

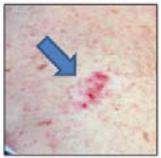
## PRESS RELEASE

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## SKIN CANCER: TO EACH TUMOR ITS PARTICULARITIES, TO EACH TUMOR ITS TREATMENT







Examples for Basal Cell Carcinoma (BCC).

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The skin cancer called basal cell carcinoma (BCC) is one of the most common types of cancer. 90 per cent of the population are at risk to develop it one day, notably because of their age and exposure to the sun UV-rays. Although common, this cancer is rarely fatal and has until now been little studied. But now geneticists at the University of Geneva (UNIGE), Switzerland, sequenced the DNA of these skin tumors, in order to determine the genes that are responsible for the cancerogenesis. Their discovery of new cancer genes that cause BCC can pave the way for new treatment methods personalized for each tumor. The results can be read in *Nature Genetics*.

Cancer is due to DNA modifications in a cell, which leads to uncontrolled cell growth. It is why, in the past few years, scientists have been using DNA sequencing of tumors more frequently to discover the genetic mutations responsible for causing cancer. Sergey Nikolaev from the UNIGE Faculty of Medicine focused on Basal Cell Carcinoma (BCC), a type of cancer that is very common in humans. Their goal was to sequence the genome of BCC tumors and compare it with the genetic profile of patients' healthy cells, to determine the genes responsible for the cancer development. To do this, they studied 293 tumors from a total of 236 patients of all ages.

## New cancer genes discovered

"It has been known for quite a while that a vast majority of BCCs are caused by abnormal activation of the so-called Sonic Hedgehog (SHh) pathway. What we did not know was that BCC is also the cancer with the highest mutation rates. In addition to SHh, numerous other genes can contribute to this skin cancer, which evidently complicates the treatment", explains Professeur Stylianos Antonarakis, who is heading the laboratory in which the research was conducted. The geneticists identified the genes MYCN, PTPN14 and LATS1 as additional drivers for the development of BCC. Their findings were confirmed through studies with patients who suffer from the so-called Gorlin syndrome and therefore possess a molecular predisposition to develop BCCs early in life. The DNA of their tumors also contained those additional cancer drivers.

But how could the geneticists from UNIGE identify those new

cancer genes? "We have developed a sophisticated statistical analysis software that permits us to localize the cancerous genes among the thousands of somatic mutations in the cancer DNA sequence", answers Sergey Nikolaev. "Once identified, we further tested their role as cancer drivers, for example by looking at molecules with and without mutations and comparing their structure and function", he adds. This was possible through the collaboration with research groups from the School of Medicine at New York University (USA), the department of Biochemistry of the University of Würzburg (Germany) and the department of Oncology of the University of Lausanne (Switzerland).

## A personalised treatment for the patient

Each tumor is rather unique because of the various genetic mutations of the genes involved. "We suggest to sequence the healthy DNA and that of the tumor of each patient to really know which genes are responsible for the development of cancer", explains Nikolaev. This precise diagnostics could improve patient care. "Cells respond differently to different ways of treatment because of the mutations of their genes. Some develop a resistance to certain drug substances, which renders the medication effect less. The additional mutations we detected could help in getting the best possible treatment results", adds Antonarakis. Even though still marginally implemented, the sequencing of tumors gradually enters into medical practice, taking into account the genetic particularities of each sick person. "The Genetic Medicine of the University Hospitals of Geneva has all the necessary infrastructure and expertise to spearhead such developments", concludes Antonarakis.

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