

### **1. ABSTRACT**

The descending contralateral movement detector (DCMD) is part of a neuronal pathway that is specifically tuned to small looming objects<sup>1</sup> and has lateral projections that synapse with motor neurons involved in collision avoidance<sup>2</sup>. DCMD responses are typically described in terms of rate coding<sup>1,3,4</sup>, yet responses to looming often display observable oscillations in mean firing rates and tight clustering of spikes in raw traces; an indication of the presence of bursting. We tested 20 locusts with 30 looming stimuli known to generate behavioural responses. We found frequent and shorter inter-spike intervals (ISIs) ranged from 1-8ms, while longer less frequent ISIs ranged from 40-50ms. A subsequent burst analysis revealed interburst frequencies of ~25Hz (within the range of the wingbeat frequency of a flying locust<sup>5,6</sup>). We propose that the DCMD employs a bimodal coding strategy to relay information regarding looming objects.



Fig 1. After making a relatively small incision exposing the ventral nerve cords, locusts (Locusta migratoria) were mounted onto a fixed platform within a flight simulator (A). Stimuli (7 cm black disc travelling at 300 cm/s aligned with the center of the right eye) were projected onto the dome screen (B) and neural recordings were taken from exposed underlying paired connectives of the ventral nerve cord anterior to the prothoracic ganglia (C and Fig. 2A); n=20 locusts and n=30 presentations.

3. LOOMING RESPONSES		
Α	B	
Fig. 2. DCMD responses to looming stimuli from previous work show evidence of bursting (A <sup>3</sup> , B <sup>8</sup> , and C <sup>9</sup> ).		

# Sensory bursts in a single motion sensitive pathway of the locust Glyn A. McMillan and John R. Gray



Fig 3. A) Visual information is transmitted to the DCMD in a 1:1 spike ratio via the lobula giant movement detector (LGMD, A). B) Extracellular recording aligned to the perceived time of collision (TOC). DCMD spikes were isolated based on threshold analysis. Increases in the stimulus subtense angle causes an increase in DCMD firing frequency that peaks before the time of collision (TOC). DCMD spike times were used for analysis and bursts (red squares in B) were identified using a burst assay (subset of data from above trace).



Fig. 4. Top) Vertical banding patterns are present in this TOC-aligned perievent raster plot (grey) representing 600 individual responses and mirrored by distinct oscillations in the mean DCMD firing rate (peristimulus time histogram (PSTH) - black line). Bottom) PSTHs show that while DCMD bursts (red line) and single spikes (blue line) increase up to 200ms before TOC, bursting dominates the response leading up to TOC.

6. ISI HISTOGRAMS AND AUTOCORRELATIONSSHOW BIMODAL DISTRIBUTIONS



- (a) Max. interval to start/maintain burst (8ms) (b) Min. number of spikes in a burst (2) (c) Min. burst duration (2ms)
- (d) Min. inter-burst interval (9ms)



interval (ISI) return map (top) has a fan-shaped distribution with a tight cluster of points at the origin (red, intraburst interval) and two other clusters along the axes (grey and blue, interburst intervals); the more scattered clusters may be the ISIs between single spikes (black). Right) Reflecting the mean DCMD PSTH, ISIs shorten as the stimulus subtense angle (blue line) increases toward TOC (red vertical line). However, the relatively constant ISI distribution in short intervals reveals a bimodal distribution of ISIs.

Fig. 4. ISI histograms and autocorrelations of DCMD responses for mean response (left), DCMD bursts (middle), and single spikes (right). The bimodal distribution of ISIs (represented more clearly in the data leading up to 200ms before TOC) are indicative of a bursting neuron showing shorter ISIs from 1-8ms and a second group with longer ISIs around 40ms. The ISIs of the bursts was relatively unimodal and occurred around 40ms. These trends are reflected in the associated autocorrelation. Overall, there was no clear trend in the distribution of single spikes.

- (or 25Hz)
- locusts<sup>10</sup>
- behaviour.

Biol 86: 87–97, 1980b. 1992.

216:3369-3380, 2013. 19:1122-1141, 1999.

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# 7. CONCLUSIONS

Evidence of bursting (particularly up to 200ms before TOC) based on distribution of spikes and autocorrelations: peak burst ISI and autocorrelation occurred around 40ms

Results suggest a behavioural implication of bursting: previous studies have shown that low DCMD firing rates that occur around 200ms before TOC may trigger avoidance steering responses in rigidly tethered

Given that the average wingbeat frequency of a flying locust is ~25 beats/s, our findings provide evidence to drive future experiments to test if DCMD bursting may have a role in gating non-rhythmical sensory input (object motion) to coordinate rhythmical modulation of wing kinematics linked to avoidance

# **8. REFERENCES**

<sup>1</sup>Schlotterer GR. Response of the locust descending movement detector neuron to rapidly approaching and withdrawing visual stimuli. Can J Zool 55:1372-1376, 1977.

<sup>2</sup>Simmons PJ. Connections between a movement-detecting visual interneurone and flight motoneurones of a locust. J Exp

<sup>3</sup>Santer RD, Rind FC, Stafford R, Simmons PJ. The role of an identified looming-sensitive neuron in triggering a flying locust's escape. J Neurophysiol 95:3391-3400, 2006.

<sup>4</sup>Simmons PJ and Rind FC. Orthopteran DCMD neuron: a reevaluation of responses to moving objects. II. Critical cues for detecting approaching objects. J Neurophysiol 68:1667-1682,

<sup>5</sup>Hatsopoulos N, Gabbiani F, and Laurent G. Elementary computation of object approach by a wide-field visual neuron. Science 270:1000-1003, 1995.

<sup>6</sup>Robertson RM and Johnson AG. Retinal image size triggers obstacle avoidance in flying locusts. Naturwiss 80:176-178, 1993. <sup>7</sup>McMillan GA, Loessin V, and Gray JR. Bilateral flight muscle activity predicts wing kinematics and 3-dimensional body orientation of locusts responding to looming objects. J Exp Biol

<sup>8</sup>Gabbiani F, Krapp HG, and Laurent G. Computation of object approach by a wide-field motion-sensitive neuron. J Neurosci

<sup>9</sup>Money TGA, DeCarlo CA, and Robertson RM. Temperaturesensitive gating in a descending visual interneuron, DCMD. J Comp Phys [A] 192:915-925, 2006.

<sup>10</sup>Matheson T, Rogers SM, Krapp HG. Plasticitiy in the visual system is correlated with a change in lifestyle of solitarious and gregarious locusts. J Neurophysiol 91: 1-12, 2004.

## **9. ACKNOWLEDGEMENTS**