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Humanitarian Demining Using an Insect Based Chemical Unmanned Aerial Vehicle

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1. Introduction

Nowadays, there are about 100 million active landmines distributed around the world as the result of earlier conflicts (Habib 2007). These cheap and simple to manufacture weapons have a long lasting effect that cause injury to the civil population even decades after the conflict has ended. Their removal is very expensive, dangerous and time consuming, and it has a high social and economical impact. Hence, although demining is a necessity for the affected populations to recover, it poses fundamental technical challenges.

At the present time, the use of animals, i.e. biological sensors, still provides the highest accuracy and safety standards in demining tasks. A number of animal species including dogs, rats and bees have been successfully trained to detect and localize landmines following the minute chemical trails of leaking explosive compounds (Fjellanger et al. 2002; Bromenshenk et al. 2003; Verhagen et al. 2006). Insects, and in particular moths, are highly optimized chemical detection systems that are extremely proficient at the detection and localization of different chemical compounds, in particular pheromones, at very long distances, i.e. several hundreds of meters. Moths use pheromone signals for sexual communication and it has been shown that males are able to detect and distinguish minute amounts of female pheromones (as little as 10⁴ molecules cm⁻³) against a background of other chemicals in very irregular and unpredictable plumes (Wyatt 2003). As a consequence, the evolutionary pressure to detect pheromones has generated specific neural and behavioral adaptations to deal with this specific problem. Nonetheless, the chemical search task is not reduced to a unique olfactory process but is a multi-modal task that includes the integration of complex behavioral strategies with visual, olfactory and wind sensing information (Kennedy & Marsh 1974; Ludlow 1982; Charlton & Cardé. 1990).

In this chapter we analyze the relationship between the chemical detection and localization problem and its biological solution, and we will show how our understanding of the biological solution can be exploited to construct efficient autonomous chemo-sensing Unmanned Aerial Vehicles (cUAV). Firstly, we describe a blimp-based technology for a cUAV. Subsequently, we investigate the computational and behavioral principles underlying the opto-motor system of the fly and the locust, and we show that relying solely

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on vision, biologically constrained neuronal models of the fly visual system suffice for course stabilization and altitude control of a blimp-based UAV. Then, we augment this system with a collision avoidance model based on the Lobula Giant Movement Detector neuron of the Locust. A number of chemical search experiments are described with a mobile robot in a controlled wind tunnel environment. In these experiments, a number of chemical mapping strategies and behaviorally and biologically constrained models derived from the moth are tested and their performance is assessed. Finally, some of these solutions are evaluated in outdoor chemical detection, mapping and localization tasks using a cUAV. We show that our insect based approach that combines detailed biologically constrained models of the fly, locust and moth provides for a robust system and can constitute a viable approach towards the detection and localization of explosives and therefore be used for humanitarian demining. Moreover, our insect based cUAV demonstrates that the detailed understanding of biological solutions to real-world problems can provide for novel and robust artificial systems.

2. Technology

Sensor Technology

The effectiveness of the landmine detection process depends fundamentally on the sensor technology used and the target mine. The most commonly used sensors are metal detectors, electromagnetic, acoustic and seismic methods, and biological sensors (Bruschini & Gros 1998; Habib 2002; Gooneratne et al. 2004). However, the current technology is still too limited to deal with the great variety of mines available, and usually is rather specific to a particular kind of explosives or mines.

Aiming at mimicking the best sensors known so far, here we propose the use of an artificial nose sensor sensitive to a wide range of volatile compounds in combination with the current understanding of the best studied chemical detection system, the male moth (Pyk et al. 2006; Bermúdez i Badia et al. 2007a). The artificial nose sensor consists of a 6 grid array of broadly tuned thin film metal oxide chemo-sensors (Alpha MOS SA, France). The sensitivity of each of the individual sensors is controlled by variations in their dopants and semiconductor materials (Nanto & Stetter 2003; Pyk et al. 2006). These variations render variable binding properties of each sensor to chemical compounds and, hence, differential sensing capabilities. The interaction of the surface of each of the sensors with the odor molecules provokes a change in the bulk resistance of the semiconductor material, and this is then converted to voltage and measured (Nanto & Stetter 2003). To allow the release of bound compounds, the temperature of the surface of the sensor is regulated by an external circuit. Given that the compounds the sensor needs to detect are presented in the real-world in complex plume-like dynamics it is essential to also understand the dynamics of the sensor. We measured the time constants of the artificial nose sensor and showed a rise time of 2.0±0.77 s (mean±std, n=5) and a decay time of 3.1±0.84 s (mean±std, n=9) (Pyk et al. 2006). The low power consumption (of approx. 270 mW) and the lightweight and relatively high degree of miniaturization (2 x 3 x 0.38 mm) make it a suitable sensor for real-time use on an Unmanned Aerial Vehicle (UVA).

The different robotic platforms used in this project are equipped with lightweight and highresolution cameras (628 [H] x 582 [V] pixels) ("Module 3", Conrad Electronics, Germany) fitted with wide-angle lenses (2.5 mm lenses, Conrad Electronics, Switzerland). Subsequently, the obtained video stream from the cameras is broadcasted via compact PAL transmitters (SDX-21LP video transmitters on the 2.4 GHz band, produced by RF-Video, Canada). The robots use Lithium-Polymer rechargeable batteries (KOK 3270, Kokam, Kyunggi-do, Korea, www.kokam.com) that provide up to 5 times higher energy per unit of mass than regular Nickel Cadmium rechargeable batteries. On the ground station side, the processed by camera images are received and the neural simulator iar (http://iqr.sourceforge.net/) (Bernardet et al. 2002). iqr is a software for the graphical design and control of large-scale neuronal models, and their interfacing to real-world devices in real-time. Our large-scale insect based neural models communicate with the robots via a wireless radio link in the case of the indoor blimp (BIM433-F transceivers, Wireless World AG, Switzerland), and Bluetooth in the case of the mobile moth robot.

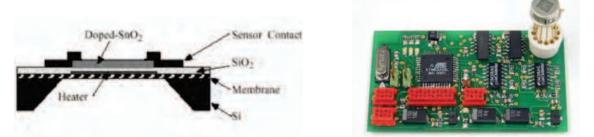


Fig. 1. The artificial nose sensor system. Left panel: schematic of each of the sensors of the 6 grid array including its main components. The size of a sensor is 0.18 x 0.2 mm (with x length), with an active area of 0.032 mm². Right panel: the full chemo-sensor package and its readout PCB. The sampling frequency of the PCB is 16.3 Hz, its dimensions 60 mm x 35 mm, and its weight 13.8 g. Adapted from Pyk et al. (2006)

The Robotic Platforms

Humanitarian demining clearly deviates from military demining in its objectives and approach. For humanitarian demining it is essential to locate and clean up every single mine in post-conflict areas for the recovery of the population. Seeking the highest accuracy and safety standards, many autonomous or remote controlled demining robots have been developed (Nicoud & Habib 1995; Nonami et al. 2000; Gonzalez de Santos et al. 2002; Marques et al. 2002; Santana & Barata 2005). Ideally, a robotic platform suitable to work in human non-accessible or unspecified environments with the highest degree of user safety is desired. Therefore, we propose the use of Unmanned Aerial Vehicle (UAV) technology since it provides a terrain independent solution and it is unable to detonate mines during inspection. In particular, the use of a blimp-based UAVs offers us a cheaper, more stable and easier to control solution than fixed-wing or helicopter like platforms. In addition, we need to consider that the sensing platform itself should not disturb the plume structure when it wants to localize the source.

We have developed a number of robots to approach the chemical localization problem from different angles. First, we constructed a blimp-based robot designed to work within indoor environments to study course control systems based on the understanding of the neuronal principles of insect visual navigation (Fig 2, left panel). This robot was constructed from a balsa wood structure, uses a lightweight Lithium-Polymer battery, and has independent control for altitude and translation (08GS - 8mm motors, API-Portescap, La Chaux-de-

Fonds, Switzerland, www.portescap.com) (see Bermúdez i Badia et al. (2007b) for further details).

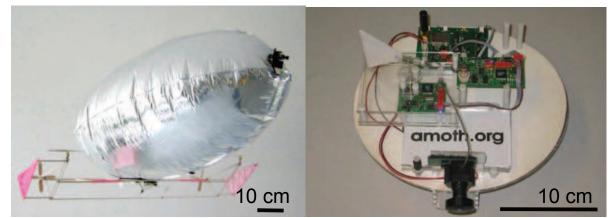


Fig. 2. The indoor artificial moth robots. Left panel: the blimp-based UAV consists of a 30 x 120 cm hull (radius x length) with a payload of about 250 g, a remote control board that provides with independent altitude and translation control, and 2 wide-angle wireless camera systems. Right panel: the ground chemosensory vehicle, of 20 cm diameter and 16 cm height, provided with, from top to bottom, a Bluetooth control board, a chemo-sensor, a wind direction sensor and a wireless camera equipped with a wide-angle lens. Right panel adapted from Pyk et al. (2006)

Subsequently, we developed a chemosensory ground vehicle as the first approach to study the odor mapping and localization problem in indoor and controlled environments (Fig 2, right panel). The robot is equipped with a chemo-sensor, a wind direction sensor and a wireless camera. The control board controls two motors, collects the sensory input from a chemo-sensor and a wind direction sensor and communicates with a ground station via a Bluetooth module. The wind direction detection sensor consists of a lightweight styrophore vane that is attached to a rotating shaft fitted with a small magnet. Then, a magnetometer is used to read out the orientation of the vane (MicroMag2, PNI Corporation, Santa Rosa, USA, www.pnicorp.com). All sensory boards are fitted with an ATmega32L microcontroller (ATmega32L, Atmel, San Jose, CA, USA, www.atmel.com) that makes local computations and interfaces the sensor with the robot infrastructure via a Two-Wire-Interface (TWI) bus. A Lithium-Polymer battery provides the robot with approximately 8 hours of autonomy. To conclude, we constructed a blimp-based outdoor chemo-sensing UAV (cUAV) to perform field experiments (Fig. 3). One of the advantages of using a blimp-based cUAV is that it can carry additional sensors for conventional control, monitoring and analysis such as a GPS, accelerometers, altimeters, cameras, etc. In this case, the cUAV has a control board that interfaces a GPS, a 3D compass, 2 altimeters and a chemo-sensory board via a common TWI bus. The cUAV exchanges sensor readings and motor commands with a ground station via a 2.4GHz communication system (AC1524 transceiver, Aerocomm, Lenexa, USA). The total weight of the electronics is 90 g. 8 Lithium Polymer cells with a capacity of 13 Ah provide up to 2 hours of run time in moderate wind speeds. Based on the sensor readings, a control layer was developed to allow for unsupervised cUAV operation (see Bermúdez i Badia et al. (2007a) for further details).

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Our custom developed cUAV consists of a PVC hull filled with helium (4.5m long, 1.2m diameter, 6m³ volume) with an approximate payload of 3 kg. Four independent DC motors are fixed on a modular and scalable carbon fiber frame of about 3 kg (Fig. 3, right panel). Each of the propellers fixed to a DC motor generates a thrust of approximately 620G. The resulting mass/power ratio turns the cUAV into a more unstable platform than conventional blimps, while faster in dynamic responses. To solve the instability problem, the motor frame is constructed around the hull as opposed to be attached exclusively to its lower part as in the case of conventional blimps (Fig. 3). The arrangement of the motors is such that the forces are applied directly at the center of mass of the cUAV, hence reducing the oscillatory behaviors that result from fast flight maneuvers. The distance between the motors used for rotation is significantly increased, meaining that greater torque forces can be generated by the same thrust. Furthermore, fins of about 0.2 m² are used as a passive stabilization mechanism. As a result, our design of the motor frame renders the cUAV more stable and with better maneuverability than standard off the shelf solutions.

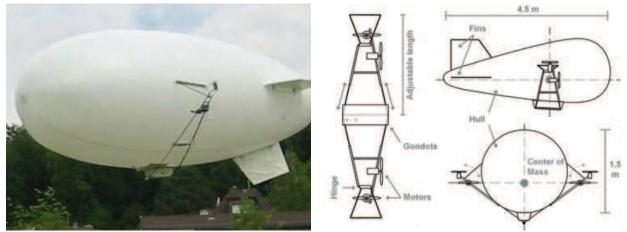


Fig. 3. The outdoor artificial moth robot. Left panel: the cUAV during autonomous flight.
Right panel: scheme of the cUAV. It includes 2 pairs of motors, a carbon fibre frame that builds around the hull of the blimp, and a hinge and a variable length mechanism that makes it adjustable to different hull sizes. Adapted from Bermúdez i Badia et al. (2007a)

The Wind Tunnel and the Odor Delivery System

In order to investigate the responses of our sensor technology to different chemical stimuli and a number of moth based localization algorithms, we constructed a wind tunnel suitable for mobile robot experiments. The wind tunnel measures $3 \times 4 \times 0.54$ m and was constructed out of wood and transparent plastic sheets. The front part of the wind tunnel was left open to let the air in. Four ventilators with adjustable wind speed were placed at the back to generate a negative pressure and create an air flow of about 0.7 m \cdot s⁻¹ from front to back, and to adjust for a uniform and symmetric velocity profile. Then, the air sucked out of the tunnel goes into an exhaust tunnel where it is removed.

As delivery system for the chemical substances we used an ultrasonic release system (Mist of Dreams, XrLight, Zhongshong City, China) that generates a rapidly evaporating mist at a rate of about 3.33 ml min⁻¹. The vaporizer was enclosed in a 40 x 30 cm chamber and the delivery from this chamber was controlled via an impeller (CGW/EDF-50, GrandWing Servo-tech C0, Ltd, Taiwan). Unless otherwise is specified, a solution of fixed concentration

of ethanol and distilled water (20% ethanol) was used for all the chemical search experiments. The delivered rate of ethanol was approximately 0.8 ml · min⁻¹.

3. Insect Based Flight Control

In order to deal with the accessibility challenge faced in humanitarian demining we propose to use a flying platform. The first property of such a system is that it can autonomously maintain a specific course and compensate for perturbations. Insects localize odour sources by using multiple sensor modalities including: chemosensing, vision, anemotaxis and mechano-sensing. It is for this reason that this particular behavior observed in the moth is referred to as opto-motor anemotactic behavior (Kennedy & Marsh 1974). Hence, for the moth, vision is as important as olfaction since they cannot orient without visual cues (Kennedy & Marsh 1974; Charlton & Cardé. 1990). In a brain of about 1mm³, insects incorporate principles for visual navigation that are not only efficient in their implementation but also robust and reliable (Posey et al. 2001). Generally in insects, about two thirds of their brain is dedicated to visual processing to support navigation (Strausfeld 1976). Visual processing is mainly done by feed-forward neural structures dedicated to extract wide filed directional optic flow that is believed to be used for flight control (Hausen 1982a; Hausen 1982b; Egelhaaf & Borst 1993a; Krapp et al. 1998; Franz & Krapp 2000; Douglass & Strausfeld 2003; Higgins et al. 2004). In later processing stages, this information is integrated and used for landing, course and altitude control and collision avoidance (Rowel 1971; David 1982; Rind & Simmons 1992; Egelhaaf & Borst 1993b; Srinivasan et al. 1996; Srinivasan et al. 2000; Tammero & Dickinson 2002; Higgins et al. 2004).

In the particular case of the moth, it is believed that vision is used to assess the wind direction by computing the optic flow caused by drifts in position generated by the wind airflow (Ludlow 1982). Moreover, vision is also known to be used to control the flight speed and set it to a constant ground speed (Kennedy & Marsh 1974; Kennedy et al. 1978).

Here we look into the flight control question in relation to the computational and behavioral principles of the opto-motor system of the fly and locust. We choose these two preparations since they are the best studied species with respect to opto-motor behaviors. An important question is to what extend these insect based principles of visually guided 3 dimensional navigation can generalize to man made flying platforms (Srinivasan et al. 1999; Franceschini et al. 2007). From the technological point of view, the idea of using insect based task oriented models for flight control appears appealing because of their computational efficiency and flexibility. Consequently, we aim at providing a flight control infrastructure based only on biologically plausible and realistic neuronal models of the insect opto-motor system that can bring in the essential capabilities for the autonomous navigation of a flying robot.

Course and Altitude Control

The visual system of insects is commonly seen as a feed forward system with clearly defined functional and anatomical divisions (Strausfeld 1976). These divisions, known as neuropils, process the visual information at different levels, from a contrast enhancement to visual motion extraction (Fig. 4). Our synthetic insect visual system models the first three neuropils (Lamina, Medulla and the Lobula complex), and starts with the capture of visual information by means of a video camera, mimicking the photo transduction that takes place in the photoreceptors. The first process on the visual input is a compression of the

luminance levels (Fig. 4a), followed by an edge enhancement (Fig. 4b). These 2 simple operations render the system more robust to changes in illumination (Dubs 1982). At the level of the Medulla we find the first visual motion sensitive neurons (Borst & Egelhaaf 1993; Douglass & Strausfeld 2003). Since the 60s, when Reichardt first proposed the, so called, correlation model, insects are believed to possess neurons that are capable to compute local directional motion, the Elementary Motion Detectors (EMD), and that they use this information for navigation purposes (Reichardt 1961) (Fig. 4c). Furthermore, at the level of the Lobula complex a number of wide field neurons have been identified in many species that provide motion information on horizontal and vertical displacement, as well as on rotations and collisions in the full visual field (Krapp et al. 1998; Gabbiani 2004). In the case of the fly, the neurons known as the Horizontal and Vertical System (HS and VS) can extract this information from the optic flow and have been suggested to be used for flight control (Egelhaaf & Borst 1993a; Franz & Krapp 2000; Tammero & Dickinson 2002).

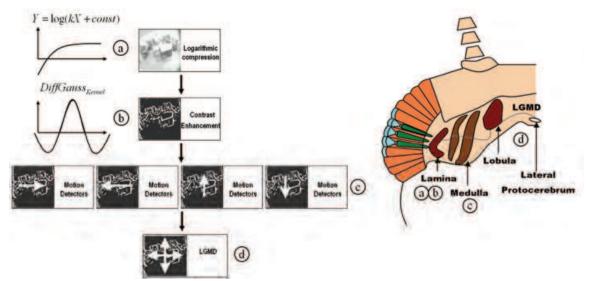


Fig. 4. The main functional and anatomical components of a prototypical insect visual system based on the locust. See text for further explanation. Adapted from Bermúdez i Badia et al. (2007b)

A course stabilization and altitude control system should make sure that the flying device follows a desired trajectory, and corrects for eventual drifts and altitude changes. Indeed we find that the HS and VS neurons of the fly exactly provide reliable information for the above mentioned functionalities (Krapp et al. 1998; Haag et al. 2004). These neurons have been widely modeled using the correlation model proposed by Reihardt to extract local directional motion information (Borst & Egelhaaf 1993; Egelhaaf & Borst 1993b; Franz & Krapp 2000; Haag et al. 2004; Bermúdez i Badia et al. 2005; Harrison 2005; Bermúdez i Badia et al. 2007b) (Fig. 5). Conceptually, the model performs a time delayed-spatial correlation to compute time dependent correlations on the normalized and enhanced input pixel values, and as such extracts local motion information (see Bermúdez i Badia et al. (2007b) for model details). Subsequently, all the correlations computed per pixel are summed to extract wide field motion information (Fig. 5). Therefore, depending on the two parameters that define the spatio-temporal correlation (δ and spatial shift), wide field neurons can be modeled to

be sensitive to different kinds of optic flow such as the ones generated for horizontal and vertical displacements during flight (Fig. 5).

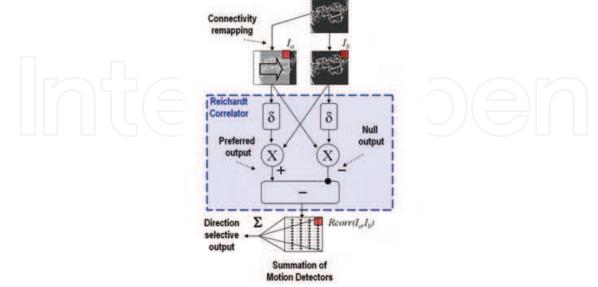


Fig. 5. The functional principles underlying a model of the Horizontal System neuron that extracts directional motion information from the optic flow based on the, so called, Reichardt correlator. In this particular case, the model is maximally sensitive to rightward motion (preferred direction). δ represents a delay, x the multiplication operation and – the subtraction operation. See text for further explanation. Adapted from Bermúdez i Badia et al. (2007b)

To detect rotations and altitude changes from the optic flow of the video stream broadcasted from the cameras of the indoor UAV, wide field HS and VS neurons are modelled. In order to map displacements detected by the HS and VS systems we used a P control system (proportional) that executed a motor command proportional to the responses of the HS and VS neurons, and in the opposite direction. The P system was chosen since it is the simplest solution and the one that makes the least assumptions on the computations that an insect brain could perform.

A blimp has a very prototypical and inconvenient dynamics caused by the added mass and Coriolis forces that prevent it from following a straight course. In order to evaluate the performance of our altitude and course control systems, we implemented them in the neural simulator **iqr** and we recorded synchronously the motor commands, UAV position and neuronal responses of our model. The effects caused by the added mass and Coriolis forces seem to disappear after applying our model to the UAV (Fig. 6, left panel). Several test flights were performed with the UAV to quantify the performance of the model. A maximum off-course deviation of 15° with respect to the perfect trajectory, with a mean deviation of 7.05°, was measured. During all these tests, a mean velocity of 0.62 m \cdot s⁻¹ was maintained. In addition, we used the same method to quantify the performance of the altitude control system to support a constant altitude. The examination of the responses of the UAV shows that the motor compensation forces of the altitude control system are tightly coupled to the variations of the altitude of the UAV (Fig. 6, right panel). Hence, we observe that the altitude control system is compensating the upwards and downwards

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displacements of the UAV with strong motor responses proportional to the vertical displacement detected by the neural model (Bermúdez i Badia et al. 2007b).

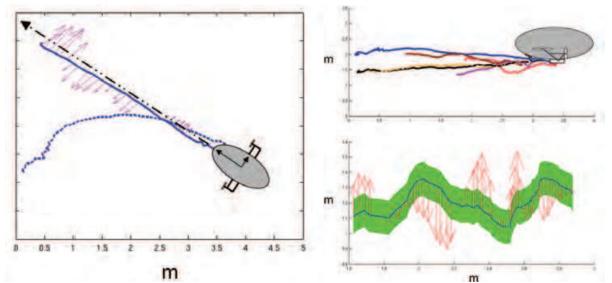


Fig. 6. UAV behaviour under the control of the proposed biologically based neuronal models for course stabilization. Left panel: top view comparison of a trace of the blimp-based UAV without the course control system (dashed line) and a trace of the UAV controlled by the neuronal models. Top right panel: side view of 6 UAV traces under the control of the altitude control system. Bottom right panel: side view showing the compensation motor forces (arrows) and the standard deviation (filled area) of an example trace. Adapted from Bermúdez i Badia et al. (2007b)

Collision Avoidance

A collision avoidance system is a must for any autonomous navigation system. Many sensors, from ultrasonic, infrared and laser sensors to vision have been applied to this purpose, although no general solution has been found (Fox et al. 1997; Surmann et al. 2003; Harrison 2005). The neural correlate of collision avoidance has been identified in many insect species, and due to its impressive performance and accessibility it has been subject of abundant research (Rowel 1971; Gabbiani 2002; Tammero & Dickinson 2002; Krapp & Gabbiani 2005). The best studied collision avoidance system in insects is the one of the Locust, known as the Lobula Giant Movement Detector (LGMD), which relies exclusively on vision. The LGMD has been shown to robustly signal collisions with objects independent of their size, texture, shape and approaching angle (Gabbiani et al. 1999; Gabbiani 2001; Gabbiani 2002; Gabbiani 2004).

Previous studies shave shown that a system that extracts visual expansion, i.e. looming sensitive, based on Reichardt's correlation can account for many aspects of the responses generated by the LGMD at the same time it is a suitable system for a robot implementation (Indiveri 1998; Blanchard et al. 2000; Bermúdez i Badia & Verschure 2004; Harrison 2005; Bermúdez i Badia et al. 2007b).

Relying on the same principles as the course and altitude control models, the local computation of motion, our LGMD model computes motion in a radial outward fashion (Fig. 7, left panel). This alignment of EMDs makes the system looming sensitive. Looming is greater when it is caused by faster approaching objects or by closer ones. Thus, the level of

the looming signal is used to trigger collision detections as opposed to be used to estimate distance-to-contact or time-to-contact. All the neural responses of the specifically arranged EMDs are integrated by the excitatory pre-synaptic fan of the LGMD and inhibited by a global motion signal (Rowel 1971; O'Shea & Williams 1974; O'Shea & Rowell 1976) (Fig. 7, right panel). This connectivity converts the LGMD neuron into a looming sensitive system (excitatory pathway) with a normalization signal that regulates the looming sensitivity of the system with respect to the properties of the visual input (inhibitory pathway). The regulatory signal is crucial since more activity at the input level generates more spurious activity at the EMD level, which may lead to random correlations and therefore to more false positives. After the integration of the activity of the inhibitory and excitatory synaptic connections, a threshold operation is used to decide when the looming signal is high enough to be considered a collision (see Bermúdez i Badia et al. (2007b) for model details).

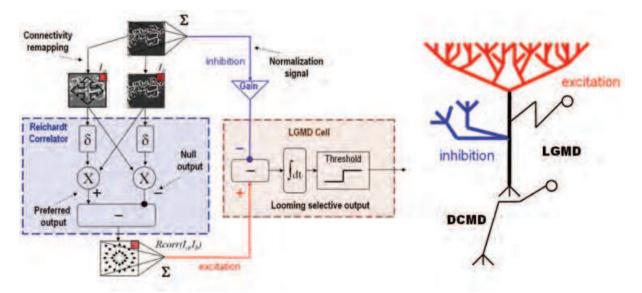


Fig. 7. The functional principles of our model of the Lobula Giant Movement Detector, that extracts expanding motion information from optic flow, and the morphology of its biological counterpart. δ represents a delay, x the multiplication operation and – the subtraction operation. See text for further explanation. Adapted from Bermúdez i Badia et al. (2007b)

To successfully avoid collisions, the UAV not only has to reliably detect approaching objects but also do it at a prudent distance. This is also critical for fast moving vehicles, vehicles with slow reaction times or when inertial forces play an important role. To test the performance of the proposed model of the LGMD, we implemented it in **iqr** and employed it to trigger avoidance manoeuvres on the indoor UAV whenever a collision was detected. The avoidance manoeuvre was triggered in the opposite direction of the collision detection, and with a rotation speed proportional to the amplitude of the looming measure provided by the model

The tests were run in an empty room (~ 5×4 m) equipped with curtains with random black filled squares to provide visual cues. An analysis of the system showed robust collision detection and a correlation between detection distance and translational speed where later responses were observed for high speeds, being in the worst case at a distance of about 1.75

m. In addition, the peak response always occurs before the collision takes place, largely independent of the approaching speed (Bermúdez i Badia et al. 2007b). This system allowed for successful autonomous navigation of the UAV with a minimal number of collisions (~ 90% correct detections) (Fig. 8). Most missed collisions occurred at very shallow angles where the cameras and their optics do not capture sufficient visual information.

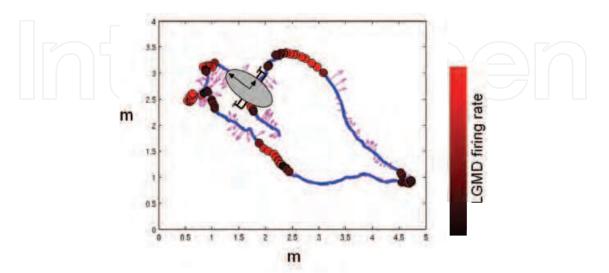


Fig. 8. Trace example for one minute of flight controlled by our LGMD neural model. The blue line represents the position of the UAV within the test area, the arrows represent the motor commands of the course stabilization model, and the filled dots indicate when a collision is detected. The intensity of the red colour is proportional to the amplitude of the model's response. Adapted from Bermúdez i Badia et al. (2007b)

4. Localization Strategies

Usually, the study of mine clearance is reduced to the investigation and development of sensor technology that can detect a number of mines. However, the ultimate sensor technology does not yet exist among the technologies of potential use to this problem. Moreover, demining does not only depend on the selection of the appropriate sensor technology but more aspects play a role. Often overlooked, the efficiency of the clearance process largely depends on the chosen localization strategy. This becomes more relevant when the mined areas are very large or unknown, or when clearance is urgently required. Thus, there is a need for an efficient explosive localization strategy that is not only able to locate possible explosive artifacts but that performs this task in an efficient way.

Our target model, the moth, has developed the appropriate sensor neural structures to be extremely sensitive in the detection of chemical signals (Kennedy & Marsh 1974). In addition, via evolution it has developed a behavioral strategy to optimize success rate when tracing a particular odor in a plume of complex dynamics and within complex odor blends (Kennedy et al. 1978; Baker & Kuenen 1982; Baker 1990). Therefore, the understanding of the origin of the high sensitivity of moths to pheromones is very valuable when studying optimal odor encoding and processing mechanisms, and developing a real-time mine localization robot. Hence, the objective is to learn the key elements from the neural substrate

of the systems of the moth involved in the odor localization task, and to present the according biologically constrained models of our cUAV.

Mapping of an Area

The first and most straight forward approach for localization is to perform a mapping of the area to be examined. The objective of mapping an area is to generate a density plot in which sensor measures are associated to specific locations in space. There are many ways to generate such maps, by means of numerous static sensors located at specific points in space, by means of a dynamic scanning of the full area, a random search within the area, etc. The resulting maps contain information that, independent of the sensor technology used, can be used to derive the probability of having an explosive at specific x and y coordinates. Interestingly enough, this approach has been also used to locate explosives with bees. Bees trained to detect certain chemical compounds were released in a mined area, and then the measured density of bees \cdot m⁻² was used to estimate the position of the mines (Bromenshenk et al. 2003).

Mapping using multiple sensors and parallel measures speeds up the process but it becomes unfeasible when the sensor technology is very expensive. Alternatively, the scanning of an area may lead to the best results but requiring a long time to explore the entire surface, becoming then impracticable for very large areas. A trade off would be to use a mapping strategy that maximizes the area coverage per unit of time. In particular, a random search strategy applied on larger areas explores initially a larger surface per unit of time but then it requires an infinite amount of time to explore the entire area. Thus, depending on the time and surface area constraints, a strategy has to be chosen to maximize coverage, i.e. probability of mine detection (Bermúdez i Badia et al. 2007a).

The mapping of known areas or environments can also be used as a way to characterize the sensitivity, reliability and dynamic responses of our sensor technology. This is a necessary step to assess the limitations of the sensor technology and therefore its applicability to a particular task, in this case the autonomous chemical search by means of a cUAV. The first question that arises is whether our metal thin oxide sensor technology can reliably measure a chemical plume and reconstruct it. In order to assess that, we constructed a 3 x 4 m wind tunnel in which we generated a chemical plume using an ultrasonic delivery system with a 9.4% solution of ethanol in distilled water and a wind speed of 0.7 m \cdot s⁻¹ (see section 2).

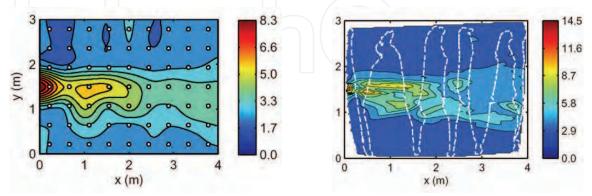


Fig. 9. Wind tunnel chemical mapping experiments by means of static (left panel) and dynamic (right panel) measurements of a chemical plume. Both maps are generated by the sensor responses to a 9.4% solution of ethanol in distilled water delivered at position (0 m, 1.5 m). The map was smoothened using a bi-cubic filter and divided

into 11 contours. The dots represent the sensor location, the trace the position of a remote controlled mobile robot, and the color bar the sensor response strength.

Adapted from Pyk et al. (2006) rder to investigate the effect of the samplin

In order to investigate the effect of the sampling strategy on the accuracy of a chemical map we performed windtunnel experiments comparing static and dynamic smapling approaches. The chemical plume generated in the wind tunnel was sequentially measured during 2 min using a single sensor placed at 9 times 7 equally spaced locations which were interpolated to generate a map (Fig. 9, left panel). The resulting map shows that the static mapping of a controlled chemical plume can be done using our sensor technology, clearly displaying the ethanol distribution in the wind tunnel. Moreover, as a control experiment, the wind tunnel was remapped using the same procedure but this time with a distilled water plume, this demonstrated that the sensor was responding exclusively to the ethanol content of the plume (see for Pyk et al. (2006) further details).

However, it is not clear how time-averaged static measurements generalize to measurements performed by sensors in behaving artifacts. Therefore, we repeated the experiment using our mobile robot equipped with a thin metal oxide chemo-sensor. In this case, the robot was manually driven scanning the wind tunnel from back to front at a speed of $10 \text{ cm} \cdot \text{s}^{-1}$. Its position was tracked using a custom made visual tracking system called AnTS. The map generated using the dynamical measurements is consistent with the previous one, and shows that the chemo-sensor provides a rapid and reliable measurement of the ethanol concentration while the robot is moving. Hence, we can conclude that this technology seems suitable for an on board implementation in our cUAV that performs dynamic mapping (Fig. 9, right panel).

A Moth Behavior Based Localization Strategy

The problem of odor localization is considerably more complex than the one of mapping since the chemical cues are carried by filamentous plumes of a complicated structure and dynamics that are unpredictable and follow complex patterns (Murlis 1986). Localization, as opposed to mapping, can be less time consuming and does not necessarily require the exploration of the complete area.

In nature we can find a large amount of animals that solved the localization problem of odor cues as an effective way of chemical communication (Kennedy & Marsh 1974; Thorp & Ammerman 1978; Johnston 2003). Female moths release extremely low concentrations of sex attractant chemical signals (pheromones) that male moths are capable of tracing over large spatial scales (hundreds of meters). Therefore, the understanding of the mechanisms underlying moth chemical communication would allow human made robots to reproduce their accuracy, sensitivity and strategies adapted to a number of applications such us fire detection, environmental monitoring, demining, etc.

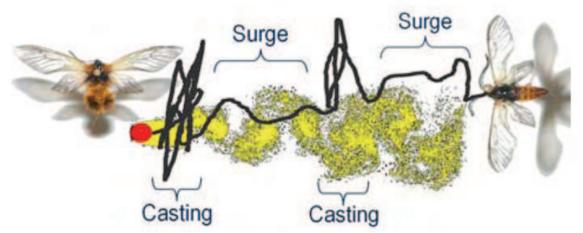


Fig. 10. An actual moth trace (black line) following upwind a pheromone plume (in yellow) released by a female moth (left). The male moth displays two stereo-typical behaviors, surge up-wind and casting cross wind. The red dot represents the location of the release point

The remarkable performance of the male moth in the chemical location task has been largely attributed to the alternation of two behaviors that help it to sample the pheromone plume. The surge behavior consists of a relatively straight upwind flight believed to be triggered by the contact with a pheromone filament. Instead, the casting behavior is a zigzagging crosswind movement triggered in this case by the absence of pheromone contact (Kennedy et al. 1978; Kennedy 1983; Baker 1990; Olberg 1993; Balkovsky & Shraiman 2002).

Based on the moth chemical search behavior, many models have been proposed and tested in both simulations and robot experiments (Kennedy et al. 1978; Kennedy 1983; Baker 1990; Olberg 1993; Balkovsky & Shraiman 2002). We have proposed a model augmented with the previously introduced opto-motor course stabilization and the collision avoidance system based on the LGMD neuron of the locust. This model of the opto-motor anemotactic behavior of the moth is based on the surge and casting behavioral modes observed in the moth, and the switches between them are triggered by the contact with the targeted chemical compound. Additionally, the LGMD collision avoidance system overwrites any of those behaviors triggering an avoidance maneuver whenever an imminent collision is detected (Fig. 11). As a result, the model requires of a chemo-sensor, wind direction information for the upwind and crosswind flight (anemometer), and a partial insect visual system to detect collisions (wireless camera system). All these sensor systems are integrated in our mobile platform (Fig. 2, right panel), and a biologically plausible neural implementation of the opto-motor and chemical search model, consisting of 7082 neurons, aggregated in 97 groups, and 180 connections with 11887 synapses, is simulated in real-time using iqr (see Pyk et al. (2006) for further details).

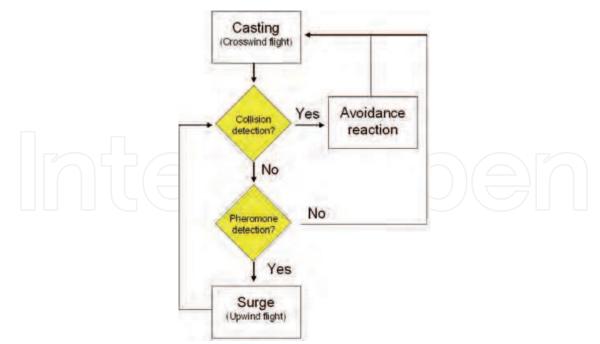


Fig. 11. Block diagram of the moth opto-motor anemotactic behavior model augmented with a collision avoidance system. Adapted from Pyk et al. (2006)

The previous mapping experiments described earlier demonstrated that the chemo-sensor technology proposed provides a functional signal for the detection of ethanol. Further, the wind vane sensor provides the wind direction to the chemical search model. We tested the performance of the proposed moth behavioral model in our wind tunnel in two test and two control conditions. 37 trials were performed and for each trial the robot was placed around 3.5 m downwind from the source on an arbitrary y-coordinate.

Firstly, the model was tested without the collision avoidance system to evaluate the performance of the chemo-sensing mobile robot for an ethanol plume. Two chemical concentrations were tested (17 trials on 9.4% and 20 trials on 23.5%) of water ethanol solution. In all the cases the robot was able to locate the ethanol delivery point, and it required of a median time of 74.2 s (Fig. 12, top left panel) (see table 1 for details). The search times did not differ significantly between the two conditions (Wilcoxon rank sum test for equal medians), being the performance not dependent on the chemical concentration. We observed that the casting mode can be distinguished from the surge mode without difficulty because of the zigzag pattern, lower sensor readings and smaller upwind displacement (Fig. 12, top left panel).

Ethanol source	Mode	Median [s]	Percentile 10% [s]	Percentile 90% [s]
9.4%, n=17	Casting	39.90	28.37	45.72
	Surge	33.10	17.91	103.66
	Total	74.97	62.35	121.89
23.5%, n=20	Casting	32.32	16.42	61.73
	Surge	31.88	14.23	136.53
	Total	67.83	45.50	174.37

Table 1. Performance analysis of the chemical search neural model. Median, 10% percentileand 90% percentile total search time for the two plume conditions

Secondly, the experiment was repeated with an obstacle and the collision avoidance system enabled. The robot behavior demonstrated a successful integration of the moth anemotactic chemical search model with the insect based collision avoidance model, being capable of finding the odor source even in the presence of objects obstructing the direct path (Fig. 12, top right panel).

Finally, two control experiments were performed in the absence of a chemical plume and in the absence of wind flow (Fig. 12, bottom). In the first case, the robot was unable to detect the ethanol plume and therefore displayed exclusively the casting behavior (Fig. 12, bottom right panel). In the second case, the robot not only could not detect the chemical trail but it was unable to measure the wind direction, and thus unable to find the source direction. This resulted in an erratic behavior without a clear movement direction or target direction (Fig. 12, bottom left panel). This demonstrates that both airflow and chemical stimuli are required for the robot to complete the localization behavior.

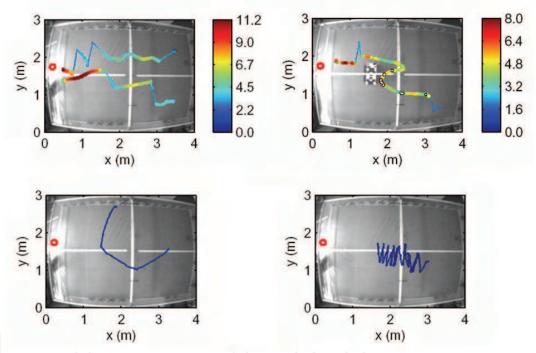


Fig. 12. Prototypical robot trials for the chemical search experiments using the moth behavior based localization model. Top left panel: two trials of the chemical search model with the collision avoidance system disabled. Top right panel: example trial of the chemical search model augmented with the collision avoidance system in the presence of obstacles. Bottom left panel: example trial of the chemical search model in the absence of airflow. Bottom right panel: example trial of the chemical search model in presence of airflow and absence of chemical plume. The thick robot trace indicates surge mode whereas the thin one indicates casting mode. The color of the trace represents the amplitude of the sensor readings of the robot in that particular position. The white dots indicate collision detection and the red dot indicates the position of the ethanol delivery system. Adapted from Pyk et al. (2006)

Consequently, the moth based search behavior model seems to be a comparatively faster odor localization strategy than mapping. Moreover, its success rate demonstrates not only the reliability of this model to successfully localize the odor source under turbulent conditions, but also the matching between chemical plume dynamics and sensor readings, and its integration and enhancement with the LGMD collision avoidance model.

A moth Neural Based Localization Strategy

It has been very difficult to assess more detailed aspects of moth chemotaxis directly because of the impossibility of visualizing a plume without interfering with the flight behavior of the moth. Hence, it is still not quantitatively established whether the moth responds to a chemical gradient, filament contact or uses a more complicated behavioral strategy. However, experiments have been performed where a male moth was equipped with a third antenna with a wireless transmission system to approximate what it would sense (Kuwana et al. 1999). Although the results of this approach were technologically interesting and challenging, it is insufficient for a proper characterization of the relationship between the stimuli the moth is exposed to and the behavior it displays since it does not directly measure from the real antennae.

As shown by our behavior based model, a simple strategy switching between upwind (surge) and crosswind flight (casting) can be very successful in solving the localization problem. Nevertheless, the moth developed some neural systems to deal with the specific characteristics of the structure of the pheromone plume that are not exploited in our behavior based strategy. For instance, it is known that the temporal structure of the stimulus is encoded in the responses of the nervous system of the insect, and that this structure is crucial to keep the moth flying upwind in the direction of the source of pheromone (Murlis & Jones 1981; Vickers & Baker 1994; Mafra-Neto 1995; Kuwana et al. 1999; Quero et al. 2001; Justus 2002). In this case, the frequency of the odour filaments has been shown to have a strong impact on the behavior of the moth, where the moth appears to be tuned to respond maximally to a specific detection frequency (Willis & Baker 1984; Vickers & Baker 1994; Vickers 1996).

The question arises whether a model based exclusively on phenomenological observations of the moth can fully account for its behavior without taking into account the neural substrate that regulates it. Hence, we aim at constraining our localization model with the current knowledge of the odor processing stages in the moth and identified neuronal mechanisms.

Our neural based model relies on two important hypotheses, the use of *stereo information* and the *pheromone frequency dependency*. The first hypothesis is supported by the, so called, flip/flop neurons (Kanzaki et al. 1989). These are, so called, Descending Neurons of the protocerebrum that arborize in the Lateral Accessory Lobe, and that show a bi-stable high and low frequency response. The activity of these, so called, flip/flop neurons is directly correlated with the body orientation and zigzagging of the moth while tracing a pheromone trail. Moreover, the orientation switches appear to be caused by the difference of pheromone concentration in the anteanne (Olberg 1993; Mishima & Kanzak 1998; Wada & Kanzaki 2005). Therefore, olfactory stereo information is used in our model to trigger the orientation changes whenever the sensor reading difference of two chemo-sensors is above threshold (Fig. 13).

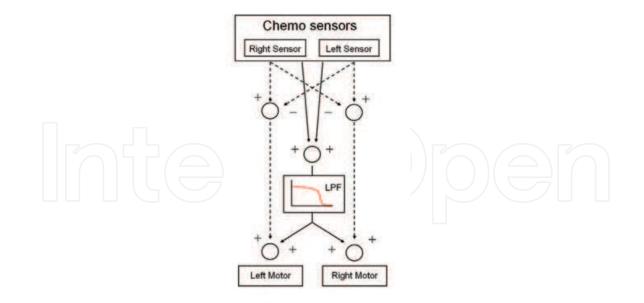


Fig. 13. Scheme of the proposed neural based model for the control of chemical search. The left and right chemo-sensor responses are continuously compared to re-orient the robot in the direction of the maximum concentration. The Low Pass Filter of the sum of the sensor readings (LPF) is used to drive the two motors of the robot at the same speed to obtain an upwind translation. Adapted from Bermúdez i Badia et al. (2007a)

The second corner stone of the model is based on the Macro Glomerular Complex (MGC) of the Antennal Lobe (AL) of the moth, a glomerulus that evolved to solely process the pheromone signals (Christensen & Hildebrand 1987; Kanzaki et al. 1989; Hansson et al. 1991; Christensen et al. 1993; Christensen et al. 1995). Approximately 85% of the neurons of the MGC display transient responses correlated with the pheromone signal, but are only able to resolve odor pulses of up to a few Hz (Christensen & Hildebrand 1987; Christensen et al. 1993; Lei & Hansson 1999). Surprisingly, this frequency response range is similar to the characteristics of the plume pulsing experiments that show that moths display a faster upwind displacement for pulsed pheromone signals (Murlis & Jones 1981; Murlis 1992). Thus, a Low Pass Filter (LPF) of the readings of the chemical sensors is used to trigger translation displacement (independent of the heading direction) (Fig. 13).

As opposed to the previous behavior based model, in this case the wind direction information is not used to direct the robot upwind but only to prevent the robot from an undesired downwind displacement.

To quantify the performance of the neural based model we used the following criteria: from an evolutionary point of view one can assume that the most advantageous strategy has to trade off some aspects such as the consumption of energy, the search time, the number of competitors, the number of female moths and, of course, the success rate. Therefore, and given the amount of male moths that can receive the chemical cues of a single female, we assume that the optimal search strategy has to mainly balance search time with success rate, i.e. the longer a moth takes to search the more chances it has to find the female, but it is also more likely that another male moth will have found it before while the probability to be detected by a predator will increase. Hence, in the subsequent experiments we compared the accuracy and search time of a modified version of the behavior based model, which uses a single chemo-sensor, with the neural based model, which uses two chemo-sensors mounted on our mobile robot. Inspired by the theoretical studies of Balkovsky et al., the casting mode of the behavior based model was modified from having a fix crosswind length to increasing it over time, increasing consequently the probability of the robot intercepting a pheromone filament over time since it explores a larger area (Balkovsky & Shraiman 2002) (Fig. 11).

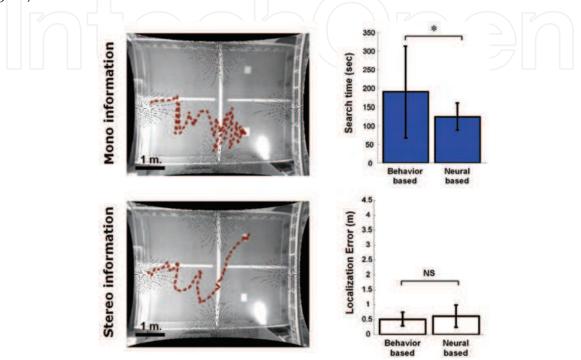


Fig. 14. Comparative behavior analysis of the two proposed moth based chemical search models. Left panel: Example traces for the behavior based (mono) and neural based (stereo) chemical search models. Right top panel: Search time comparison between models. Right bottom panel: Localization error comparison between models. A total of 100 test runs were performed for the two conditions. Bars indicate the data variance. Adapted from Bermúdez i Badia et al. (2007a)

In order to compare the performance of both models, a total of 100 robot test runs were done starting randomly at the rear left or right side of the wind tunnel (Fig. 14, white areas in the left panel). Then a chemical plume was created using a water/ethanol solution of 20% ethanol (see section 2). The performance of both moth based models was evaluated measuring the total search time and the localization error. The accuracy of the behavior and neural based models, measured as the distance between the ethanol delivery system and the robot end position, is of about 0.5 m, with no significant difference (2 sample t-test, p > 0.05) (fig. 14, bottom right panel). Nevertheless, when we take into account the total search time we observe that the neural based model outperforms significantly (reducing the average search time by more than one minute which is 31% of the mean task duration) the performance of the behavior based model (2 sample t-test, p < 0.05) (Fig. 14, top right panel). Thus, the neural based model is significantly faster than the behavior based yet displaying the same localization precision. If we compare the mean search time of the neural based

model (approx. 2 min.) to the time required for the robot to scan the whole wind tunnel (approx. 5 min) it is more than twice faster. Moreover, this difference will increase even further for greater areas, since scanning search time directly depends on the area size to be explored.

Outdoor cUAV Experiments

In order to test the different models studied in indoor controlled environments in more realistic scenarios we constructed an outdoor chemo-sensing UAV (see section 2). As starting point, we equipped the outdoor cUAV with a single chemo-sensor, a GPS, an altimeter, a gyroscope and a compass. These traditional sensing technologies were added for monitoring and analysis purposes. Then, a standard Proportional Integral Derivative (PID) control software was developed to allow for autonomous and remote navigation of the flying platform (see Bermúdez i Badia et al. (2007a) for further details). Besides the robot and sensor technology, the performance of the proposed mine detection technology in large and unspecified terrains does not only depend on the localization strategies but also on the scheduling and coordination of a number of these cUAVs that are applied to the demining problem. Therefore, we propose to use a control layer that permits the scheduling of a fleet of cUAVs to coordinate the search in large environments. A base station divides the target area into a number of sub-regions in which each of the cUAV performs an autonomous flight. For that, the cUAVs make use of their GPS to ensure that they do not surpass the limits of the region assigned to them.

The first outdoor scenario used to test our cUAV was a university football field (40x80 m.) in which we placed a single odor delivery source. For these experiments, the ultrasonic delivery system released a solution of 40% ethanol in water at a rate of 200 ml \cdot h⁻¹. The weather station reported an average wind speed of 3 km \cdot h⁻¹ during the experiments.

In these experiments, the platform was successfully tested to perform a number of autonomous mapping strategies (random search and scanning). These experiments demonstrated the reliability of the hardware and software control infrastructure of the cUAV, and the feasibility of fully autonomous chemical localizations based on predefined strategies (see Bermúdez i Badia et al. (2007a) for further details).

Nevertheless, the chemical source localization becomes impracticable by a single cUAV when search is required over very large areas. Thus, we tested the possibility of scheduling a cUAV fleet based on non-overlapping divisions of the area to explore. Inside each of these regions, the cUAVs perform an autonomous chemical mapping. Therefore, we scheduled our outdoor cUAV to perform a chemical mapping using a random search strategy on a predefined area, and we re-scheduled to another adjacent area after it realized the first mapping (Fig. 15). These experiments show that the cUAV is able to sequentially examine non-overlapping areas autonomously and create a consistent chemical map of the sensed ethanol distribution in the football field (Fig. 15). Moreover, the data provided by the reconstructed map suffices to detect and locate the position of the ethanol delivery system, although false positives can occur.

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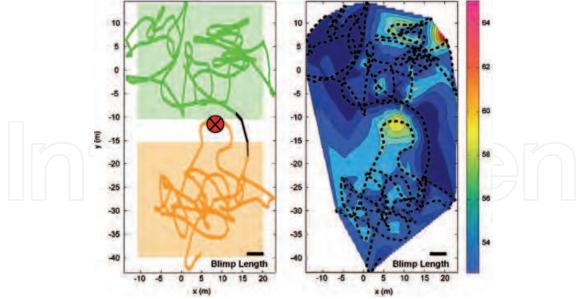


Fig. 15. Autonomous cUAV scheduling and mapping experiments using a random search strategy. Left panel: the cUAV explores sequentially the two search areas (green and orange filled areas). The red circle represents the position of the odor delivery system, and the color lines the trace of the cUAV. Right panel: reconstructed ethanol distribution map of the explored area using a bi-cubic filter and divided into 11 contours. The dashed line represents the cUAV trace. The sensor reading amplitude and the blimp dimensions are indicated. Adapted from Bermúdez i Badia et al. (2007a)

5. Conclusions

In this project we investigated the use of a chemo-sensing Unmanned Aerial Vehicle (cUAV) for humanitarian demining tasks. In this chapter we have shown a wide spectrum of developments oriented towards the autonomous and non-supervised mine detection on large scale and unspecified environments by means of a blimp-based chemo-sensing UAV equipped with a broadly tuned metal thin oxide chemo-sensor. These developments include the design and construction of a number of indoor robots (mobile and flying robots), the first outdoor chemo-sensing UAV, novel insect based flight control methods and a number of moth based localization strategies. The central assumption of our work is that we can develop advanced and effective demining systems by basing them on insect solutions to flight control, navigation and chemical communication.

First, new methods for the autonomous flight control of flying robots have been developed and tested using an indoor blimp-based UAV. These methods, based on the Horizontal and Vertical System neurons of the fly, include a course stabilization and altitude control system. The robot tests demonstrate the feasibility of this technology and that the principles of the neural substrate of the fly visual system can successfully control the course and the altitude of the UAV. Then, the indoor UAV system has been augmented with a collision avoidance system that captures the computational principles of the Lobula Giant Movement Detector neuron of the locust. The UAV tests confirmed the collision avoidance neural model to be robust and reliable during a free indoor flight in a controlled environment.

In addition, the feasibility of the proposed chemo-sensor technology has been proven first by static and dynamic chemical mapping experiments of a controlled environment, then by two moth based odor localization experiments, and finally by a mapping and scheduling experiment of an uncontrolled outdoor environment using the chemo-sensing UAV. This outdoor platform is shown to be very well suited to the mine detection task since it is terrain independent, does not trigger mines, and can perform either fully autonomous chemical search or it can be manually driven.

Our results corroborate the viability of the proposed technology for demining in real-world scenarios. Further, the results support our biologically based approach where the knowledge of biological systems is used to solve challenging engineering problems. This is contrasted to a "biologically inspired" approach that does not require its solutions to satisfy standard scientific norms of validity and or plausibility. Given our results we suggest that a biologically based approach is more effective since it is both advancing our understanding of nature and the construction of real-world artefacts.

All the biological systems considered in this study are of great interest for robotics and computation because of their simplicity (implementation wise) and effectiveness. Research with robots permits to study those systems by making complex models of the different subsystems involved in the flight control and moth chemical search, and test them in controlled and in more realistic and complex scenarios. Moreover, these new models that are developed contain implicit hypothesis on unknown neural mechanisms of the moth, and as such robotics can contribute to a better understanding of biological systems at the same time novel and robust solutions are developed. Hence, the convergent validation of physiology, anatomy and behavior advances our understanding of the insect solution to chemical communication.

We have explored the main components required to build a fully synthetic moth robot. However the full integration of some of these system parts is not yet completed. In future work, we expect to be able to test the different moth based chemical localization strategies using the outdoor cUAV in more complex outdoor environments. Further, the biologically based flight control neural models have been shown to deliver a rotation and altitude change related measure, thus in the future we plan them to replace the altimeter and gyroscope sensors used in the outdoor cUAV.

In addition, the limit of the sensitivity of the thin metal oxide chemo-sensing technology, and in particular the applicability to the particular case of the detection of leaking explosives has to be further assessed. It has been shown that the sensitivity of this type of technology can go up to few parts per million (ppm), providing a wide range sensitivity that allows for identification of organic compounds (Neubecker et al. 1997; Eickhoff et al. 2003). Nevertheless, the chemical traces of some explosives such as TNT are shown to be made of few parts per billion (ppb) (Rose et al. 2005). Despite this problem, the particularities of the sensor technology used for demining do not alter the important contribution of the biologically based approach, the studied search strategies, the flight control mechanisms, and the terrain independent platform approach.

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9. References

- Baker, T. C. (1990). Upwind flight and casting flight: complimentary phasic and tonic systems used for location of a sex pheromone sources by male moths, *Proceedings of Proc. of the 10th International Symposium on Olfaction and Taste*, pp. 18-25, Oslo, Norway.
- Baker, T. C. & L. P. S. Kuenen (1982). Pheromone source location by flying moths: a supplementary non-anemotactic mechanism. *Science* Vol. 216, 424-427.
- Balkovsky, E. & B. I. Shraiman (2002). Olfactory search at high Reynolds number. *Proc. Natl. Acad. Sci. USA* Vol. 99, 20, 12589-93.
- Bermúdez i Badia, S., U. Bernardet, A. Guanella, P. Pyk & P. F. M. J. Verschure (2007a). A Biologically Based chemo-sensing UAV for Humanitarian Demining. *International Journal of Advanced Robotic Systems* Vol. 4, 2, 187-198.
- Bermúdez i Badia, S., P. Pyk & V. P.F.M.J. (2007b). A fly-locust based neuronal control system applied to an unmanned aerial vehicle: the invertebrate neuronal principles for course stabilization, altitude control and collision avoidance. *The International Journal of Robotics Research* Vol., 26, 759-772.
- Bermúdez i Badia, S., P. Pyk & P. F. M. J. Verschure (2005). A Biologically Based Flight Control System for a Blimp-based UAV, *Proceedings of International Conference on Robotics and Automation (ICRA05)*, pp., Barcelona, Spain.
- Bermúdez i Badia, S. & P. F. M. J. Verschure (2004). A Collision Avoidance Model Based on the Lobula Giant Movement Detector of the Locust., *Proceedings of Proc. of the International Joint Conference on Neural Networks* 2004 (IJCNN'04), pp. 1757, Budapest, Hungary.
- Bernardet, U., M. Blanchard & P. F. M. J. Verschure (2002). IQR: a distributed system for real-time real-world neural simulation. *Neurocomputing* Vol. 44-46, 1043-1048.
- Blanchard, M., F. C. Rind & P. F. M. J. Verschure (2000). Collision avoidance using a model of the locust LGMD neuron. *Robotics and Autonomous Systems* Vol. 30, 1-2, 17-38.
- Borst, A. & M. Egelhaaf (1993). Detecting visual motion: theory and models. *Rev Oculomot Res* Vol. 5, 3-27.
- Bromenshenk, J. J., H. C.B., R. A. Seccomb, S. D. Rice, R. T. Etter, B. S.F.A., P. J. Rodacy, J. A. Shaw, N. L. Seldomridge, L. H. Spangler & J. J. Wilson (2003). Can Honey Bees Assist in Area Reduction and Landmine Detection? *Journal of Mine Action* Vol., http://maic.jmu.edu/journal/7.3/focus/bromenshenk/bromenshenk.htm.
- Bruschini, C. & B. Gros (1998). A Survey of Research on Sensor Technology for Landmine Detection. *Journal of Humanitarian Demining* Vol. 2, 1.
- Charlton, R. E. & R. T. Cardé. (1990). Orientation of male gypsy moths, Lymantria dispar (L.), to pheromone sources: the role of olfactory and visual cues. *J. Insect Behav.* Vol. 3, 443-469.
- Christensen, J., M. Pedersen, B. Aasted & S. Alexandersen (1995). Purification and characterization of the major nonstructural protein (NS-1) of Aleutian mink disease parvovirus. *J Virol* Vol. 69, 3, 1802-9.

- Christensen, T. A. & J. G. Hildebrand (1987). Male-specific, sex pheromone-selective projection neurons in the antennal lobes of the moth Manduca sexta. *J Comp Physiol [A]* Vol. 160, 5, 553-69.
- Christensen, T. A., B. R. Waldrop, I. D. Harrow & J. G. Hildebrand (1993). Local interneurons and information processing in the olfactory glomeruli of the moth Manduca sexta. *J Comp Physiol* [A] Vol. 173, 4, 385-99.
- David, C. T. (1982). Compensation for height in the control of groundspeed by Drosophila in a new `barber's pole' wind tunnel. *J. Comp. Physiol. A* Vol. 147, 485 -493.
- Douglass, J. K. & N. J. Strausfeld (2003). Anatomical organization of retinotopic motionsensitive pathways in the optic lobes of flies. *Microsc Res Tech* Vol. 62, 2, 132-50.
- Dubs, A. (1982). The spatial integration of signals in the retina and lamina of the fly compound eye under different conditions of luminance. *Journal of Comparative Physiology A: Neuroethology, Sensory, Neural, and Behavioral Physiology* Vol. 146, 3, 321-343.
- Egelhaaf, M. & A. Borst (1993a). A look into the cockpit of the fly: visual orientation, algorithms, and identified neurons. *J Neurosci* Vol. 13, 11, 4563-74.
- Egelhaaf, M. & A. Borst (1993b). Motion computation and visual orientation in flies. *Comp Biochem Physiol Comp Physiol* Vol. 104, 4, 659-73.
- Eickhoff, M., J. Schalwig, G. Steinhoff, O. Weidemann, L. Görgens, R. Neuberger, M. Hermann, B. Baur, G. Müller, O. Ambacher & M. Stutzmann (2003). Electronics and sensors based on pyroelectric AlGaN/GaN heterostructures - Part B: Sensor applications. *Physica Status Solidi* Vol. 0, 6, 1908-1918.
- Fjellanger, R., E. Andersen & I. McLean (2002). A training program for filter-search mine detection dogs. *International Journal of Comparative Psychology* Vol. 15, 277-286.
- Fox, D., W. Burgard & S. Thrun (1997). The dynamic window approach to collision avoidance. *Robotics & Automation Magazine* Vol. 4, 1, 23-33.
- Franceschini, N., F. Ruffier & J. Serres (2007). A bio-inspired flying robot sheds light on insect piloting abilities. *Curr Biol* Vol. 17, 4, 329-35.
- Franz, M. O. & H. G. Krapp (2000). Wide-field, motion-sensitive neurons and matched filters for optic flow fields. *Biol Cybern* Vol. 83, 3, 185-97.
- Gabbiani, F., H. G. Krapp & G. Laurent (1999). Computation of object approach by a widefield, motion-sensitive neuron. *J Neurosci* Vol. 19, 3, 1122-41.
- Gabbiani, F., Krapp, H. G., Hatsopoulos, N., Mo, C. H., Koch, C. and Laurent, G. (2004). Multiplication and stimulus invariance in a looming-sensitive neuron. *J Physiol Paris* Vol. 98, 1-3, 19-34.
- Gabbiani, F., Krapp, H. G., Koch, C. and Laurent, G. (2002). Multiplicative computation in a visual neuron sensitive to looming. *Nature* Vol. 420, 6913, 320-4.
- Gabbiani, F., Mo, C.and Laurent, G. (2001). Invariance of angular threshold computation in a wide-field looming-sensitive neuron. *J Neurosci* Vol. 21, 1, 314-29.
- Gonzalez de Santos, P., E. Garcia, J. Estremera & M. A. Armada (2002). SILO6: Design and configuration of a legged robot for humanitarian demining, *Proceedings of IARP*, *Workshop on Robots for Humanitarian Demining*, pp., Vienna, Austria.
- Gooneratne, C., S. C. Mukhopadhyay & G. Sen Gupta (2004). A review of Sensing Technologies for Landmine Detection: Unmanned Vehicle Based Approach, *Proceedings of The second International Conference on Autonomous Robots and Agents*, pp. 401-407, Palmerston North, New Zealand.

- Haag, J., W. Denk & A. Borst (2004). Fly motion vision is based on Reichardt detectors regardless of the signal-to-noise ratio. *Proc Natl Acad Sci U S A* Vol. 101, 46, 16333-8.
- Habib, M. K. (2002). Mine Clearance Techniques and Technologies for Effective Humanitarian Demining. *Journal of Mine Action* Vol., 6.1.
- Habib, M. K. (2007). Humanitarian Demining: Reality and the Challenge of Technology -The State of the Arts. *International Journal of Advanced Robotic Systems* Vol. 4, 2, 151-172.
- Hansson, B. S., T. A. Christensen & J. G. Hildebrand (1991). Functionally distinct subdivisions of the macroglomerular complex in the antennal lobe of the male sphinx moth Manduca sexta. *J Comp Neurol* Vol. 312, 2, 264-78.
- Harrison, R. R. (2005). A biologically-inspired analog IC for visual collision detection. *IEEE Transactions on Circuits and Systems* Vol. I 52, 2308-2318.
- Hausen, K. (1982a). Motion sensitive interneurons in the optomotor system of the fly. I. The horizontal cells: Structure and signals. *Biol. Cybern* Vol., 45, 143-156.
- Hausen, K. (1982b). Motion sensitive interneurons in the optomotor system of the fly. II. The horizontal cells: Receptive field organization and response characteristics. *Biol. Cybern* Vol., 46, 67-79.
- Higgins, C. M., J. K. Douglass & N. J. Strausfeld (2004). The computational basis of an identified neuronal circuit for elementary motion detection in dipterous insects. *Vis Neurosci* Vol. 21, 4, 567-86.
- Indiveri, G. (1998). Analog VLSI Model of Locust DCMD Neuron Response for Computation of Object Approach. *PROGRESS IN NEURAL PROCESSING* Vol., 10, 47-60.
- Johnston, R. E. (2003). Chemical communication in rodents: from pheromones to individual recognition. *Journal of Mammalogy* Vol. 84, 4, 1141-1162.
- Justus, K. A., Schofield, S.W., Murlis, J., and Cardé, R.T. (2002). Flight behaviour of Cadra cautella males in rapidly pulsed pheromone plumes. *Physiol. Entomol* Vol. 27, 58-66.
- Kanzaki, R., E. A. Arbas, N. J. Strausfeld & J. G. Hildebrand (1989). Physiology and morphology of projection neurons in the antennal lobe of the male moth Manduca sexta. *J Comp Physiol* [A] Vol. 165, 4, 427-53.
- Kennedy, J. S. (1983). Zigzagging and casting as a programmed response to wind-borne odour : a review. *Physiol Ent* Vol. 8, 109-120.
- Kennedy, J. S. & D. Marsh (1974). Pheromone-regulated anemotaxis in flying moths. *Science* Vol. 184, 140, 999-1001.
- Kennedy, J. S., D. M. Marsh & A. R. Ludlow (1978). Anemotactic zigzagging flight in male moths stimulated by pheromone. *Physiological Entomology* Vol. 3, 221-240.
- Krapp, H. G. & F. Gabbiani (2005). Spatial distribution of inputs and local receptive field properties of a wide-field, looming sensitive neuron. *J Neurophysiol* Vol. 93, 4, 2240-53.
- Krapp, H. G., B. Hengstenberg & R. Hengstenberg (1998). Dendritic structure and receptivefield organization of optic flow processing interneurons in the fly. J Neurophysiol Vol. 79, 4, 1902-17.
- Kuwana, Y., S. Nagasawa, I. Shimoyama & R. Kanzaki (1999). Synthesis of the pheromoneoriented behaviour of silkworm moths by a mobile robot with moth antennae as pheromone sensors. *Biosensors Bioelectronics* Vol. 14, 195-202.
- Lei, H. & B. S. Hansson (1999). Central processing of pulsed pheromone signals by antennal lobe neurons in the male moth Agrotis segetum. *J Neurophysiol* Vol. 81, 3, 1113-22.

- Ludlow, A., Perry, J., David, C., Kennedy, J. and Wall, C. (1982). A reappraisal of insect flight towards a distant source of wild-borne odor. *Journal of Chemical Ecology* Vol. 8, 1207-1215.
- Mafra-Neto, A., and Carde, R.T. (1995). Influence of plume structure and pheromone concentration on upwind flight of caudra cautella males. *Physiol. Entomol.* Vol. 20, 117-133.
- Marques, L., M. Rachkov & A. T. de Almeida (2002). Mobile pneumatic robot for demining, Proceedings of International Conference on Robotics and Automation, pp. 3508-3513.
- Mishima, T. & R. Kanzak (1998). Coordination of flipflopping neural signals and head turning during pheromone-mediated walking in a male silkworm moth Bombyx mori. *J Comp Physiol A* Vol. 183, 273-282.
- Murlis, J. (1986). The structure of odour plumes. Mechanisms in insect olfaction. T. L. Payne, M. C. Birch & C. E. J. Kennedy. Oxford, UK, Clarendon Press: 27-38.
- Murlis, J., Elkinton, J.S. and Carde, R.T. (1992). Odor plumes and how insects use them. *Annu. Rev. Entomol.* Vol. 37, 505-32.
- Murlis, J. & C. Jones (1981). Fine scale structure of odor plumes in relation to insect orientation to distant pheromone and other attractant sources. *Physiol. Ent.* Vol. 6, 71-86.
- Nanto, H. & J. R. Stetter (2003). Introduction to Chemosensors. Handbook of Machine Olfaction. T. C. Pearce, S. S. Schiffman, H. T. Nagle & J. W. Garnder. Weinheim, Wiley-VCH: 79-103.
- Neubecker, A., T. Pompl, T. Doll, W. Hansch & I. s. Eisele (1997). Ozone-enhanced molecular beam deposition of nickel oxide (NiO) for sensor applications. *Thin Solid Films* Vol. 310, 1-2, 19-23.
- Nicoud, J. D. & M. K. Habib (1995). The Pemex-B autonomous demining robot: perception and navigationstrategies, *Proceedings of International Conference on Intelligent Robots and Systems*, pp. 419-424, Pittsburgh, PA, USA.
- Nonami, K., Q. J. Huang, D. Komizo, N. Shimoi & H. Uchida (2000). Humanitarian Mine Detection Six-
- Legged Walking Robot, *Proceedings of 3rd International Conference on Climbing and Walking Robots*, pp. 861-868, Madrid, Spain.
- O'Shea, M. & C. H. F. Rowell (1976). The neuronal basis of a sensory analyzer, the acridid movement detector system. II. Response decrement, convergence, and the nature of the excitatory afferents to the fan-like dendrites of the LGMD. *J. Exp. Biol.* Vol., 65, 289-308.
- O'Shea, M. & J. L. D. Williams (1974). The anatomy and output connection of a locust visual interneurone; the lobula giant movement detector (LGMD) neurone. *J Comp Physiol* [*A*] Vol., 91, 257-266.
- Olberg, R. M. (1993). Pheromone-triggered flip-flopping interneurons in the ventral nerve cord of the silkworm moth, Bombyx mori. *J Comp Physiol* [A] Vol., 152, 297-307.
- Posey, K. L., L. B. Jones, R. Cerda, M. Bajaj, T. Huynh, P. E. Hardin & S. H. Hardin (2001). Survey of transcripts in the adult Drosophila brain. *Genome Biol* Vol. 2, 3, RESEARCH0008.
- Pyk, P., S. Bermúdez i Badia, U. Bernardet, P. Knuesel, M. A. Carlsson, J. Gu, E. Chanie, B. S. Hansson, T. C. Pearce & P. F. M. J. Verschure (2006). An artificial moth: Chemical

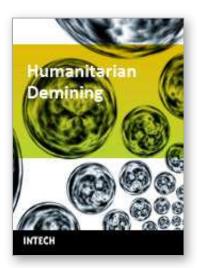
source localization using a robot based neuronal model of moth optomotor anemotactic search. *Autonomous Robots* Vol. 20, 3, 197-213.

- Quero, C., H. Y. Fadamiro & T. C. Baker (2001). Responses of male helicoverpa zea to single pulses of sex pheromone and behavioural antagonist. *Physiological Entomology* Vol. 26, 106-115.
- Reichardt, W. E. (1961). *Autocorrelation, a principle for evaluation of sensory information by the central nervous system,* John Wiley, New York.
- Rind, F. C. & P. J. Simmons (1992). Orthopteran DCMD neuron: a reevaluation of responses to moving objects. I. Selective responses to approaching objects. *J Neurophysiol* Vol. 68, 5, 1654-66.
- Rose, A., Z. Zhu, C. F. Madigan, T. M. Swager & V. Bulovic (2005). Sensitivity gains in chemosensing by lasing action in organic polymers. *Nature* Vol. 434, 7035, 876-9.
- Rowel, C. H. F. (1971). The orthopteran descending movement detector (DMD) neurones: a characterisation and review. *Z Vergl. Physiol* Vol., 73, 167-194.
- Santana, P. & J. Barata (2005). Unmanned Helicopters Applied to Humanitarian Demining, *Proceedings of Emerging Technologies and Factory Automation*, pp. 729-738.
- Srinivasan, M., S. Zhang, M. Lehrer & T. Collett (1996). Honeybee navigation en route to the goal: visual flight control and odometry. *J Exp Biol* Vol. 199, Pt 1, 237-44.
- Srinivasan, M., S. W. Zhang, J. S. Chahl, E. Barth & S. Venkatesh (2000). How honeybees make grazing landings on flat surfaces. *Biol Cybern* Vol. 83, 3, 171-83.
- Srinivasan, M. V., J. S. Chahl, W. K., S. Venkatesh, M. G. Nagle & S. W. Zhang (1999). Robot navigation inspired by principles of insect vision *Robotics and Autonomous Systems* Vol. 26, 2, 203-216.
- Strausfeld, N. J. (1976). Atlas of an Insect Brain, Springer-Verlag, New York.
- Surmann, H., A. Nüchter & J. Hertzberg (2003). An autonomous mobile robot with a 3D laser range finder for 3D exploration and digitalization of indoor environments. *Robotics and Autonomous Systems* Vol. 45, 3-4, 181-198.
- Tammero, L. F. & M. H. Dickinson (2002). Collision-avoidance and landing responses are mediated by separate pathways in the fruit fly, Drosophila melanogaster. *J Exp Biol* Vol. 205, Pt 18, 2785-98.
- Thorp, J. H. & K. S. Ammerman (1978). Chemical Communication and Agonism in the Crayfish, Procambarus acutus. *American Midland Naturalist* Vol. 100, 2, 471-474.
- Verhagen, R., F. Weetjens, C. Cox, B. Weetjens & M. Billet (2006). Rats to the Rescue: Results of the First Tests on a Real Minefield. *Journal of Mine Action* Vol. 9.2, http://www.maic.jmu.edu/journal/9.2/RD/verhagen/verhagen.htm.
- Vickers, N. J., and Baker, T.C. (1996). Latencies of behavioral response to interception of filaments of sex pheromone and clean air influence flight track shape in Heliothis virescens (F) males. *J. Comp. Physiol.* [A] Vol., 178, 831-847.
- Vickers, N. J. & T. C. Baker (1994). Reiterative responses to single strands of odor promote sustained upwind flight and odor source location by moths. *Proc. Natl. Acad. Sci. USA* Vol. 91, 13, 5756-60.
- Wada, S. & R. Kanzaki (2005). Neural control mechanisms of the pheromone-triggered programmed behavior in male silkmoths revealed by double-labeling of descending interneurons and a motor neuron. *J Comp Neurol* Vol. 484, 2, 168-82.

- Willis, M. A. & T. C. Baker (1984). Effects of intermittent and continuous pheromone stimulation on the flight behaviour of the oriental fruit moth, Grapholita molesta. *Physiological Entomology* Vol. 9, 341-358.
- Wyatt, T. D. (2003). *Pheromones and Animal Behaviour Communication by Smell and Taste,* University of Oxford.







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United Nation Department of Human Affairs (UNDHA) assesses that there are more than 100 million mines that are scattered across the world and pose significant hazards in more than 68 countries. The international Committee of the Red Cross (ICRC) estimates that the casualty rate from landmines currently exceeds 26,000 persons every year. It is estimated that more than 800 persons are killed and 1,200 maimed each month by landmines around the world. Humanitarian demining demands that all the landmines (especially AP mines) and ERW affecting the places where ordinary people live must be cleared, and their safety in areas that have been cleared must be guaranteed. Innovative solutions and technologies are required and hence this book is coming out to address and deal with the problems, difficulties, priorities, development of sensing and demining technologies and the technological and research challenges. This book reports on the state of the art research and development findings and results. The content of the book has been structured into three technical research sections with total of 16 chapters written by well recognized researchers in the field worldwide. The main topics of these three technical research sections are: Humanitarian Demining: the Technology and the Research Challenges (Chapters 1 and 2), Sensors and Detection Techniques for Humanitarian Demining (Chapters 3 to 8), and Robotics and Flexible Mechanisms for Humanitarian Demining respectively (Chapters 9 to 16).

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