Professor Neumann has six publications in **JAMA and journals of the JAMA group**: one in 1995, one in 2002, one in 2004, one in 2005, one in 2017, and one in 2019.

The publication in 1995 is from the time when the indications for genetic testing for carriership of germline mutations predisposing for MEN 2 was unclear.

In 2002 there was a comment published as a Letter to the Editor regarding biochemical analyses and imaging tests for the diagnosis of pheochromocytoma.

In 2004 a first study on prevalence and penetrance of germline mutations of the genes SDHB and SDHD in patients with pheochromocytoma and paraganglioma was published.

In 2005 a first study on prevalence and penetrance of germline mutations of the gene SDHC pheochromocytoma and paraganglioma was published.

In 2017 we characterized with colleagues from an international cooperation the phenotype of patients with mutations in the genes SDHA, TMEM127, MAX, and SDHAF2.

In 2019 a large scale international study on bilateral pheochromocytoma guided by Professor Neumann was published in JAMA Network Open.

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