

Bio-molecular cryptography for protecting nano-network transmissions in healthcare applications

Alessandra Rizzardi*, Giuseppe Piro^{†‡}, Sabrina Sicari^{*§}, Luigi Alfredo Grieco^{†‡}, Alberto Coen-Porisini*

*DISTA, Università degli Studi dell'Insubria, via O. Rossi 9 - 21100 Varese (Italy)

[†]DEI, Politecnico di Bari, via Orabona 4, 70125 Bari (Italy)

[‡]NIT, Consorzio Nazionale Interuniversitario per le Telecomunicazioni

[§]Corresponding author

Email: {sabrina.sicari; alessandra.rizzardi; alberto.coenporisini}@uninsubria.it

giuseppe.piro@poliba.it, alfredo.grieco@poliba.it

Abstract—Nano-technologies and communications at the nano-scale emerged as new approaches for data collection in several application domains, such as healthcare, bio-medicine, agro-food, industry, and military/defense. They usually entail a huge amount of continuous and real-time information to be processed from nano-devices, thus paving the way for the spreading of an innovation paradigm, represented by the Internet of Nano-Things (IoNT). The reliability of the envisioned IoNT applications still represents an open issue, due to complexity of protecting the transmitted data at the nano-scale. In this context, this work proposes to adopt bio-molecular cryptography to secure data exchanges among the nano-devices in nano-network, which leverages both molecular diffusion and electromagnetic-based communication protocols. Bio-molecular cryptography is essentially based on the DNA and proteins' properties. Starting from such premises, a cryptographic algorithm is developed and its performance is evaluated on a reference IoNT scenario in the healthcare domain, along with the study of the behavior of different routing algorithms.

Keywords—*Internet of Nano-Things, Nano-Networks, Cryptography, Security*

I. INTRODUCTION

In the last years, a novel networking paradigm entered heterogeneous application contexts, such as healthcare, bio-medicine and biomedical engineering, agro-food, industry, and military/defense domains [1]. Such an approach includes the presence of nano-technologies and it is known as *Internet of Nano-Things (IoNT)*. IoNT conceives the interconnection of nano-scale devices with existing communication architectures and, ultimately, with the Internet [2], by means of commonly used smart devices (i.e., gateways, smartphones, etc.). The final aim is to acquire useful data from the environment where the nano-devices are placed in, process them, and share the results in the form of services to the end-users. Note that IoNT essentially extends the functionalities of the well-known Internet of Things (IoT) concept to the nano-scale [3].

In general, *nano-devices* are able to perform both sensing and actuation tasks. Some of them may also act as *nano-routers*, thus becoming able to forward data to other devices, such as an access point, a sink node, a smartphone, and so on. A *nano-device* is expected to have limited capabilities. In such a scenario, which includes electromagnetic transmission and wi-fi communications, security and privacy emerge as

critical issues, mainly in presence of sensitive information [4]. Possible threats maDy include people's privacy violation (e.g., eavesdropping of medical information, alteration of vital parameters, home habits, localization), causes of pollution, food poisoning. Therefore, the design of a secure IoNT architecture must be carried out in order to: (i) ensure that information are efficiently protected against manipulation by non-authorized parties; (ii) guarantee data confidentiality, integrity, and availability; (iii) be accepted by users, which could not trust such a kind of technology. Furthermore, the solution must be as lightweight as possible, due to the aforementioned resources' constraints of *nano-devices*.

To support data confidentiality and integrity, the interaction between *nano-devices* and *nano-routers* must be properly protected by means of specific security functionalities. As pointed out in [5], invasion vectors can gain unauthorized access to the nano-network, exploiting system vulnerabilities, installing different types of malware, and launching cyber attacks. To this end, this paper proposes the adoption of bio-molecular end-to-end cryptography for encrypting the information acquired from the *nano-devices* and forwarded by the *nano-routers*, with the mediation of *nano-controllers*, which exploit molecular diffusion as a communication mean. The conceived algorithm is based on the DNA and proteins' properties [6]. Few mechanisms are until now proposed by the scientific community concerning the cryptographic operations at the nano-scale. This work aims to at least partially fill this gap by proposing a bio-molecular approach towards the realization of more complex cryptographic algorithms, which can be further developed and optimized in the next future. In fact, DNA cryptography represents an open research topic [7] [8] [9] [10] since many years. The impact of the investigated approach is evaluated through a case study in the healthcare domain; different routing algorithms have been taken into account, in order to analyze the solution, able to minimize the computational and communication overhead introduced by the new added cryptography algorithm.

The rest of the paper is organized as follows. Section II discusses the state of the art regarding nano-technology applications and security. Section III presents the architecture of an IoNT network, followed by Section IV, which is about the proposed bio-molecular cryptographic algorithm. Section V includes the performance evaluation of the presented approach. Section VI provides the conclusions and some hints

for future research directions.

II. RELATED WORK

The field of security and privacy in nano-technology is not still mature enough. Moreover, the interconnection and interoperability of the nano-devices with existing communication networks and paradigms require the design and development of new IoNT infrastructures and standards.

The authors of [11] coined the term biochemical cryptography to emphasize the need of new security and cryptographic solutions, in the field of nano-communications. In fact, actual approaches might be not applicable due to antenna size and channel limitations, as well as limited available memory and processing capabilities of the so-called nano-machines. Instead, in [12], the authors present an architecture for trusted remote sensing in *Public Physical Unclonable Function (PPUF)*, based on nano-technology. The proposed security protocol consists of authentication and time-stamping, in order to counteract statistical guessing attack. DNA origami is proposed in [13] to secure communications by means of some bio-molecular cryptography principles, based on protein binding-based steganography and self-assembled braille-like patterns.

The authors of [14] propose a DNA based cryptographic method, which consists of a symmetric algorithm based on the pseudo DNA cryptography and molecular biology. Splicing and padding techniques are combined with complementary rules to make the algorithm secure by means of an additional layer of security than conventional cryptographic techniques. The work presented in [15] proposes to improve the key strength by combining the genetic algorithm with the Diffie-Hellman key exchange algorithm; while a C++ algorithm for DNA cryptography is envisioned in [16], even if their feasibility is neither evaluated nor tested. However, important information is provided concerning the speed and storage of DNA computing. In particular, the authors state that a conventional computer can compute at the rate of 100 millions of instructions per second (MIPS) approximately, but experimentally it is found that DNA strands combinations are generated by combining DNA strands on computing at the rate of 109 MIPS or 100 times faster than a fastest computer. Instead, media storage requires 10-12 cubic nano-meter to store 1 bit, while DNA needs only 1 bit per cubic nano-meter, so very large amounts of data can be stored in compact volume.

The work in [17] proposes a modified lightweight DNA-based encryption method which employs the randomization nature of the DNA sequences to generate the encryption key: such an encryption key will be exploited to make the encryption process more simple and fast to accommodate the computations of IoT devices. Experiments have been conducted on encryption images processing. A similar evaluation is described in [18], where a lightweight DNA-based encryption method and the elliptic curve encryption (ECC) are coupled together to secure IoT communications. Instead, in [19], secure data storage is provided in the form of DNA sequences; more in detail, a DNA-based Cryptographic Security Framework incorporates a robust key agreement protocol and encryption algorithm, which has been integrated into CloudSim simulator. The claimed target domain is health cloud data, even if experiments do not present an e-health scenario.

As emerged from the analysis reported above, none of available solutions is specifically targeted to IoNT. Many works describe some methods related to DNA cryptography, but none of them verifies their behavior in a running nano-network, considering relevant network conditions, such as the routing protocol, the number of devices, and so on. The authors of [20] focuses on drug delivery in the human body, in a scenario where the drug molecule is protected with the help of protein cryptography and genetic signature. MATLAB is used for simulations, while this paper aims to represent a starting point for the future design and development of complex and effective cryptographic solutions by means of ad-hoc simulators, to provide security and privacy functionalities in different IoNT contexts.

III. IONT ARCHITECTURE

Nano-devices are equipped with nano-scale components, able to perform basic and specific tasks at nano-level, such as: sensing a “simple” information, actuating a “simple” action, computing basic operations (e.g., simple cryptographic functions), storing data with limited memory capacity, and communicating in short range. An IoNT system should be autonomous: once deployed, nano-devices must be able to self-act inside the environment. The environment hosting the nano-network should be closed (i.e., the network area must be delimited). It is worth to remark that, thanks to the nano-scale size of the devices, they can be massively deployed in a non-invasive way across a multitude of biological or environmental contexts, such as the human body. The reference architecture is depicted in Figure 1. It embraces:

- A *smart device*, which is not affected by power-constraints or resources’ limitations. It is responsible for collecting the information provided by the nano-network, reporting the activity of the IoNT system, and possibly taking some actions in response to the outcomes of the monitoring activities; the smart device is usually connected to the Internet.
- The *nano-routers*, which are in charge of forwarding data, when received by other nano-routers or nano-devices, to other nano-routers or to the smart device. The information exchange takes place by means of electromagnetic waves.
- The *nano-devices*, which are able to sense information from the environment where they are placed in, and, eventually, actuate some actions in response to specific commands received by nano-routers. Nano-devices send, following a hop-by-hop schema, the acquired data towards nano-routers by means of electromagnetic waves, which guarantee very high transmission throughput (i.e., in the order of Tbps) [21]; hence nano-devices encode messages through electromagnetic waves, sent in the terahertz band [22].
- The *nano-controllers*, which are particular nano-devices that transmit and receive information by means of molecular communications; their main goal is to add a control on the reliability of nano-devices’ behavior, as will be explained in Section IV-C.

A. Nano-Sim architecture and routing strategies

Nano-Sim is integrated within the *NS-3* simulation framework [23], and it has been originally conceived for modelling baseline IoNT architectures, where the communication

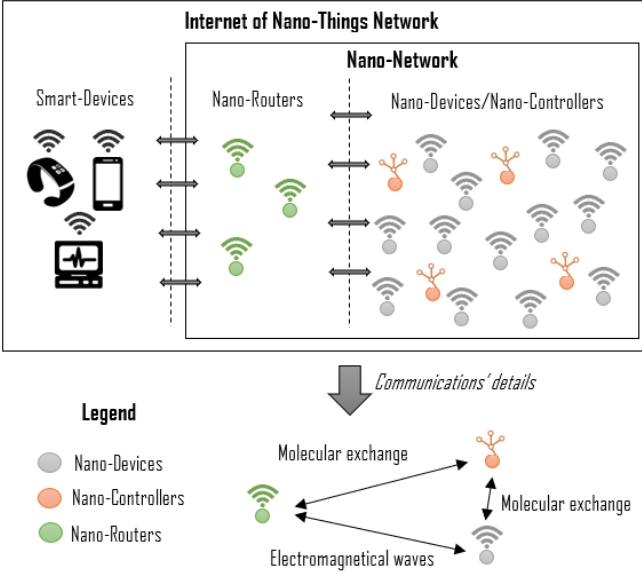


Fig. 1. The reference IoNT architecture

among nodes is handled, at the nano-scale, by means of electromagnetic waves generated in the terahertz band. *Nano-Sim* identifies each nano-component by means of a unique *Dev-ID*. The most important entities are: *Message Processing Unit*, *Network Layer*, *MAC (Media Access Control) Layer* and *Physical Interface (PHY)*.

The task of generating and processing messages is delegated to the *Message Processing Unit*. Such a module periodically (i.e., following a constant bit-rate) creates packets of fixed length, customizable by the developer. Each packet also contains some headers, including: (i) the *Dev-ID* of the owner/source of the data; (ii) the *Dev-ID* of the sender; (iii) the *Dev-ID* of the nano-device, which represents the next hop; (iv) an identifier of the packet (i.e., *Pack-ID*); (v) the *Time To Live (TTL)*. Once created a new message, the *Message Processing Unit* sends it through the protocol stack towards the *PHY*. When a new packet is successfully received by the network channel thorough the *MAC Layer*, it is further delivered to the *Network Layer* that will check the destination of the message itself. Then, the *Network Layer* routes the packet to the final recipient.

Due to the fact that nano-devices communicate with the *PHY* through very limited transmission ranges, multi-hop transmissions between the sender and the recipient are required. As a consequence, the adopted routing protocol must efficiently manage the multi-hop connections. *Nano-Sim* provides a flexible *Network Layer*, which supports two possible routing strategies:

- The *Selective Flooding Routing (SFR)* implies the sharing of the received packet with all the devices within its transmission range. In order to avoid bandwidth waste, it is important to avoid duplicates. Hence, every message can be uniquely identified by the couple (*Pack-ID*, source *Dev-ID*); then, if each node keeps in memory the couples (*Pack-ID*, source *Dev-ID*) associated with the last n messages received, each node will be not able to transmit packets already received.
- The *Random Routing (RR)* randomly selects the next nano-device from a set of neighbors. Hence, each node

knows the devices within its transmission range; for such a reason it must cooperate with a specific MAC strategy, which is *Smart-MAC*, in this work. Also, the *RR* algorithm stores the information related to the last packets received, in order to avoid loops within the nano-network.

Both the routing protocols (i.e., *SFR* and *RR*), in order to speed up the delivery of messages towards the smart device, require that the nano-devices preferably try to send the information to a nano-router, if available, as the next hop. Instead, due to nano-devices' energy constraints, the most promising modulation scheme for nano-communications is the *Time Spread On-Off Keying (TS-OOK)* [24]. TS-OOK offers two important advantages: (i) it does not require synchronization of nano-devices before starting packets' transmission; (ii) it allows sharing among multiple devices. TS-OOK parameters, such as the signal duration, the transmission, the frequency and the pulse energy can be optimized on the basis of the application scenario, as explained in Section V-A. Note that, since the most suitable transmission techniques for the nano-networks are based on impulse communications, the asynchronous MAC algorithms are the best candidates for a nano-network, because they allow the packets' transmission without the need for nodes to compete for the channel. Furthermore, thanks to the low computing requirements, asynchronous MAC schemes are easier to implement on nano-devices with limited resources. In line with such premises, two different asynchronous strategies have been defined at the MAC level:

- The *Transparent-MAC* transmits packets from the *Network Layer* to the *Physical Interface (PHY)* without performing any kind of check.
- The *Smart-MAC* does not immediately transmit the packets from the *Network Layer* to the *PHY*, but it stores them in a dedicated queue; before starting the transmission, the MAC layer takes advantage from an hand-shake procedure to discover all the nano-devices available in the transmission range of the nano-device owing the message. If at least one nano-device is found, the package is then sent to the *PHY*. If the *RR* algorithm is active and the next hop has not been selected yet, the choice will be delegated to the MAC layer. In case the sending nano-device has no "neighbors", the MAC layer will apply a random back-off delay, before restarting the hand-shake procedure. Such a delay is uniformly distributed in the interval (*MIN back-off time*, *MAX back-off time*), according to the application context.

IV. SECURITY FUNCTIONALITIES

The envisioned security functionalities and the threats model are detailed in this section, after briefly presenting the concepts behind the bio-molecular cryptography.

A. Bio-molecular cryptography

Bio-molecular cryptography is based on the use of biological information of living beings, which turn out to be unique and, therefore, suitable to be used to encrypt sensitive information. In this paper we will focus on the use of *Deoxyribose Nucleic Acid (DNA)* and its biological properties. The design of DNA-based bio-molecular encryption is a technique

that needs parallel processing and bio-molecular computing capacity in order to convert short messages, starting from a hexadecimal numeric system or from ASCII code. This work considers the bio-molecular properties of DNA sequences with the aim of generating keys that allow the protection of sensitive data contained in a packet, transmitted within the reference IoNT network. DNA processing techniques were conceived thanks to the work of Leonard Max Adleman [25]. The different phases will be set up to manipulate and encode the data, starting from the alignment of DNA with other biological languages up to protein building process. Protein formation from sequences of DNA goes through a well-defined process, and includes the addition of several catalysts and enzymes, while still ensuring the DNA integrity. DNA represents a double helix sequence of nucleotides, which determine the code of each gene, made up of four basic elements: *Adenine (A)*, *Guanine (G)*, *Cytosine (C)*, *Thymine (T)*. Although there are only four basic elements in the sequence, their arrangement is purely random and billions of combinations exist. The calculation, made using a DNA sequence, is called *DNA Computing*. Several issues, such as support on mobile devices for sending text, communication and video simultaneously [26], have been solved thanks to the use of parallel computing methods. James Dewey Watson [27] has succeeded to combine traditional cryptography techniques with DNA sequences and introduced a new concept of hybrid security, which can be now extended to the nano-technologies domain.

The complementary strands that generate the DNA are triplets of nucleotide codons represented as follows:

AGG CTC AAG TCC TAG
TCC CAG TTC AGG ATC

If DNA strands are mapped to numbers, alphabetic letters or other attributes, they can be used for coding and decoding information, as well as for digital data storage [28].

B. DNA algorithm in the IoNT network

DNA synthesis represents one of the central dogmas of biology and is the key to understand and operate within the bio-molecular encryption domain. The synthesis process involves a transition from DNA sequences to *mRNA* sequences finally going to get proteins. When two DNA strands are separated by an enzyme, a new strand, named *Messenger RNA (mRNA)*, is formed, and it is complementary to the DNA strand. The RNA filament is formed by mapping DNA sequences (A, T, C and G) with RNA complementary sequences, which consist of U (Uracil), A, G and C [29]. Figure 2 shows how basic DNA elements are copied in an mRNA strand, by replacing codon T with codon U. Finally, the combination of codons belonging to the mRNA determines the aminoacids' order to build a protein cell, whose scheme is sketched in Figure 3.

Information are secured through the use of the DNA sequence that characterizes the biological material containing the nano-device, by mapping the message with the basic elements of DNA, properly setup. In the proposed algorithm, a DNA strand is simulated by means of the following mapping between binary code and portions of DNA strand:

- 00 → "first portion of DNA strand"
- 01 → "second portion of DNA strand"
- 10 → "third portion of DNA strand"
- 11 → "fourth portion of DNA strand"

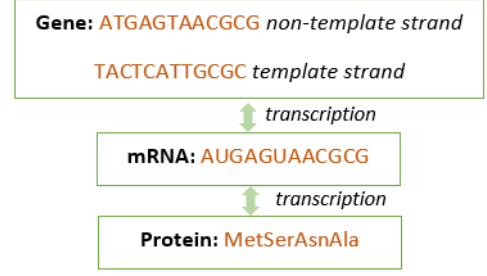


Fig. 2. Transformation process from mRNA to aminoacids

		Second letter →					
First letter →		U	C	A	G		
	U	UUU } Phe UUC } UUA } UUG } Leu	UCU } UCC } Ser UCA } UCG }	UAU } Tyr UAC } UAA } Stop UAG }	UGU } Cys UGC } UGA } Stop UGG } Trp	U C A G	Third letter →
	C	CUU } CUC } Leu CUA } CUG }	CCU } CCC } Pro CCA } CCG }	CAU } His CAC } CAA } Gln CAG }	CGU } CGC } Arg CGA } CGG }	U C A G	
	A	AUU } AUC } Ile AUA } AUG } Met	ACU } ACC } Thr ACA } ACG }	AAU } Asn AAC } AAA } Lys AAG }	AGU } Ser AGC } AGA } Arg AGG }	U C A G	
	G	GUU } GUC } Val GUA } GUG }	GCU } GCC } Ala GCA } GCG }	GAU } Asp GAC } GAA } Glu GAG }	GGU } GGC } Gly GGA } GGG }	U C A G	

Fig. 3. Coding table for DNA/RNA codons

The four different DNA portions are assigned by the nano-network taking at each time different portions of the currently involved DNA sequence; in this way, when a new data or data set must be transmitted, different portions are selected. Their nature depends on the particular IoNT context considered. In fact, if the analyzed field is that of healthcare, we can suppose that the sequences can be directly extracted from the individuals and kept different for all the people monitored by the same healthcare structure. Note that it is important that such sequences are known to whom must decrypt the information. Other fundamental aspects in the encryption process are: (i) the length of such sequences, since it affects the robustness of the security algorithm (i.e., it represents the length of the encryption key); (ii) the two seeds (i.e., *seed*₁ and *seed*₂) owned by each nano-device.

P _{DNA}	00	01	10	11
AAAA...	TGGA...	CTAA...	...	...
GAAA...	ACGT...	...	...	...
...	...	...	...	...

Fig. 4. Map for DNA associations

Several steps are required before reaching the cipher-text. The first step of the algorithm is characterized by the acquisition of sensitive data and their conversion into ASCII code and, subsequently, in binary code [30]. Once generated the binary code B_{code} , the DNA elements will be used to convert the message into usable DNA sequences, starting

from the aforementioned seeds. When a DNA portion P_{DNA} is extracted, then it is mapped with a corresponding DNA sequence P'_{DNA} , taken from a pre-defined map (containing all the possible combinations of DNA portions with the same length of P_{DNA}), associated with the current nano-network, in the position $[seed_1, \text{couple of binary values (i.e., 00, 01, 10, 11)}]$ from $B1_{code}$. The two seeds are in the range $[0 : (length_{P_{DNA}})!]$ and we assume that P_{DNA} and P'_{DNA} have the same length, even if this feature is not strictly mandatory. A scheme of the map is depicted in Figure 4. The subsequent DNA portion, namely $P2_{DNA}$, is mapped to the sequence positioned in $[seed_2, \text{couple of binary values } B2_{code}]$, and so on, until the end of B_{code} sequence. The two seeds change each time a data or a set of data which must be encrypted, by increasing their value of one unit $\% length_{P_{DNA}}$, in order to fit the range of the columns' number in the map. Once the DNA sequence is obtained, the biological synthesis can begin, which includes: (i) transformation of DNA into mRNA; (ii) mapping of the mRNA strand into aminoacids according to Figure 5.

Ala	A	GCU, GCC, GCA, GCG	Leu	L	UUA, UUG, CUU, CUC, CUA, CUG
Arg	R	CGU, CGC, CGA, CGG, AGA, AGG	Lys	K	AAA, AAG
Asn	N	AAU, AAC	Met	M	AUG
Asp	D	GAU, GAC	Phe	F	UUU, UUC
Cys	C	UGU, UGC	Pro	P	CCU, CCC, CCA, CCG
Gln	Q	CAA, CAG	Ser	S	UCU, UCC, UCA, UCG, AGU, AGC
Glu	E	GAA, GAG	Thr	T	ACU, ACC, ACA, ACG
Gly	G	GGU, GGC, GGA, GGG	Trp	W	UGG
His	H	CAU, CAC	Tyr	Y	UAU, UAC
Ile	I	AUU, AUC, AUA	Val	V	GUU, GUC, GUA, GUG
start		AUG, GUG	stop		UAG, UGA, UAA

Fig. 5. Lookup-table for the mapping of 20 "ordinary aminoacids"

Once completed the last step, the set of amino acids, which concur to the formation of proteins, is obtained. In order to complete the encryption of the obtained message, other status changes are needed: (i) conversion of the message formed by aminoacids into ASCII code; (ii) conversion to binary code. The result is the encrypted message, which is now ready for its transmission towards the nano-routers, along with the information associated by the nano controllers (see Section IV-C). No decryption is needed within the IoNT network itself, following an end-to-end approach in guaranteeing the confidentiality, the integrity, and the availability of the transmitted information from their acquisition by a nano-device to their reception to the smart device. Figure 6 shows the flow chart related to the status changes of the encryption algorithm and the reverse decoding process.

C. Interaction with nano-controllers

Molecular exchanges do not suffer of proximity attacks, since no electromagnetic waves are generated, but only molecular reactions. Hence, nano-controllers, which interact with nano-routers and nano-devices through molecular exchanges, are responsible for adding a control on the reliability of the nano-network during the time. Nano-controllers are properly configured in order to react in presence of nano-devices and nano-routers [31] [32] [33]. More in detail, each nano-controller is randomly disposed within the nano-network and is configured, before network deployment, with a unique identifier id_{nc} . Nano-controllers periodically generate

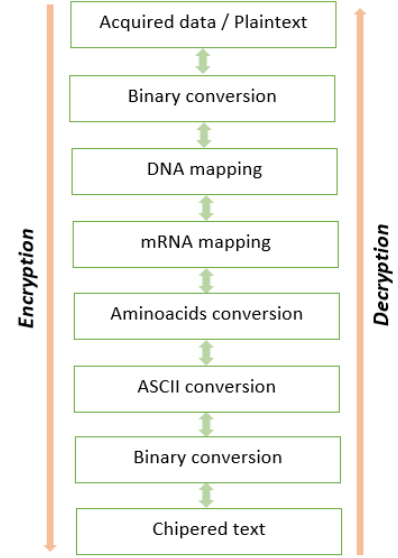


Fig. 6. Scheme of coding and decoding processes

a concatenation between the identifier and a sequence number seq , along with a lightweight symmetric key (key_{nc}). Such a couple of information (i.e., $[id_{nc}seq, key_{nc}]$) is transmitted encoded by the nano-controller itself and can only be decoded by nano-routers and nano-devices. In the former case, the nano-devices will use the gathered information to add a security token to the data to be sent to the nearer nano-router. In the latter case, the nano-routers will use the received token for assessing the reliability of the data received by the nano-devices. Further details on such a mechanism are available in [34].

D. Threat model

Considering the terahertz communications among nano-devices and among nano-devices and nano-routers, remote attacks, may occur. Although such a kind of attack does not require a physical contact, the attacker should be in the proximity of the nano-network, in order to reach the short-range signals emitted by nano-devices. It is not yet clear if an external entity may alter the behavior of nano-devices or if only communications may be affected. Undoubtedly, the nano-network could be a victim of spoofing and eavesdropping attacks. In this paper, we deal with the aforementioned security issues by means of the bio-molecular cryptography and molecular exchanges. Confidentiality, data integrity and access control are guaranteed thanks to unique mapping among the message and the DNA sequences, which are strictly related to the owner of the data acquired by the nano-devices, to the seeds and to the map used for DNA mapping. A further control on data reliability is also provided by nano-controllers, which add a security token to the transmitted information. As a consequence, even if a non authorized entity is able to trace the encrypted messages (e.g., MITM attack), it cannot decrypt the packets' content, since it should also be able to exact the DNA combination used to map the data in clear (transformed into binary code). Moreover, the decryption process takes place outside the nano-network system, not inside the biological system, in a protected environment (e.g., the smart device).

In case of packet sniffing, the attacker could recognize the number of transmitted packets, the messages' size, and the data transmission frequency, but no useful information can be gathered regarding the encryption/decryption algorithm. Furthermore, the nano-network, thanks to both the adopted routing algorithms, results to be fault tolerant; hence, if a nano-device depletes its resources and stops working, the packets are surely transmitted to the neighbour devices, thus guaranteeing their transmission to the final recipient.

Also, interactions among nano-routers and smart device must be protected from violations, as presented in [34], where a negotiation procedure between each nano-router and the smart device to obtain the proper session key to be used for the communication exchange, following the well-known certified *Diffie-Hellman* scheme, is detailed. In this way, the smart device and the nano-routers can compute the session key, via an ordinary/standard *Key Derivation Function (KDF)*, thus preventing both replay and *Man-In-The-Middle (MITM)* attacks. Finally, the smart device offers an interface between the nano-network and the rest of the Internet. Seamlessly, data confidentiality and integrity checks must be put in action, by means of conventional security protocol already used in typical IoT deployments [35].

V. PERFORMANCE ASSESSMENT

Despite the recent advances in the nano-technology field, evaluating the behavior of nano-networks through experimental tests is still difficult. In line with the current scientific literature, the system's performance are evaluated through computer simulations. The proposed encryption algorithm has been implemented in C++ programming language, inside *NanoSim*; all the measurements and collected data are obtained by means of a MacbookPRO with the following features: (i) 2.5 GHz quad-core Intel Core i7 processor; (ii) 16 GB RAM memory; (iii) 500 GB flash hard disk; (iv) macOS Catalina operating system. Since *NanoSim* is mainly able to provide networking related key performance indices, to assess the complexity of DNA computing related operations, another tool has been adopted, similarly named *NanoSim*¹. Implemented in Python, it provides pre-loaded data-sets, which can be freely downloaded from the tool's dedicated website and configured inside it. Instead, for modelling molecular exchanges, *N3Sim* [36] has been integrated.

A. Implemented scenarios and parameter settings

The study considers 30 mm long artery, with a diameter set to 1 mm [37]. Nano-devices are uniformly distributed in this bounded environment). Nano-devices periodically send the acquired information towards the nano-routers, across multi-hop paths. Nano-routers forwards the received data to a smart device by using the IEEE 802.11 wireless technology. Such a scenario perfectly fits within the e-health context, which is, for a few years, involved in the adoption of nano-technologies [38]. Examples of applications are drug delivery, gene delivery, molecular diagnostics, imaging, cardiac therapy, dental care, orthopedic applications, and so on. Regarding the protocol suite adopted for the electromagnetic-based communication inside *NanoSim*, three simulation's setups are considered, combining different routing logic (*routing-type*) and MAC (*mac-type*) procedures, as summarized in

Table I; while Table II shows the parameter settings for two different scenarios. Note that data-link and routing strategies are baseline protocols already implemented in *NanoSim*.

Specific information, concerning the complexity of DNA computing operations, are obtained from a separate analysis, conducted by means of *NanoSim*. The first step of *NanoSim* process is read characterization, which provides a comprehensive alignment-based analysis and generates a set of read profiles serving as the input to the next step, which is the simulation stage, where DNA reactions take place. The authors of the simulator [39] measured that the runtime of *NanoSim* scales up linearly with the number of reads and the memory requirement depends on the length of the reference sequence. For example, *NanoSim* requires 4 minutes and 39 seconds and a peak memory usage of 120 MB for processing 20,000 reads. Since we simulate a packet generation time interval equals to 0.5 seconds and simulation time is set to 5 seconds, then we expect to produce 10 reads per nano-device. As a consequence, the time required for processing the total amount of read approximately is 0,14 seconds per nano-device; while the required storage should not exceed 0.06 MB per nano-device. The delay value guarantees the efficiency of the IoNT network, also in presence of DNA computing operations, as shown in the following; while we assume that information are not persistently stored in nano-devices once transmitted, due to the excessive storage occupancy which could be generated during the time.

TABLE I. SIMULATION SETUP

Simulation	routing-type	mac-type
<i>Sim</i> ₁	SFR	Transparent-MAC
<i>Sim</i> ₂	SFR	Smart-MAC
<i>Sim</i> ₃	RR	Smart-MAC

TABLE II. SIMULATION SCENARIOS

Scenario	n. nano-devices	n. nano-routers	n. smart devices
<i>Sce</i> ₁	50	10	1
<i>Sce</i> ₂	200	50	1

Each simulation is repeated 5 times with a different initial seed, which affects the random placement of nano-devices within the artery. An overview of parameter settings, derivated from the study in [40], is reported in Table III.

B. Obtained results

The message processing overhead reports the delay introduced by the new added bio-molecular cryptography algorithm. Note that the delay in generating the encrypted message is evaluated taking into account the average time (in milliseconds), required for the execution of the encryption algorithm for each nano-device.

From Figure 7 the first analysis which can be carried out concerns the simulation *Sim*₂ applied to scenario *Sce*₁; in fact, the algorithm would seem to optimally work with the combination *Selective Flooding Routing - Smart-MAC*. Coming back to the same data, but considering the scenario *Sce*₁, a peak is immediately notice, as the time needed to encrypt the data is greater than those of all other simulations. Finally, simulation *Sim*₂ is not suitable for a system with the

¹NanoSim tool, <https://www.bcgsc.ca/resources/software/nanosim>

TABLE III. SIMULATION PARAMETERS

General parameters	Value
Simulation time	5 s
Number of seeds	5
Artery size	30 mm x 1 mm x 1 mm
Number of nano-devices	[50, 200]
Number of nano-routers	[10, 50]
Number of smart-devices	1
Details related to the electromagnetic-based communication channel	Value
Packet size	128 bytes
Packet generation time interval	0.5 s
Pulse energy	100 pJ
Pulse duration	100 fs
Pulse interval time	10 ps
Modulation scheme	TS-OOK
Backoff interval	[0 ns, 100 ns]
TTL	100
Details related to the molecular communication channel	Value
Modulation scheme	OOK
Nano-controllers communication range	0.02 nm
Diffusion coefficient	1.0 nm^2/ns
Control packet generation time interval in the molecular channel	0.5 s
Brownian Motion Factor	0.5
Inertia Factor	0.5

presence of many nano-devices, due to the fact that the delay in generating the encrypted message follows an exponential trend, which is proportional to the number of deployed nodes. Now, we consider simulations Sim_1 and Sim_3 . In a scenario including a reduced number of devices (e.g., Sce_1), the use of the bio-molecular encryption algorithm, combined with *SFR* or *RR*, and *Transparent-MAC* and *Smart-MAC*, respectively, substantially requires the same time. However, by increasing the amount of nano-devices involved in the network (i.e., Sce_2), we immediately note that the *RR* algorithm needs more processing time than the *SFR* algorithm. In fact, the hand-shake procedure applied by *Smart-MAC* requires much processing time with respect to the *Transparent-MAC* procedure, due to the fact that each nano-device, before sending a packet, must always check which other nano-devices are in their transmission range at that moment. More nano-devices are present in the network, longer the time required by the packet routing algorithm is, and this consequently causes a slowdown of the network.

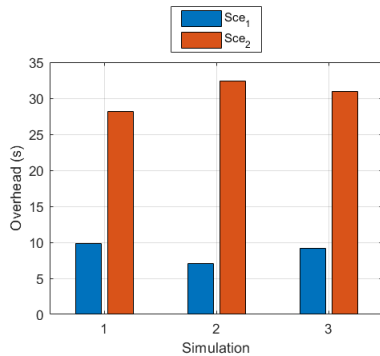


Fig. 7. Average delay of encrypted message generation

Figure 8 analyzes the average delay generated by a single nano-device in the nano-network to execute the whole mes-

sage processing, which includes encryption algorithm and transmission time. As the devices deployed in the network increase (i.e., Sim_2), the time that each nano-device requires to encrypt the data to be transmitted significantly increases, making the algorithm poorly performing in large IoNT networks, due to the fact that the nano-device is simultaneously involved in forwarding operations. On the other hand, in a scenario characterized by a reduced number of nano-devices, *SFR* combined with *Smart-MAC* turns out to be the best choice. Focusing the attention on simulation Sim_1 , we can see that, as the number of sensors deployed in the network increases, the time that each of them requires to process data results more optimized. Finally, simulation Sim_3 presents a proportional trend with respect to the number of nano-devices involved in the system; this is due to the fact that *Smart-MAC* maps each device to check if it is within the transmission range of the device that wants to transmit the packet.

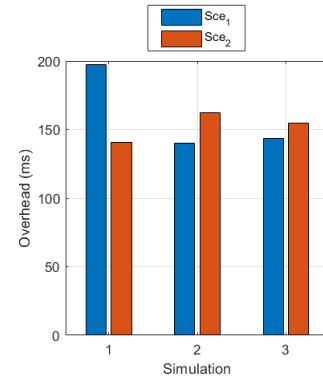


Fig. 8. Average delay of data processing per nano-device

Table IV summarizes the analysis carried out in relation to the delay, considering the execution of the whole application and the average delay generated per nano-device.

VI. CONCLUSIONS

IoNT paradigm is planned to be integrated in various edge devices, such as cell-phones, sensors, vehicles, household-appliances, etc. Hence, security functionalities represent significant open research problems. The paper introduced an end-to-end bio-molecular cryptography algorithm within an IoNT network. The proposed approach is based on the unique DNA coding properties, which offers many advantages, such as ultra-high storage density, ultra-low energy consumption, and the potential of ultra-large-scale parallel computing to realize the envisioned cryptographic functions. Simulation results revealed significant outcomes for evaluating the impact of the proposed secure approach in the IoNT context, in presence of different number of nano-devices and routing protocols. A future interesting aspect to investigate is how to simulate different kinds of attacks and violation attempts towards the IoNT system.

REFERENCES

- [1] H. E. El-Din and D. Manjaiah, "Internet of nano things and industrial internet of things," in *Internet of Things: Novel Advances and Envisioned Applications*. Springer, 2017, pp. 109–123.
- [2] I. F. Akyildiz and J. M. Jornet, "The internet of nano-things," *IEEE Wireless Communications*, vol. 17, no. 6, 2010.

TABLE IV. SUMMARY OF DELAY ASSESSMENT

Sim/Delay	Global encryption delay	Per nano-device encryption delay
Sim_1	Acceptable increase, proportionally to the number of sensors	Decrease in the per nano-device incidence due to the insignificant increase in the total delay as the devices in the network increase
Sim_2	Not suitable for a system with a high number of nano-devices	Optimal only in the case of few nano-devices
Sim_3	Acceptable increase proportionally to the number of sensors	Acceptable increase proportionally to the number of sensors

- [3] I. F. Akyildiz, F. Brunetti, and C. Blázquez, “Nanonetworks: A new communication paradigm,” *Computer Networks*, vol. 52, no. 12, pp. 2260–2279, 2008.
- [4] A. D. Maynard, “Nanotechnology: assessing the risks,” *Nano Today*, vol. 1, no. 2, pp. 22–33, 2006.
- [5] N. Akhtar and Y. Perwej, “The internet of nano things (iont) existing state and future prospects,” *GSC Advanced Research and Reviews*, vol. 5, no. 2, pp. 131–150, 2020.
- [6] Y. Niu, K. Zhao, X. Zhang, and G. Cui, “Review on dna cryptography,” in *Bio-inspired Computing: Theories and Applications: 14th International Conference, BIC-TA 2019, Zhengzhou, China, November 22–25, 2019, Revised Selected Papers, Part II 14*. Springer, 2020, pp. 134–148.
- [7] G. Xiao, M. Lu, L. Qin, and X. Lai, “New field of cryptography: Dna cryptography,” *Chinese Science Bulletin*, vol. 51, no. 12, pp. 1413–1420, 2006.
- [8] B. Anam, K. Sakib, M. Hossain, and K. Dahal, “Review on the advancements of dna cryptography,” *Proceedings of International Conference on Software, Knowledge, Information Management and Applications (SKIMA)*, 10 2010.
- [9] T. Mahjabin, A. Olteanu, Y. Xiao, W. Han, T. Li, and W. Sun, “A survey on dna-based cryptography and steganography,” *IEEE Access*, vol. 11, pp. 116423–116451, 2023.
- [10] L. Chu, Y. Su, X. Yao, P. Xu, and W. Liu, “A review of dna cryptography,” *Intelligent Computing*, vol. 4, p. 0106, 2025.
- [11] F. Dressler and F. Kargl, “Towards security in nano-communication: Challenges and opportunities,” *Nano communication networks*, vol. 3, no. 3, pp. 151–160, 2012.
- [12] J. B. Wendt and M. Potkonjak, “Nanotechnology-based trusted remote sensing,” in *Sensors*. IEEE, 2011, pp. 1213–216.
- [13] Y. Zhang, F. Wang, J. Chao, M. Xie, H. Liu, M. Pan, E. Kopperger, X. Liu, Q. Li, J. Shi *et al.*, “Dna origami cryptography for secure communication,” *Nature communications*, vol. 10, no. 1, pp. 1–8, 2019.
- [14] C. S. Sreeja, M. Misbahuddin, and N. P. Mohammed Hashim, “Dna for information security: A survey on dna computing and a pseudo dna method based on central dogma of molecular biology,” in *International Conference on Computing and Communication Technologies*, 2014, pp. 1–6.
- [15] E. Vidhya and R. Rathipriya, “Key generation for dna cryptography using genetic operators and diffie-hellman key exchange algorithm,” *Computer Science*, vol. 15, no. 4, pp. 1109–1115, 2020.
- [16] A. Das, S. K. Sarma, and S. Deka, “Data security with dna cryptography,” in *Transactions on Engineering Technologies*. Springer, 2021, pp. 159–173.
- [17] B. AL-Shargabi and A. Dar Assi, “A modified lightweight dna-based cryptography method for internet of things devices,” *Expert Systems*, vol. 40, no. 6, p. e13270, 2023.
- [18] S. Aqeel, A. S. Khan, I. A. Abbasi, F. Algarni, and D. Grzonka, “Enhancing iot security with a dna-based lightweight cryptography system,” *Scientific Reports*, vol. 15, no. 1, p. 13367, 2025.
- [19] I. Qiqieh, J. Alzubi, and O. Alzubi, “Dna cryptography based security framework for health-cloud data,” *Computing*, vol. 107, no. 1, p. 35, 2025.
- [20] P. K. Bulasara and S. R. Sahoo, “A robust and secure drug delivery with single transmitter and dual symmetrical receivers in an internet of bio-nano things,” *IEEE Internet of Things Journal*, vol. 11, no. 14, pp. 25 074–25 087, 2024.
- [21] I. Akyildiz and J. M. Jornet, “Nano communication networks,” *Nano Communication Networks*, vol. 1, pp. 3–19, 2010.
- [22] J. M. Jornet and I. F. Akyildiz, “Channel modeling and capacity analysis for electromagnetic wireless nanonetworks in the terahertz band,” *IEEE Transactions on Wireless Communications*, vol. 10, no. 10, pp. 3211–3221, 2011.
- [23] G. Piro, L. A. Grieco, G. Boggia, and P. Camarda, “Simulating wireless nano sensor networks in the ns-3 platform,” in *IEEE 27th International Conference on Advanced Information Networking and Applications Workshops (WAINA)*. IEEE, 2013, pp. 67–74.
- [24] J. M. Jornet and I. F. Akyildiz, “Information capacity of pulse-based wireless nanosensor networks,” in *8th Annual IEEE Communications Society Conference on Sensor, Mesh and Ad Hoc Communications and Networks*. IEEE, 2011, pp. 80–88.
- [25] L. M. Adleman, “Molecular computation of solutions to combinatorial problems,” *Science*, pp. 1021–1024, 1994.
- [26] A. Azfar, K.-K. R. Choo, and L. Liu, “A study of ten popular android mobile voip applications: Are the communications encrypted?” in *47th Hawaii International Conference on System Sciences*. IEEE, 2014, pp. 4858–4867.
- [27] J. D. Watson, F. Crick *et al.*, “A structure for deoxyribose nucleic acid,” 1953.
- [28] L. M. Adleman, P. W. Rothenmund, S. Roweis, and E. Winfree, “On applying molecular computation to the data encryption standard,” *Journal of Computational Biology*, vol. 6, no. 1, pp. 53–63, 1999.
- [29] S. Lloyd and Q. O. Snell, “Sequence alignment with traceback on reconfigurable hardware,” in *International Conference on Reconfigurable Computing and FPGAs*. IEEE, 2008, pp. 259–264.
- [30] C. Kreibich and J. Crowcroft, “Efficient sequence alignment of network traffic,” in *Proceedings of the 6th ACM SIGCOMM Conference on Internet Measurement*, 2006, pp. 307–312.
- [31] M. Pierobon and I. F. Akyildiz, “A physical end-to-end model for molecular communication in nanonetworks,” *IEEE Journal on Selected Areas in Communications*, vol. 28, no. 4, 2010.
- [32] B. Atakan, O. B. Akan, and S. Balasubramaniam, “Body area nanonetworks with molecular communications in nanomedicine,” *IEEE Communications Magazine*, vol. 50, no. 1, 2012.
- [33] L. Chouhan and M.-S. Alouini, “Interfacing of molecular communication system with various communication systems over internet of every nano things,” *IEEE Internet of Things Journal*, vol. 10, no. 16, pp. 14 552–14 568, 2023.
- [34] S. Sicari, A. Rizzardi, G. Piro, A. Coen-Porisini, and L. Grieco, “Beyond the smart things: towards the definition and the performance assessment of a secure architecture for the internet of nano-things,” *Computer Networks*, vol. 162, p. 106856, 2019.
- [35] S. Sicari, A. Rizzardi, L. A. Grieco, and A. Coen-Porisini, “Security, privacy and trust in internet of things: The road ahead,” *Computer networks*, vol. 76, pp. 146–164, 2015.
- [36] I. Llatser, D. Demiray, A. Cabellos-Aparicio, D. T. Altılar, and E. Alarcon, “N3sim: Simulation framework for diffusion-based molecular communication nanonetworks,” *Simulation Modelling Practice and Theory*, vol. 42, pp. 210–222, 2014.
- [37] G. Piro, K. Yang, G. Boggia, N. Chopra, L. A. Grieco, and A. Alo-mainy, “Terahertz communications in human tissues at the nanoscale for healthcare applications,” *IEEE Transactions on Nanotechnology*, vol. 14, no. 3, pp. 404–406, 2015.
- [38] S. Sahoo, S. Parveen, and J. Panda, “The present and future of nanotechnology in human health care,” *Nanomedicine: Nanotechnology, Biology and Medicine*, vol. 3, no. 1, pp. 20–31, 2007.
- [39] C. Yang, J. Chu, R. L. Warren, and I. Birol, “Nanosim: nanopore sequence read simulator based on statistical characterization,” *Giga-Science*, vol. 6, no. 4, p. gix010, 2017.
- [40] C. Funck, F. B. Laun, and A. Wetscherek, “Characterization of the diffusion coefficient of blood,” *Magnetic resonance in medicine*, vol. 79, no. 5, pp. 2752–2758, 2018.