



Paul Komminoth

Kontakt

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Publikationen (7)

Savic S, Soltermann A, Zweifel R, Zettl A, von Gunten M, Singer G, Rössle M, Jasarevic Z, Letovanec I, McKee T, Komminoth P, Jochum W, Fleischmann A, Diebold J, Cathomas G, Eppenberger-Castori S, Berezowska S, Bubendorf L. PD-L1 testing of non-small cell lung cancer using different antibodies and platforms: a Swiss cross-validation study. *Virchows Arch* 2019; 475:67-76.

Zulewski H, Dettmer M, Minder A, Clerici T, Triponez F, Oertli D, Weidner S, Steinert H, Haldemann A, Christ E, Bilz S, Giovanella L, Komminoth P. Multidisciplinary approach for risk-oriented treatment of low-risk papillary thyroid cancer in Switzerland. *Swiss Med Wkly* 2019; 149:w14700.

Zhong Q, Diebold J, McKee T, Jochum W, Kashofer K, Hofman P, Zischka M, Moch H, Rechsteiner M, Rogel U, Vassella E, Wagner U, Kurt H, Molinari F, Cathomas G, Komminoth P, Barman-Aksözen J, Schneider-Yin X, Rey J, Wild P. Multi-laboratory proficiency testing of clinical cancer genomic profiling by next-generation sequencing. *Pathol Res Pract* 2018; 214:957-963.

Diebold M, Kohler M, Oezkan F, Darwiche K, Berezowska S, Theegarten D, Grobholz R, Jochum W, Komminoth P, Bubendorf L, Haile S, Hottinger S, Soltermann A, Franzen D. Prognostic value of MIB-1 proliferation index in solitary fibrous tumors of the pleura implemented in a new score - a multicenter study. *Respir Res* 2017; 18:210.

Collaud S, Tischler V, Atanassoff A, Wiedl T, Komminoth P, Öhlschlegel C, Weder W, Soltermann A. Lung neuroendocrine tumors: correlation of ubiquitinylation and sumoylation with nucleo-cytosolic partitioning of PTEN. *BMC cancer* 2015; 15:74.

Mihic-Probst D, Zhao J, Saremaslani P, Baer A, Öhlschlegel C, Paredes B, Komminoth P, Heitz P. CGH analysis shows genetic similarities and differences in atypical fibroxanthoma and undifferentiated high grade pleomorphic sarcoma. *Anticancer Res* 2004; 24:19-26.

Maier M, Schmid C, Galeazzi R, Krull I, Heitz P, Locher T, Schmid S, Saremaslani P, Komminoth P, Brändle M, Perren A. A novel succinate dehydrogenase subunit B gene mutation, H132P, causes familial malignant sympathetic extraadrenal paragangliomas. *The Journal of clinical endocrinology and metabolism* 2004; 89:362-7.

Projekte (0)

Keine Resultate gefunden.

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