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Antisense oligonucleotide treatment for human astrocytoma

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Active antisense oligonucleotides directed against the insulin-like growth factor type I receptor (IGF-IR/AS ODN) have shown potential as an antitumour agent in animal studies. In the April issue of [Journal of Clinical Oncology](#) David Andrews and colleagues from [Thomas Jefferson University Hospital](#), Philadelphia, describe their novel implantation of IGF-IR/AS ODN-treated cells in patients with astrocytoma.

Andrews *et al.* studied 12 patients treated by surgery for malignant astrocytoma. Glioma cells collected at surgery were treated *ex vivo* with an IGF-IR/AS ODN and reimplanted in the rectus sheath within 24 hours of craniotomy. Examination of the IGF-IR/AS ODN-treated cells after a 24-hour *in situ* incubation showed that only 2% were intact, and none of the intact cells was viable in culture thereafter. IGF-IR was down regulated to 10% after *ex vivo* antisense treatment (*J Clin Oncol* 2001, **19**:2189-2200).

In addition, clinical and radiographic improvements were observed in eight of 12 patients. This suggests that *ex vivo* IGF-IR/AS ODN treatment of autologous glioma cells can induce apoptosis and an *in vivo* host response without side effects.

References

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2. Thomas Jefferson University Hospital, [<http://www.jeffersonhospital.org/show.asp?durki=3858>]