

MEMORY-BASED IMMUNE NETWORK FOR MULTI-ROBOT COOPERATION

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ABSTRACT

In this paper, basic biological immune systems and their responses to external elements to maintain an organism's health state are fully described. The relationship between immune systems and multi-robot systems are also discussed. Our proposed algorithm is based on immune network theories that have many similarities with the multi-robot systems domain. The paper describes a memory-based immune network that enhance a robot's action-selection process and can obtain an overall a quick group response. The algorithm which is named as *Immune Network T-cell-regulated—with Memory* (INT-M) is applied to the dog and sheep scenario. Simulation experiments are being conducting on the Player/Stage platform and experimental data are being evaluated.

INTRODUCTION

Usually mobile robots need to interact and engage with one another in order to achieve assigned tasks more efficiently. These autonomous multi-robot systems would be highly beneficial in assisting humans to complete suitable tasks. In such systems, distributed intelligence is highly needed in the team whereby decisions are processed in each individual robots (Parker 1998). Furthermore, these robots would need to have the mechanism to cooperate so that they would achieve the assigned task (Cao et al. 1997).

Biological systems are examples of distributed information processing that are capable of solving problems in living organisms in a distributed manner. Some of these biological systems have neural networks in the brain that is capable of processing information through impulses at the synapses, genetic systems in constructing the organism genes and immune systems which protect and

maintain the homeostatic state of the living organism. Biological immune systems are particularly interesting, not only because they have no central processing but also exhibit cooperative capability among the antibodies in maintaining the internal stable environment of the body.

This leads to the advances in research on Artificial Immune Systems (AIS) and the application of AIS in engineering fields particularly in Multi-Robot Systems (MRS) domain (Cao et al. 1997, Li et al. 2007, Parker 1998). Situations faced by multi-robot systems require real-time processing and response. Furthermore, such situations would also require these systems to be robust to changes in the environment and some unexpected events, such as failure of robots in the team. Thus, mimicking the biological immune system is appropriate.

This paper proposes a memory-enhanced immune system algorithm to achieve cooperative behavior in a team of robots. Using the algorithm inspired by the immune network theory, the robots have the capability of performing their mission in a dynamically changing environment. The proposed algorithm is applied to the dog and sheep scenario (Li et al. 2007, Schultz et al. 1996). Simulation experiments are arranged to investigate the proposed algorithm using the above scenario.

The proposed approach would be suitable in various application domains such as military, disaster rescue operation and even service robotics in domestic environment. These example applications usually require several robots and have continuously changing environments. The test scenario can be extended for the chosen application domain.

BACKGROUND

This section explains the principle of the biological immune response and the Idiotypic Network Hypothesis which describe the cooperative behaviour achieved by

immune systems in vertebrate organisms. This is followed by the generic relation between immune systems and multi-robot systems.

Biological Immune Systems

Immune system is a system that eliminates foreign substances from an organism’s body. These foreign substances such as bacteria, fungi or virus cells that can harm the host are called pathogens. When such substance activates an immune response it is called *antigen*, which stimulates the system’s antibody generation. Each antigen has a unique set of identification on its surface called *epitope*. These antigenic determinants are where the host’s antibodies would attach to by using their *paratope*, as shown in Figure 1. *Antibodies* are cells in the immune system that kill antigens in order to maintain the host homeostatic state—i.e. balancing the body’s health status.

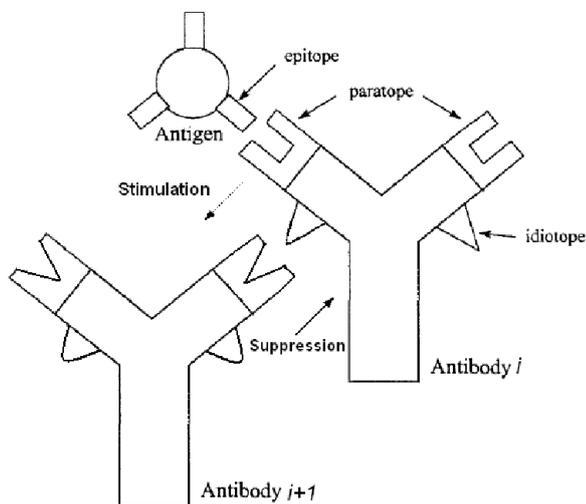


Figure 1: Antigen-antibody binding and Jerne’s Idiotypic Network Theory

The immune system can be divided into two general categories, innate immunity and adaptive immunity. Innate immunity is the first line of defense of the immune system. Generic pathogens that can be recognized and killed by the innate immunity cells would not be able to harm the host further. However, certain disease carrying antigens would bypass this defense mechanism because the innate immunity does not adapt to antigens that originate from various types of illnesses. The adaptive immunity would then play its role through the use of *lymphocytes* which are generally known as white blood cells. Lymphocytes have two main types, *T-cells* that mainly help in recognizing antigen cells and *B-cells* that mainly produce antibodies to fight specific antigens. In humans, T-cells are primarily produced in the thymus while B-cells are produced in bone marrows. These innate and adaptive immune responses make up effective

and important defense mechanism for living organisms.

Immune Response

The immune response basically can be viewed in six phases of recognition and activation, as seen in Figure 2. Pathogen is digested by Antigen Presenting Cells (*APCs*) where it is broken down into *peptides* (de Castro and Timmis 2002). These peptides will then bind to Major Histocompatibility Complex (*MHC*) molecules, then presented on the APC surface. T-cells recognize these different APC receptors and thus become activated. They divide and release *lymphokines* that transmit chemical signals to stimulate other immune system components to take action. B-cells would then travel to the affected area and be able to recognize the antigen. This would activate the B-cells which then mature into *plasma cells*. Plasma cells are the ones which release specific antibody molecules that neutralize the particular pathogens.

This immune response cycle results in the host’s immunity against the antigen which triggers it, thus having protection in future attacks (de Castro and Timmis 2002). Prominent characteristics of the immune system is that there is no central control of the lymphocytes in fighting antigens that invade the host and the system’s adaptability in responding to various kind of antigens. The B-cells cooperatively merge at the affected area and produce appropriate antibodies for that particular situation. This phase of immune response exhibits cooperative behavior of the related cells.

Idiotypic Network Hypothesis

Studies in immunology have suggested that antibodies are not isolated but they ‘communicate’ with each other. Each type of antibody has its specific *idiotope*, an antigenic determinant as shown in Figure 1. Jerne (1984) who is an immunologist proposed the Idiotypic Network Hypothesis (also known as Idiotypic Network Theory) which views the immune system as a large-scale closed system consisting of interaction of various lymphocytes (i.e. B-cells). Referring to Figure 1, idiotope of antibody *i* stimulates antibody *i+1* through its paratope. Antibody *i+1* views that idiotope (belonging to antibody *i*) simultaneously as an antigen. Thus, antibody *i* is suppressed by antibody *i+1*. These mutual stimulation and suppression chains between antibodies form a controlling mechanism for the immune response (de Castro and Timmis 2002).

Farmer et al. (1987) proposed differential equations of Jerne’s idiotypic network theory. These equations consist of antibodies’ stimulus and suppression terms, antigen-antibody affinity, and cell’s natural mortality rate (Farmer et al. 1987). This large-scale closed system

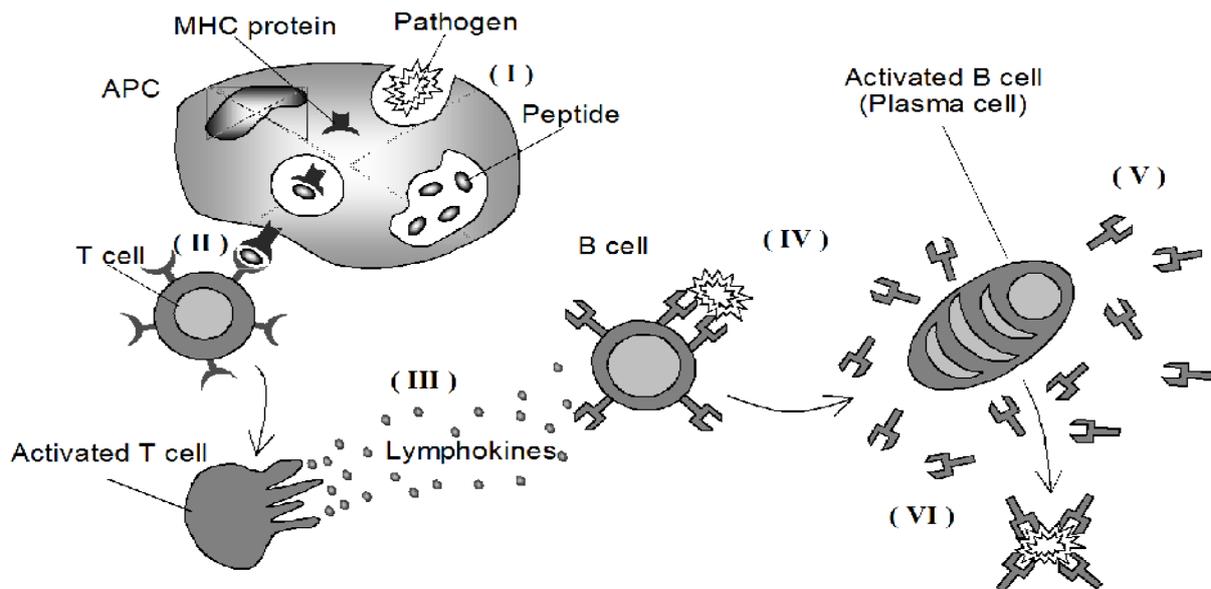


Figure 2: Basic biological immune systems response (de Castro and Timmis 2002)

interaction is the main mechanism that can be used for cooperation of multi-robot systems.

Immune Systems and MRS

The relationship of the immune systems with multi-robot systems is evident where obstacles, robots and their responses are antigens, B-cells and antibodies respectively. Table 1 lists the obvious parallel of MRS and immune systems terminologies.

Table 1: Immune Systems and MRS relationship

Immune Systems	Multi-Robot Systems
B-cell	Robot
Antigen	Robot's Environment
Antibody	Robot's action
T-cell	Control parameter
Plasma cell	Excellent robot
Inactivated cell	Inferior robot
Immune network	Robots interaction
Stimulus	Adequate robot stimulation
Suppression	Inadequate robot stimulation

Immune network theory as previously described is suitable as a basis for emulating cooperative behavior in a multi-robot environment. This is because the immune network uses affinity measures that are dependent on other cells concentration and location in determining the next action. Other than that, multi-robot systems require recognition ability of obstacles and other robots, which is parallel to the immune system recognition and activation phase of an immune response. Obviously, in immune network the processing of information is done

in real-time and in a distributed manner—as what a multi-robot system requires.

IMMUNE NETWORK BASED MULTI-ROBOT COOPERATION

Sun et al. (2001) have proposed a model based on Farmer's immune network equation that involves T-cells as control parameter which provides adaptation ability in group behaviour.

The group control or coordination phase is done in a distributed manner via local communication between nearby robots. When a robot encounters other robot and both have the same or similar strategy, this strategy is stimulated; if not, the strategy is suppressed. This facilitates the group to self-organize towards a common action which is optimal for the local environment. If a robot is stimulated beyond a certain threshold—which makes it an excellent robot, its behaviour is regarded as adequate in the system such that it can transmit its strategy to other inferior robots. This is a metaphor of the plasma cell in the biological immune systems.

The advantage of adding the T-cell model is that the system adapts quickly to the environment by recovery of antibody concentration to the initial state, when antigens have successfully been removed. Thus, the system is more adaptable to environmental changes.

Our proposed approach is based on Sun et al. (2001) work, with the extension of Memory ability so that quick responses can be achieved in certain relevant situation.

IMMUNE NETWORK WITH MEMORY

In biological immune response, there is a Clonal Selection process, whereby various B-cells try to identify the antigen. Once the appropriate B-cell is selected, it is activated and multiplied (i.e. proliferate) so that adequate immune response could be mounted later. The activated B-cells will proliferate and differentiate into Plasma cells that will secrete specific antibodies and memory cells which will be in the host body for quite a long time (de Castro and Timmis 2002). These memory cells will act as catalysts in mounting a quick immune response to the same antigen in the future.

The INT-M Model

In order to improve the approach by Sun et al. (2001), a specific memory mechanism is proposed in order to retain the appropriate action for relevant environment condition. This mechanism is introduced when the newly sensed environment is similar to the previous environment. Thus, a quick action-selection process can be executed without the need of re-evaluating the new situation.

The approach is aptly named as *Immune Network T-cell-regulated—with Memory* (INT-M) as it involves modelling the memory part of the biological immune systems. The general algorithm is shown in algorithm 1 which is an extension of Sun et al. (2001). The algorithm being displayed is for each robot in the group, and uses Equations (2), (3) and (4).

$$A(t) = \sum_{j=0}^{N-1} (m_{ij} - m_{ji}) s_j(t-1) \quad (1)$$

$$S_i(t) = S_i(t-1) + \left(\alpha \frac{A(t)}{N} + \beta g_i - c_i(t-1) - k_i \right) s_i(t-1) \quad (2)$$

$$s_i(t) = \frac{1}{1 + \exp(0.5 - S_i(t))} \quad (3)$$

$$c_i(t) = \eta(1 - g_i(t)) S_i(t) \quad (4)$$

In equations (2) and (3), $S_i(t)$ is the stimulus value of antibody i where $i, j = 0 \dots N$, N is the number of antibody types. m_{ij} is the mutual stimulus of antibody i and j , which is referred to in Table 2. g_i is the affinity of antibody i and antigen, which can arbitrarily be assigned using a function. $s_i(t)$ is the concentration of antibody i . The difference with Farmer et al. (1987) immune network equation is that $s_j(t)$ is not the concentration of self-antibody, but that of other robot's antibody obtained by communication.

Table 2: Mutual stimulus coefficient, m_{ij}

robot $i \setminus$ robot j	Ab₀	Ab₁	Ab₂	Ab₃
Aggregation, Ab₀	1	-0.4	-0.2	-0.4
Search, Ab₁	-0.4	1	-0.4	-0.2
Dispersion, Ab₂	-0.2	-0.4	1	-0.4
Homing, Ab₃	-0.4	-0.2	-0.4	1

Equation (4) is the T-cell model whereby $c_i(t)$ is the concentration of T-cell which controls the concentration of antibody i . α , β , and η are constants, whereby α and β are parameters of response rate of other robot and the environment (antigen) respectively. In biological immune systems, helper T-cells activate B-cells when antigen invades, and suppressor T-cells prevent the activation of B-cells when the antigen has been eliminated thus ensuring that the system adapts quickly to the environment by recovery of antibody concentration to the initial state.

Equations (5) and (6) are the functions and its corresponding values for the upper ($\bar{\tau}$) and lower ($\underline{\tau}$) thresholds in determining whether a robot becomes an excellent (i.e. plasma cell) or an inferior (i.e. inactivated cell) robot.

$$\bar{\tau} = \frac{1}{1 + e^{-0.5}} = 0.622 \quad (5)$$

$$\underline{\tau} = \frac{1}{1 + e^{0.5}} = 0.378 \quad (6)$$

SIMULATION

In this research we investigate shepherding behavior of robots. Shepherding behavior is similar to a flocking behavior but having agents/robots outside of the flock guiding or controlling the members (Lien et al. 2004). Figure 3 shows the screenshot of the dog and sheep scenario.

In a dog and sheep problem, a few dogs try to guide a few sheep to the grazing site (also called the safety zone) without going beyond the borders (Schultz et al. 1996). Dogs are required to cooperate in shepherding the sheep which are moving away from the dogs or wandering randomly inside the area. The objective is to prevent the sheep from going out of the grazing site while having partial information of what is happening in the area.

This problem is highly dynamic and obviously requires the robots to have real-time processing of partial information of the environment. The robot dogs use the proposed immune-inspired approach in cooperating with one another while the robot sheep have basic avoidance and flocking behaviours.

Algorithm 1 Immune Network T-cell-regulated—with Memory (INT-M)

Require: $t = 0$, $S_i(0) = s_i(0) = 0.5$ for $i = 0 \dots N - 1$,
 N is number of actions

Ensure: retain previous Ab if robot is not inferior within similar environment, execute Ab_{max}

$Ab_{max} \leftarrow Ab_1$
robot \leftarrow inferior
environment \leftarrow similar

loop

Execute Ab_{max}

{robot is activated (normal) or excellent}

if robot \neq inferior **then**

{environment sensed is *similar* to previous}

if $g_i(t) \approx g_i(t - 1)$ **then**

$S_i(t) \leftarrow S_i(t - 1)$

$s_i(t) \leftarrow s_i(t - 1)$

$c_i(t) \leftarrow c_i(t - 1)$

else

environment \leftarrow changed

end if

end if

{robot is inferior or environment has changed}

if (robot=inferior) || (environment=changed) **then**

for $i \leftarrow 0$ to $N - 1$ **do**

Calculate $S_i(t)$

Calculate $s_i(t)$

Calculate $c_i(t)$

end for

if $S_i(t) > \bar{\tau}$ **then**

robot \leftarrow excellent

else if $S_i(t) < \underline{\tau}$ **then**

robot \leftarrow inferior

if robot encounter $robot_{excellent}$ **then**

for all i **do**

receive Ab_i

renew $s_i(t)$

end for

end if

end if

end if

if Ab_i has $max(s_i(t))$ **then**

$Ab_{max} \leftarrow Ab_i$

end if

$t \leftarrow t + 1$

end loop

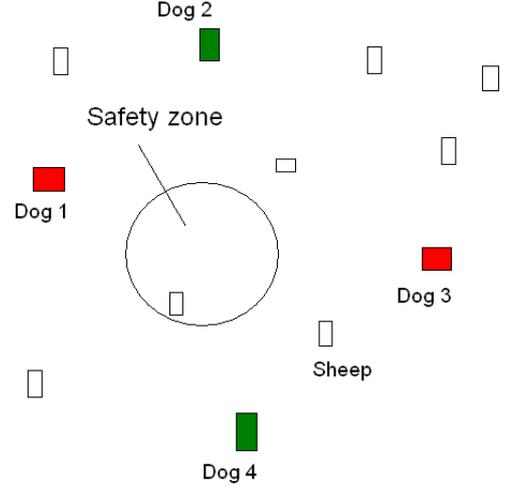


Figure 3: The Dog and Sheep problem environment

The proposed approach as described in algorithm 1 is applied to the dog and sheep problem and adjusted where necessary. The Player/Stage simulation platform (Gerkey et al. 2003) on a Fedora Core 6 Linux operating system is being used to test the proposed algorithm. Figure 4 shows a sample screenshot of the simulation platform. Experimental data is currently being collected to analyze the behaviours of the simulated robots.

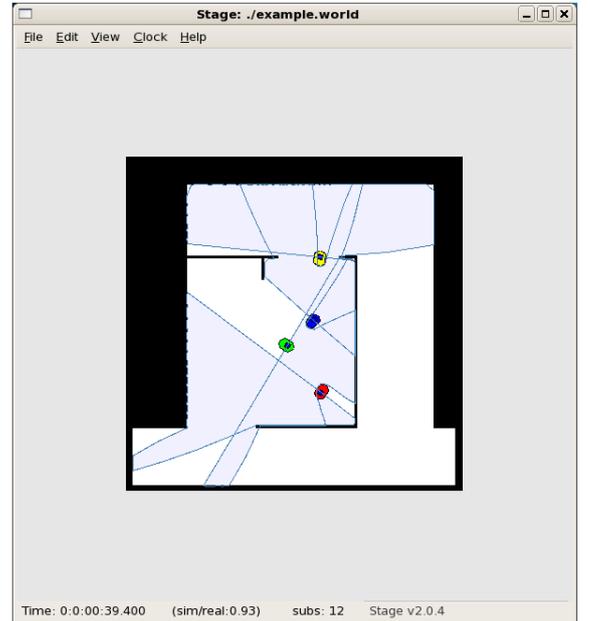


Figure 4: The Player/Stage simulation platform

A distinct part of this study is that we are looking into the memory-based immune network cooperation approach by the robots (i.e. dogs) in maintaining the herd (i.e. sheep). This utilizes the advantage of mem-

ory in the action-selection phase and affects the resulting dynamic behaviour of both the robot dogs and the robot sheep. The dog-sheep scenario partly contains the pursuit-evasion problem and can be further applied to other robot coordination problems, such as robot formation and the likes.

CONCLUSIONS

In this paper a memory-based immune system inspired approach for cooperation in multi-robot systems has been proposed. We have described the basic concepts and mechanisms of biological immune systems, and argued that the immune network is a suitable analogy for multi-robot cooperation problem. We have also proposed a multi-robot cooperation algorithm—the INT-M model, and applied to the dog-sheep test scenario. An experimental simulation environment has been setup to evaluate the proposed approach and algorithm. The approach can be easily extended to other application domains which require several agents (robots) to work cooperatively in a distributed way in a dynamic environment.

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