

PROPOSED APPROACH FOR IMPROVING RNA CRYPTO-KEY BASED ON POLYNOMIAL CONVOLUTION

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Abstract - Random Number Generators are fundamental tools for cryptography protocols and algorithms. The basic problems that face any crypto key generator are randomness, correlations and distribution of the state of key sequence. This paper proposed a new method to enhance RNA crypto key generation. It has been implemented by extending the crypto key by applying polynomial convolution technique which extracts the mask filter from the same RNA key sequence depending on the start and end codon properties. This will provide another high level of extension and generate random-strength crypto key. The proposal approach could pass through the statistical measurements successfully and achieved high rate of randomness (approximated to 96%).

Keywords – Cryptography, Key generation, RNA, Bimolecular Computation, Polynomial Convolution.

I. INTRODUCTION

Random numbers generations are a major tool in many applications of cryptography such as key generation [1]. The security of cryptography methods basically depends on the robustness of encryption method and secrecy of key [3]. It is so important to provide generators which are capable to produce amount of secure random numbers such as polynomial Feedback Shift Registers (LFSRs) [2]. Bimolecular Computation Methods (BMC) have been developed for a various field of operations on DNA and RNA strands. BMC methods have solved hard combinatorial search problems such as Travelling Salesman Problem (TSP), breaking the Data Encryption Standard (DES) and for cryptography key generation [4]. This approach will propose a new method to improve RNA crypto key generation basically by depending on the concept of Polynomial convolution 1D, this technique will add second level of extension to achieve a desired level of secrecy and randomness In [5], A. Hassan proposed a method for key generation based on the concept of translating RNA to protein chain. Their approach first determines a key sequence size, for example 9 byte, then it produces crypto key with extended length suitable with the length of secret message. Their proposed approach consists of several steps contain (Mapping, Extended, Complementing and Rotation). The generated key was acceptable according to statistical tests of randomness.

II. DNA AND RNA TRANSCRIPTION

DNA is a double stranded sequence of four nucleotides; the four nucleotides that compose a strand of DNA are as follows: Adenine (A), Guanine (G), Cytosine (C), and Thymine (T), they are often called *bases*. The chemical structure of DNA (the famous double-helix) was discovered by James Watson and Francis Crick in 1953. It consists of a particular bond of two linear sequences of bases. This bond follows a property of complementarity: Adenine bonds with Thymine (A-T) and vice versa (T-A), Cytosine bonds with Guanine (C-G) and vice versa (G-C). This is known as Watson-Crick complementarity. Each DNA strand has two different ends that determine its polarity: the 3' end, and the 5' end. The double helix is an anti-parallel (two strands of opposite polarity) bonding of two complementary strands [6].

III. RNA CRYPTOSYSTEM

RNA-Crypto System (shortly RCS) is a private key algorithm to encrypt data, in particular from the observation of RNA behavior and some of its properties. The RNA sequences have some sections called Introns. Introns, derived from the term "intra-genic regions", are non-coding sections of precursor mRNA (pre-mRNA) or other RNAs, that are removed (spliced out of the RNA) before the mature RNA is formed.

Once the introns have been spliced out of a pre-mRNA, the resulting mRNA sequence is ready to be translated into a protein. The corresponding parts of a gene are known as introns as well. It uses the presence of Introns in the RNA-Crypto System output as a strong method to access to the secret key to code the messages. In the RNA-Crypto System algorithm, the introns are sections of the ciphered message with non-coding information as well as in the precursor mRNA [7].

are Equipment, Middle-ware and Introduction [1]. Equipment is made up of sensors and actuators, the Middle-ware provides stockpiling and processing apparatuses and finally, the introduction provides the elucidation devices open on different steps. It is not plausible the processing of information that has been collected from a great number of sensors, which set mindful Middle-ware arrangements are suggested for enabling a sensor for choosing the most critical data to be handled [24].

Characteristically speaking, the design of the internet of things offers no suitable edge to achieve the fundamental actions

IV. POLYNOMIAL VECTOR CONVOLUTION

In mathematics and some particular functional analysis, convolution is a mathematical operation on two functions *g* and *f*, to produce a third function that is typically showed as a modified version of one of the original two functions in which one of the original functions is translated. Convolution is similar to cross-correlation. It has many applications in probability, statistics, computer vision, image and signal processing [9]. Segment is made out of a four-round structure utilizing two non-direct substitution boxes *S7* and *S9*. The structure is compacted in two rounds utilizing parallelism. The unequal division of *FI* is because of the way that objective elements of odd size are for the most part superior to those of even size from the perspective of provable security against direct and differential cryptanalysis. *S7* and *S9* have been planned in a way that stays away from direct structures in *FI*. This reality has been affirmed by measurable testing. *S*-boxes (*S7* and *S9*) are actualized in combinational rationale despite the fact that they could be executed by "look-into tables" to diminish the extent of our usage. As a result of the parallelism, just two part of each *S7* and *S9* is required for the calculation of *A5*. Along these lines we utilize each *S7* and *S9* part 120 times.

Let *m* = length (*u*) and *n* = length (*v*). Then *w* is the vectors of length *m+n-1* whose *k*th element is:

$$W(k) = \sum_j u(j) v(k - j + 1) \tag{1}$$

The sum is over all the values of *j* that lead to legal subscripts for *u*(*j*) and *v*(*k-j+1*), specifically *j* = max(1, *k+1-n*):1:min(*k*, *m*). When *m* = *n*, this gives

$$\begin{aligned} w_1 &= u_1 * v_1 \\ w_2 &= u_1 * v_2 + u_2 * v_1 \\ w_3 &= u_1 * v_3 + u_2 * v_2 + u_3 * v_1 \\ &\dots\dots\dots \\ w_n &= u_1 * v_n + u_2 * v_{n-1} + \dots + u_n * v_1 \end{aligned}$$

V. IMPROVEMENT OF RNA-CRYPTO-KEY

This approach discusses some drawbacks that have been observed by using the previous approach of RNA key generation, and we try to find an improved version for addressing those mistakes stated as follows:

- Translation of mRNA strand based on start and end codons, investigated from which by calculating the number of these confined codons to apply rotate shift on RNA sequence.

Benefit from mRNA translation just for applies "Rotate Shift on RNA sequence" may be considered as weakness investment for this property, it looks like "simple permutation".

In this approach, extended RNA-Crypto-key basically depends on the "Polynomial convolution technique", contributes to

provide self-extension with variable RNA length by treating RNA confined codons as mask filter for convolution applied with RNA key chain. The length of resultant key will be approximated to the summation of the length of mask filter and the length of RNA chain. It is worth mentioning, that mask filter has variable length according to confined codons number in RNA key chain, so the output of extended RNA-Crypto-key also has variable length with a high rate of randomness. This approach has two levels of RNA key extension, the first one comes from RNA codons table and the second one by applying Polynomial convolution technique. Algorithm (1) shows main steps of the proposal approach.

Algorithm (1): Improved RNA-Crypto-Key

Input:
Initial key (characters, numbers), size (9, 12, 15, ... etc/ byte), No. Iterations

Output:
Improving RNA random key with variable expanded size

Begin

Step1: Convert initial key sequence to binary sequence

Step2: Coding each 2bit from the message binary sequence to RNA 4base using table (1).

Step3: Split RNA strand into group of codons (3 nitrogenous bases)

Step4: Extend each codon in RNA sequence by selecting another codon that belong to the same amino acid and appending them, according to table (II).

Step5: Read RNA strand until finding the AUG that is used to begin protein synthesis, count the number codons and stop when finding end codon is (UAA or UAG or UGA)

Step6: Extract Self-RNA-mask filter starting from AUG to (UAA or UAG or UGA)

Step7: Apply rotate right shift on RNA strand based on the number of confined codons between start and end codon

Step8: Apply (Polynomial-Convolution) technique between Self-RNA-mask filter and RNA strand based on equation (1)

Step9: Convert New-RNA key to binary sequence and generate final crypto-key

End

TABLE (1): CONVERT BIT SEQUENCE INTO MRNA NUCLEOTIDES

Bit Sequence	mRNA Base
00	A
01	U
10	C
11	G

TABLE (2): CODING AMINO ACID GROUPS INTO BIT SEQUENCE

Decimal code	RNA code	Binary code based (6 bit)
0	UUU	000000
1	UUC	000001
2	UUA	000010
3	UUG	000011
4	CUU	000100
5	CUC	000101
6	CUA	000110
7	CUG	000111
8	AUU	001000
9	AUC	001001
10	AUA	001010
11	AUG	001011
12	GUU	001100
13	GUC	001101
14	GUA	001110
15	GUG	001111
16	UCU	010000
17	UCC	010001
18	UCA	010010
19	UCG	010011
20	CCU	010100
21	CCC	010101
22	CCA	010110
23	CCG	010111
24	ACU	011000
25	ACC	011001
26	ACA	011010
27	ACG	011011
28	GCU	011100
29	GCC	011101
30	GCA	011110
31	GCG	011111
32	UAU	100000
33	UAC	100001
34	UAU	100010
35	UAC	100011
36	CAU	100100
37	CAC	100101
38	CAA	100110
39	CAG	100111
40	AAU	101000
41	AAC	101001
42	AAA	101010
43	AAG	101011
44	GAU	101100
45	GAC	101101
46	GAA	101110
47	GAG	101111
48	UGU	110000
49	UGC	110001
50	UGU	110010
51	UGG	110011
52	CGU	110100
53	CGC	110101
54	CGA	110110
55	CGG	110111
56	AGU	111000
57	AGC	111001
58	AGA	111010
59	AGG	111011
60	GGU	111100
61	GGC	111101
62	GGA	111110
63	GGG	111111

Note: Highlight green code on in Table (2) represents starts and Red represents end codons in mRNA strand

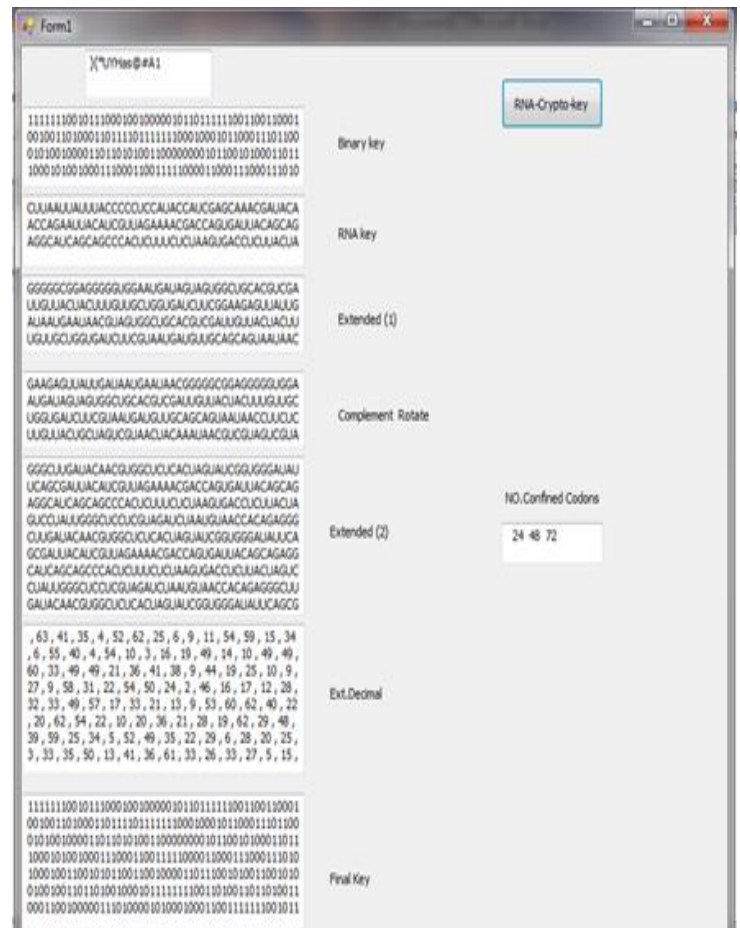


Fig. 1. Interface of proposed program using c#.net

VI. EXPERIMENT A RESULT AND DISCUSSION

This section illustrates the results that are obtained from the implementation of the proposed approach which is explained in the previous sections, these results can be presented as follows: Basic Statistical measurements of randomness have been done on the improved RNA as shown in Table (3). These statistical tests such as, frequency, serial, poker and runs are used to evaluate randomness and distributed properties of several RNA crypto- key samples [11]. Randomness tests were applied on various key sizes such as 9, 12, 72, 96 and 144 bits or bytes. Randomness is used to check random and distributed properties of several RNA crypto- key samples. Useful statistical tests are four basic tests, including: *Frequency test*, *Serial test*, *Poker test*, *Runs test* [10]. The output of tests must be compared with passes values to decide if the outputs of randomness tests are good or not. Randomness tests are applied on different key sizes: 72, 96 and 144 bits.

TABLE (2): CODING AMINO ACID GROUPS INTO BIT SEQUENCE

Initial key size (bit)	Improved RNA key at 3itr. size(bit)	Frequency Pass <= 3.84	Serial Pass <= 5.99	Poker Pass <=11.1	Run Pass <=22.362
96	3324	0.123	0.544	7.321	5.161
	2190	0.154	1.119	4.754	3.368
	2388	0.342	3.182	10.407	15.164
72	1758	2329	6.254	7.146	6.387
	2088	3.065	3.571	8.321	18.462
	1824	0.967	1.430	7.394	12.737
144	3306	0.756	2.49	10.89	8.406
	3204	1.709	4.77	6.262	6.824
Average		1.18	2.92	7.81	10.21

As shown in table (3), Improved RNA -key is achieved to high rate of randomness (96% percentage of tests that is passed through statistics) when compared with previous method of RNA-key generation in table (4) that shows just Average of many tests. While Improved -key size is may be much larger than 40%.

TABLE (4): RANDOMNESS STATISTICAL TESTS OF PREVIOUS RNA-CRYPTO-KEY (BY AVERAGE)

Initial Key	Previous RNA key at 3 itr. Size (bit)	Frequency Pass <= 3.84	Serial Pass <=5.99	Poker Pass <=11.1	Run Pass <=22.362
96	1920	27.5	65.9	92.4	58.2

Figure (2) illustrates Chart that shows the difference between the key expansion rates of the improved and traditional RNA key method through several iterations.

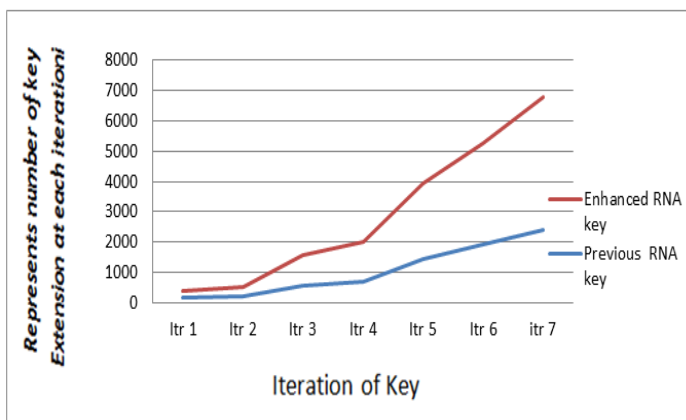


Fig. 2. Comparison in term of key expansion between traditional and proposed approaches

VII. CONCLUSION

This paper describes an efficient method for Improving RNA-crypto-key generation using Polynomial Convolution. It successfully provides an extra level of extension and security for RNA-crypto-key when compared with the previous approach. Robust features of this proposed method could be as follows:

- RNA-crypto-key expansion basically depends on several factors such as (start and end codons for translation, iteration numbers, and self-expansion based on codon table).
- Polynomial convolution provided more than 40% expansion on traditional RNA₂
- Improved RNA-crypto-key provides high rate of security randomness achieved 96% on other traditional RNA key generation method.
- Size of convolution mask filter may have a variable length based on start and end codon of each RNA strand and encoding of initial key.

VIII. FUTURE WORK

Improving -RNA algorithm considered as one of the optimization techniques which are used for solving any NP-complete problem such as path planning, four colors mapping, TSP, CPU scheduling problems.

IX. APPENDIX(1)

This section illustrates the implementation of the proposed approach which is programmed by using Visual c#.net 2015 as shown in Fig (2). The following example explains in details steps of proposed algorithm work:

- Initial key (12 byte)=)(*UYHas@#A1

Step1. Binary coding (96 bits)=

```
(111111001011100010010000010 110 11111100110
011000100100 1101000 11011110111111000 10001 01100 01
110110001010010000110110 10100110 0 0 0
000010110010100011011100010100100011100011001111100
001100011100011101010001001100101011001100100001101
11001010011001010010010011011010010001011111110011
010011011010011000110010000011101000010100010001100)
```

Step2. RNA Coding (48 nucleotide)=

```
(CUUAAUUUUUACCCCUCCAUAACCAUCGAGCAAAC
GAUACAA CCAGA)
```

Step3. Extended (1) based on codon table(96 nucleotide)=

```
(GAAGAGUUUUGAUAAUGAAUAACGGGGGCGG
AGGGGGU GGA
AUGAUAGUAGUGGCUGCACGUCGAUUGUUACUACUU
UGUUGCUGGUGAUCUUCG)
```

Step4. Complement RNA & Rotate=

```
(GGGGGCGGAGGGGGUGGAAUGAUAGUAGUGGCUGC
ACGUCGAUUGUUACUACUUUGUUGCUGGUGAUCUUC
GGAAGAGUUUUGAUAAUGAAUAAC)
```

Step5. Extended (2) based on Polynomial Convolution (Note: mask filter length = 24)

(GGGCUUGAUACAACGUGGCUCUCACUAGUAUCGGUG
GGAUUAUCAGCGAUUACAUCGUUAGAAAACGACCAG
UGAUUACAGCAGAGGCAUCAGCAGCCCACUCUUUCU
CUAAGUGACCUCUUACUAGUCCUUAUUGGGCUCCUCG
UAGAUCUAAUGUAACCACAGA)

Step6. Random Key Generation (330 bit) =

(1111110010111000100100000101101111100110011000100
10011010001101111011111100010001011000111011000101
00100001101101010011000000010110010100011011100010
100100011100011001111100001100011100011101010001001
100101011001100100001101110010100110010100100100110
11010010001011111110011010011011010011000110010000
011101000010100010001100)

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