

## 研究成果の刊行に関する一覧表

## 書籍

著者氏名	論文タイトル名	書籍全体の編集者名	書籍名	出版社名	出版地	出版年	ページ
呉 繁夫	「非ケトーシス型高グリシン血症」		先天異常症候群辞典	日本臨床社		印刷中	

## 雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Kure S, Kanako K, Kudo T, Kanno K, Aoki Y, Suzuki Y, Shinka T, Sakata Y, Narisawa K, and Matsubara Y.	Chromosomal localization, structure, single nucleotide polymorphisms, and expression of human H-protein gene of the glycine cleavage system ( <i>GCSH</i> ), a candidate gene for nonketotic hyperglycinemia.	J Hum Genet			In press
Takayanagi M, Kure S, Tada K, Sakata Y, Kurihara Y, Ohya Y, Kajita M, Tada K, Matsubara Y, Narisawa K.	Human glycine decarboxylase ( <i>GLCD</i> ) gene and its processed type pseudogene: Structures, expression, and identification of a large deletion in a family with nonketotic hyperglycinemia.	Human Genet	106	298-305.	2000
Kure S, Hou DC, Suzuki Y, Yamagishi A, Hiratsuka M, Fukuda T, Sugie H, Kondo N, Matsubara Y, Narisawa K.	Glycogen storage disease type Ib without neutropenia.	J Pediatr	137	253-256	2000

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Kudo T, Ikeda K, Kure S, Matsubara Y, Oshima T, Kawase T, Narisawa K, Takasaka T.	New common mutation in the Connexin 26 gene (GJB2) in childhood deafness in the Japanese population.	Am J Med Genet	90	141-145	2000
Fujii K, Matsubara Y, Akanuma J, Takahashi K, Kure S, Suzuki Y, Imaizumi M, Iinuma K, Sakatsme O, Rinaldo P, Kuniaki Narisawa.	Mutation detection by TaqMan-allele specific amplification: its application to the molecular diagnosis of glycogen storage disease type Ia and medium-chain acyl-CoA dehydrogenase deficiency.	Hum Mutaion	15	189-196.	2000
Mizugaki M, Hiratsuka M, Agatsuma Y, Matsubara Y, Fujii K, Kure S, Narisawa K.	Rapid detection of CYP2C18 genotypes by real-time fluorescence polymerase chain reaction.	J Pharm Pharmacol	52	199-205.	2000
Akanuma J, Nishigaki T, Matsubara Y, Fujii K, Takahashi K, Kure S, Suzuki Y, Ohura T, Miyabayashi S, Oagawa E, Iinuma K, Inui K, Okada S, Narisawa K.	Molecular diagnosis of 50 Japanese patients with glycogen storage disease type Ia: three mutations in the glucose-6-phosphatase gene account for 96% of mutant alleles.	Am J Med Genet	91	107-112	2000
Takahashi K, Akanuma J, Matsubara Y, Fujii K, Kure S, Suzuki Y, Wataya K, Sakamoto O, Aoki Y, Ohura T, Miyabayashi S, Narisawa K.	Heterogeneity of mutations in the glucose-6-phosphatase gene in Japanese patients with glycogen storage disease type Ia.	Am J Med Genet	92	90-94	2000

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Senoo M, Matsubara Y, Fujii K, Nagasaki Y, Hiratsuka M, Kure S, Uehara S, Okamura K, Narisawa K.	Adenovirus-mediated in utero gene transfer in mice and guinea pigs: tissue distribution of recombinant adenovirus determined by quantitative TaqMan PCR assay.	Mol Genet Metab	69	269-276	2000
Yong X, Aoki Y, Li X, Sakamoto O, Hiratsuka M, Gibson KM, Kure S, Narisawa K, Matsubara Y, Suzuki Y.	Haplotype analysis suggests that the two predominant mutations in Japanese patients with holocarboxylase synthetase deficiency are founder mutations.	J Hum Genet	45	358-362	2000

## 別紙 5

## 研究成果の刊行に関する一覧表

## 書籍

著者氏名	論文タイトル名	書籍全体の 編集者名	書 籍 名	出版社名	出版地	出版年	ページ
Tanaka T, Hashizume K, Kunimoto M, Maeda T, Hodozuka A, Nakai H.	Multiple subpial transection versus callosal section in the treatment of experimentally induced cortical focal seizures In	H O Luders and Y Comair	Epilepsy Surgery, 2 <sup>nd</sup> Edition	Lippincott Williams &Wilkins	Maryland	2001	801-806

## 雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Tanaka T, Hashizume K, Sawamura A, Yoshida K, Takano k, Hodozuka A, Nakai H.	Influence of basal ganglia upon seizure propagation of the kainic acid-induced focal epilepsy	Epilepsia	41(Suppl 7)	26-27	2000
Hashizume K, Kiriyaama K, Kunimoto M, Maeda T, Tanaka T, Miyamoto A, Miyokawa N, Fukuhara M	Correlation of EEG, neuroimaging and histopathology in an epilepsy patient with diffuse cortical dysplasia	Child's Nervous System	16(2)	75-79	2000
Hashizume, K, Tanaka T	Antiepileptic effect of nefiracetam on kainic acid-induced limbic seizure in rats	Epilepsy Research	39(3)	221-228	2000
Hashizume, K, Tanaka T, T. Fujita, S. Tanaka	Generalized seizures induced by an epileptic focus in the mesencephalic reticular formation: Impact on the understanding of the generalizing mechanism	Stereotact Funct Neurosurg	74	153-160	2000

## 研究成果の刊行に関する一覧表

## 書籍

著者氏名	論文タイトル名	書籍全体の 編集者名	書 籍 名	出版社名	出版地	出版年	ページ

## 雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
M. Kawamoto, K. Ohno, Kuriyama, T. Kubo, and K. Sato	Developmental changes in GABA transporter (GAT1, GAT3) mRNA expressions in the rat olfactory bulb.	Dev. Brain Res	126	137-145	2001
M. Utsumi, K. Ohno, H. Onchi, K. Sato and M. Tohyama	Differential expression patterns of three glutamate transporters (GLAST, GLT1 and EAAC1) in the rat main olfactory bulb	Mol. Brain Res.	in press		
C. Kanaka, K. Ohno, A. Okabe, K. Kuriyama, T. Itoh, A. Fukuda, and K. Sato	The differential expression patterns of messenger RNAs encoding K-C1 cotransporters (KCC1,2) and Na-K-2Cl cotransporter (NKCC1) in the rat nervous system.	Neuroscience	in press		