## Differentiation between osteomyelitis and malignant tumor infiltration into the mandibular bone using [<sup>18</sup>F]FDG PET

Yasufumi Hara<sup>1</sup>, Kazunori Terasaki<sup>2</sup>, Masanori Shozushima<sup>3</sup> and Yoshiki Sugiyama<sup>1</sup>

<sup>1</sup>Div. of Oral and Maxillofacial Surgery, <sup>3</sup>Div. of Dental Radiology, Department of Reconstructive Oral and Maxillofacial Surgery, Iwate Medical University 19-1 Uchimaru, Morioka, Iwate 020-8505, Japan

> <sup>2</sup>Cyclotron Research Center, Iwate Medical University 348-58 Tomegamori, Takizawa, Iwate 020-0603, Japan

## Abstract

Introduction

When clinical findings similar to osteomyelitis are noted in the course observation after oral cancer treatment, it is difficult to differentiate between the recurrence of the cancer or osteomyelitis caused by radiation.

**Objectives** 

This study was performed to evaluate the usefulness of PET using  $^{18}\mathrm{F}\text{-}\mathrm{FDG}$  for such differentiation.

## Materials and Methods

The subjects were 4 patients who had been diagnosed with carcinoma of the tongue or gingiva and received treatment. In the postoperative course observation, <sup>18</sup>F-FDG PET was performed to differentiate between osteomyelitis and the recurrence of cancer. Twenty cases of carcinoma of the gingiva with jaw bone infiltration were selected as the control.

Results

The mean SUV was  $5.6 \pm 1.2$  in osteomyelitis, and  $11.2 \pm 5.7$  in gingival carcinoma. No significant difference in the mean was noted between the two groups, although clear differentiation was impossible. However, in cases in which SUV is over 7, it is estimated that the possibility of cancer is markedly high. <u>Conclusions and discussion</u>

1) Although the mean SUV  $_{max}$  was higher in bone infiltration by malignant tumors than in osteomyelitis, no significant difference was noted. However, in cases in which SUV is over 7, it is estimated that the possibility of malignant tumors is markedly high.

2) From these findings, it is considered that <sup>18</sup>F-FDG accumulates not only in cancer cells and inflammation-activated macrophages and lymphocytes, but also in bone metabolism-related osteoclasts and osteoblasts.