

# 'Game-changing' immunotherapy doubles head and neck cancer survival

October 9 2016

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Killer T cells surround a cancer cell. Credit: NIH

An immunotherapy drug has been hailed as a potential 'game changer' after being found to greatly improve survival for patients with relapsed head and neck cancer - a disease which is notoriously difficult to treat.

Nivolumab became the first treatment to extend survival in a phase III clinical trial for [patients](#) with head and [neck cancer](#) in whom [chemotherapy](#) had failed - and it did so with fewer side-effects than existing therapeutic options.

More than double the number of patients taking nivolumab were alive after one year as those treated with chemotherapy, reported the major international trial, published today (Sunday) in the *New England Journal of Medicine*.

There are currently no other [treatment options](#) that improve the survival of patients with cisplatin-resistant relapsed or metastatic head and neck cancers. This group of patients are expected to live less than six months.

The trial was led in the UK by Professor Kevin Harrington of The Institute of Cancer Research, London, and The Royal Marsden NHS Foundation Trust, and involved 20 research organisations from around the world. It was funded by Bristol Myers Squibb.

Of the 361 patients in the trial, 240 with relapsed or metastatic head and neck cancer were allocated to receive nivolumab and 121 to one of three different chemotherapies. UK patients received the chemotherapy drug docetaxel, which is the only treatment approved for advanced head and neck cancer by NICE.

After one year of the study, 36 per cent of patients treated with nivolumab were still alive compared with 17 per cent for the comparator arm.

Median survival for patients on nivolumab was 7.5 months, compared with 5.1 months for chemotherapy.

The survival benefit was more pronounced in patients whose tumours

had tested positive for human papillomavirus (HPV). These patients survived an average of 9.1 months with nivolumab and 4.4 months with chemotherapy.

HPV-negative patients survived an average of 7.5 months with nivolumab and 5.8 with chemotherapy.

Importantly, fewer patients experienced serious side-effects from taking nivolumab than with conventional treatment - only 13 per cent compared with 35 per cent of patients who received chemotherapy.

Patients given chemotherapy reported feeling physically, socially and emotionally worse off, whereas those who were given nivolumab remained stable during the course of treatment.

Professor Harrington will be presenting some of the findings at the European Society for Medical Oncology 2016 Congress in Copenhagen, simultaneously with publication.

Nivolumab will still have to go through approval by the European Medicines Agency and NICE before it is available for head and neck cancer patients on the NHS.

UK trial lead Professor Kevin Harrington, Professor of Biological Cancer Therapies at The Institute of Cancer Research, London, and Consultant at The Royal Marsden NHS Foundation Trust, said:

"Nivolumab could be a real game-changer for patients with advanced head and neck cancer. This trial found that it can greatly extend life among a group of patients who have no existing treatment options, without worsening quality of life.

"Once it has relapsed or spread, head and neck cancer is extremely

difficult to treat. So it's great news that these results indicate we now have a new treatment that can significantly extend life, and I'm keen to see it enter the clinic as soon as possible."

Professor Paul Workman, Chief Executive of The Institute of Cancer Research, London, said:

"Nivolumab is one of a new wave of immunotherapies that are beginning to have an impact across cancer treatment. This phase III clinical trial expands the repertoire of nivolumab even further, showing that it is the first treatment to have significant benefits in relapsed head and neck cancer.

"We hope regulators can work with the manufacturer to avoid delays in getting this drug to patients who have no effective treatment options left to them."

Provided by Institute of Cancer Research

Citation: 'Game-changing' immunotherapy doubles head and neck cancer survival (2016, October 9) retrieved 5 May 2023 from <https://medicalxpress.com/news/2016-10-game-changing-immunotherapy-neck-cancer-survival.html>

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