

神戸大学大学院医学研究科 神経内科学分野 業績集 2017

さらに詳細は[分子脳科学のHP](#)も参照して下さい

原著論文：

2017 年 英文

WNK1/HSN2 founder mutation in patients with hereditary sensory and autonomic neuropathy: A Japanese cohort study.

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Xeroderma pigmentosum clinical practice guidelines revision committee.

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The Ror1 receptor tyrosine kinase plays a critical role in regulating satellite cell proliferation during regeneration of injured muscle.

Kamizaki K, Doi R, Hayashi M, Saji T, Kanagawa M, Toda T, Fukada SI, Ho HH, Greenberg ME, Endo M, Minami Y.

J Biol Chem. 2017; 292(38): 15939-51.

Corrigendum: Novel missense mutation in DLL4 in a Japanese sporadic case of Adams-Oliver syndrome.

Nagasaka M, Taniguchi-Ikeda M, Inagaki H, Ouchi Y, Kurokawa D, Yamana K, Harada R, Nozu K, Sakai Y, Mishra SK, Yamaguchi Y, Morioka I, Toda T, Kurahashi H, Iijima K.

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Novel missense mutation in DLL4 in a Japanese sporadic case of Adams-Oliver syndrome.

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総説、解説：

2017 年 和文

【パーキンソン病の新展開-発症の分子機構と新規治療】 分子機構解明の新しい展開 ゲノムワイド関連解析 (GWAS)からの新展開

戸田 達史, 佐竹 渉

医学のあゆみ 262 巻 6 号 Page 617-621 (2017.08)

【認知症 1,000 万人時代を目前に控えて-最新の診断,マネジメント,そして分子標的治療へ】 認知症診療の最前線 兵庫県ならびに神戸市における認知症診療体制の現状

古和 久朋

内科 120 巻 2 号 Page 225-228 (2017.08)

神経筋電気診断 Case of the Issue(No.12) 「首下がり症候群」への筋電図によるアプローチ 体幹部の筋電図検査の重要性について

関口 兼司

臨床神経生理学 45 巻 4 号 Page 190-197 (2017.08)

【最新遺伝医学研究と遺伝カウンセリング シリーズ 2 最新精神・神経遺伝医学研究と遺伝カウンセリング】 (第 1 章)総論 孤発性疾患のリスク遺伝子の発見 ゲノムワイド関連解析の現状、進化と今後

佐竹 渉, 戸田 達史

遺伝子医学 MOOK 別冊最新精神・神経遺伝医学研究と遺伝カウンセリング Page 39-44 (2017.04)

【パーキンソン病-基礎・臨床の最新情報-】 総論 パーキンソン病の原因遺伝子

戸田 達史

日本臨床 75 巻 1 号 Page 21-27 (2017.01)