学会記事

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En block resection of the temporomandibular joint is relatively difficult because it islocated at inferior aspect of the middle cranial fossa of the temporal bone. Technique for dissection are limited by the need to avoid undue damage to the body surface of fresh cadavers for pathological examination. In our presentation, we will introduce technique that we use for dissection of the temporomandibular joint in human fresh cadavers at our institution. The techniques will be discussed with respect to their application for various study method.

13.Studies on distribution and characterization of proteoglycans (PGs) in porcine Temporomandibular Joint (TMJ) Tomoaki Shibuya $^{1)}$, Kouji Kino $^{1)}$, Junji Kobayashi $^{1)}$, Akiko Kobayashi $^{1)}$, Teruo Amagasa $^{1)}$ and Tohru Takagi $^{2)}$ $^{1)}$ Dept. of Oral Maxillofacial Surgery, Faculty of Den tistry, Tokyo Medical and Dental Univ.

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Distribution of porcine TMJ PGs are different com position in TMJ Disk, retrodiskal tissue and superficial layer of condyle PGs. In this studies, Glycosaminoglycans were assaied by cellulose acetate mem brane electrophoresis. The core proteins were assaied by western-blotting. A high molecular weight PG resembled the Aggrecan and two low moleculaer weight PGs resembled Biglycan and Decorin.

14. Thermal analysis of decomposition of collagen at high temperature

Toshiro Sakae¹⁾, Yukie Sato^{2),} Yukishige Kozawa^{1),} Hiroyuki Yamamoto²⁾

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It is well known that denaturing temperature of collagen has been extreamly studied. We found that at the higher temperature range collagen decomposed and showed a variety of decomposition pattern in the thermal analysis curves. Possibility of applying thermal analysis technique for study of collagen is discussed.

15.Bone formation induced by the implantation of bone marrow cell collagen matrix complexes in vivo.Morimichi Mizuno, Daiji Kobayashi, Eichi Tsuruga,Yoshinori KubokiDepartment of Biochemistry, School of Dentistry, Hokkaido University

The purpose of this study is to induce bone by the implantation of bone marrow cell-collagen matrix. Bone marrow cells differentiated to osteoblasts on collagen matrix. When cell-collagen complexed were implanted to nude mice, bone was formed and it contained bone marrow.

16. Immunohistochemical localization of TGF-b1 in the process of BMP induced heterotopic osteogenesis Toshiyuki Kawakami, Noriyuki Takei, Masayuki Kanatani, and Shigeo Eda

Department of Oral Pathology, Matsumoto Dental College Using ddY mice, immunohistochemical localizations

of TGF-b1 in the early phase of BMP induced heterotopic osteogenesis were examined. In one week specimen, TGF-b1 was detected in chondrocytes and the matrices, and in two weeks appeared in osteoblasts and the matrices. In three weeks case, the peptide was shown in osteoblasts and osteocytes, and newly formed bone marrow tissues. The positive reaction remained only in a part of the matrices in four weeks specimen.

17. Expression of Bone Matrix Proteins during Ectopic

Bone Formation Induced by Bone Morphogenetic Protein Noriyuki Nagai¹⁾, Manabu Kanyama^{1)2),} Hidetugu Tujigiwa¹⁾ Yuzou Ishiwar¹⁾, and Hironobu Konouchi¹⁾

- 1) Dept. of Oral Pathology, Okayama Univ. Dental School.
- ²⁾ Dept. of Fixed Prosthodontics, Okayama Univ. Dental School mRNA expression of bone matrix proteins was investigated during the formation of ectopic bone induced by bone morphogenetic protein. Partially purified bovine BMP was implanted into the dorsal subcutaneous tissues of Wistar strain rats. The specimens were removed at 1, 2, 3, 5, 7, 10, and 14 days after implantation. Total RNA was extracted from all specimens, and mRNAs of osteopontin, osteonectin, osteocalcin, and type I collagen were evaluated by Northern blotting. The gene expression of bone matrix proteins was found to be in accordance with the bone formation process.
- 18. Electron microscopic study of ectopic bone formation induced by BMP

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Partially purified bovine bone morphogenetic protein (BMP) and type collagen as carrier were combined and lyophilized. The freeze-dried pellets were implanted into the dorsal subcutaneous tissues of 3 week old Wistar strain rats. The specimens were removed at various periods and embedded in Epon 812. The none-decalcified specimens were observed by elec-