.