

## **Clinical Trial Now Recruiting Dogs With Histiocytic Sarcoma**

Last August the Morris Animal Foundation announced a new clinical trial **funded by the BMDCA** to test the effectiveness of a new treatment - Trametinib - for histiocytic sarcoma:

<https://www.morrisanimalfoundation.org/article/new-clinical-trial-assess-canine-cancer-treatment-histiocytic-sarcoma>

**That trial is now accepting patients at the following four universities.** Each has websites describing what dogs may participate, and what dogs are not eligible.

- University of Florida

<https://research.vetmed.ufl.edu/phase-ii-open-label-non-randomized-multicenter-clinical-trial-of-trametinib-for-dogs-with-histiocytic-sarcoma/>

- Virginia Tech

<https://research.vetmed.vt.edu/clinical-trials/current-studies/trametinib.html>

- Michigan State University

<https://cvm.msu.edu/hospital/veterinarians/clinical-trials/phase-ii-open-label-non-randomized-multicenter-clinical-trial-of-trametinib-for-dogs-with-histiocytic-sarcoma>

- University of Wisconsin *website to be available soon.*

The drug and physical exam rechecks and any routine tests or imaging needed at those recheck visits are free of cost and must be completed at the trial site.

### **Eligibility criteria:**

- \* Dogs of any age, weight, sex, or breed with a cytological or histological diagnosis of histiocytic sarcoma
- \* Dogs with measurable tumors (primary or metastatic)
- \* Dogs with an estimated life expectancy of >4 weeks
- \* Participation in the study will be offered when conventional therapy is no longer effective or is declined by the dog's client-caregiver.
- \* Signed informed consent and consent to necropsy will be obtained at time of enrollment.
- \* No prior chemotherapy within a minimum 2-week period
- \* No prior radiation therapy within a minimum 1-month period
- \* Dogs with adequate organ and marrow function and without serious systemic disorders that could compromise the patient's health
- \* Dogs with a prior or concurrent cancer whose prognosis or treatment does not have the potential to interfere with the safety or efficacy assessment of the investigational regimen

**Pat Long,  
BMDCA Health Committee Chair**