



**Hartmut P H Neumann**

**Contact**

Hartmut P H Neumann

## Publications (4)

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.

Jörger M, Köberle D, Neumann H, Gillessen Sommer S. Von Hippel-Lindau disease--a rare disease important to recognize. *Onkologie* 2005; 28:159-63.

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.

---

Kantonsspital St.Gallen

Rorschacher Strasse 95

CH-9007 St.Gallen

T: +41 71 494 11 11

[support.forschung@kssg.ch](mailto:support.forschung@kssg.ch)