



DNA Computation

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Abstract:

The process capability of living systems has intrigued researchers for years. Primarily, the main target has been on implementing aspects of living systems in process devices. Laptop literal peoples expand their hand to the biologist and chemist to explore the potential for computation of biological molecules line deoxyribonucleic acid (DNA) and polymer (RNA) that square measure info carrying molecules. during this context, polymer computation is largely a group of specially designated polymer strands whose mixtures can end in the answer to some issues. Current DNA-based circuits square measure non-homogenous, consisting of separate combinatorial gates with corresponding static process talents. This analysis focuses on any developing DNA-based methodologies to mimic digital information manipulation. Uniform logic style principles square measure introduced, giving birth the foundations of dynamic process whereby data is parsed through a digital circuit comprised of DNA-based logic gates. The final word aim would be to facilitate shift-and-add calculation. A lot of immediate applications concern information tamper-proofing, serving security desires whose progress was antecedent restricted. a unique computer circuit style supported chemical reactions is conferred within which observance of double stranded sequences indicates a truth analysis. Circuits square measure obfuscated by removing of physical sequence connections, permitting client-specific representative strands for input sequences, sterilization the input sequence strands over time, and ranging the input sequence length.

Keywords: DNA, DNA-based logic gates, Finger Printing, Logic gates, DNA-based shift register, Gates-input, DNase.

1.INTRODUCTION

Traditional silicon-based electronic equipment is vulnerable to security attacks as a consequence of the static nature of its style. Metrics utilised to judge security area unit typically supported the 'good feeling' of engineers instead of empirical proof. Assessments typically lead to statements like "it is much not possible to access the memory from the outside" or "it is not possible to access the info bus that carries the key from storage to the processor if they're all on an equivalent piece of semiconducting material." This diminishes the complete subject of security to a mere form instead of scientific proof. The truth is, once a static circuit is obtained by Associate in Nursinging offender, it's a matter of your time before one will reverse engineer its configuration.

True tamper-proof security should satisfy 3 principal requirements:

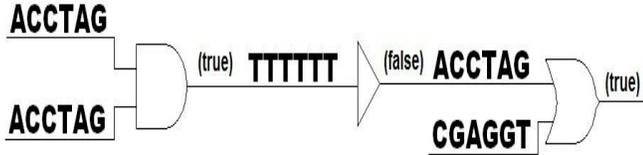
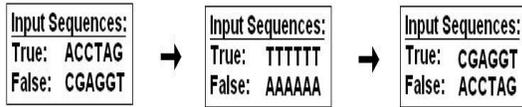
- (1) Resist static attacks that involve direct penetration of memory cells.
- (2) Resist dynamic attacks that decide to access info because it is moves from memory to the process unit.
- (3) Resist dynamic attacks that decide to access info throughout process.

To bypass such meddling, circuit should be dynamic naturally. We tend to argue that DNA-based logic circuits, once the technology matures, could give revolutionary solutions to tamper proofing. A DNA-based style permits electronic equipment to be supported organic chemistry and environmental stimuli. Distinct parts, like those ascertained in

CMOS circuits, area unit non-existent. Meddling would therefore have new which means, presumably preventing it altogether supported correct scientific observations. After construction of AN info storage methodology with information validation and random range generation electronic equipment, subsequent logical objective is to develop a DNA-based register. A register may be a primary part of the machine processor that permits computation at a gate level followed by shifting of the knowledge to the continuing gate. Merely moving the info by itself has no machine that means. A register needs the combination of each logic and shifting; thereby making an entire process unit that performs serial calculations on an input stream of data. Thus, its development is vital to the continuing advancement of dynamic computing. With every of those new theories introduced, we tend to move nearer to the sensible applications afforded by dynamic computing. As a primary step, DNA-based methodologies are developed to mimic existing silicon-based technologies in information manipulation, specifically info storage, random range generation, and a register. The remainder of the paper treats the subsequent. Active DNA-based logic gates as we tend to understand them to be created area unit introduced. A discussion of techniques used to alter gate style, resulting in tamper proofing, is then bestowed. The power to form present ternary logic gates exploitation the planned gate style is delineate. this is often followed by a discussion of however the planned gates might be joined to construct circuits. A biological approach to shifting is then bestowed, proceeded by a technique by that DNA sequences that function gate inputs might be quickly hold on. Finally, circuit fabrication concerns area unit mentioned.

II. SYSTEM ARCHITECTURE

DNA prefers to be in double stranded form; single stranded DNA sequences naturally migrate towards complementary sequences to form double stranded complexes.



2.1. Gate input

Each DNA-based logical operation input is delineated by one stranded DNA sequence, with the need that the sequence representing a “true” analysis is complementary to the sequence representing a “false” analysis for one gate. For instance, one might assign ACCTAG to represent “true” and CTAGGT representing “false,” as CTAGGT is that the reverse complement of ACCTAG. This allows sequence assignment to be dynamic in nature. A user might every which way assign a replacement set of representative sequences for every gate during a circuit. Contemplate the circuit comprised of 3 DNA-based logic gates in Figure one. The primary gate uses the sequences bestowed higher than, wherever ACCTAG represents “true” and CTAGGT represents “false.” Once evaluating the primary gate, the user dynamically changes the representative input sequences, wherever TTTTTT currently represents “true” and AAAAAA currently represents “false.” Finally, the user reuses the primary set of sequences, however reverses the assignment such CTAGGT currently represents “true” and ACCTAG currently represents “false.”

2.2. Gate Result

DNA’s preference to be in double stranded kind allows ancient logic operations to be performed. For every various DNA-based gate style, a preset mixture is equipped containing a selected single stranded sequence to induce the suitable chemical process, called the bottom mixture. If the gate input sequence provided is complementary to the bottom sequence, then the corresponding double stranded polymer sequence can kind. Thus, the presence or absence of a double stranded sequence is employed to judge gate output; the presence of a double stranded sequence represents a “true” analysis whereas its absence represents a “false” analysis. Fluorescent labels are often accustomed sight the presence or absence of the double stranded sequence. During this method, fluorescent molecules are connected to the ester sequence, and absorb and emit light-weight at a selected wavelength. Thus, by attaching the fluorescent molecule to 1 of the strands of the double stranded sequence, the double stranded sequence are often detected.

III. Proposed System

The following are the gates that will be use full in determining the result.

As we know that the nucleotides present in DNA are ATGC. Base pairs occur when nitrogenous bases make hydrogen bonds with each other. Each base has a specific partner:

guanine with cytosine, adenine with thymine (in DNA) or adenine with uracil (in RNA).

3.1. NOT Gate

In digital logic, an inverter or NOT gate is a logic gate which implements logical negation. Just one input is equipped to the gate, and therefore the output is that the corresponding complementary sequence. As a result of the output ought to assess “true” solely within the presence of a “false” input, the bottom mixture provided to the gate contains the representative “true” sequence. DNAase is equipped to destroy any single stranded sequences. If a double stranded sequence is discovered, then the result's “true”; otherwise, the result's “false.” Contemplate the instance conferred in Figure a pair of, wherever the sequence TTTTTT represents a “true” input and AAAAAA represents a “false” input. The bottom mixture would so contain the sequence TTTTTT. If the input sequence is “false,” then AAAAAA can bind with the provided TTTTTT sequence to make a double stranded sequence. DNAase can haven't any impact on the sequences, and therefore the double stranded sequence are going to be discovered, representing a “true” analysis.

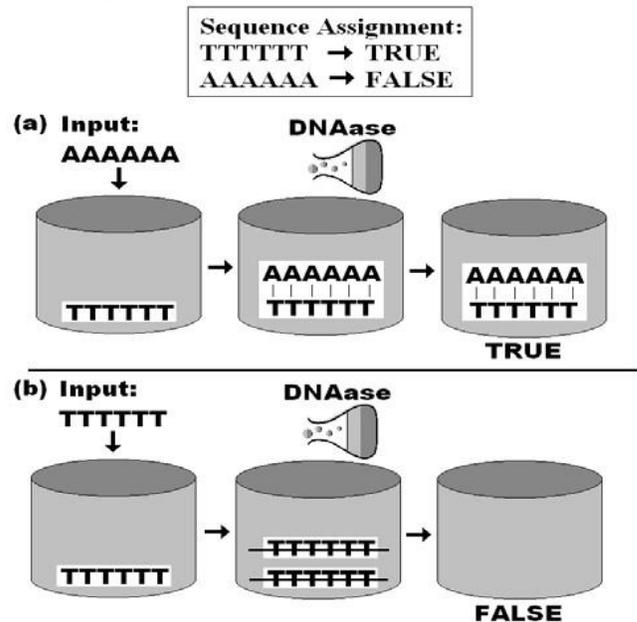


Fig 3.a : NOT gate.

3.2. OR Gate

The logic gate evaluates “true” if a minimum of one amongst the gate inputs area unit “true.” Introducing the “false” sequence as (a) One true and one false input ends up in a double stranded sequence representing a truth analysis. Conversely, (b) 2 true inputs and (c) 2 false inputs don't produce a double stranded sequence. The base mixture would force a minimum of one amongst the inputs be “true” to create a double stranded sequence. DNAase can destroy any single stranded sequence within the mixture. If a double stranded sequence is discovered, then the result's “true;” otherwise, the result's “false.” Contemplate the instance in Figure four wherever the sequence TTTTTT represents a “true” input and also the sequence AAAAAA represents a “false” input. If each of the input sequences area unit “true” TTTTTT sequences, then one amongst the sequences can mix with the bottom “false” AAAAAA sequence to supply a double stranded

sequence. DNAase can destroy the remaining input sequence and also the double stranded sequence can lead to a “true” analysis.

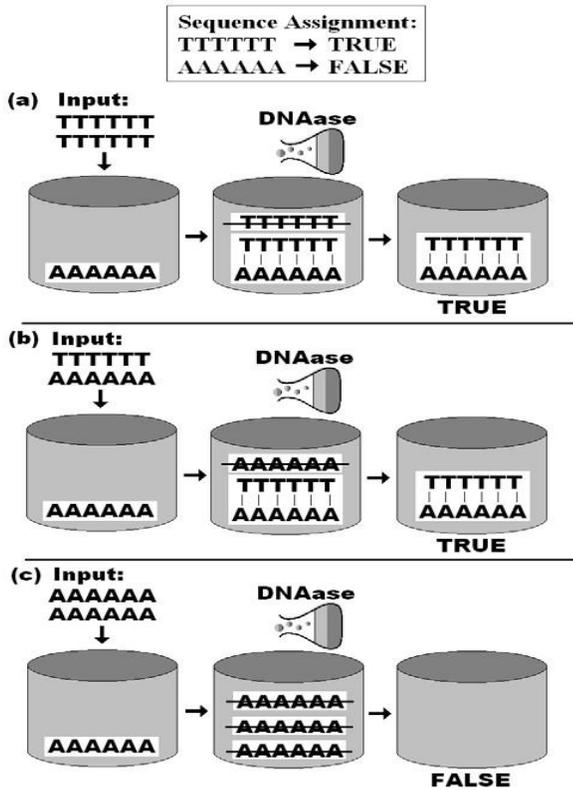


Fig 3.b : OR gate

3.3. XOR Gate

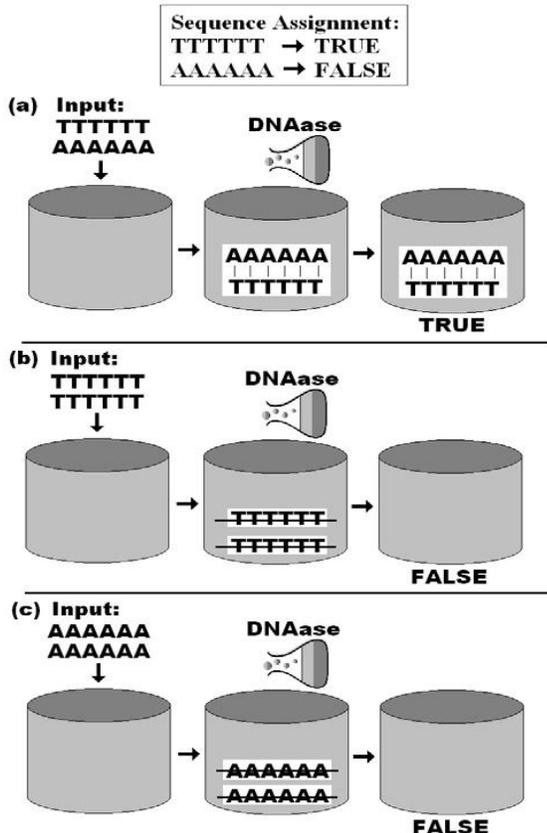


Fig 3.c : XOR gate

The gate evaluates “true” as long as precisely one amongst the input sequences evaluates “true.” With binary inputs, XOR is outlined as evaluating “true” if input values are opposite. In DNA-based logic gates, the gate is that the most oversimplified style in this no base sequence has to be provided to the gate. Opposite input sequences are complementary and can bind along to create a double stranded sequence. If the inputs don't seem to be opposite, then the sequences won't be ready to bind to 1 another and DNAase can destroy each input sequences. If a double stranded sequence is ascertained, then the result's “true;” otherwise, the result's “false.”

3.4. NAND Gate

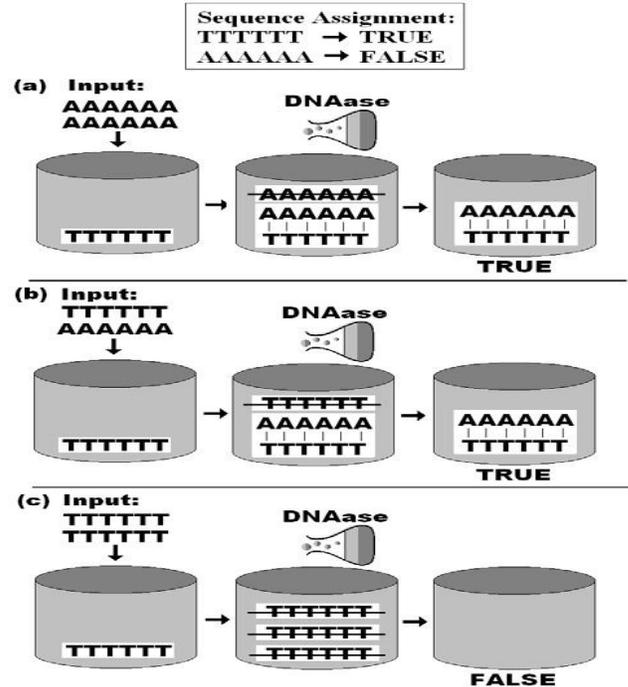


Fig 3.d : NAND gate

The NAND gate evaluates “true” if inputs aren't each “true,” in different words, if there's a minimum of one false input. The DNA-based NAND computer circuit is analogous to the logic gate conferred antecedent, except the bottom sequence contains the sequence representing “true” instead of “false.” Thus, a minimum of one in every of the inputs should be “false” so as to create a double stranded sequence. DNAase can destroy any single stranded sequence within the mixture. If a double stranded sequence is determined, the result's “true;” otherwise, it evaluates to “false.” continued with the instance higher than, if each of the input sequences square measure “false” AAAAAA sequences, then one can mix with the reference “true” TTTTTT sequence to provide a double stranded molecule. DNAase can destroy the remaining input sequence and also the double stranded sequence can lead to a “true” analysis. If one input sequence is “false” and also the different input sequence is “true,” the “false” AAAAAA input sequence. each (a) 2 false inputs and (b) one true input and one false input lead to the formation of a double stranded sequence representing a truth analysis. (c) but, 2 true inputs don't produce a double stranded sequence can mix with the “true” TTTTTT sequence to provide the mandatory double stranded sequence. DNAase can then destroy the remaining “false” sequence and also the gate can still lead to a “true” analysis.

Finally, if each of the input sequences square measure “true” TTTTTT sequences, then neither of the sequences are going to be ready to mix with the bottom “true” sequence. DNAase can destroy all sequences within the mixture, leading to a “false” analysis of the gate.

3.5. For rest of Gates

NOT, XOR, OR, and NAND represent four of the seven commonest mathematical logic gates. From these four DNA-based logic gates, one will simply devise a DNA-based illustration for all alternative mathematical logic gates. take into account the 3 remaining digital logic gates of the seven commonest – AND, NOR, and XNOR. The gate, that evaluates “true” only if each inputs area unit “true,” is formed by applying the NOT gate to the output of the NAND circuit. The NOR gate, that evaluates “true” once each inputs area unit “false,” is formed by applying the NOT gate to the results of the OR circuit. Finally, the XNOR gate, that evaluates “true” once each inputs area unit constant, is formed by applying the NOT gate to 1 of the inputs, then applying the XOR gate to the result and also the alternative input. just like the preceding gate styles, the presence of a double stranded sequence indicates a “true” analysis of the gate, whereas the absence of a double stranded sequence indicates a “false” analysis of the gate.

IV. Input vs. output

The benefits of DNA-based logic gates don't seem to be restricted to the reduction within the range of the gates supported the illustration of a further output state; it conjointly permits circuits to be compressed. The planned DNA based computer circuit output analysis relies exclusively on the presence or absence of the double stranded molecule. Thus, a myriad of input sequences will be condensed into one gate mixture. for instance, a series of gates will be integrated into one DNA-based OR gate.

INPUT 1	INPUT 2	OUTPUT
A	A	Identical
A	C	Different
A	G	Different
A	T	Complementary
C	A	Different
C	C	Identical
C	G	Complementary
C	T	Different
G	A	Different
G	C	Complementary
G	G	Identical
G	T	Different
T	A	Complementary
T	C	Different
T	G	Different
T	T	Identical

The presence of one “true” sequence within the mixture can end in the formation of the double stranded molecule irrespective of the magnitude of inputs gift. maybe the advantages of DNA based computer circuit style lies not in mimicking the formal logic of their digital counterparts, however in fashioning a brand-new set of logical operations enabled by the ternary logic structure combined with the DNA-based style.

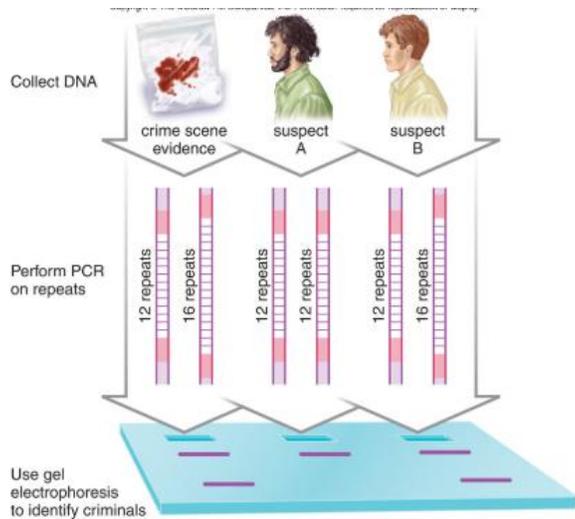
V. DNA Fingerprinting

DNA procedure, conjointly referred to as polymer typewriting, polymer identification, procedure, genotyping, or identity testing, in biological science, technique of analytic and distinguishing variable components at intervals the base-pair sequence of polymer (deoxyribonucleic acid).

DNA procedure could be a laboratory technique want to establish a link between biological proof and a suspect during a criminal investigation. A DNA sample taken from against the law scene is compared with a DNA sample from a suspect. If the 2 DNA profiles area unit a match, then the proof came from that suspect. Conversely, if the 2 DNA profiles don't match, then the proof cannot have return from the suspect. DNA procedure is additionally wont to establish paternity. The procedure for making a genetic fingerprint consists of 1st getting a sample of cells, like skin, hair, or blood cells, that contain polymer. The polymer is extracted from the cells and refined. In Jeffreys's original approach, that was supported fragment length polymorphism (RFLP) technology, the polymer was then cut at specific points on the strand with proteins called restriction enzymes. The enzymes made fragments of variable lengths that were sorted by inserting them on a gel and so subjecting the gel to an electrical current (electrophoresis): the shorter the fragment, the additional quickly it emotional toward the positive pole (anode). The sorted double-stranded polymer fragments were then subjected to a blotting technique within which they were split into single strands and transferred to a nylon sheet. The fragments underwent radiography within which they were exposed to polymer probes—pieces of artificial polymer that were created hot which certain to the minisatellites. a chunk of film was then exposed to the fragments, and a dark mark was made at any purpose wherever a hot probe had become connected. The resultant pattern of marks might then be analysed. The assay developed by Jeffreys has been supplanted by approaches that area unit supported the employment of the enzyme chain reaction (PCR) and alleged microsatellites (or short bicycle repeats, STRs), that have shorter repeat units (typically a pair of to four base pairs in length) than minisatellites (10 to over one hundred base pairs in length). PCR amplifies the required fragment of deoxyribonucleic acid (e.g., a selected STR) repeatedly over, making thousands of copies of the fragment. it's an automatic procedure that needs solely tiny amounts of deoxyribonucleic acid as beginning material and works even with part degraded deoxyribonucleic acid. Once AN adequate quantity of deoxyribonucleic acid has been created with PCR, the precise sequence of ester pairs in an exceedingly section of deoxyribonucleic acid is determined by exploitation one amongst many biomolecular sequencing strategies. machine-controlled instrumentality has greatly enhanced the speed of deoxyribonucleic acid sequencing and has created offered several new sensible applications, as well as pinpointing segments of genes that cause genetic diseases, mapping the human order, engineering drought-resistant plants, and manufacturing biological medication from genetically altered microorganism.

An early use of deoxyribonucleic acid process was in legal disputes, notably to assist solve crimes and to see paternity. The technique was challenged, however, over considerations regarding sample contamination, faulty preparation procedures, and incorrect interpretation of the results. Additionally, RFLP needed massive amounts of

high-quality deoxyribonucleic acid that restricted its application in forensics.



Rhetorical deoxyribonucleic acid samples often area unit degraded or area unit collected post-mortem, which implies that lower-quality and subject to manufacturing less-reliable results than samples that are obtained from a living individual. a number of the considerations with deoxyribonucleic acid process, and specifically the employment of RFLP, subsided with the event of PCR- and STR-based approaches.

VI.DNA profiling

Modern-day DNA profiling is also called STR analysis and relies on microsatellites rather than the minisatellites used in DNA fingerprinting. Microsatellites, or short tandem repeats (STRs), are the shorter relatives of minisatellites usually two to five base pairs long. Like minisatellites they are repeated many times throughout the human genome, for example ‘TATATATATATA’. DNA profiles are very useful in forensics because only a tiny sample of human material left behind after a crime may be sufficient to identify someone. DNA profiling can be used to help confirm whether two people are related to one another and is commonly used to provide evidence that someone is, or is not, the biological parent of a child.

VII. Conclusion

Traditional silicon-based electronic equipment is liable to security attack as a consequence of the static nature of its style. True tamper-proof security needs circuits be dynamic naturally. we have a tendency to argue that DNA-based logic circuits, once the technology matures, might offer revolutionary solutions to tamper proofing. As DNA paradigms will be developed to represent their digital equivalents, we have a tendency to move nearer to the sensible applications afforded by dynamic computing.

As a primary step, DNA-based methodologies are developed to mimic existing silicon-based shift registers, demonstrating however data will be passed through a circuit comprised of DNA-based logic gates. A unique computer circuit style supported chemical reactions is given during which observance of double stranded sequences indicate a truth analysis. Circuits are obfuscated by removing physical sequence connections,

permitting client specific representative strands for input sequences, fixing the input sequence strands over time, and ranging the input sequence length. Shifting on the input stream to dissect individual inputs is accomplished through simulated various conjunction of DNA sequences.

VIII. References

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