Dendritic Cell Algorithm*

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Abstract

The Clever Algorithms project aims to describe a large number of Artificial Intelligence algorithms in a complete, consistent, and centralized manner, to improve their general accessibility. The project makes use of a standardized algorithm description template that uses well-defined topics that motivate the collection of specific and useful information about each algorithm described. This report describes the Dendritic Algorithm using the standard algorithms template.

Keywords: Clever, Algorithms, Description, Optimization, Dendritic, Cell, Algorithm

1 Introduction

The Clever Algorithms project aims to describe a large number of algorithms from the fields of Computational Intelligence, Biologically Inspired Computation, and Metaheuristics in a complete, consistent and centralized manner [1]. The project requires all algorithms to be described using a standardized template that includes a fixed number of sections, each of which is motivated by the presentation of specific information about the technique [2]. This report describes the Dendritic Cell Algorithm using the standard algorithms template.

2 Name

Dendritic Cell Algorithm, DCA

3 Taxonomy

The Dendritic Cell Algorithm belongs to the field of Artificial Immune Systems, and more broadly to the field of Computational Intelligence. The Dendritic Cell Algorithm is the basis for extensions such as the Deterministic Dendritic Cell Algorithm (dDCA) [4]. It is generally related to other Artificial Immune System algorithms such as the Clonal Selection Algorithm, and the Immune Network Algorithm.

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4 Inspiration

The Dendritic Cell Algorithm is inspired by the Danger Theory of the mammalian immune system, and specifically the role and function of dendritic cells. The Danger Theory was proposed by Matzinger and suggests that the roles of the acquired immune system is to respond to signals of danger, rather than discriminating self from non-self [9, 10]. The theory suggests that antigen presenting cells (such as helper T-cells) activate an alarm signal providing the necessarily costimulation of antigen-specific cells to respond. Dendritic cells are a type of cell from the innate immune system that respond to some specific forms of danger signals. There are three main types of dendritic cells: 'immature' that collect parts of the antigen and the signals, 'semimature' that are immature cells that internally decide that the local signals represent safe and present the antigen to T-cells resulting in tolerance, and 'mature' cells that internally decide that the local signals represent danger and present the antigen to T-cells resulting in a reactive response.

5 Strategy

The information processing objective of the algorithm is to prepare a set of mature dendritic cells (prototypes) that provide context specific information about how to classify normal and anomalous input patterns. This is achieved as a system of three asynchronous processes of 1) migrating sufficiently stimulated immature cells, 2) promoting migrated cells to semi-mature (safe) or mature (danger) status depending in their accumulated response, and 3) labeling observed patterns as safe or dangerous based on the composition of the sub-population of cells that respond to each pattern.

6 Procedure

Algorithm 1 provides pseudo-code for training a pool of cells in the Dendritic Cell Algorithm, specifically the Deterministic Dendritic Cell Algorithm. Mature migrated cells associate their collected input patterns with anomalies, whereas semi-mature migrated cells associate their collected input patterns as normal. The resulting migrated cells can then be used to classify input patterns as normal or anomalous. This can be done through sampling the cells and using a voting mechanism, or more elaborate methods such as a 'mature context antigen value' (MCAV) which is $\frac{M}{Ag}$ (where M is the number of mature cells with the antigen and Ag is the sum of the exposures to the antigen by those mature cells), which gives a probability of a pattern being an anomaly.

7 Heuristics

- The Dendritic Cell Algorithm is not specifically a classification algorithm, it may be considered a data filtering method for use in anomaly detection problems.
- The canonical algorithm is designed to operate on a single discrete, categorical or ordinal
 input and two probabilistic specific signals indicating the heuristic danger or safety of the
 input.
- The danger and safe signals are problem specific signals of the risk that the input pattern is an anomaly or is normal, both typically $\in [0, 100]$.
- The danger and safe signals do not have to be reciprocal, meaning they may provide conflicting information.

Algorithm 1: Pseudo Code for the Dendritic Cell Algorithm.

```
Input: InputPatterns, iterations_{max}, cells_{num}, MigrationThresh_{bounds}
    Output: MigratedCells
 1 ImmatureCells \leftarrow InitializeCells (cells_{num}, MigrationThresh_{bounds});
 2 MigratedCells \leftarrow 0;
 3 for i = 1 to iterations_{max} do
        P_i \leftarrow \texttt{SelectInputPattern}(\texttt{InputPatterns});
        k_i \leftarrow (Pi_{danger} - 2 \times Pi_{safe});
 5
        cms_i \leftarrow (Pi_{danger} + Pi_{safe});
 6
        foreach Cell_i \in ImmatureCells do
 7
            UpdateCellOutputSignals(Cell_i, k_i, cms_i);
 8
            StoreAntigen(Cell_i, Pi_{antigen});
 9
            if Celli_{lifespan} \leq 0 then
10
                ReInitializeCell(Cell_i);
11
            else if Celli_{csm} \geq Celli_{thresh} then
12
                RemoveCell(ImmatureCells, Cell_i);
13
14
                ImmatureCells \leftarrow CreateNewCell(MigrationThresh_{bounds});
                if Celli_k < 0 then
15
                    Celli_{type} \leftarrow \mathsf{Mature};
16
17
                else
                     Celli_{type} \leftarrow Semimature;
18
                end
19
                MigratedCells \leftarrow Cell_i;
20
            end
\mathbf{21}
        end
22
23 end
24 return MigratedCells;
```

- The system was designed be used in real-time anomaly detection problems, not just static problem.
- Each cells migration threshold is set separately, typically $\in [5, 15]$

8 Code Listing

Listing 1 provides an example of the Dendritic Cell Algorithm implemented in the Ruby Programming Language, specifically the Deterministic Dendritic Cell Algorithm (dDCA). The problem is a contrived anomaly-detection problem with ordinal inputs $\in [0, 50]$, where values that divide by 10 with no remainder are considered anomalies. Probabilistic safe and danger signal functions are provided, suggesting danger signals correctly with P(danger) = 0.70, and safe signals correctly with P(safe) = 0.95.

The algorithm is an implementation of the Deterministic Dendritic Cell Algorithm (dDCA) as described in [11, 4], with verification from [7]. The algorithm was designed to be executed as three asynchronous processes in a real-time or semi-real time environment. For demonstration purposes, the implementation separated out the three main processes and executed the sequentially as a training and cell promotion phase followed by a test (labeling phase).

```
def random_vector(search_space)
return Array.new(search_space.length) do |i|
search_space[i][0] + ((search_space[i][1] - search_space[i][0]) * rand())
end
end
```

```
6
    def construct_pattern(class_label, domain, p_safe, p_danger)
7
     set = domain[class_label]
8
     selection = rand(set.size)
9
     pattern = {}
10
     pattern[:class_label] = class_label
11
     pattern[:input] = set[selection]
12
     pattern[:safe] = (rand() * p_safe * 100)
13
     pattern[:danger] = (rand() * p_danger * 100)
14
     return pattern
15
    end
16
17
    def generate_pattern(domain, p_anomaly, p_normal)
18
     pattern = nil
19
     if rand() < 0.5
20
21
       pattern = construct_pattern("Anomaly", domain, 1.0-p_normal, p_anomaly)
       puts ">Generated Anomoly [#{pattern[:input]}]"
22
23
     else
24
       pattern = construct_pattern("Normal", domain, p_normal, 1.0-p_anomaly)
25
     end
26
     return pattern
27
28
    def initialize_cell(thresh, cell={})
29
     cell[:lifespan] = 100.0
30
31
     cell[:k] = 0.0
     cell[:cms] = 0.0
32
     cell[:migration_threshold] = thresh[0] + ((thresh[1]-thresh[0]) * rand())
33
34
     cell[:antigen] = {}
     return cell
35
    end
36
37
    def store_antigen(cell, input)
38
     if cell[:antigen][input].nil?
39
40
       cell[:antigen][input] = 1
41
42
       cell[:antigen][input] += 1
     end
43
44
    end
45
    def expose_cell(cell, cms, k, pattern, threshold)
46
     cell[:cms] += cms
47
     cell[:k] += k
48
     cell[:lifespan] -= cms
49
     store_antigen(cell, pattern[:input])
50
51
      if cell[:lifespan] <= 0</pre>
       initialize_cell(threshold, cell)
52
     end
53
    end
55
   def can_cell_migrate?(cell)
56
    return (cell[:cms]>=cell[:migration_threshold] and !cell[:antigen].empty?)
57
    end
58
59
   def expose_all_cells(cells, pattern, threshold)
60
61
     migrate = []
62
     cms = (pattern[:safe] + pattern[:danger])
63
     k = pattern[:danger] - (pattern[:safe] * 2.0)
64
     cells.each do |cell|
       expose_cell(cell, cms, k, pattern, threshold)
65
       if can_cell_migrate?(cell)
66
         migrate << cell
67
         cell[:class_label] = (cell[:k]>0) ? "Anomaly" : "Normal"
68
```

```
end
69
70
      end
      return migrate
71
72
73
    def train_system(domain, max_iter, num_cells, p_anomaly, p_normal, threshold)
74
75
      immature_cells = Array.new(num_cells){ initialize_cell(threshold) }
76
      migrated = []
      max_iter.times do |iter|
77
        pattern = generate_pattern(domain, p_anomaly, p_normal)
78
        migrants = expose_all_cells(immature_cells, pattern, threshold)
79
        migrants.each do |cell|
80
          immature_cells.delete(cell)
81
          immature_cells << initialize_cell(threshold)</pre>
82
          migrated << cell
83
84
        puts "> iter=#{iter} new=#{migrants.size}, migrated=#{migrated.size}"
85
86
      end
87
      return migrated
88
    end
89
    def classify_pattern(migrated, pattern, response_size)
90
91
      input = pattern[:input]
      num_cells, num_antigen = 0, 0
92
      migrated.each do |cell|
93
        if cell[:class_label] == "Anomoly" and !cell[:antigen][input].nil?
94
95
          num cells += 1
96
          num_antigen += cell[:antigen][input]
97
        end
98
      end
      mcav = num_cells.to_f / num_antigen.to_f
99
      return (mcav>0.5) ? "Anomaly" : "Normal"
100
    end
101
102
    def test_system(migrated, domain, p_anomaly, p_normal, response_size)
103
      correct = 0
104
105
      100.times do
        pattern = construct_pattern("Normal", domain, p_normal, 1.0-p_anomaly)
106
107
        class_label = classify_pattern(migrated, pattern, response_size)
        correct += 1 if class_label == "Normal"
108
109
      end
      puts "Finished testing Normal inputs #{correct}/#{100} (#{correct}%)"
110
      correct = 0
111
      100.times do
112
        pattern = construct_pattern("Anomaly", domain, 1.0-p_normal, p_anomaly)
113
114
        class_label = classify_pattern(migrated, pattern, response_size)
        correct += 1 if class_label == "Anomaly"
115
      end
116
      puts "Finished testing Anomaly inputs #{correct}/#{100} (#{correct}%)"
117
118
119
    def run(domain, max_iter, num_cells, p_anomaly, p_normal, threshold, response_size)
120
      migrated = train_system(domain, max_iter, num_cells, p_anomaly, p_normal, threshold)
121
      test_system(migrated, domain, p_anomaly, p_normal, response_size)
122
    end
123
124
125
    if __FILE__ == $0
126
      domain = {}
      domain["Normal"] = Array.new(50){|i| i}
127
      domain["Anomaly"] = Array.new(5){|i| (i+1)*10}
128
      domain["Normal"] = domain["Normal"] - domain["Anomaly"]
129
      p_{anomaly} = 0.70
130
      p_normal = 0.95
131
```

```
iterations = 100
num_cells = 20
threshold = [5,15]
response_size = 10

run(domain, iterations, num_cells, p_anomaly, p_normal, threshold, response_size)
end
```

Listing 1: Deterministic Dendritic Cell Algorithm (dDCA) in the Ruby Programming Language

9 References

9.1 Primary Sources

The Dendritic Cell Algorithm was proposed by Greensmith, Aickelin and Cayzer describing the inspiring biological system and providing experimental results on a classification problem [6]. This work was followed shortly by a second study into the algorithm by Greensmith, Twycross, and Aickelin, focusing on computer security instances of anomaly detection and classification problems [8].

9.2 Learn More

The Dendritic Cell Algorithm was the focus of Greensmith's thesis, which provides a detailed discussion of the methods abstraction from the inspiring biological system, and a review of the technique's limitations [3]. A formal presentation of the algorithm is provided by Greensmith et al. [7]. Greensmith and Aickelin proposed the Deterministic Dendritic Cell Algorithm (dDCA) that seeks to remove some of the stochastic decisions from the method, reduce the complexity and to make it more amenable to analysis [4]. Stibor, et al. provide a theoretical analysis of the Deterministic Dendritic Cell Algorithm, considering the discrimination boundaries of single dendrite cells in the system [11]. Greensmith and Aickelin provide a detailed overview of the Dendritic Cell Algorithm focusing in the information processing principles of the inspiring biological systems as a book chapter [5].

10 Conclusions

This report described the Dendritic Cell Algorithm using the standard algorithms template. The description of the Deterministic Dendritic Cell Algorithm in the literature was found to be incomplete and ambiguous. As such, the current implementation of the algorithm in the Ruby programming language, while it executes, is considered incomplete. Similarly, the pseudo code listing may too require amendment. It is suggested that the author of the approach be contacted and as such, remains an area for further work.

11 Contribute

Found a typo in the content or a bug in the source code? Are you an expert in this technique and know some facts that could improve the algorithm description for all? Do you want to get that warm feeling from contributing to an open source project? Do you want to see your name as an acknowledgment in print?

Two pillars of this effort are i) that the best domain experts are people outside of the project, and ii) that this work is (somewhat) wrong by default. Please help to make this work less wrong by emailing the author 'Jason Brownlee' at jasonb@CleverAlgorithms.com or visit the project website at http://www.CleverAlgorithms.com.

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