

GROUNDBREAKING NEW IMMUNOTHERAPY FOR CANINE OSTEOSARCOMA

ELIAS Cancer Immunotherapy (ECI[®]) is the only combination adoptive T cell therapy available in veterinary medicine



WHAT IS IT?

ECI[®] is a vaccine primed form of adoptive T cell therapy, which has been developed for the treatment of canine osteosarcoma and, potentially, other types of canine cancers. Instead of genetically manipulating T cells or harvesting these cells directly from tumor tissue, ECI[®] uses autologous attenuated cancer cell vaccines manufactured from surgically removed host cancer tissue. These autologous vaccines are used to condition host T cells to the cancer antigens. The vaccine-primed mononuclear cells are later harvested via apheresis, and expanded and activated *ex vivo* prior to manufacture a killer T cell infusion that targets cancer cells and causes apoptosis.

ECI[®] STUDY RESULTS

Study	ECI-OSA-01 (completed) <i>n</i> =14	ECI-OSA-04 (ongoing) <i>n</i> =100
Safety	<p>Reported AEs were generally mild to moderate:</p> <ul style="list-style-type: none"> • Vaccine: 14 reports of Grade 1 injection site reactions; one report of Grade 1 vomiting • T cell infusion: 7 reports of Grades 1-2, and one manageable Grade 3 pyrexia (fever) <p>Treatment found to be safe and tolerable</p>	<ul style="list-style-type: none"> • Multi-center, randomized trial • Control group is carboplatin • Results expected in late 2022
Efficacy	<p>Outcomes observed:</p> <ul style="list-style-type: none"> • CR and PR with remission of metastatic disease • Slow DP progression in some patients <ul style="list-style-type: none"> ◦ Median Survival Time = 415 days ◦ 5 dogs alive 719-946 days at study end ◦ 4 of those 5 with no evidence of disease 	



How will you make a difference for your canine patients?

Ideal candidates for ECI® are dogs with newly diagnosed appendicular osteosarcoma (limb intact) that have no signs of metastatic disease at diagnosis

RESULTS OBSERVED IN HUMANS

The protocol for canine osteosarcoma is adapted from the human version of this immunotherapy that was used in Phase 1/2 clinical trials. It has demonstrated efficacy in human Phase 2 trials against stage IV renal carcinoma and recurrent malignant melanoma.

Exciting Progress

In April 2020, this immunotherapy was given FastTrack designation by the FDA for development in humans. ELIAS' canine osteosarcoma data was a key component in achieving this milestone.

FIND OUT HOW TO BRING ECI® TO YOUR PATIENTS



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ECI® IS AVAILABLE FOR CANINE OSTEOSARCOMA PATIENTS NOW

ECI® is authorized by the USDA for sale to licensed veterinarians prior to full approval.

- Administered to more than 200 dogs through clinical trials or commercial use
- Available in 23 states and expanding; authorized in all 50 states and District of Columbia
- 17 apheresis centers participating (and expanding)

Contact ELIAS Animal Health to learn how your hospital can offer ECI® and bring new hope to your patients and their families.

ECI® is an experimental autologous prescription product available for sale to veterinarians under 9 CFR 103.3. Safety and efficacy have not been established.

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