Alternative DNA Security using BioJava

Mircea-Florin Vaida¹, Radu Terec¹, Lenuta Alboaie²

¹Technical University of Cluj-Napoca, Faculty of Electronics, Telecommunications and Information Technology, Departament of Communications, 26 – 28 Gh. Baritiu, 400027, Cluj-Napoca, Romania, Phone: (+40) 264 401810, Mircea.Vaida@com.utcluj.ro, RaduTerec@gmail.com

²Alexandru Ioan Cuza University of Iasi, Romania, Faculty of Computer Science, Berthelot, 16, Iasi, Romania, adria@infoiasi.ro

Abstract. This paper presents alternative security methods based on DNA. From the alternative security methods available, a DNA algorithm was implemented using symmetric coding in BioJava and MatLab. As results, a comparison has been made between the performances of different standard symmetrical algorithms using dedicated applications. In addition to this, we also present an asymmetric key generation and DNA security algorithm. The asymmetric key generation algorithm starts from a password phrase. The asymmetric DNA algorithm proposes a mechanism which makes use of more encryption technologies. Therefore, it is more reliable and more powerful than the OTP DNA symmetric algorithm.

Keywords: DNA security, BioJava, asymmetric cryptography.

1 Introduction

With the growth of the information technology (IT) power, and with the emergence of new technologies, the number of threats a user is supposed to deal with grew exponentially. For this reason, the security of a system is essential nowadays. It doesn't matter if we talk about bank accounts, social security numbers or a simple telephone call. It is important that the information is known only by the intended persons, usually the sender and the receiver.

In the domain of security, to ensure the confidentiality property two main approaches can be used: that of symmetrical and asymmetrical cryptographic algorithms. Cryptography consists in processing plain information [1], [2], applying a cipher and producing encoded output, meaningless to a third-party who does not know the key. Symmetrical algorithms use the same key to encrypt and decrypt the data, while asymmetric algorithms use a public key to encrypt the data and a private key to decrypt it. By keeping the private key safe, you can assure that the data remains safe. The disadvantage of asymmetric algorithms is that they are computationally intensive. Therefore, in security a combination of asymmetric and symmetric algorithms is used.

In the future it is most likely that the computer architecture and power will evolve. Such systems might drastically reduce the time needed to compute a cryptographic key. As a result, security systems need to find new techniques to transmit the data securely without relying on the existing pure mathematical methods.

We therefore use alternative security concepts [9]. The major algorithms which are accepted as alternative security are the elliptic, vocal, quantum and DNA encryption algorithms. Elliptic algorithms are used for portable devices which have a limited processing power, use a simple algebra and relatively small ciphers.

The quantum cryptography is not a quantum encryption algorithm but rather a method of creating and distributing private keys. It is based on the fact that photons send towards a receiver changing irreversibly their state if they are intercepted. Quantum cryptography was developed starting with the 70s in Universities from Geneva, Baltimore and Los Alamos.

In [18] two protocols are described, BB84 and BB92, that, instead of using general encryption and decryption techniques, verify if the key was intercepted. This is possible because once a photon is duplicated, the others are immediately noticed. However, these techniques are still vulnerable to the Man-in-the-Middle and DoS attack.

DNA Cryptography is a new field based on the researches in DNA computation [4] and new technologies like: PCR (Polymerase Chain Reaction), Microarray, etc. DNA computing has a high level computational ability and is capable of storing huge amounts of data. A gram of DNA contains 10^{21} DNA bases, equivalent to 10^{8} terabytes of data. In DNA cryptography we use existing biological information from DNA public databases to encode the plaintext [7], [12].

The cryptographic process can make use of different methods. In [9] the *one-time pads* (OTP) algorithms are described, which is one of the most efficient security algorithms, while in [15] a method based on the DNA splicing technique is detailed. In the case of the *one-time pad* algorithms, the plaintext is combined with a secret random key or *pad* which is used only once. The pad is combined with the plaintext using a typical modular addition, or an XOR operation, or another technique. In the case of [15] the start codes and the pattern codes specify the position of the introns, so they are no longer easy to find. However, to transmit the spliced key, they make use of public-key secured channel.

Additionally, we will describe an algorithm which makes use of asymmetric cryptographic principles. The main idea is to avoid the usage of both purely mathematical symmetric and asymmetric algorithms and to use an advanced asymmetric algorithm based on DNA. The speed of the algorithm should be quite high because we make use of the powerful parallel computing possibilities of the DNA. Also, the original asymmetric keys are generated starting from a user password to avoid their storage.

This paper is structured in 5 sections. In section 2 we present some general aspects about the genetic code. In section 3 we show 2 algorithms for the symmetric DNA implementation, a MatLab implementation and one realized in BioJava. We will

also expose the limitation imposed by these platforms. In section 4 we describe an advanced asymmetric DNA encryption algorithm. We will conclude this paper in section 5 where a comparison between the obtained results is made and the conclusions and possible continuations of our work are presented.

2 General aspects about Genetic code

There are 4 nitrogenous bases used in making a strand of DNA. These are adenine (A), thymine (T), cytosine (C) and guanine (G). These 4 bases (A, T, C and G) are used in a similar way to the letters of an alphabet. The sequence of these DNA bases will code specific genetic information [7].

In our previous work we used a one-time pad, symmetric key cryptosystem [19]. In the OTP algorithm, each key is used just once, hence the name of OTP. The encryption process uses a large non-repeating set of truly random key letters. Each pad is used exactly once, for exactly one message. The sender encrypts the message and then destroys the used pad. As it is a symmetric key cryptosystem, the receiver has an identical pad and uses it for decryption. The receiver destroys the corresponding pad after decrypting the message. New message means new key letters. A cipher text message is equally likely to correspond to any possible plaintext message. Cryptosystems which use a secret random OTP are known to be perfectly secure.

By using DNA with common symmetric key cryptography, we can use the inherent massively-parallel computing properties and storage capacity of DNA, in order to perform the encryption and decryption using OTP keys. The resulting encryption algorithm which uses DNA medium is much more complex than the one used by conventional encryption methods.

To implement and exemplify the OTP algorithm, we downloaded a chromosome from the open source NCBI GenBank. As stated, in this algorithm the chromosomes are used as cryptographic keys. They have a small dimension and a huge storage capability. There is a whole set of chromosomes, from different organisms which can be used to create a unique set of cryptographic keys. In order to splice the genome, we must know the order in which the bases are placed in the DNA string.

The chosen chromosome was "*Homo sapiens FOSMID clone ABC24-1954N7 from chromosome 1*". It's length is high enough for our purposes (37983 bases).

GenBank offers different formats in which the chromosomal sequences can be downloaded:

- GenBank,
- GenBank Full,
- FASTA,
- ASN.1.

We chose the FASTA format because it's easier to handle and manipulate. To manipulate the chromosomal sequences we used BioJava API methods, a framework for processing DNA sequences. Another API which can be used for managing DNA sequences is offered by MatLab. Using this API, a dedicated application has been implemented [10].

In MatLab, the plaintext message was first transformed in a bit array. An encryption unit was transformed into an 8 bit length ASCII code. After that, using functions from the Bioinformatics Toolbox, each message was transformed from binary to DNA alphabet. Each character was converted to a 4-letter DNA sequence and then searched in the chromosomal sequence used as OTP, [19].

Next, we will present an alternative implementation which makes use of the BioJava API.

The core of BioJava is actually a symbolic alphabet API, [20]. Here, sequences are represented as a list of references to singleton symbol objects that are derived from an alphabet. The symbol list is stored as often as possible. The list is compressed and uses up to four symbols per byte.

Besides the fundamental symbols of the alphabet (A, C, G and T as mentioned earlier), the BioJava alphabets also contain extra symbol objects which represent all possible combinations of the four fundamental symbols. The structure of the BioJava architecture together with its most important APIs is presented below:

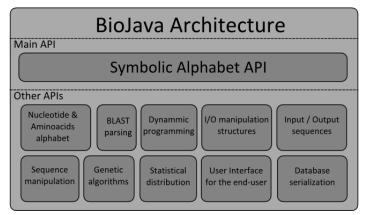


Figure 1. The BioJava Architecture

By using the symbol approach, we can create higher order alphabets and symbols. This is achieved by multiplying existing alphabets. In this way, a codon can be treated as nothing more than just a higher level alphabet, which is very convenient in our case. With this alphabet, one can create views over sequences without modifying the underlying sequence.

In BioJava a typical program starts by using the sequence input/output API and the sequence/feature object model. These mechanisms allow the sequences to be loaded from a various number of file formats, among which is FASTA, the one we used. The obtained results can be once more saved or converted into a different format.

3 DNA Cryptography Implementations

In this chapter we will start by presenting the initial Java implementation of the symmetric OTP encryption algorithm, [19]. We will then continue by describing the corresponding BioJava implementation and some drawbacks of this symmetric algorithm.

3.1. Java implementation

Due to the restrictions that limit the use of JCE, the symmetric cryptographic algorithm was developed using OpenJDK, which is based on the JDK 6.0 version of the Java platform and does not enforce certificate verification. This algorithm involves three steps: key generation, encryption and decryption.

In this algorithm, the length of the key must be exactly the same as the length of the plaintext. In this case, the plaintext is the secret message, translated according to the following substitution alphabet: 00 - A, 01 - C, 10 - G and 11 - T. Therefore, the length of the key is three times the length of the secret message. So, when trying to send very long messages, the length of the key would be huge. For this reason, the message is broken into fixed-size blocks of data. The cipher encrypts or decrypts one block at a time, using a key that has the same length as the block.

The implementation of block ciphers raises an interesting problem: the message we wish to encrypt will not always be a multiple of the block size. To compensate for the last incomplete block, padding is needed. However, this DNA Cipher will not use a standard padding scheme but a shorter version (a fraction) of the original key. The only mode of operation implemented by the DNA Symmetric Cipher is ECB (Electronic Code Book). ECB mode has the disadvantage that the same plaintext will always encrypt to the same ciphertext, when using the same key.

As we mentioned, the DNA Cipher applies a double encryption in order to secure the message we want to keep secret. The first encryption step uses a substitution cipher.

For applying the substitution cipher a HashMap object was used. HashMap is a java.util class that implements the Map interface. These objects associate a value to a specified unique key in the map. Each character of the secret message is represented by a combination of 3 DNA bases.

The result after applying the substitution cipher is a string containing characters from the DNA alphabet (A, C, G and T). This will further be transformed into a byte array, together with the key. The exclusive or operation (XOR) is then applied to the key and the message in order to produce the encrypted message.

When decrypting an encrypted message, it is essential to have the key and the substitution alphabet. While the substitution alphabet is known, being public, the key is kept secret and is given only to the addressee. Any malicious third party won't be able to decrypt the message without the original key.

For the decryption, the received message is XOR-ed with the secret key which results in a DNA-based text. This text is then broken into groups of three characters and with the help of the reverse map each such group will be replaced with the corresponding letter. The reverse map is the inverse of the one used for translating the original message into a DNA message. This way the receiver is able to read the original secret message.

A powerful implementation should consider medical analysis of a patient. In [8] an improved DNA algorithm is proposed.

3.2 BioJava Implementation

In this approach, we use more steps to obtain the DNA code starting from the plaintext. For each character from the message we wish to encode, we first apply the *get_bytes()* method which returns an 8bit ASCII string of the character we wish to encode. Further, we apply the *get_DNA_code()* method which converts the obtained 8 bit string, corresponding to an ASCII character, into DNA alphabet. The function returns a string which contains the DNA-encoded message.

The *get_DNA_code()* method is the main method for converting the plaintext to DNA encoded text. For each 2 bits from the initial 8 bit sequence, corresponding to an ASCII character, a specific DNA character is assigned: 00 - A, 01 - C, 10 - G and 11 - T. Based on this process we obtain a raw DNA message.

Table 1. DNA encryption test sequence

```
Plaintext message: "test"
ASCII message: 116 101 115 116
Raw DNA message: "TCACGCCCTATCTCA"
```

The coded characters are searched in the chromosome chosen as session key at the beginning of the communication. The raw DNA message is split into groups of 4 bases. When such a group is found in the chromosome, its base index is stored in a vector. The search is made between the first characters of the chromosome up to the 37983th. At each new iteration, a 4 base segment is compared with the corresponding 4 base segment from the raw DNA message. So, each character from the original string will have an index vector associated, where the chromosome locations of that character are found.

The *get_index()* method effectuates the parsing – the comparison of the chromosomal sequences and creates for each character an index vector. To parse the sequences in the FASTA format specific BioJava API methods were used.

BioJava offers us the possibility of reading the FASTA sequences by using a FASTA stream which is obtained with the help of the SeqIOTools class. We can pass through each of the sequences by using a *SequenceIterator* object. These sequences are then loaded into an *Sequence* list of objects, from where they can be accessed using the *SequenceAt()* mrthod.

In the last phase of the encryption, for each character of the message, a random index from the vector index is chosen. We use the *get_random()* method for this purpose. In this way, even if we would use the same key to encrypt a message, we would obtain a different result because of the random indexes.

Since the algorithm is a symmetric one, for the decryption we use the same key as for encryption. Each index received from the encoded message is actually pointing to a 4 base sequence, which is the equivalent of an ASCII character.

So, the *decode()* method realizes following operations: It will first extract the DNA 4 base sequences from the received indexes. Then, it will convert the obtained raw DNA message into the equivalent ASCII-coded message. From the ASCII coded message we finally obtain the original plaintext. And with this, the decryption step is completed.

The main vulnerability of this algorithm is that, if the attacker intercepts the message, he can decode the message himself if he knows the coding chromosomal sequence used as session key.

4 BioJava asymmetric algorithm description

In this chapter we will present in detail an advanced method of obtaining DNAencoded messages. It relies on the use of an asymmetric algorithm and on key generation starting from a user password.

We will also present a pseudo-code description of the algorithm.

4.1 Asymmetric key generation

Our first concern when it comes to asymmetric key algorithms was to develop a way in which the user was no longer supposed to deal with key management authorities or with the safe storage of keys. The reason behind this decision is fairly simple: both methods can be attacked. Fake authorities can pretend to be real key-management authorities and intruders may breach the key storage security. By intruders we mean both persons who have access to the computer and hackers, which illegally accessed the computer.

To address this problem, we designed an asymmetric key generation algorithm starting from a password. The method has some similarities with the RFC2898 symmetric key derivation algorithm [21]. The key derivation algorithm is based on a combination of hashes and the RSA algorithm. Below we present the basic steps of this algorithm:

• Step 1: First, the password string is converted to a byte array, hashed using SHA256 and then transformed to BigInteger number. This number is transformed in an odd number, *tmp*, which is further used to apply the RSA algorithm for key generation.

• Step 2: Starting from *tmp* we search for 2 random pseudo-prime number p and q. The relation between *tmp*, p and q is simple: p < tmp < q. To spare the computational power of the device, we do not compute traditionally if p and q are prime but make primality tests.

• A primality test determines the probability according to which a number is prime. The sequence of the primality test is the following: First, trial

divisions are carried out using prime numbers below 2000. If any of the primes divides this BigInteger, then it is not prime. Second, we perform base 2 strong pseudo-prime test. If this BigInteger is a base 2 strong pseudo-prime, we proceed on to the next step. Last, we perform the strong Lucas pseudo-prime test. If everything goes well, it returns true and we declare the number as being pseudo-prime.

• Step 3: Next, we determine Euler totient: phi = (p - 1) * (q - 1); and $n = p^{*}q$;

• Step 4: Next, we determine the public exponent, *e*. The condition imposed to *e* is to be coprime with *phi*.

• Step 5: Next, we compute the private exponential, d and the CRT (Chinese Reminder Theorem) factors: dp, dq and qInv.

• **Step 6**: Finally, all computed values are written to a suitable structure, waiting further processing.

• The public key is released as the public exponent, *e* together with *n*.

• The private key is released as the private exponent, d together with n and the CRT factors.

The scheme of this algorithm is presented below:

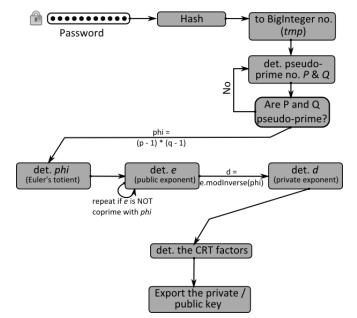


Figure 2. Asymmetric RSA compatible key generation

In comparison with the RFC2898 implementation, here we no longer use several iterations to derive the key. This process has been shown to be time consuming and provide only little extra security. We therefore considered it safe to disregard it.

The strength of the key-generator algorithm is given by the large pseudo-prime numbers it is using and of course, by the asymmetric algorithm. By using primality tests one can determine with a precision of 97 - 99% that a number is prime. But most importantly, the primality tests save time. So, the average computation time, including appropriate key export, for the whole algorithm is 143 ms. After the generation process was completed, the public or private key can be retrieved using the static *ToXmlString* method.

Next, we will illustrate the algorithm through a short example. Suppose the user password is "DNACryptography". Starting from this password, we compute its hash with SHA256. The result is shown below. This hashed password is converted into the BigInteger number *tmp*. Starting from it, and according to the algorithm described above, we generate the public exponent e and the private exponent d.

Table 2. Asymmetric DNA encryption test sequence

We conducted several tests and the generated keys match the PKCS #5 specifications. Objects could be instantiated with the generated keys and used with the normal system-build RSA algorithm.

4.2 Asymmetric DNA algorithm

The asymmetric DNA algorithm proposes a mechanism which makes use of three encryption technologies. In short, at the program initialization, both the initiator and its partner generate a pair of asymmetric keys. Further, the initiator and its partner negotiate which symmetric algorithms to use, its specifications and of course, the codon sequence where the indexes of the DNA bases will be looked up. After this initial negotiation is completed, the communication continues with normal message transfer. The normal message transfer supposes that the data is symmetrically encoded, and that the key with which the data was encoded is asymmetrically encoded and attached to the data. This approach was first presented in [17].

Next, we will describe the algorithm in more detail and also provide a pseudocode description for a better understanding. **Step 1:** At the startup of the program, the user is asked to provide a password phrase. The password phrase can be as long or as complicated as the user sees fit. The password phrase will be further hashed with SHA256.

Step 2: According to the algorithm described in section 4.1, the public and private asymmetric keys will be generated. Since the pseudo-prime numbers p and q are randomly chosen, even if the user provides the same password for more sessions, the asymmetric keys will be different.

Step 3: The initiator selects which symmetric algorithms will be used in the case of normal message transfer. He can choose between 3DES, AES and IDEA. Further, he selects the time after which the symmetric keys will be renewed and the symmetric key length. Next, he will choose the codon sequence where the indexes will be searched. For all this options appropriate visual selection tools are provided.

Step 4: The negotiation phase begins. The initiator sends to its partner its public key. The partner responds by encrypting his own public key with the initiators public key. After the initiator receives the partner's public key, he will encrypt with it the chosen parameters. Upon receiving the parameters of the algorithms, the partner may accept or propose his own parameters. In case the initiators parameters are rejected, the parties will chose the parameters which provide the maximum available security.

Step 5: The negotiation phase is completed with the sending of a test message which is encrypted like any regular message would be encrypted. If the test message is not received correctly by any of the two parties or if the message transfer takes too much time, the negotiation phase is restarted. In this way, we protect the messages from tampering and interception.

Step 6: The transmission of a normal message. In this case, the actual data will be symmetrically encoded, according to the specifications negotiated before. The symmetric key is randomly generated at a time interval t. The symmetric key is encrypted with the partner's public key and then attached to the message. So, the message consists in the data, encrypted with a symmetric key and the symmetric key itself, encrypted with the partner's public key. We chose to adopt this mechanism because symmetric algorithms are faster than asymmetric ones. Still, in this scenario, the strength of the algorithm is equivalent to a fully asymmetric one because the symmetric key is encrypted asymmetrically. The procedure is illustrated below:

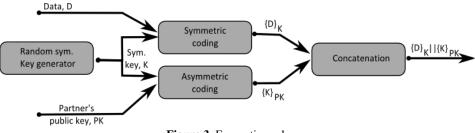


Figure 3. Encryption scheme

Next, the obtained key will be converted into a byte array. The obtained array will be converted to a raw DNA message, by using a substitution alphabet. Finally, the raw DNA message is converted to a string of indexes and then transmitted.

The decryption process is fairly similar. The user converts the index array back to raw DNA array and extracts the ASCII data. From this data he will decipher the symmetric key used for that encryption, by using its private key. Finally the user will obtain the data by using the retrieved symmetric key. At the end of the communication, all negotiated data is disregarded (symmetric keys used, the asymmetric key pair and the codon sequence used).

5 Conclusions and compared results

In this chapter we will present the results we obtained for the symmetric algorithm implementation along with the conclusions of our present work.

Our first goal was to compare the time required to complete the encryption/ decryption process. We compared the execution time of the DNA Symmetric Cipher with the time required by other classical encryption algorithms. We chose a random text of 360 characters, in string format which was applied to all tests.

The testing sequence is:

 Table 3. Testing sequence

```
k39pc3xygfv(!x|jl+qo|9~7k9why(ktr6pkiaw|gwnn&aw+be|r|*4u+rz$
wm)(v_e&$dz|hc7^+p6%54vp*g*)kzlx!%4n4bvb#%vex~7c^qe_d745h40i
$_2j*6t0h$80!c~9x4^2srn81x*wn9&k%*oo_co(*~!bfur7tl4udm!m4t+a
|tb%zho6xmv$6k+#1$&axghrh*_3_zz@0!05u*|an$)5)k+8qf0fozxxw)_u
pryjj7_|+nd_&x+_jeflua^peb_+%@03+36w)$~j715*r)x(*bumozo#s^j
u)6jji@xa3y35^$+#mbyizt*mdst&h|hbf6o*)r2qrwm10ur+mbezz(1p7$f
```

To be able to compute the time required for encryption and decryption, we used the public static *nanoTime()* method from the *System* class which gives the current time in nanoseconds. We called this method twice: once before instantiating the Cipher object, and one after the encryption. By subtracting the obtained time intervals, we determine the execution time.

It is important to understand that the execution time varies depending on the used OS, the memory load and on the execution thread management. We therefore measured the execution time on 3 different machines:

- System 1: Intel Core 2 Duo 2140, 1.6 GHz, 1 Gb RAM, Vista OS
- System 2: Intel Core 2 Duo T6500, 2.1 GHz, 4 Gb RAM, Windows 7 OS
- System 3: Intel Dual Core T4300, 2.1 GHz, 3 Gb RAM, Ubuntu 10.04 OS

Next, we present the execution time which was obtained for various symmetric algorithms in the case of the first, second and the third system, for different cases:

Analysis results for Vista OS							
DES	Encryption	50	26	1.03	0.81	0.84	0.84
	Decryption	1.63	0.35	0.33	0.32	0.34	0.36
AES	Encryption	80	26	0.92	0.95	0.88	0.54
	Decryption	27	2.09	0.30	22.26	0	0.14
Blowfish	Encryption	65	10.91	25	24	0.15	1.45
	Decryption	3	1.87	1.72	29	1.09	1
3DES	Encryption	82	24	2.41	25	2.12	1.42
	Decryption	1.56	1.42	26	1.23	1.41	0.66
BIO sym.	Encryption	4091	4871	4875	4969	4880	4932
algorithm	Decryption	6.29	4.19	4.19	4.19	4.19	4.19

Table 4. Results obtained for System 1

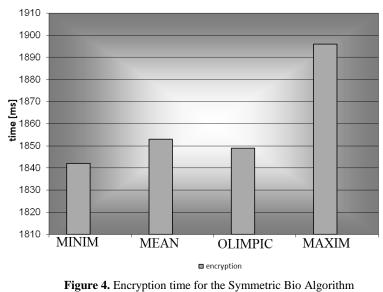
 Table 5. Results obtained for System 2

Analysis results for Windows 7							
DES	Encryption	34	1.43	1.09	1.2	1.73	1.19
	Decryption	0.75	0.37	0.44	0.42	0.38	0.37
AES	Encryption	28	1.3	1.16	0.07	1.77	0.82
	Decryption	0.12	0.14	2.09	0.9	2.09	0.16
Blowfish	Encryption	22	28.4	6.2	4	1.6	2.83
	Decryption	2.24	2.21	1.8	1.8	1.8	1.71
3DES	Encryption	41	6.59	2.78	2.62	2.69	2.12
	Decryption	1.12	1.78	1.24	1.74	1.48	1
BIO sym.	Encryption	3970	3884	3887	3901	3900	3910
algorithm	Decryption	4.19	4.19	4.19	2.09	4.19	2.09

Table 6. Res	ults obtained	for S	vstem 3	3
--------------	---------------	-------	---------	---

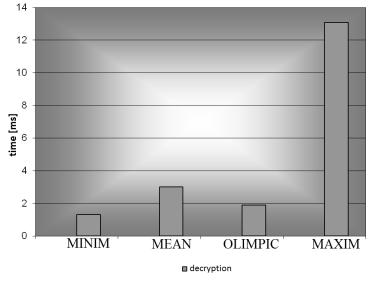
Analysis results for Ubuntu 10.04							
DES	Encryption	12.64	0.9	0.61	0.59	0.61	0.56
	Decryption	1.24	0.45	0.44	0.45	0.43	0.41
AES	Encryption	0.66	0.6	0.63	0.63	0.62	0.63
	Decryption	0.66	0.71	0.64	0.64	0.19	0.19
Blowfish	Encryption	37.07	32	19	13	15	14
	Decryption	0.81	0.77	0.81	0.58	0.74	0.59
3DES	Encryption	14	11	17.7	10.21	10.11	13
	Decryption	0.77	0.79	0.78	0.6	0.6	0.6
BIO sym.	Encryption	1896	1848	1857	1846	1850	1850
algorithm	Decryption	2.62	13.1	1.83	1.31	1.57	2.62

Below, we will illustrate the maximum, mean, olimpic (by eliminating the absolute minimum and maximum values) and minimum encryption and decryption time for the Symmetric Bio Algorithm.



Encryption Time for the Symmetric Bio Algorithm





Decryption time for the Symmetric Bio Algorithm

Figure 5. Decryption time for the Symmetric Bio Algorithm

First of all, we can notice that the systems 1 and 2 (with Windows OS) have larger time variations for the encryption and decryption processes. The third system, based on the Linux platform, offers a better stability, since the variation of the execution time is smaller.

As seen from the figures and tables above, the DNA Cipher requires a longer execution time for encryption and decryption, comparatively to the other ciphers. We would expect these results because of the type conversions which are needed in the case of the symmetric Bio algorithm. All classical encryption algorithms process array of bytes while the DNA Cipher is about strings. The additional conversions from string to array of bytes and back make this cipher to require more time for encryption and decryption then other classic algorithms.

However, this inconvenience should be solved with the implementation of full DNA algorithms and the usage of Bio-processors, which would make use of the parallel processing power of DNA algorithms.

In this paper we proposed an asymmetric DNA mechanism that is more reliable and more powerful than the OTP DNA symmetric algorithm. As future developments, we would like to make some test for the asymmetric DNA algorithm and increase its execution time.

Acknowledgments. This work was supported by CNCSIS–UEFISCSU, project number PNII – IDEI 1083/2007-2010.

References

1. David Hook, Beginning Cryptography with Java, Wrox Press, (2005)

2. Kahn D., The codebrakers McMillan, New York, (1967)

3. M. Schena, Microarray analysis Wiley-Liss, July (2003)

4. L. M. Adleman, Molecular computation of solution to combinatorial problems, *Science*, 266, 1021-1024, (1994)

5. B. Schneier, Applied cryptography: protocols, algorithms, and source code in C, John Wiley & Sons Inc, (1996)

6. Java Cryptography Architecture. Sun Microsystems.

http://java.sun.com/j2se/1.4.2/docs/guide/security/CryptoSpec.html (2011)

7. Genetics Home Reference. U.S. National Library of Medicine.

http://ghr.nlm.nih.gov/handbook/basics/dna. (2011)

8. T. Hodorogea, Mircea-F. Vaida, Blood Analysis as Biometric Selection of Public Keys, 7 th International Carpathian Control Conference ICCC'2006, Ostrava – Beskydy, Czech Republic, May 29-31, pp. 675-678, (2006)

9. Ashish Gehani, Thomas LaBean, John Reif, *DNA-Based Cryptography*. s.l.: DIMACS Series in Discrete Mathematics and Theoretical Computer Science, Vol. 54, and *Lecture Notes in Computer Science, Springer*, (2004) 10. Olga Tornea, Monica Borda, Tatiana Hodorogea, Mircea-Florin Vaida, Encryption System with Indexing DNA Chromosomes Cryptographic Algorithm, IASTED International Conference on Biomedical Engineering (BioMed 2010), 15-18 Feb., Innsbruck, Austria, paper 680-099, pp. 12-15, (2010)

11. R. K. Wilson, The sequence of Homo sapiens FOSMID clone ABC14-50190700J6, submitted to http://www.ncbi.nlm.nih.gov, (2009)

12. DNA Alphabet. VSNS BioComputing Division. http://www.techfak.uni-

bielefeld.de/bcd/Curric/PrwAli/node7.html#SECTION0007100000000000000000, (2011)

13. Wagner, Neal R., The Laws of Cryptography with Java Code. [PDF], (2003).

14. B. Schneier, Description of a New Variable-Length Key, 64-Bit Block Cipher (Blowfish): Springer-Verlag, andf, Fast Software Encryption, Cambridge Security Workshop Proceedings (1993).

15. S. T. Amin, M. Saeb, S. El-Gindi, A DNA-based Implementation of YAEA Encryption Algorithm, IASTED International Conference on Computational Intelligence, San Francisco, pp. 120-125, (2006)

 BioJava - http://java.sun.com/developer/technicalArticles/javaopensource/biojava/ (2011)
 Nicolas Nobelis, Karima Boudaoud, Michel Riveill – "Une architecture pour le transfert électronique sécurisé de document", PhD Thesis, Equipe Rainbow, Laboratories I3S – CNRS, Sophia-Antipolis, France, (2008)

18. Piya Techateerawat, A Review on Quantum Cryptography Technology, International Transaction Journal of Engineering, Management & Applied Sciences & Technologies, Vol. 1, pp. 35-41, (2010)

 Mircea-Florin Vaida, Radu Terec, Olga Tornea, Chiorean Ligia, Alexandra Vanea, DNA Alternative Security, Advances in Intelligent Systems and Technologies Proceedings ECIT2010 – 6th European Conference on Intelligent Systems and Technologies, Iasi, Romania, October 07-09, pp. 1-4, (2010)

20. R.C.G. Holland; T. Down; M. Pocock; A. Prlić; D. Huen; K. James; S. Foisy; A. Dräger; A. Yates; M. Heuer; M.J. Schreiber - "BioJava: an Open-Source Framework for Bioinformatics", Bioinformatics (2008)

21. RSA Security Inc. Public-Key Cryptography Standards (PKCS) – "PKCS #5 v2.0: Password-Based Cryptography Standard", (2000)