Table S8 – Results of proportional hazards regression analysis of secondary endpoints for detection of differences between treatment groups in the FAS and PPS2 populations

Endpoint	FAS			PPS2		
Endpoint	Hazard ratio	95% CI	P	Hazard ratio	95% CI	P
Time to study withdrawal						
All	0.96	0.54 – 1.71	.89	0.82	0.41 – 1.64	.57
LVOTO	3.14	1.13 – 8.73	**.028***	1.82	0.48 – 6.80	.39
No LVOTO	0.55	0.28 – 1.10	.089	0.63	0.29 – 1.38	.25
Time to cardiac death, euthanasia due to cardiac disease, and treatment failure						
All	0.87	0.35 – 2.18	.77	0.92	0.32 – 2.59	.87
LVOTO	2.04	0.29 – 14.49	.48	2.28	0.14 – 36.48	.56
No LVOTO	0.70	0.25 – 1.95	.50	.80	0.27 – 2.39	.69
Time to 1st escalation of furosemide dose						
All	0.77	0.39 – 1.52	.45	0.80	0.37 – 1.72	.56
LVOTO	1.78	0.54 – 5.87	.35	1.82	0.48 – 6.80	0.38
No LVOTO	0.56	0.25 – 1.22	.14	0.57	0.23 – 1.37	0.21
Time to furosemide >10 mg/kg/d						
All	1.25	0.33 – 4.75	.75	0.79	0.19 – 3.23	.74
LVOTO	n.d.	n.d.	n.d	n.d.	n.d.	n.d.
No LVOTO	0.61	0.14 – 2.73	.52	0.51	0.11 – 2.31	.39

Notes: *, includes 3 cats that were withdrawn from the study due to worsening of LVOTO on D1. Due to low number of observations (1 in each treatment group) data on time to arterial thromboembolism are not displayed.

Abbreviations: PPS2, per protocol set 2; CI, confidence interval; LVOTO, left ventricular outflow tract obstruction; n.d., not determined due to the low number of observations.