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STUDY OF THE FLUORINE AND BORON-10 CONTAINING COMPOUNDS TOWARD MRI AND **12 pt. bold ALL CAPS**

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Yoshihide Hattori¹, Yoshihiro Yamaguchi¹, Hitoshi Yamamoto², Tomohiro Asano³, Mitsunori Kiriha⁴, Underline for a speaker Teaki Wakamiya¹

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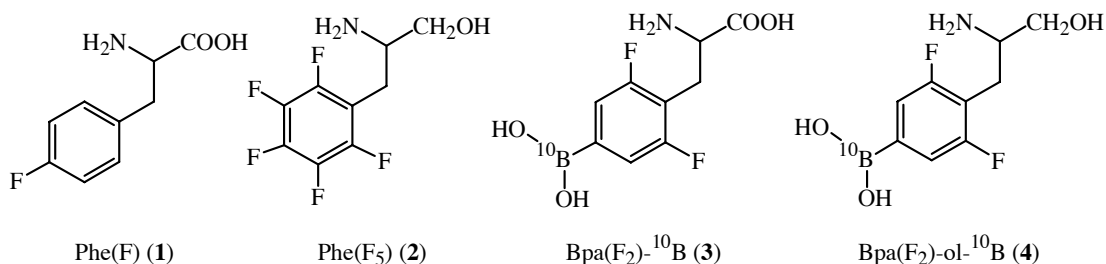
¹Faculty of Science and Technology, Kinki University, Higashi-osaka, Osaka 577-8502, Japan, ²Graduate School of Science, Osaka University, Toyonaka, Osaka 560-0043, Japan, ³College of Agriculture, Univer **12 pt. (18 pt. single-spaced)** Osaka 599-8231, Japan, ⁴Department of Neurosurgery, Aino Junior College Hospital, Ibaraki, Osaka 567-0018, Japan

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Dipeptides containing 3-(4-fluorophenyl)alanine [Phe(F)] (1) seem to be transferred into some kinds of tumor cells through the oligopeptide transporter. Furthermore, in a previous study **No subtitles** containing 3-(2,3,4,5,6-pentafluorophenyl)alanine [Phe(F₅)] (2) was certified to be detectable by ¹⁹F NMR up to μM order concentration. These facts suggest that magnetic resonance imaging (M **12 pt. typeface with 1.5-line spacing (18 pt. fixed).** of the Phe(F₅)-containing peptides internalized into the tumor cells may be acceptable as a promising means for diagnosis of cancer.

From the standpoint of the treatment of brain cancer or melanoma, the boron neutron capture therapy (BNCT) based on the interaction of ¹⁰B isotope and neutron has been highly noted in recent years [1]. In **SPECIMEN** and BNCT, we designed and synthesized the m and boron-10 atoms such as 3-(4-borono **[B]** (3) and 3-(4-borono-2,6-difluorophenyl)alaninol {[Bpa(F₂)-¹⁰B]-ol} (4). In the present paper we focus on ¹⁹F NMR measurement and tumor cell killing effect of various compounds containing both fluorine and boron-10 atoms.

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[1] Soloway, H. A., Tjarks, W., Barnum A. B., Ronø F. Barth F. R., Codogni, M. I., Wilson, J. G. (1998) *Chem. Rev.*, **98**, 1515-1527. **12 pt. (18 pt. single-spaced)**

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