

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
東元健、副 島英伸	Beckwith-Wiedemann症候群と小児腫瘍	監修：佐々木裕之、編集：中尾光善、中島欽一	遺伝子医学MOOK25 エピジェネティクスと病気	メディカルドゥ	大阪	2013	195-201

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
Court F, Tayama C, Romanelli V, Martin-Trujillo A, Iglesias-Platas I, Okamura K, Sugahara N, Simón C, Moore H, Harness JV, Keirstead H, Sanchez-Mut JV, Kaneki E, Lapunzina P, Soejima H, Wake N, Esteller M, Ogata T, Hata K, Nakabayashi K, Monk D.	Genome-wide parent-of-origin DNA methylation analysis reveals the intricacies of the human imprintsome and suggests a germline methylation independent establishment of imprinting.	Genome Res	24(4)	554-69	2014
Ohnishi K, Semi K, Yamamoto T, Shimizu M, Tanaka A, Mitsunaga K, Okita K, Osafune K, Arioka Y, Maeda T, Soejima H, Moriwaki H, Yamamoto S, Woltjen K, Yamada Y.	Premature termination of reprogramming <i>in vivo</i> leads to cancer development through altered epigenetic regulation.	Cell	156(4)	663–677	2014
Higashimoto K, Joza K, Kosho T, Matsubara K, Fuke T, Yamada D, Yatsuki H, Maeda T, Ohtsuka Y, Nishioka K, Joh K, Koseki H, Ogata T, Soejima H.	A novel de novo point mutation of the OCT-binding site in the <i>IGF2/H19</i> -imprinting control region in a Beckwith-Wiedemann syndrome patient.	Clin Genet			published online:4 December, 2013
Rumbajan JM, Maeda T, Souzaki R, Mitsuui K, Higashimoto K, Nakabayashi K, Yatsuki H, Nishioka K, Harada R, Aoki S, Kohashi K, Oda Y, Hata K, Saji T, Taguchi T, Tajiri T, Soejima H, Joh K.	Comprehensive analyses of imprinted differentially methylated regions reveal epigenetic and genetic characteristics in hepatoblastoma.	BMC Cancer	13	608	2013

Miyazaki H, Higashimoto K, Yada Y, A. Endo T, Sharif J, Komori T, Matsuda M, Koseki Y, Nakayama M, Soejima H, Handa H, Koseki H, Hirose S, Nishioka K.	Ash1l methylates Lys36 of histone H3 independently of transcriptional elongation to counteract Polycomb silencing.	PLoS Genet	9(11)	e1003897	2013
Soejima H, Higashimoto K.	Epigenetic and genetic alterations of the imprinting disorder Beckwith-Wiedemann syndrome and related disorders.	J Hum Genet	58(7)	402-409	2013
大場隆、片渕秀隆、副島英伸。	間葉性異形成胎盤Placental mesenchymal dysplasia (PMD)の診断と原因遺伝子	病理と臨床	32(5)	535-540	2014
前田寿幸、東元健、副島英伸。	Beckwith-Wiedemann症候群とSilver-Russell症候群。	小児科臨床	66(増刊号)	1308-1314	2013
Yamaguchi M, Tashiro H, Motohara K, Ohba T, Katabuchi H.	Primary strumal carcinoid tumor of the ovary: A pregnant patient exhibiting severe constipation and CEA elevation.	Gynecol Oncol	4	9-12	2013
大場隆、坂口勲、片渕秀隆。	「プロメテウス 婦人科がん最新医療」絨毛性疾患 3.絨毛性疾患の診断 (2)画像診断	産婦人科の実際	62(12)	2027-2031	2013