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Publications (3)

Erlic Z, Fottner C, Forrer F, Plöckinger U, Petersenn S, Zabolotny D, Kollukch O, Yaremchuk S, Januszewicz A, Walz M, Eng C, Neumann H, Klose S, Opocher G, Rybicki L, Peçzkowska M, Golcher H, Kann P, Brauckhoff M, Müssig K, Muresan M, Schäffler A, Reisch N, Schott M, Fassnacht M, European-American Pheochromocytoma Study Group. Clinical predictors and algorithm for the genetic diagnosis of pheochromocytoma patients. *Clin Cancer Res* 2009; 15:6378-85.

Schiavi F, Välimäki M, Kawecki A, Szutkowski Z, Schipper J, Walz M, Pigny P, Bauters C, Willet-Brozick J, Baysal B, Januszewicz A, Eng C, Opocher G, Neumann H, Forrer F, Walter M, Boedeker C, Bausch B, Peçzkowska M, Gomez C, Strassburg T, Pawlu C, Buchta M, Salzmann M, Hoffmann M, Berlis A, Brink I, Cybulla M, Muresan M, European-American Paraganglioma Study Group. Predictors and prevalence of paraganglioma syndrome associated with mutations of the SDHC gene. *JAMA* 2005; 294:2057-63.

Neumann H, Apel T, Treier M, Reineke M, Walz M, Hoang-Vu C, Brauckhoff M, Klein-Franke A, Klose P, Schmidt H, Maier M, Peçzkowska M, Szmigielski C, Glaesker S, Manz T, Munk R, Bausch B, McWhinney S, Bender B, Gimm O, Franke G, Schipper J, Klisch J, Althoefer C, Zerres K, Januszewicz A, Eng C, Smith W, Freiburg-Warsaw-Columbus Pheochromocytoma Study Group. Germ-line mutations in nonsyndromic pheochromocytoma. *N Engl J Med* 2002; 346:1459-66.

Projects (0)

No results found.

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