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CONTEXT & 01 PROBLEM







CPPS & IIOT SECURITY



Antao, L., Pinto, R., Reis, J., & Gonçalves, G. (2018, April). Requirements for testing and validating the industrial internet of things. In ICSTW (pp. 110-115). IEEE. Pinto, R., Rossetti, R. J., & Gonçalves, G. (2016, July). Wireless sensor network simulation for fault detection in industrial processes. In SIMULTECH (pp. 1-6). IEEE.





\$3.1 BILLION

cyber attacks are launched against IoT devices each month.



The worldwide spending on IoT security is expected to reach this year

- The security of CPPS is crucial to the development and acceptance of the technology. As the IoT environment becomes complex, the resulting security issues present more challenges than any other existing network system.
- The limited computational power of COTS devices in CPPS restricts the usage of heavy security solutions.
- □ Security solution must be lightweight and Self-X* compliant.



- Prevention alone is no good, since no system can be full-prove to all attack vectors, especially considering zero-day attacks and inside threats.
- The autonomous and complex nature of CPPS does not comply with traditional IT security solutions.
- Intrusion Detection Systems may be used as a 2nd layer of protection.

Organizations use and rely on a security prevention tool

70%

Since 2018, the number of security incidents involving insiders has increased by

47%





Of security technology is expected to rely on artificial intelligence by 2023



55%

of enterprises plan to increase their cybersecurity spendings this year



- By focusing on ML detection solutions, IDS is limited in model training, online detection and real-time data processing.
- Develop of security solutions in CPPS is a complex and error-prone process. Security features should be included in the CPPS already in the design stage.
- Avoid the training phase, batch data processing and offline detection.
- Follow an Model-based Engineering approach using an industrial standard, such as the IEC 61499.



VALUE Proposition











- Since attackers are becoming more and more underhand in their techniques, which are increasingly sophisticated, data breaches are inevitable.
- □ It is extremely difficult to fix all the possible vulnerabilities of a system and stop all intrusions/attacks to happen.
- Prevention won't stop inside attackers.



📐 IDS TAXONOMY





SELF-IMMUNITY

 In the context of Autonomic Computing, a system is Self-immune when it recovers its safety properties in the face of an external (malicious) action. Ultimately, safety properties are no longer compromised when the same action occurs, acquiring tolerance over time.

All Self-immune systems are self-healing, but the reverse is not true.



APPLICATION OF AIS IN SMS

- AIS are attracting researcher attention in Autonomic Systems, Control Systems, Islanding, A-IDS, Fault Diagnosis and Operations Research areas.
- □ The main motivations are:
 - □ Handle large volumes of data & manage very complex systems
 - Generate Self-behaving systems and adaptability to changes
 - □ MAS-oriented architecture for dynamic, heterogeneous and distributed environments
- □ The industry adoption of the AIS technology is very slow. The average TRL of bio-inspired solutions in industry is between 3 and 4 (proof-of-concept and validation in lab), while the LOA is between 2 and 3 (partially/limited autonomy). Possible reasons may be:
 - □ The strong stochastic nature of the solutions makes them hard to assess, explain, and use in different contexts.
 - **Cost** & effort of the industrial systems and infrastructure transformation.





INSPIRED Protection

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IMMUNE SYSTEM

The immune system is a network of biological processes that protects an organism from diseases. It detects and responds to a wide variety of pathogens (viruses, bacteria, cancer cells...), distinguishing them from the organism's own healthy tissue.

There are two major subsystems of the immune system. The innate immune system provides a preconfigured response to broad groups of situations and stimuli. The adaptive immune system provides a tailored response to each stimulus by learning to recognize molecules it has previously encountered. Both use molecules and cells to perform their functions.



IMMUNE CELLS - LEUKOCYTES



IMMUNE PROCESS OVERVIEW

→ Negative Selection Algorithm

- \rightarrow Clonal Selection Algorithm
- → Artificial Immune Network
- → Danger Theory (Dendritic Cell Algorithm)



NEGATIVE Selection Algorithm

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a) Immature T cells with various antigen receptors are produced in bone marrow and migrate to the thymus.

b) In the thymus, cells that recognize self undergo apoptosis. Cells that don't recognize any self antigen are allowed to live.

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NEGATIVE SELECTION ALGORITHM

Algorithm 1: General Negative Selection Algorithm Detector Set Generation		
Input: S_{self} set of points that are normal		
Output: D detector set		
<pre>while not stop_condition() do</pre>		
2	$self_detected = True;$	
3	while self_detected do	
4	$d = generate_new_detector();$	
5	foreach s _i in S _{self} do	
6	if $detect(d, s_i)$ then	
7	break;	
8	end	
9	$self_detected = False;$	
10	end	
11	end	
12	$D \leftarrow d$	
13 end		
14 I	14 return D	



CLONAL Selection Algorithm



d.1) some cells differentiate to short-lived plasma B cells which release free antibodies to fight or inhibit the intruder. These B cells die shortly after.

d.2) other cells will differentiate into long-lived memory cells that will readily react to future occurrences of antigen with a similar signature

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CLONAL SELECTION ALGORITHM

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	Al	gorithm 2: General Clonal Selection Algorithm	
	Input: N population size		
	n	number of antibodies to clone	
	a	number of new antibodies to be generated each iteration	
	0	Dutput: P best population	
	1 F	$P \leftarrow generate_antibodies(N)$	
	2 V	<pre>while not stop_condition() do</pre>	
	3	foreach p in P do	
	4	update_affinity(p)	
	5	end	
	6	$P' \leftarrow select(P,n)$ foreach p' in P' do	
	7	$C \leftarrow clone(p')$	
	8	end	
	9	foreach c in C do	
	10	hypermutate(c)	
	11	end	
	12	foreach c in C do	
	13	update_affinity(c)	
	14	end	
	15	$P \leftarrow insert(C,n)$	
	16	$P_{new} \leftarrow generate_antibodies(d)$	
	17	$P \leftarrow replace(P, Pr)$	
	18 end		
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ARTIFICIAL IMMUNE NETWORK



Algorithm 3: A specific AIN implementation for generating an ARB population, based on AINE

Input: A input data (antigens) N population size NAT network affinity threshold R_m mutation rate N_c number of clones on each ARB Output: SARB set of all ARBs 1 $S_{ARB} \leftarrow initialize_population()$ 2 $C_{ARB} = \{\}$ 3 while True do foreach arb in SARR do 4 foreach a in A do 5 present_antigen(p) 6 end 7 8 end foreach arb in SARB do 9 update stimulation(arb) 10 11 end foreach arb in SARB do 12 allocate_b_cells(arb) 13 end 14 if stop_condition() then 15 16 break end 17 $C_{ARB} = clone_and_mutate(S_{ARB})$ foreach arb in C_{ARB} do 18 $S_{ARB} \leftarrow arb$ 19 end 20 21 end 22 return SARB

DANGER Theory

a) immature dendritic cells (iDC) navigate throughout the body, collecting antigens they encounter without descriminating whether the antigens come from self or non self.

b.1) iDCs also collect signals that originate from tissue cells. Cells that are damaged can chaotically release their contents which can be identified as danger signals by the neighboring iDC.

b.2) even if an organism doesn't harm the tissue, the iDC still collects these antigens, while also recognizing safe signals, such as those from programmed cell death (apoptosis)



c) after collecting a sufficient amount of danger or safe signals, the iDC matures into either a mature DC (mDC) or a semi mature DC (smDC), for dangerous or safe contexts, respectively. In both cases the DCs present their collected antigen to T cells. T cells that recognize the antigens proceed to initiate an immune response for mDCs, or tolerate the presented antigens for smDC.

DENDRITIC CELL ALGORITHM

Algorithm 4: DCA algorithm implemented in this section. This algorithm is adapted from J. Greensmith Begin End Input: Ag input data (antigens) N DC population size c number of times any antigen will be presented to distinct DCs tmigration migration threshold after which DC matures Initialize DCs t_{mcay} MCAV threshold above which the antigen is considered anomalous Population Output: Classification of antigens as normal or anomalous 1 foreach ag in Ag do $signals = compute_signals(ag)$ Antigen is 2 Sample Input Antigen is Sample Antigens Normal $DC_{sample} = sample_random(c)$ Signals Abnormal 3 foreach dc in DCsample do 4 /* update csm and k of each cell and store the antigen in Yes their repertoire Update expose(dc, ag, signals) 5 Cumulative if $dc.csm > t_{migration}$ then 6 **Output Signals** if dc.k > 0 then 7 No /* this will increment the count of mDC for all MCAV > At antigens that this cell had */ 8 mature(dc.MATURE) Upload DC Internal state else 9 mature(dc, SEMIMATURE) 10 11 end Yes 12 reset(dc) Calculate MCAV 13 end DC Lifespan > 0 14 end 15 end 16 No 17 foreach ag in Ag do $mcav = \frac{mDC_count(ag)}{mcav}$ 18 $mcav = \frac{c}{t_{mcav}}$ if $mcav > t_{mcav}$ then 19 Danger Yes Mature DC DC Migration classify(ag, OUTLIER) 20 Context? 21 else classify(ag,NORMAL) 22 23 No end Semi-Mature DC 24 end

FUTURE PERSPECTIVES



Enable Self-Immunity in industrial systems

- Enable automatic security features adjustment according to needs of the system to be protected.
- Achieve autonomous reconfiguration of the industrial Edge devices in case of an attack/failure detected.

□ Enhance the detection approach & performance

- Relating process host-related data with network-based data for a meaningful detection approach.
- Decentralized detection approach to enable a global self-awareness of the entire industrial system to be protected.

D Automatic security requirement validation, solution adjustment and deployment

- Assess the application context and, based on the specific requirements, adjust the detection technique and deploy the solution.
- **G** Scalability, to be used in multiple application scenarios and business segments.

THANKS!



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