

Solving SATISFIABILITY with Molecular Algorithms

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Abstract

Molecular computation uses techniques from molecular biology and combinatorial chemistry to perform computation. We explore, via simulation, three distinct molecular algorithms for solving SATISFIABILITY. The simulation measures the number of molecular operations each algorithm needs to solve SATISFIABILITY. The test input consists of a set of random 3-SAT instances distributed over a range of clause-variable ratios ($\alpha = [0.2, 14.0]$).

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Chapter 1

Introduction

Molecular computation uses interactions between genetic molecules, such as DNA or RNA, to perform computational tasks. We provide an experimental system for simulating three molecular algorithms. In this chapter we discuss the advantages of molecular computation versus standard computation. This discussion includes an introduction to the simulation of molecular algorithms. We conclude the chapter with an overview of the contents of this report.

1.1 Introduction to molecular computation

NP problems, such as SATISFIABILITY, may be verified in polynomial time with the aid of a short (i.e. of length polynomial in the length of the input) proof called a *witness*; NP problems may be solved by checking all possible *witness candidates*. In a standard computational environment, brute force search checks all witness candidates in exponential time.

Molecular computation requires exponential space in order to represent all witness candidates. This combinatorial space of witness candidates can be filtered in polynomial time by parallel molecular operators.

A Boolean formula ϕ is said to be in Conjunctive Normal Form (CNF) if ϕ consists of a conjunctive set of Boolean disjunctive clauses. (Throughout this report we use CNF instances ϕ with n variables, m clauses, and k literals.) We have, e.g.,

$$\phi = C_1 \wedge C_2 \wedge \cdots \wedge C_m,$$

where each clause C_i contains k disjunctive Boolean literals

$$C_i = (v_1 \vee v_2 \vee \cdots \vee v_k).$$

A witness for a SATISFIABILITY instance is a Boolean assignment to the variables that makes the formula true. Such a witness can be represented as a vector B in $\{0, 1\}^n$, where

n is the number of variables in ϕ , as follows. Let $\{v_1, \dots, v_n\}$ be the variables of ϕ , for each i in $\{1, \dots, n\}$, the i th element of B is denoted B_i and represents an assignment of true or false to v_i . A witness candidate for a SATISFIABILITY instance may be verified in polynomial time with $\text{CHECKSAT}(\phi, B)$ (Algorithm 1.1 below).

Algorithm 1.1: $\text{CHECKSAT}(\phi, B)$

```

for each clause  $C$  in  $\phi$ 
   $test\_clause \leftarrow False$ 
  do { for each literal  $\ell$  in  $C$ 
    do { if  $B$  satisfies  $\ell$ 
      then  $test\_clause \leftarrow True$ 
    if  $test\_clause = False$ 
      then return ( $False$ )
  return ( $True$ )
}

```

Algorithm 1.1: $\text{CHECKSAT}(\phi, B)$ iterates over each of the clauses C in the CNF instance ϕ . The bit-vector B encodes a witness candidate for the CNF instance ϕ . The $test_clause$ variable gets set to $False$, assuming that the clause cannot be satisfied. If the clause can be satisfied with the input configuration B , then the algorithm continues. If each of the m clauses can be satisfied, then $\text{CHECKSAT}(\phi, B)$ returns $True$; otherwise the algorithm returns $False$.

$\text{CHECKSAT}(\phi, B)$ may be used as a subroutine in a brute force SATISFIABILITY solver. Algorithm 1.2 provides pseudocode for a brute force SATISFIABILITY solver $\text{BRUTE}\text{SAT}(\phi)$.

In this project, we consider molecular algorithms to solve SATISFIABILITY. Molecular algorithms permit parallelism on a massive scale [1, 16]. Molecular operations, such as *append* or *extract*, can perform in parallel on all of the string contents of a test tube [1, 16, 13]. In Chapter 3, we explore techniques from combinatorial chemistry to generate combinatorial sets [16, 10, 13].

Algorithm 1.2: BRUTESAT(ϕ)

```
// Input  $\phi$  consists of  $n$  variables.  
// Bit-vector  $B$  represents a witness candidate.  
  
for each  $B \in \{0, 1\}^n$   
do { if CHECKSAT( $\phi, B$ )  
      then return (SATISFIABLE)  
return (UNSATISFIABLE)
```

Algorithm 1.2: BRUTESAT(ϕ) tests a maximum of 2^n Boolean witness candidates, using the CHECKSAT(ϕ, B) algorithm. If the witness candidate B satisfies the input instance ϕ , then the algorithm returns SATISFIABLE; otherwise the algorithm returns UNSATISFIABLE.

Algorithm 1.3: EXTRACTSAT(ϕ)

```
// Input  $\phi$  consists of  $n$  variables.  
  
 $T \leftarrow \text{COMBINATORIALGENERATE}(n)$   
for each clause  $C$  in  $\phi$   
  do  $\begin{cases} T_C \leftarrow \emptyset \\ \text{for each literal } \ell \text{ in } C \\ \quad \text{do } \begin{cases} T \leftarrow \text{purify}(T) \\ T_T \leftarrow \text{extract}(T, \ell) \\ T_C \leftarrow \text{mix}(T_C, T_T) \end{cases} \\ T \leftarrow T_C \end{cases}$   
  if  $T = \emptyset$   
    then return (UNSATISFIABLE)  
  return (SATISFIABLE)
```

Algorithm 1.3: EXTRACTSAT(ϕ) collects satisfying Boolean literals from each clause in ϕ . Initially, EXTRACTSAT(ϕ) constructs a combinatorial space T using the subroutine COMBINATORIALGENERATE(n) which we introduce in Chapter 3. The initial space T contains string configurations representing all potential witness candidates for ϕ . The space T gets filtered down to witness all clauses. These potential solutions are incrementally mixed into the tube T_C for each clause.

Let us consider Algorithm 1.3 as a simplified version of Lipton’s algorithm [16, 13]. The EXTRACTSAT(ϕ) function provides an introductory view of a molecular algorithm. EXTRACTSAT differs in how the algorithm validates each candidate. The brute force algorithm, BRUTESAT, generates sequentially an exponential number of witness candidates. On the other hand, exponential witness canidates with EXTRACTSAT get filtered in parallel.

1.2 Simulation of molecular SATISFIABILITY solvers

We consider three molecular algorithms for solving SATISFIABILITY: Lipton’s [16], Ogihara and Ray’s [19, 20], and a new algorithm, introduced here, that we call the ‘Distribution’ algorithm. Lipton’s algorithm begins with a combinatorial space of all n -bit witness candidates and filters the combinatorial space so that only those that satisfy the input formula remain. Ogihara and Ray’s algorithm constructs a space of witness candidates using heuristic search. The Distribution algorithm expands a set of witnesses with non-conflicting literals from each clause. Chapters 3 and 4 discuss the implementation of these algorithms.

This project introduces a SATISFIABILITY solver framework for molecular algorithms, which we call ‘Molecular Simulation’. This system provides standard operations for molecular computation which we introduce in Chapter 2. It also records runtime metrics, including counts of molecular operators, memory footprints, and execution times. These metrics let us analyze the algorithmic performance of each molecular algorithm.

Molecular Simulation automates execution of DIMACS CNF instances. It measures key properties for a set of randomly generated 3-SAT instances. The 3-SAT instances span discrete clause-variable ratios from 0.2 to 14.0 in increments of 0.2, creating a sweep of SATISFIABILITY instances. This experimental setup generates SATISFIABILITY problem instances with both SATISFIABLE and UNSATISFIABLE configurations.

1.3 Report Overview

In the following chapters, we describe molecular algorithms for solving SATISFIABILITY. We begin Chapter 2 with an introduction to gene sequencing technologies and molecular biology. We define molecular operations for operating on DNA or RNA. Next, we introduce SATISFIABILITY as a language and as a Boolean circuit.

Chapters 3 and 4 introduce each of the three molecular algorithms for solving SATISFIABILITY. In Chapter 3, we discuss Lipton’s [16, 13] and Ogihara and Ray’s [19, 20, 26] algorithms for SATISFIABILITY. The chapter concludes with a discussion of existing simulation frameworks and physical implementations of these molecular algorithms. Chapter 4 introduces the Distribution algorithm.

Chapters 5 and 6 discuss the project implementation. In Chapter 5, we introduce our software, Molecular Simulation, for simulating molecular algorithms. Chapter 6 describes

the experimental workflow for importing SATISFIABILITY instances for each of the three molecular algorithms we study.

Chapter 7 provides a discussion of algorithm performance based test results. Chapter 8 concludes with a summary of contributions of this project and future directions for molecular computation.

Chapter 2

Background

This chapter provides a background on molecular computation techniques. We begin with an introduction to nanotechnology and then provide an example of how information is encoded using molecular matter. Following this example, we introduce Adleman’s molecular operators for solving an instance of HAMILTONIAN PATH. The operators provide an instruction set for molecular computation, and provide the primitives for constructing molecular algorithms.

In the second half of this chapter, we provide an introduction to SATISFIABILITY. We define SATISFIABILITY as a circuit. We then view SATISFIABILITY as a language. We also discuss practical matters related to efficiently evaluating SATISFIABILITY, such as how to encode input and output, and how to classify instances of SATISFIABILITY in the tests that we perform.

2.1 On nanotechnology and construction of molecules

Richard Feynman founded the field of nanotechnology in his 1959 talk “There’s Plenty of Room at the Bottom” [7]. Examples of applied nanotechnology include the manufacturing of graphene [24] and DNA nanopores [17]. Graphene consists of a planer arrangement of carbon atoms that provides desirable physical and electrical properties [24]. DNA nanopores use graphene to create a physical channel for reading genetic sequences [11]. Gene sequencing technologies provide an example of applied nanotechnology [11, 15, 21].

Smaller and cost-effective DNA sequencers provide the ability to read the contents of a gene. Benchtop sequencers [15, 21] allow doctors to treat patients at the genome level from their office. Life Technologies and Oxford Nanopore offer gene sequencers based on solid-state semiconductor technology [15, 21].

2.1.1 Genetic encoding schemes

Microbiology studies the interactions among organic molecules. In this project, we explore the use of applied genetics as a means for generalized computation. Molecular computation encodes data as sequences of DNA or RNA.

Arbitrary encodings that represent mappings from variables to physical oligonucleotides may have undesirable structure and functionality. Conventional techniques from molecular computation employ variable mappings from a library of oligonucleotides.

An *oligonucleotide* is a short string of genetic information. There are several configurations for DNA and RNA; these include +RNA, -RNA, +DNA, -DNA, \pm RNA, \pm DNA, and +mRNA [2]. The polarity of DNA denotes the direction of the genetic information. ‘+DNA’ is denoted 5’—3’ and ‘-DNA’ is denoted 3’—5’. We focus on +DNA and -DNA as the substrate for computational states. The computational states, in our setting, encode candidate witnesses for SATISFIABILITY.

Table 2.1: A mapping of the integers [0, 4] with arbitrary oligonucleotide definitions.

Integer	Oligonucleotide	Reverse-complement
0	5’—TCTCCC—3’	3’—AGAGGG—5’
1	5’—AAACCC—3’	3’—TTTGGG—5’
2	5’—GGTAAA—3’	3’—CCATTT—5’
3	5’—CCCTCC—3’	3’—GGGAGG—5’
4	5’—CTTTTC—3’	3’—GAAAAG—5’

Suppose for example that we would like to encode the sequence of integers $S = [1, 3, 4, 3, 2, 0]$ as an equivalent oligonucleotide representation with the definitions in Table 2.1. Gene sequencing tools permit one to read and decode data according to Table 2.1. The resulting oligonucleotide O is defined as

$$O = 5' - \text{AAACCC} \cdot \text{CCCTCC} \cdot \text{CTTTTC} \cdot \text{CCCTCC} \cdot \text{GGTAAA} \cdot \text{TCTCCC} - 3'.$$

Molecular computation uses oligonucleotides for both storing and operating on a problem state. These operations include matching and replication. Although this report describes artificial processes, DNA in natural settings undergoes the same transformations that we exploit here. Interactions between genetic molecules are the fundamental mechanism for generic computation with oligonucleotides.

In the following chapters, we describe molecular algorithms for SATISFIABILITY. In the next section, we introduce techniques from Adleman’s molecular toolbox [1].

2.2 Adleman's molecular toolbox for solving HAMILTONIAN PATH

In 1994, Leonard Adleman performed the first molecular computation using recombinant DNA in a bench laboratory setting [1]. This experiment solved a six vertex instance of HAMILTONIAN PATH, an NP-complete problem. In this section, we describe the techniques used in this experiment. We provide definitions for the following operations from Adleman's molecular toolbox: append, extract, mix, split, and purify.

Definition 2.2.1. HAMILTONIAN PATH

Given an undirected graph G , does there exist a path that visits every vertex exactly once?

Adleman uses oligonucleotides for defining each vertex for encoding a graph. His scheme for encoding a graph's vertices shares a similar definition from our example of encoding a sequence of integers, given in Table 2.1. Representing edges requires a reverse-complement oligonucleotide, which connects the suffix of the vertex v_i with the prefix of v_j . Let us consider an example. Let

$$\begin{aligned} v_1 &= 5' - \text{ATCTTT} - 3' \\ v_2 &= 5' - \text{CCTATA} - 3'. \end{aligned}$$

From the definition of v_1 and v_2 , we can construct an edge $e_{1,2}$ as

$$e_{1,2} = 3' - \text{AAAGGA} - 5'.$$

Appending v_2 to v_1 is accomplished by first attaching the edge $e_{1,2}$ to the vertex v_1

$$\begin{aligned} &5' - \text{ATC-TTT} - 3' \\ &\quad 3' - \text{AAAGGA} - 5'. \end{aligned}$$

Next we attach v_2 to the resulting complex, yielding

$$\begin{aligned} &5' - \text{ATCTTT-CCTATA} - 3' \\ &\quad 3' - \text{AAAGGA} - 5'. \end{aligned}$$

Finally the edge may be removed and we have the sequence

$$v_1 \cdot v_2 = 5' - \text{ATCTTT} \cdot \text{CCTATA} - 3'.$$

The sequence $v_1 \cdot v_2$ represents the path v_1 to v_2 , and can be obtained with the *append* operation. A test tube T stores witness candidates. The tube T starts as an empty tube. To solve HAMILTONIAN PATH, we introduce equimolar portions of each oligonucleotide vertex for a starting configuration, using the *mix* operation.

Definition 2.2.2. Mix combines n test tubes of information.

$$T \leftarrow \text{mix}(T_1, \dots, T_n)$$

The output consists of a single set $T = T_1 \cup \dots \cup T_n$.

A small initial set may be amplified using *polymerase chain reaction* (PCR). PCR thermocycles the contents of the tube to replicate the contents. Introducing each vertex representation to the contents randomly generates all potential paths. A set of DNA configurations are generated to represent the set of all witness candidates for Hamiltonian Paths in a graph instance. This set of DNA configurations will be filtered to only include configurations that witness Hamiltonian Paths in G .

Append attaches a string to each string contained in a test tube. *Split* portions a tube into multiple portions. In Chapter 3, we will use split-mix synthesis as a means for generating a combinatorial space.

Definition 2.2.3. Append concatenates each element in T with the oligonucleotide s .

$$T' \leftarrow \text{append}(T, s)$$

Definition 2.2.4. Split portions T into two tubes.

$$[T', T''] \leftarrow \text{split}(T)$$

Each of the resulting tubes, T' and T'' , contain the same representative elements of T .

The initial and terminal conditions for the graph get fulfilled by extracting, from the tube T , only paths that begin with V_{in} and end with V_{out} . Extracting only strings from T that match these conditions constrain the number of potential strings to only those that satisfy the conditions of the graph instance.

Definition 2.2.5. Extract separates all oligonucleotides from T containing the sequence s .

$$T' \leftarrow \text{extract}(T, s)$$

The output consists of a set T' of those oligonucleotides containing s .

The tube T consists of possible encodings that have the correct starting and ending vertices. We select only strings of length n , where n is the number of vertices in G , to ensure that all vertices get traversed. This can be performed using *gel electrophoresis*, a technique for sorting molecules by mass.

Next, we ensure that each vertex occurs exactly once. Introducing reverse-complement oligonucleotides for each vertex to the set of witness candidates binds to the respective

vertex. If a vertex occurs multiple times in a path, then the string representation gets discarded. This process ensures each vertex corresponds to a potential Hamiltonian Path.

Once all of the vertices have been filtered, we check T using *detect* to determine if any valid paths remain. If valid paths exist, then the oligonucleotide from T may be read for the path assignment.

Definition 2.2.6. Detect determines if any encodings are present in T .

$$\text{detect}(T)$$

The output consists of ‘true’ or ‘false’ for $T \neq \emptyset$ or $T = \emptyset$, respectively.

In the implementation of a simulation system, we avoid redundant string representations with the *purify* operation. This is a synthetic version of PCR. Purify balances the space representation of molecules with a uniform distribution.

Definition 2.2.7. Purify provides a uniform distribution from the contents of T as T' .

$$T' \leftarrow \text{purify}(T)$$

2.3 Definition of SATISFIABILITY

Definition 2.3.1. SATISFIABILITY

$$\text{SATISFIABILITY} = \{\langle \phi \rangle \mid \phi \text{ is a satisfiable Boolean formula}\}[23].$$

Cook and Levin introduced independently the canonical instance of an NP-complete language SATISFIABILITY [4, 14]. An NP-complete language is one that is in NP and NP-hard. An NP-hard language is a decision problem that can be reduced in polynomial time from any NP language [23]. NP-hard problems includes the HALTING PROBLEM [23], which asks if a program on a given input can terminate. Witnesses for SATISFIABILITY or any other NP problem are of length polynomial in the length of the input ϕ .

Standard forms for SATISFIABILITY include Boolean CNF, k -CNF, and k -SAT problem definitions.

Definition 2.3.2. CNF is a Boolean formula that consists of the conjunction of sets of disjunctive literals.

Definition 2.3.3. k -CNF is a CNF Boolean formula where each disjunctive clause contains k literals.

Definition 2.3.4. k -SAT is a problem variant of SATISFIABILITY where the satisfiable instances are exactly k -CNF satisfiable.

One way to validate a SATISFIABILITY instance is to input witness candidates to a circuit. Let us consider a circuit for a SATISFIABILITY instance having three levels. This circuit consists of n inverters, m **OR** gates, and one **AND** gate with m -fan-in. This circuit behaves according to the internal wiring of the input CNF instance ϕ . Figure 2.1 contains a schematic for SATISFIABILITY.

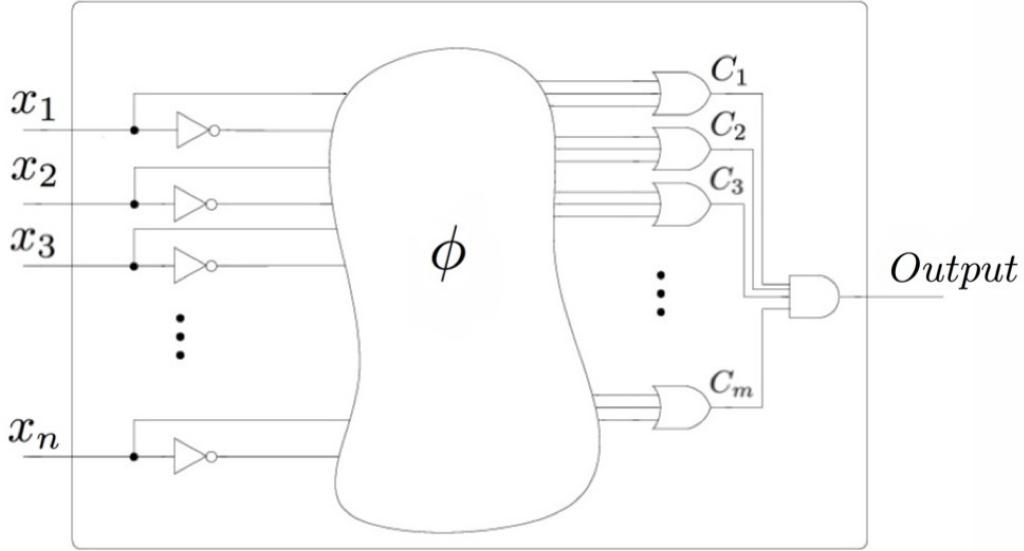


Figure 2.1: A circuit describing SATISFIABILITY.

The realization of SATISFIABILITY as a circuit reveals two aspects of this problem. SATISFIABILITY can be implemented as a circuit with the number of gates proportional to the problem size. The worst case verification for all 2^n possible witness candidates may be performed with the circuit in Figure 2.1. This circuit consists of the hardware equivalent version of the SATISFIABILITY validator CHECKSAT, as shown in Chapter 1.

2.4 Evaluating SAT Solvers

In this section, we describe the standards for encoding the SATISFIABILITY problem that we adopt. The standards come from the SATISFIABILITY Competition [5, 22].

Next, we introduce a problem instance classification scheme for SATISFIABILITY. Classification of SATISFIABILITY problem instances include random, combinatorial, and industrial SAT instances [22]. Our experiment in Chapter 6 generates random k -SAT inputs.

2.4.1 Input and output

The SAT Competition ranks implementations of solvers for evaluating SATISFIABILITY [22]. SAT solvers are evaluated on three categories of input: industrial, combinatorial, and random instances. The input and output standards for SATISFIABILITY allow common benchmarks for SAT solvers. We conform to the standards of this competition <http://www.satcompetition.org/>.

Input

DIMACS CNF provides a standard input for SATISFIABILITY [5]. The format permits sharing of existing SATISFIABILITY benchmarks by encoding SATISFIABILITY in conjunctive normal form (CNF). We provide an example of this encoding in Section 5.6.

Output

SAT Competition output consists of the status of a DIMACS CNF input instance [22]. The status is provided for the instance as either SATISFIABLE, UNSATISFIABLE, or UNKNOWN. If an instance can be determined as satisfiable, then a witness satisfying the instance gets included with the output status. We provide an example along with custom output logging in Section 5.7.

2.4.2 Metrics for classifying SATISFIABILITY

SAT phase transition and SAT backbones are two metrics for classifying SATISFIABILITY. We will use these metrics in the next section for defining a collection of random k -SAT instances.

The ratio of m clauses to n variables $\alpha = m/n$ provides a characterization where phase transitions may occur in the space of all k -CNF formula [6, 12]. The SAT phase transition is a region where both satisfiable and unsatisfiable instances are likely [12]. This region frequently separates trivially satisfiable and frequently over-constrained unsatisfiable SATISFIABLE problem instances. SATISFIABILITY instances with low α are frequently under-constrained and trivially satisfiable; those instances with high α are frequently over-constrained and trivially unsatisfiable [12].

Definition 2.4.1. SAT *backbones* are the variable assignments present in all of the satisfying assignments to a SATISFIABILITY problem instance [27].

SAT backbones contain a set of variables that occur in all satisfiable witnesses for an input instance. If there are no such variables in the set of all witnesses for a problem instance, then the set is empty.

2.4.3 SATISFIABILITY instances

There are several methods for generating SATISFIABILITY instances. We consider three classes [22]: random assignment, combinatorial problems, and industrial applications.

Random SAT

A random k -SAT instance [25], for fixed m and n , is one drawn uniformly from the set of all k -CNF formulas having m clauses and n variables.

Hard combinatorial SAT

Combinatorial problem instances are well known difficult benchmark cases. These instances include games and graph theoretic problems represented as SATISFIABILITY.

Industrial SAT

Industrial processes apply SATISFIABILITY to solve real world problems, including circuit layout, planning, logistics, circuit fault testing, and many other industrial NP-complete problems. Industrial SAT applications often apply heuristics and approximation techniques to relax the problem. This allows approximate solutions to be computed efficiently with respect to time.

Chapter 3

Existing molecular algorithms for SATISFIABILITY

In this chapter, we discuss two molecular algorithms for SATISFIABILITY. These algorithms construct sets of all witnesses for SATISFIABILITY instances. Lipton’s algorithm requires a combinatorial space of all witness candidates to be constructed and then filters out invalid ones. Ogihara and Ray’s algorithm constructs a set of witness candidates throughout execution. Following the description, we explore the physical implementations of—and simulation frameworks for—these algorithms.

Table 3.1: Components of Boolean literals and equivalent literal representations.

Literal	Variable (v)	Polarity (P)	DIMACS ($\pm v$)	Condensed (v_P)
x_1	1	T	1	1_T
$\neg x_1$	1	F	-1	1_F
x_n	n	T	n	n_T
$\neg x_n$	n	F	$-n$	n_F

The algorithm definitions and example traces use the literal conventions listed in Table 3.1. Table 3.1 lists components of a literal (variable and polarity), along with equivalent forms (DIMACS and a condensed representation).

In Chapter 1, witness candidates for SATISFIABILITY were represented as a bit-vector B . Consider the equivalent representation for witness candidates in Figure 3.1.

$$\begin{aligned}
B &= [0, 1, 0, 1] \\
L &= \{\neg x_1, x_2, \neg x_3, x_4\} \\
D &= \text{SFTFT} \\
Z &= \text{S-1+2-3+4}
\end{aligned}$$

Figure 3.1: The bit-vector $B = [0, 1, 0, 1]$ can be represented as the set of literal assignments $L = \{\neg x_1, x_2, \neg x_3, x_4\}$. A directed polarity string (D) with initial sequence (S), followed by a sequence of literals provides $D = \text{SFTFT}$. A directed integer string (Z) consists of an initial sequence (S) followed by DIMACS literal assignments $Z = \text{S-1+2-3+4}$.

We use the directed string notation (representation D in Figure 3.1) as shorthand for a directed oligonucleotide. The directed string representation D can be indexed by the variable v using the condensed literal from Table 3.1. In Lipton’s algorithm, literal configurations for a variable v get extracted directly (v_T or v_F).

Ogihara and Ray’s algorithm extracts satisfying literal configurations from an ordered clause (a, b, c) . We use the condensed literal notation v_P to indicate the assignment (either T or F) for the literal v . We use the literal configuration v_N to indicate any other non-satisfiable literal assignment to v .

In the discussion of the Distribution algorithm in Chapter 4, we use the directed integer notation (representation Z in Figure 3.1) as shorthand for a directed sequence of integers.

Along with our shorthand representations (D and Z) from Figure 3.1, we include a small combinatorial library corresponding to positive and negative literals. Table 3.2 lists a mapping of literals to oligonucleotides used for literal matching and assignment.

Table 3.2: Positive and negative literal assignment and matching oligonucleotides.

Literal	Assignment	Matching
x_1	5'-TTT-3'	3'-AAA-5'
$\neg x_1$	5'-TTA-3'	3'-AAT-5'
x_2	5'-CTT-3'	3'-GAA-5'
$\neg x_2$	5'-ATT-3'	3'-TAA-5'
x_3	5'-ATG-3'	3'-TAC-5'
$\neg x_3$	5'-GTT-3'	3'-CAA-5'
x_4	5'-TCT-3'	3'-AGA-5'
$\neg x_4$	5'-CCT-3'	3'-GGA-5'
x_5	5'-ACT-3'	3'-TGA-5'
$\neg x_5$	5'-GCT-3'	3'-CGA-5'
Start	5'-TTG-3'	3'-AAC-5'

The examples throughout this report use this combinatorial library designed with codons. Larger combinatorial libraries [13] take multiple considerations for the genetic encoding, including uniform melting temperature and non-self complementary sequences. Codons provide redundant encodings that we exploit for a small library.

3.1 Lipton’s algorithm for SATISFIABILITY

Introduced in 1995 by Richard Lipton [16], this algorithm filters satisfiable witnesses from a combinatorial space of all witness candidates. Lipton’s algorithm is analogous to a conventional brute-force search for all witnesses of a SATISFIABILITY instance.

Lipton’s algorithm (Algorithm 3.2) first constructs a combinatorial space T containing oligonucleotide configurations for all potential witness candidates. The algorithm iterates over each of the clauses C in ϕ .

From each clause C , each of the literals contained within each clause C get filtered to satisfiable witnesses. The contents of T get filtered by incrementally extracting those literals that satisfy the current clause. The next iteration filters witnesses that satisfy the previous clauses from T and the literal contents from C . The algorithm terminates with a set of witnesses T for the CNF input ϕ . If ϕ is unsatisfiable, then $T = \emptyset$.

Algorithm 3.2: LIPTON'S ALGORITHM(ϕ)

```
 $T \leftarrow \text{COMBINATORIAL GENERATE}(n)$ 
for each clause  $C$  in  $\phi$ 
  do  $\begin{cases} T_c \leftarrow \emptyset \\ \text{for each literal } v \text{ in } C \\ \quad \text{do } \begin{cases} \text{if } v \text{ is a positive literal} \\ \quad \text{then } \begin{cases} T_P \leftarrow \text{extract}(T, v_T) \\ T_c \leftarrow \text{mix}(T_P, T_c) \end{cases} \\ \text{else } \begin{cases} T_N \leftarrow \text{extract}(T, v_F) \\ T_c \leftarrow \text{mix}(T_N, T_c) \end{cases} \end{cases} \\ T \leftarrow \emptyset \\ T \leftarrow \text{mix}(T, T_C) \\ T \leftarrow \text{purify}(T_c) \end{cases}$ 
return (detect( $T$ ))
```

Algorithm 3.2: LIPTON'S ALGORITHM iterates over each of the m clauses. The contents of T_C grows incrementally with configurations from T that satisfy the literal v . Once the entire clause C has been evaluated, T_C contains configurations that witness the observed conditions. The contents of T_C are stored as T for the next clause; T now contains configurations that witness all previous clauses. Once complete, the tube T contains all witnesses for ϕ , if any witnesses exist.

3.1.1 Description of Lipton's algorithm

The function COMBINATORIAL GENERATE (Algorithm 3.3) implements the split-mix synthesis technique [9, 10]. COMBINATORIAL GENERATE returns a tube T_{comb} consisting of oligonucleotides that represent all 2^n distinct witness candidates. The tube T_{comb} begins with an initial medium denoted by \mathbf{S} . An iterative loop extends T_{comb} using split-mix synthesis. Each split corresponds with appending the tubes with a true (T) and false (F) assignment. The two tubes are mixed and purified to contain equimolar portions of each witness candidate.

Algorithm 3.3: COMBINATORIAL GENERATE(n)

```

 $T_{comb} \leftarrow \{\mathbf{S}\}$ 
for  $v \leftarrow 1$  to  $n$ 
  do  $\begin{cases} [T_1, T_2] \leftarrow \text{split}(T_{comb}) \\ T_1 \leftarrow \text{append}(T_1, v_T) \\ T_2 \leftarrow \text{append}(T_2, v_F) \\ T_{comb} \leftarrow \text{mix}(T_1, T_2) \end{cases}$ 
 $T_{comb} \leftarrow \text{purify}(T_{comb})$ 
return ( $T_{comb}$ )
  
```

Algorithm 3.3: COMBINATORIAL GENERATE constructs a combinatorial space consisting of 2^n molecular configurations in polynomial time.

Let us consider an example execution of COMBINATORIAL GENERATE with $n = 2$. The tube T_{comb} begins as an empty tube. A start configuration \mathbf{S} initiates the tube T_{comb} with a medium for combinatorial synthesis. We begin with the initial contents

$$T_{comb} = \{\mathbf{S}\},$$

$$T_{comb} = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}.$$

Iteration $v = 1$:

First, split the contents of T_{comb} . We have

$T_1 = \{\mathbf{S}\}$ $T_1 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} - 3'\}$	$T_2 = \{\mathbf{S}\}$ $T_2 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} - 3'\}.$
--	---

Next, append each of the tubes with a positive (T) and negative (F) assignment for the literal v_1 . We have

$T_1 = \{\mathbf{ST}\}$ $T_1 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTT}}_{x_1} - 3'\}$	$T_2 = \{\mathbf{ST}, \mathbf{SF}\}$ $T_2 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTA}}_{\neg x_1} - 3'\}.$
---	--

Mix the contents of T_1 and T_2 to form T_{comb} for the next iteration. We have

$T_{comb} = \{\mathbf{ST}, \mathbf{SF}\}.$ $T_{comb} = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTT}}_{x_1} - 3',$ $5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTA}}_{\neg x_1} - 3'\}.$
--

Iteration $v = 2$:

Split the contents of T_{comb} . We have

$T_1 = \{\mathbf{ST}, \mathbf{SF}\}$ $T_1 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTT}}_{x_1} - 3',$ $5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTA}}_{\neg x_1} - 3'\}$	$T_2 = \{\mathbf{ST}, \mathbf{SF}\}$ $T_2 = \{5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTT}}_{x_1} - 3',$ $5' - \underbrace{\mathbf{TTG}}_{\text{Start}} \cdot \underbrace{\mathbf{TTA}}_{\neg x_1} - 3'\}.$
--	---

Next, append each of the tubes with a positive (T) and negative (F) assignment for the literal v_2 . We have

$$T_1 = \{\text{STT}, \text{SFT}\}$$

$$T_1 = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

$$T_2 = \{\text{STF}, \text{SFF}\}$$

$$T_2 = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

Mix the contents of T_1 and T_2 to form T_{comb} for the final iteration. The algorithm COMBINATORIAL GENERATE returns the following tube

$$T_{comb} = \{\text{STT}, \text{SFT}, \text{STF}, \text{SFF}\},$$

$$T_{comb} = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

COMBINATORIAL GENERATE generates all witness candidates for a SATISFIABILITY instance. Lipton's algorithm filters, from a combinatorial space T , configurations that represent witnesses for the input ϕ .

3.1.2 Detailed trace of Lipton's algorithm

Appendix B lists a detailed execution trace for Lipton's algorithm.

3.2 Ogihara and Ray's algorithm for SATISFIABILITY

Ogihara and Ray's algorithm (Algorithm 3.4) consist of a breadth-first evaluation of clauses from a CNF formula [19, 20]. The algorithm constructs a set of witness candidates based on a parse of a 3-CNF formula. In this section, we describe the preconditions and execution of Ogihara and Ray's algorithm.

Algorithm 3.4: OGIHARA AND RAY'S ALGORITHM(ϕ)

```

// Input  $\phi$  consists of  $n$  variables.
// Each clause  $C$  contains ordered literals  $(a, b, c)$ .
// Extract literals using the condensed notation  $v_P$ , where:
//    $v_P$  matches the literal, and  $v_N$  matches the negated literal.

 $T \leftarrow \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$ 
for  $v \leftarrow 3$  to  $n$ 
   $[T_P, T_N] \leftarrow \text{split}(T)$ 
  for each clause  $C$  in  $\phi$ 
     $(a, b, c) \leftarrow C$ 
    if  $v_T = c$ 
      then  $\begin{cases} T_{P1} \leftarrow \text{extract}(T_N, a_P) \\ T_{N1} \leftarrow \text{extract}(T_N, a_N) \\ T_{P2} \leftarrow \text{extract}(T_{N1}, b_P) \\ T_N \leftarrow \text{mix}(T_{P1}, T_{P2}) \\ T_N \leftarrow \text{purify}(T_N) \end{cases}$ 
    do  $\begin{cases} \text{if } v_F = c \\ \text{then } \begin{cases} T_{P1} \leftarrow \text{extract}(T_P, a_P) \\ T_{N1} \leftarrow \text{extract}(T_P, a_N) \\ T_{P2} \leftarrow \text{extract}(T_{N1}, b_P) \\ T_P \leftarrow \text{mix}(T_{P1}, T_{P2}) \\ T_P \leftarrow \text{purify}(T_P) \end{cases} \end{cases}$ 
     $T_P \leftarrow \text{append}(T_P, v_T)$ 
     $T_N \leftarrow \text{append}(T_N, v_F)$ 
     $T \leftarrow \text{mix}(T_P, T_N)$ 
     $T \leftarrow \text{purify}(T)$ 
  return ( $\text{detect}(T)$ )

```

Algorithm 3.4: OGIHARA AND RAY'S ALGORITHM evaluates each subsequent variable and determines possible assignments. The possible assignments for the variables a and b get extracted if c matches the current variable v . Effectively pruning only potential solutions. These potential solutions T_P and T_N get appended with the positive or negative string assignments. The algorithm continues until each variable gets evaluated. The remaining space T contains all solutions for the CNF instance ϕ after the algorithm terminates.

3.2.1 Description of Ogihara and Ray's algorithm

Ogihara and Ray's algorithm begins with four initial witness candidates, we have

$$T = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$$

$$T = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$$

$$5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3',$$

$$5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$$

$$5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

For each clause $C = (x_i \vee x_j \vee x_k)$, we have that $1 \leq i < j < k \leq n$. As an example, let us evaluate the clause

$$C = x_1 \vee \neg x_2 \vee \neg x_3.$$

On the first iteration, we compare the third ordered literal c with x_3 . Since $c = \neg x_3$, extract configurations that satisfy $a \vee b$.

$T_P = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$ $T_P = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\},$	$T_N = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$ $T_N = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3',$ $5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$
--	--

From T , select configurations that satisfy $a_P = a_T$

$$T_{P1} = \{\text{STT}, \text{STF}\}.$$

$$T_{P1} = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}$$

From T , Select configurations that satisfy $a_N = a_F$

$$T_{N1} = \{\text{SFT}, \text{SFF}\}.$$

$$T_{N1} = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}$$

From T_{N1} , select configurations that satisfy $b_P = b_F$

$$T_{P2} = \{\text{SFF}\},$$

$$T_{P2} = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

Mix the contents of T_{P1} and T_{P2} as the contents of T_P

$$T_P = \{\text{STT}, \text{STF}, \text{SFF}\},$$

$$T_P = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

We have the tubes

$$T_P = \{\text{STT}, \text{STF}, \text{SFF}\}$$

$$T_P = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\},$$

$$T_N = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$$

$$T_N = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}.$$

Finally append assignments that satisfy the current literal with T_P and T_F .

$$T_P = \text{append}(T_P, c_F) \\ = \{\text{STTF}, \text{STFF}, \text{SFFF}\}$$

$$T_P = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3'\}$$

$$T_N = \text{append}(T_N, c_T) \\ = \{\text{STTT}, \text{STFT}, \text{SFTT}, \text{SFFT}\}$$

$$T_N = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3'\}.$$

Mix the contents of T_P and T_N to form the set of configurations that witness the clause.

$$T = \{\text{STTF}, \text{STFF}, \text{SFFF}, \text{STTT}, \text{STFT}, \text{SFTT}, \text{SFFT}\}$$

$$\begin{aligned} T = & \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{GTT}}_{\neg x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ & 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTA}}_{\neg x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3'\}. \end{aligned}$$

3.2.2 Detailed trace of Ogihara and Ray's algorithm

Appendix B lists a detailed execution trace for Ogihara and Ray's algorithm.

3.3 Implementations of molecular SATISFIABILITY solvers

We discuss existing implementations of molecular SATISFIABILITY solvers. Physical implementations apply molecular biology techniques and actual molecules. Simulation frameworks use standard computation to simulate molecular biology techniques.

3.3.1 Physical implementations

Yoshida and Suyama implemented Ogihara and Ray's algorithm using manual molecular biology techniques [26]. This experiment solved a 3-CNF instance with four variables and 10 clauses.

Braich et al. implemented a molecular computer to filter solutions for a 3-SAT instance [3]. This experiment solved a 3-CNF instance with 20 variables and 24 clauses.

3.3.2 Simulation frameworks

Martín-Mateos et al. introduced a simulation for Lipton’s algorithm [18]. Molecular operations get implemented in **ACL2**, a Common Lisp variant. The framework for this system implemented test cases for Lipton’s algorithm.

Ogihara provides test results for an implementation of his original molecular algorithm [19]. This simulation provides a comparison to Lipton’s algorithm for practical length restrictions.

Chapter 4

A new molecular algorithm for SATISFIABILITY

This chapter introduces a new molecular algorithm for SATISFIABILITY: the Distribution algorithm distributes literals from a CNF instance into a set of non-conflicting witnesses.

4.1 Distribution algorithm for SATISFIABILITY

The Distribution algorithm (Algorithm 4.1) begins with a CNF instance ϕ . The algorithm constructs sets of witnesses stored in T_C that satisfy the current clause C along with all previously evaluated witnesses in the tube T_S . Once the Distribution completes the tube T_S contain witnesses for ϕ ; when ϕ is unsatisfiable the tube $T_S = \emptyset$.

The algorithm begins by initiating a solution tube T_S with a start sequence S . During the evaluation of m clauses¹, the tube stores witnesses for the CNF instance

$$\bigwedge_{1 \leq i \leq m} C_i.$$

The tube T_C stores witnesses for each clause. The tube T_C begins as the empty set on reading each clause C , and grows with the satisfying literal assignments.

During the evaluation of each literal, we split the contents of T_S into the tubes T_V and a copy of T_S . We use the tube T_V to filter positive and negative literal assignments into the tubes T_P and T_F , respectively. The copy of the tube T_S retains the witness candidates for each of the literals in C .

We extract from T_V witnesses candidates that conflict with the current literal in the tube T_F , and those with idempotent assignments in the tube T_P . The remaining contents

¹We consider only the parse ordering of the clauses from the CNF instance ϕ . The Distribution algorithm will resolve conflict uniquely depending on the permutations of the clauses in ϕ .

of the tube T_V , if any, contain witness candidates that can witness the current clause with the assignment of the current literal v .

The Distribution algorithm extends T_V by appending the contents with the assignment v_p in the same polarity of the current literal v . We mix the clause witnesses (T_C) with the contents of the idempotent witnesses (T_P), extended witnesses (T_V), along with existing clause witnesses in T_C .

On each iteration of clauses, the Distribution algorithm clears the contents of the tube T_S and passes the clause witnesses (T_C) to the remaining clauses. We purify the contents of T_S to create a uniform distribution of the witness candidates. Once the Distribution algorithm completes the tube T_S contains witnesses for ϕ ; if $T_S = \emptyset$ then ϕ is unsatisfiable.

Algorithm 4.1: DISTRIBUTION ALGORITHM(ϕ)

```

 $T_S \leftarrow \{S\}$ 
for each clause  $C$  in  $\phi$ 
   $T_C \leftarrow \emptyset$ 
  for each literal  $v$  in  $C$ 
    do  $\left\{ \begin{array}{l} [T_V, T_S] \leftarrow \text{split}(T_S) \\ T_N \leftarrow \text{extract}(T_V, v_N) \\ T_P \leftarrow \text{extract}(T_V, v_P) \\ T_V \leftarrow \text{append}(T_V, v_P) \\ T_C \leftarrow \text{mix}(T_C, T_P, T_V) \end{array} \right.$ 
     $T_S \leftarrow \emptyset$ 
     $T_S \leftarrow \text{mix}(T_S, T_C)$ 
     $T_S \leftarrow \text{purify}(T_S)$ 
  return (detect( $T_S$ ))

```

Algorithm 4.1: The DISTRIBUTION ALGORITHM constructs a set of witnesses for a CNF instance ϕ by clause evaluation. The literals of each clause C get distributed into the clause witness tube T_C . These witnesses contained in T_C continue to witness the evaluated portion of ϕ in the solution witness tube T_S .

4.1.1 Description of the Distribution algorithm

Let us consider an example CNF instance designed to show several aspects of the Distribution algorithm.

$$\phi = (x_1 \vee x_2) \wedge (x_1 \vee \neg x_2 \vee x_3)$$

This example demonstrates several key aspects of the algorithm. We observe how the algorithm initiates the tube T_S with literals from the first clause. The second clause contains a literal of the same (x_1) and opposite polarity ($\neg x_2$) of the literals in the first clause. The literal (x_3) demonstrates when the algorithm does match any existing literal assignments in T_S ; causing the assignment of (x_3) to be appended to all witness candidates in T_V .

With this example, we demonstrate filtering witness candidates in T_V into the tubes T_P and T_N . The tube T_N contains witness candidates that conflict with the current literal, and the tube T_P contains witnesses for the current literal. The remaining contents of T_V contain witness candidates that do not contain the current literal. We append the contents of T_V with the current literal, and store the contents of $T_V \cup T_P$ as witnesses for the current clause in the tube T_C .

After each literal from a clause has been distributed, the contents of T_C contain witnesses for both the current clause and all previously distributed clauses. The contents of T_C get stored as the tube T_S . The tube T_S stores the witness candidates for the next clause. Once each clause has been distributed, the tube T_S contains witnesses for the k -CNF instance ϕ ; if $T_S = \emptyset$, then ϕ is unsatisfiable.

Initiate the tube T_S with a start medium S .

$$T_S = \{S\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}$$

Evaluate clause $C_1 = (x_1 \vee x_2)$:

$$T_C = \{ \}$$

Select the literal x_1 from C_1 . Split the contents of T_S :

$$T_V = \{S\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}$$

$$T_S = \{S\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}$$

Extract literals from T_V that contain negative and positive assignments for x_1 .

$$T_N = \{ \}$$

$$T_P = \{ \}$$

$$T_V = \{S\}$$

Append the satisfying literal x_1 to the required variable tube T_V .

$$T_V = \{S+1\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3'\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+1\}$$

$$T_C = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3'\}$$

Select the literal x_2 from C_1 . Split the contents of T_S :

$$T_V = \{S\}$$

$$T_S = \{S\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} - 3'\}$$

Extract literals from T_V that contain negative and positive assignments for x_2 .

$$T_N = \{ \}$$

$$T_P = \{ \}$$

$$T_V = \{S\}$$

Append the satisfying literal x_2 to the required variable tube T_V .

$$T_V = \{\text{S+2}\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{\text{S+1}, \text{ S+2}\}$$

$$T_C = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

Complete the iteration of the first clause by storing the witnesses in T_C as witnesses to T_S for the evaluated instance ϕ .

$$T_S = \{\text{S+1}, \text{ S+2}\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

Evaluate clause $C_2 = (x_1 \vee \neg x_2 \vee x_3)$:

$$T_C = \{ \}$$

Select the literal x_1 from C_2 . Split the contents of T_S :

$$T_V = \{\text{S+1}, \text{ S+2}\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

$$T_S = \{\text{S+1}, \text{ S+2}\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

Extract literals from T_V that contain negative and positive assignments for x_1 .

$$\begin{aligned} T_N &= \{\} \\ T_P &= \{S+1\} \\ T_V &= \{S+2\} \end{aligned}$$

Append the satisfying literal x_1 to the required variable tube T_V .

$$\begin{aligned} T_V &= \{S+2+1\} \\ T_V &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{TTT}}_{x_1} - 3'\} \end{aligned}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$\begin{aligned} T_C &= \{S+1, S+2+1\} \\ T_C &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ &\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{TTT}}_{x_1} - 3'\} \end{aligned}$$

Select the literal $\neg x_2$ from C_2 . Split the contents of T_S :

$$\begin{aligned} T_V &= \{S+1, S+2\} \\ T_V &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ &\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\} \\ T_S &= \{S+1, S+2\} \\ T_S &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ &\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\} \end{aligned}$$

Extract literals from T_V that contain negative and positive assignments for $\neg x_2$.

$$T_N = \{\text{S+2}\}$$

$$T_P = \{\}$$

$$T_V = \{\text{S+1}\}$$

Append the satisfying literal $\neg x_2$ to the required variable tube T_V .

$$T_V = \{\text{S+1-2}\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{\text{S+1}, \text{ S+2+1}, \\ \text{ S+1-2}\}$$

$$T_C = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3'\}$$

Select the literal x_3 from C_2 . Split the contents of T_S :

$$T_V = \{\text{S+1}, \text{ S+2}\}$$

$$T_V = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

$$T_S = \{\text{S+1}, \text{ S+2}\}$$

$$T_S = \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} - 3'\}$$

Extract literals from T_V that contain negative and positive assignments for x_3 .

$$\begin{aligned}T_N &= \{\} \\T_P &= \{\} \\T_V &= \{\text{S+1}, \text{ S+2}\}\end{aligned}$$

Append the satisfying literal x_3 to the required variable tube T_V .

$$\begin{aligned}T_V &= \{\text{S+1+3}, \text{ S+2+3}\} \\T_V &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\&\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3'\}\end{aligned}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$\begin{aligned}T_C &= \{\text{S+1}, \text{ S+2+1}, \\&\quad \text{S+1-2}, \\&\quad \text{S+1+3}, \text{ S+2+3}\} \\T_C &= \{5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\&\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\&\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\&\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\&\quad 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3'\}\end{aligned}$$

Complete the iteration of the last clause by storing the witnesses in T_C as witnesses to T_S . The tube T_S contains witnesses for ϕ .

$$T_S = \{ \text{S+1}, \text{ S+2+1}, \\ \text{ S+1-2}, \\ \text{ S+1+3}, \text{ S+2+3} \}$$

$$T_S = \{ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{TTT}}_{x_1} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATT}}_{\neg x_2} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{TTT}}_{x_1} \cdot \underbrace{\text{ATG}}_{x_3} - 3', \\ 5' - \underbrace{\text{TTG}}_{\text{Start}} \cdot \underbrace{\text{CTT}}_{x_2} \cdot \underbrace{\text{ATG}}_{x_3} - 3' \}$$

4.1.2 Detailed trace of the Distribution algorithm

Appendix B lists a detailed execution trace for the Distribution algorithm.

Chapter 5

Molecular Simulation: A system for molecular computation

This chapter introduces Molecular Simulation: A system for molecular computation. We describe an overview of Molecular Simulation and its documentation, along with tools for automated execution for Molecular Simulation. This includes `Perl` execution scripts and visualization for output data. We describe example input and output for Molecular Simulation. Command line argument provides user configurable options for Molecular Simulation. Chapter 6 describes the usage of Molecular Simulation with automated execution.

5.1 Overview

Molecular Simulation implements a simulated molecular lab for operating on DNA. The present simulation implements three molecular algorithms for SATISFIABILITY. The included `Perl` scripts process DIMACS CNF input directories with invocations to Molecular Simulation.

Molecular Simulation may be executed directly or invoked with the assistance of an execution script. The system requirements to execute or design a molecular experiment are listed in this section.

This program is a simulated molecular lab for experimenting with DNA operations. Implementation of three molecular algorithms for solving SATISFIABILITY include Lipton's algorithm, Ogihara and Ray's algorithm, and the Distribution algorithm. Chapters 3 and 4 describe the background and provide pseudocode for these algorithms.

5.2 Download

Download Molecular Simulation from: <https://github.com/dncarley/MolecularSimulation>.

5.3 Requirements

This section specifies the requirements for running Molecular Simulation.

5.3.1 Hardware requirements

Molecular Simulation requires a 64-bit processor with 2 GB of RAM.

5.3.2 Software requirements

`gcc` (GNU Compiler Collection) must be installed to build Molecular Simulation.

`Perl` must be installed to automate build and execution of Molecular Simulation.

5.4 Documentation

The project website contains detailed documentation for Molecular Simulation. The documentation provides an overview of Molecular Simulation that may be used independently of Chapters 5 and 6 for getting started. The online documentation describes detailed datatype, function, and class definitions.

5.5 Tools

This project uses several tools for automating tasks and execution. In this section, we discuss tools to automate execution and visualize output from Molecular Simulation.

5.5.1 Perl utilities

The source directory includes several `Perl` scripts to assist in building and initiation of tests for Molecular Simulation. Table 5.1 documents the basic usage for build and testbench execution scripts. Each script provides detailed execution options.

Table 5.1: Perl execution commands and descriptions.

Perl script	Usage	Description
build.pl	\$ perl build.pl	Compiles Molecular Simulation and generates an executable in the directory <code>./execute/simulation</code> .
buildGenerate.pl	\$ perl buildGenerate.pl	Generates a sweep of CNF formulas over a range of k -SAT ratios. Program uses a modified random k -SAT generator from Microsoft Research.
executeMolecularSat.pl	\$ perl executeMolecularSat.pl	Executes Molecular Simulation for a directory of SATISFIABILITY instances with desired algorithms. If no options are specified, then each of the three algorithms are executed and output is generated in the same test directory.
runSimulation.pl	\$ perl runSimulation.pl	Executes <code>build.pl</code> followed by <code>executeMolecularSat.pl</code> . Any command line arguments get passed to <code>executeMolecularSat.pl</code>

5.6 Input

Input to Molecular Simulation consists of a DIMACS CNF file. The definition of the *.cnf filetype can be accessed from: <ftp://dimacs.rutgers.edu/pub/challenge/satisfiability/doc/>.

```
c comments begin with a 'c'
c
c cnf input is designated with 'p cnf'
c     followed by number of variables <n>, and clauses <m>
c
p cnf 4 3
c
c A clause is represented by a sequence of <k> integers,
c     separated by whitespace and ending with a '0'.
c Each variable is represented by the integer sequence,
c     negative polarity is represented by '-'.
c
1 2 -3 0
2 3 -4 0
-1 -3 4 0
```

5.7 Output

Output from Molecular Simulation, by default, conforms to the 2011 SAT Competition rules. The rules can be accessed from: <http://www.satcompetition.org/2011/rules.pdf>.

```
c comments begin with a 'c'
c
s SATISFIABLE
c
c A line beginning with a 's' marks the status.
c This can be either 'UNSATISFIABLE', 'SATISFIABLE', or 'UNKNOWN'.
c
v 1 2 -3 -4 0
c
c A satisfiable witness begins with a 'v' and ends with a '0'.
c     A sequence of integers, between 'v' and '0', encodes a satisfiable assignment.
```

Table 5.2 describes an extended custom output. This output reports parameters for metric performance evaluation.

Table 5.2: Molecular Simulation output logging.

Parameter	Description
c algorithmType:	Display the algorithm type: Lipton, Ogihara-Ray, Distribution
c algorithmTime:	Display the algorithm execution time in seconds.
c solutionMemory:	Display the solution space memory in Bytes.
c mixCount:	Display the number of mixes required during algorithm execution.
c extractCount:	Display the number of extracts required during algorithm execution.
c appendCount:	Display the number of appends required during algorithm execution.
c splitCount:	Display the number of splits required during algorithm execution.
c spliceCount:	Display the number of splices required during algorithm execution.
c purifyCount:	Display the number of purifications required during algorithm execution.
c numVar:	Display the number of variables in the input CNF instance.
c numClause:	Display the number of clauses in the input CNF instance.

5.8 Execution

Invocation of Molecular Simulation can be performed from the command line.

```
$ ./execute/simulation i [input] [options]
```

The [input] consists of a DIMACS CNF file. Command line [options] may be a combination of the options in Table 5.3.

Table 5.3: Command line options for Molecular Simulation. Use the Perl scripts for automated execution (See Table 6.1).

Argument	Parameters	Description
-a	d l o	Algorithm select Distribution algorithm Lipton's algorithm Ogihara and Ray's algorithm
-d		Debug
i	[input]	Input DIMACS CNF file
-w	[output]	Write output to file Output filename

5.8.1 Execution example

Suppose that we would like to execute Ogihara and Ray's algorithm for a DIMACS CNF file instance `test1.cnf` located in the directory `MolecularSimulation/testbench`. We output the results `test1-o.out` in the same directory as the input CNF.

We invoke Molecular Simulation with the following command:

```
$ ./execute/simulation i ../testbench/test1.cnf -a o -w ../testbench/test1-o.out
```

In the next chapter, we will describe the automation for a random k -SAT sweep with each of the algorithms. The provided Perl scripts are the recommended method for building and execution of Molecular Simulation.

Chapter 6

Experimental Setup

This chapter describes the use of Molecular Simulation for evaluation of a set of DIMACS CNF SATISFIABILITY instances. We discuss configuration for generation of random k -SAT instances. Further, any existing DIMACS CNF benchmark may be imported for test. Example configuration options automate the execution of Molecular Simulation. The example continues with an analysis of runtime metrics for each test instance. The next chapter describes the results from the k -SAT sweep experiment.

6.1 Setup

In this section, we describe prerequisites for executing a test bench using Molecular Simulation. Molecular Simulation requires a 64-bit architecture with a UNIX like system with `gcc` and `Perl`. The target system must meet the minimum requirements.

Building Molecular Simulation can be performed by invoking the `Perl` script `build.pl` from the command line.

```
$ perl build.pl
```

This script generates an executable `simulation` in the directory:

```
MolecularSimulation/execute
```

The next sections describe invocation of Molecular Simulation with desired options. We begin with the creation and importation of DIMACS CNF datasets.

6.2 Create dataset

We create a sweep of random k -SAT instances to observe SAT phase transition. This set consists of random 3-CNF instances at fixed $n = 20$ spanning clause-variable ratios $\alpha = m/n = [0.2, 14.0]$ in increments of 0.2. Each α ratio consists of 30 3-CNF instances.

David Wilson's `ksat.c` generates random k -SAT instances in DIMACS CNF format [25]. The program takes four arguments to create a unique DIMACS CNF instance. Invocation of the program can be performed using the following command:

```
$ ./execute/ksat k n m s > output.cnf
```

This generates `output.cnf` in DIMACS CNF format with k variables per clause n variables, m clauses, and random seed s .

We use automated Perl scripts to create a sweep of DIMACS CNF instances. Setup for a sweep configuration includes specifying a set of ratios. Invocation of the script generates a set of random k -SAT instances. The redirected output gets stored in the target directory with the previous file naming convention. We use the following command to invoke the construction of a sweep of k -SAT instances.

```
$ perl buildGenerate.pl
```

6.3 Import dataset

Datasets of DIMACS CNF input may be provided for batch processing. This includes random k -SAT instances generated from the previous section, or importing existing DIMACS CNF instances.

DIMACS CNF benchmarks are available for download from: <ftp://dimacs.rutgers.edu/pub/challenge/satisfiability/>.

6.4 Configure test

The previous chapter described a single execution of Molecular Simulation. We use automated scripts for processing datasets with each of the algorithms.

The Perl script `executeMolecularSat.pl` allows execution for a directory of DIMACS CNF input. Executing the script from the command line without arguments processes the experimental setup and saves output to the same directory.

```
$ perl executeMolecularSat.pl [options]
```

The options for `executeMolecularSat.pl` can be a combination of the options in Table 6.1.

Table 6.1: Command line options for `executeMolecularSat.pl`

Argument	Parameters	Description
-d -l -o		Distribution algorithm Lipton's algorithm Ogihara and Ray's algorithm Default: Execute all three algorithms.
-debug		Debug
-p	[CNF file path]	Specify CNF file path. Default path: data/testCNF
-f		Write output to file

6.5 Execution and collection of data

The following command builds and executes Molecular Simulation.

```
$ perl runSimulation.pl [options]
```

This command first builds Molecular Simulation with `build.pl`, and invokes Molecular Simulation with `executeMolecularSat.pl`. The `[options]` are passed directly to `executeMolecularSat.pl`. Molecular Simulation executes the default experimental setup with no `[options]` specified.

Output consists of the standard SAT Competition output appended with custom runtime metric logging. Collections of output files may be read by the data visualization program and exported into a condensed table.

6.5.1 Execution output

Molecular Simulation, by default, writes output to standard output on the console. The `-f` option saves output to a file as `[filename]-<a>.out`. The `[filename]` consists of the DIMACS CNF name and `<a>` specifies the algorithm type: `d`, `l` or `o`.

Output directed to standard output conforms to the SAT Competition rules. This output may be used during testing, or redirected to an external stream. The debug option

-debug displays detailed information about the execution. The debug option writes verbose content based on the program execution.

Reading output metrics from the saved output, as defined in Table 5.2, allows for analysis of collected data. Output files from Molecular Simulation get condensed into a Tab Separated Values (*.tsv) file. Subsequent datapoint browsing and the online view use the *.tsv file for condensed reading and transmission of data points. In the next chapter, we describe the results of the experimental setup.

Chapter 7

Results

This chapter presents results of the k -SAT execution test from the previous chapter. We consider the results of the test and analyze the algorithm metrics.

7.1 Algorithm metric comparison

This section describes the results from the simulation. We analyze the molecular operations count for append, extract, mix, purify, splice, and split. Presentation of actual computation time and required memory for the solution representation allow for comparison of algorithms.

The plots in this chapter use the shapes in Figure 7.1 for algorithm type. We superimpose a line over each plot showing the percent of satisfiable 3-CNF instances on each clause-variable ratio α for our test cases.

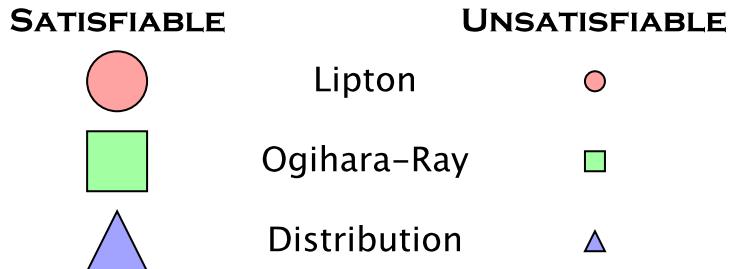


Figure 7.1: Key for output metrics. Large shapes represent satisfiable instances and small shapes represent unsatisfiable instances. Datapoints for Lipton’s algorithm are represented with red circles, Ogihara and Ray’s algorithm with green squares, and the Distribution algorithm with blue triangles.

Append concatenates two oligonucleotides. Figure 7.2 shows the number of appends for the naive implementations. Figure 7.3 shows early termination on unsatisfiable 3-CNF instances.

The Distribution algorithm uses $O(k \cdot m)$ appends. For unsatisfiable k -CNF input, the algorithm may terminate if there exist no witness candidates exist for the distribution of literals from remaining clauses.

Lipton's algorithm uses $\Theta(n)$ appends during the creation of a combinatorial space of 2^n witness candidates.

Ogihara and Ray's algorithm uses $O(n)$ appends. Unsatisfiable k -CNF input may terminate once a conflict has been detected.

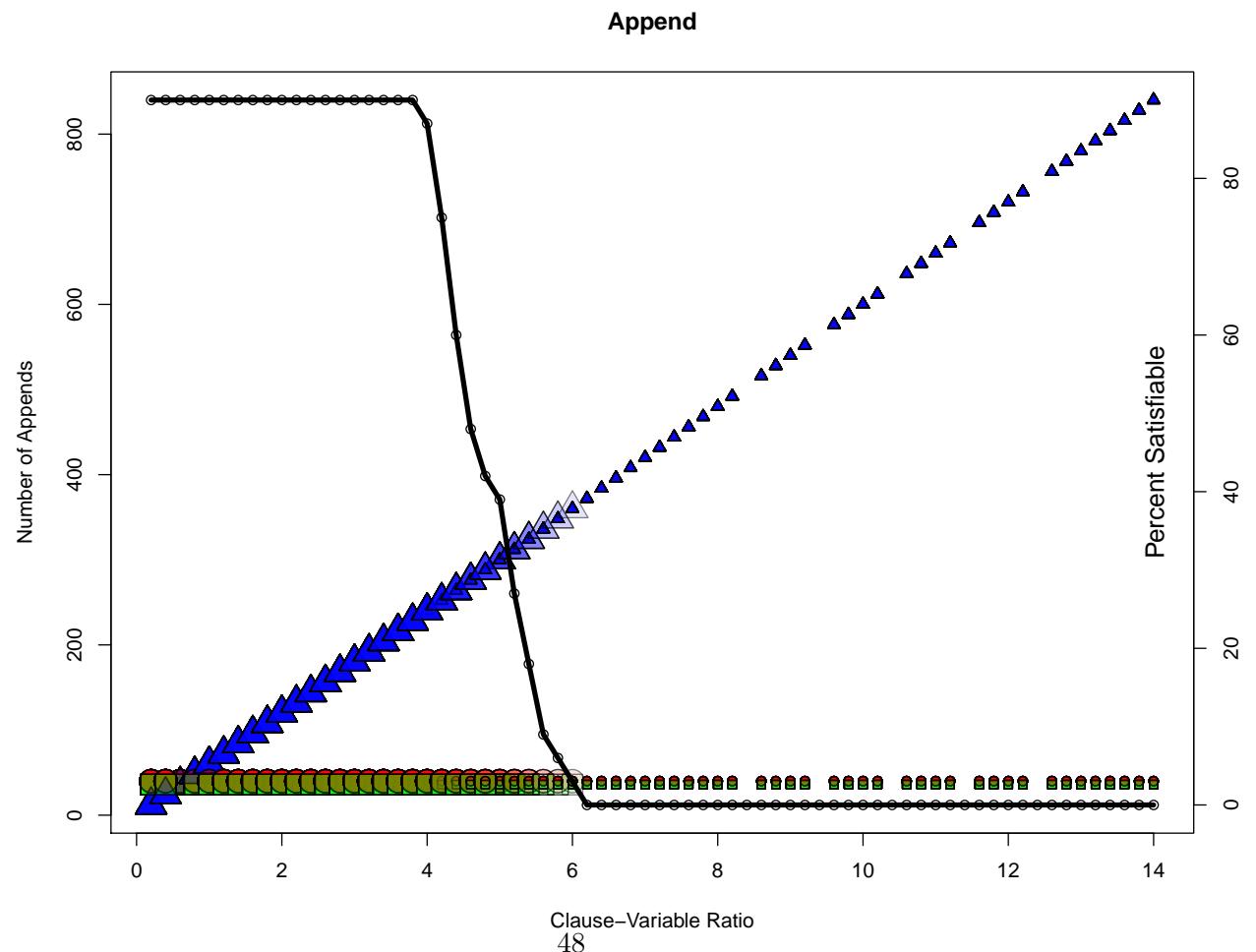


Figure 7.2: Clause to variable ratio α vs. Number of appends, with $n = 20$.

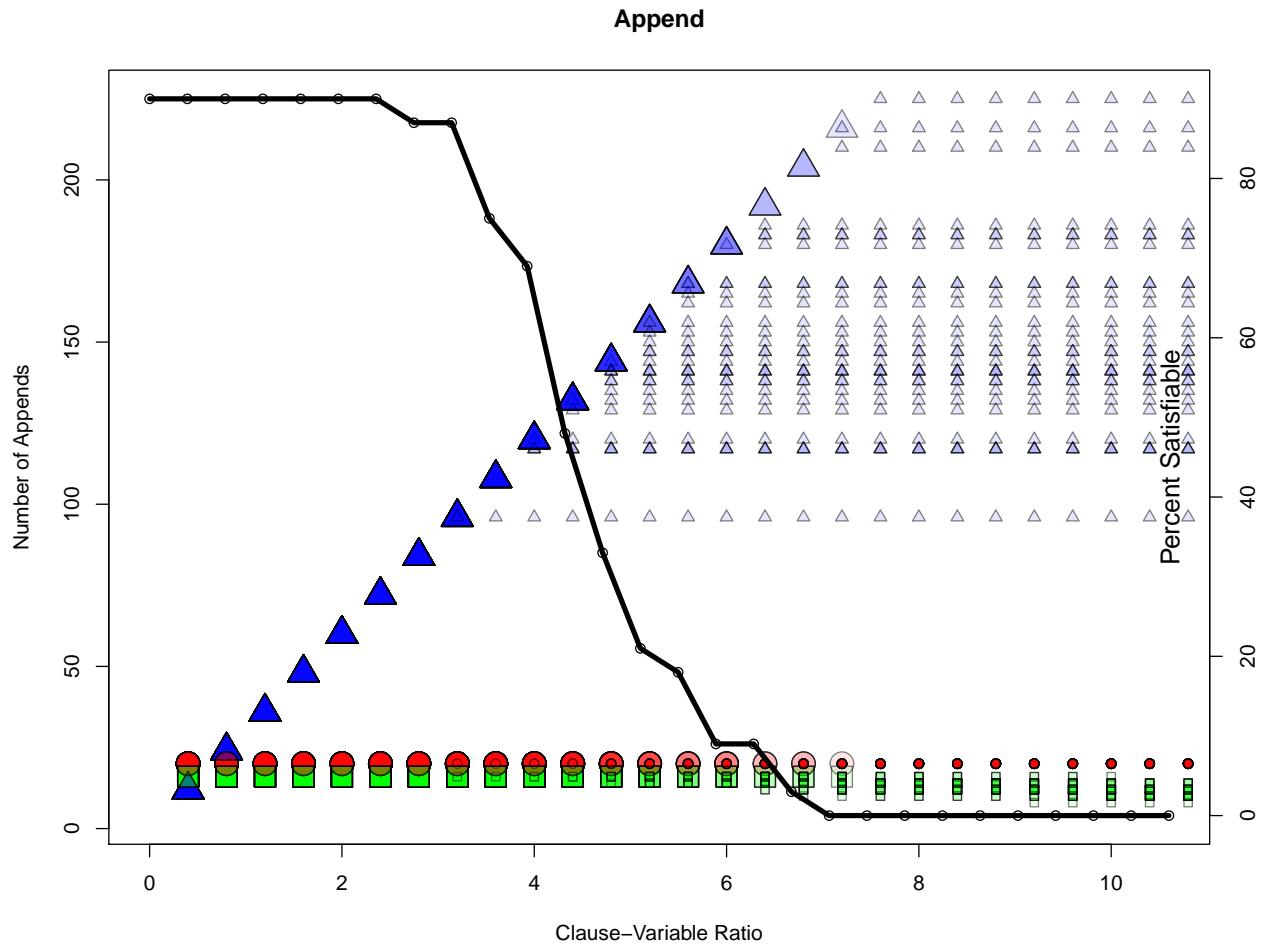


Figure 7.3: Clause to variable ratio α vs. Number of appends, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Extract filters oligonucleotides from a tube. Figure 7.4 shows the number of extracts used in the naive implementations. Figure 7.5 shows early termination on unsatisfiable 3-CNF instances.

Lipton’s algorithm and the Distribution algorithm use $O(k \cdot m)$ extracts. The Distribution algorithm varies a constant amount for maintaining the tubes T_N , T_P , and T_V .

Ogihara and Ray’s algorithm uses $O(m)$ extracts. Lipton’s algorithm shares the same complexity from the experiments since we use 3-CNF instances.

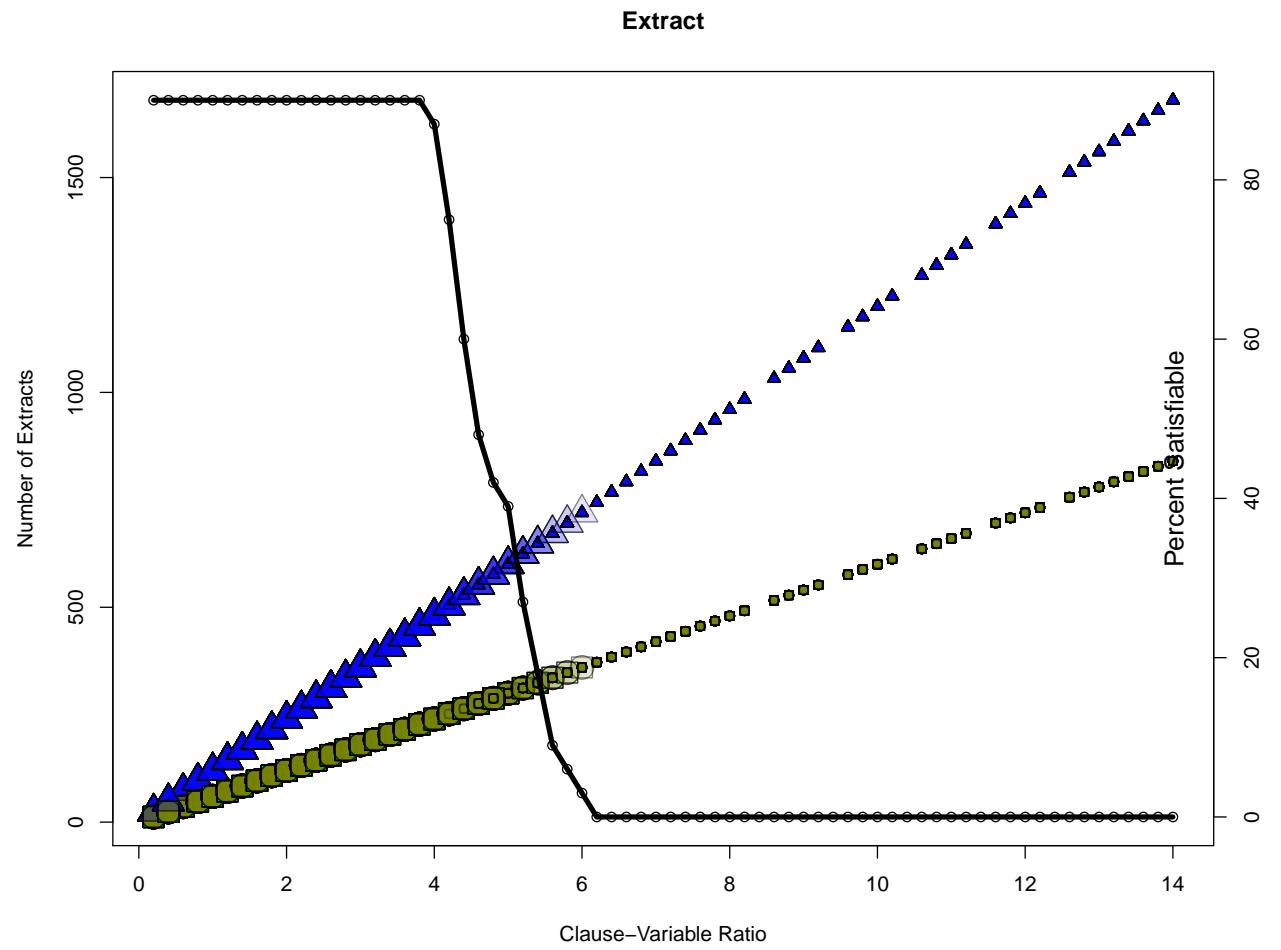


Figure 7.4: Clause to variable ratio α vs. Number of extracts, with $n = 20$.

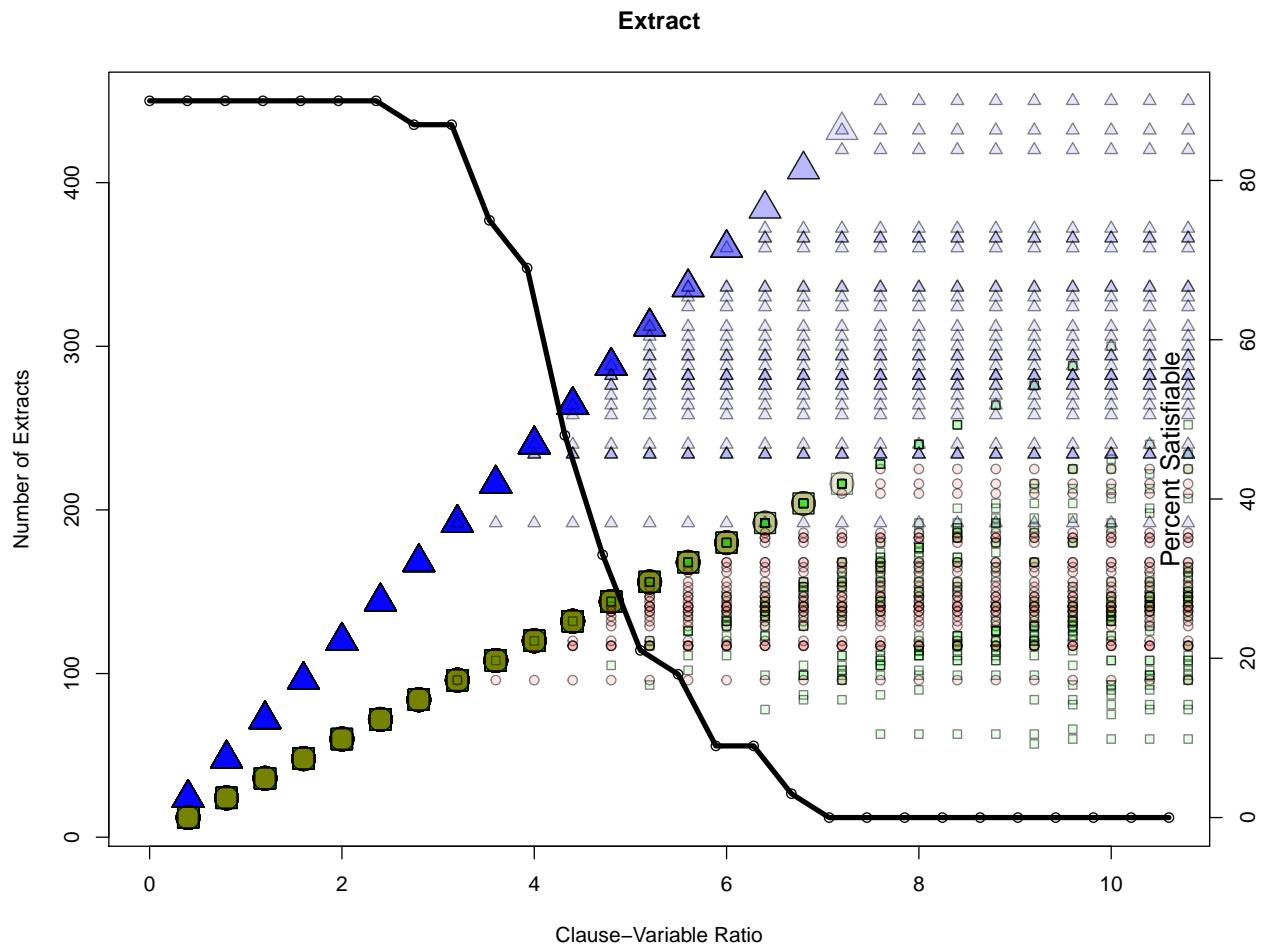


Figure 7.5: Clause to variable ratio α vs. Number of extracts, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Mix combines the contents of n tubes into a storage tube. Figure 7.6 shows the number of mixes used in the naive implementations. Figure 7.7 shows early termination on unsatisfiable 3-CNF instances.

Lipton's algorithm uses $\Theta(m(k + 1) + n)$ mix operations for satisfiable k -CNF instances. This includes preparing a combinatorial space with the COMBINATORIAL GENERATE subroutine.

The Distribution algorithm uses $O(k \cdot m)$ mixes for distributing each clause with k literals into a set of witnesses satisfying each clause. This set of witnesses get mixed into the tube satisfying all distributed clauses.

Ogihara and Ray's algorithm expands on clauses matching in the third ordered literal. The algorithm uses $O(m + n)$ mixes. This includes a maximum of m mixes for each clause, and n mixes for extending witnesses for ϕ .

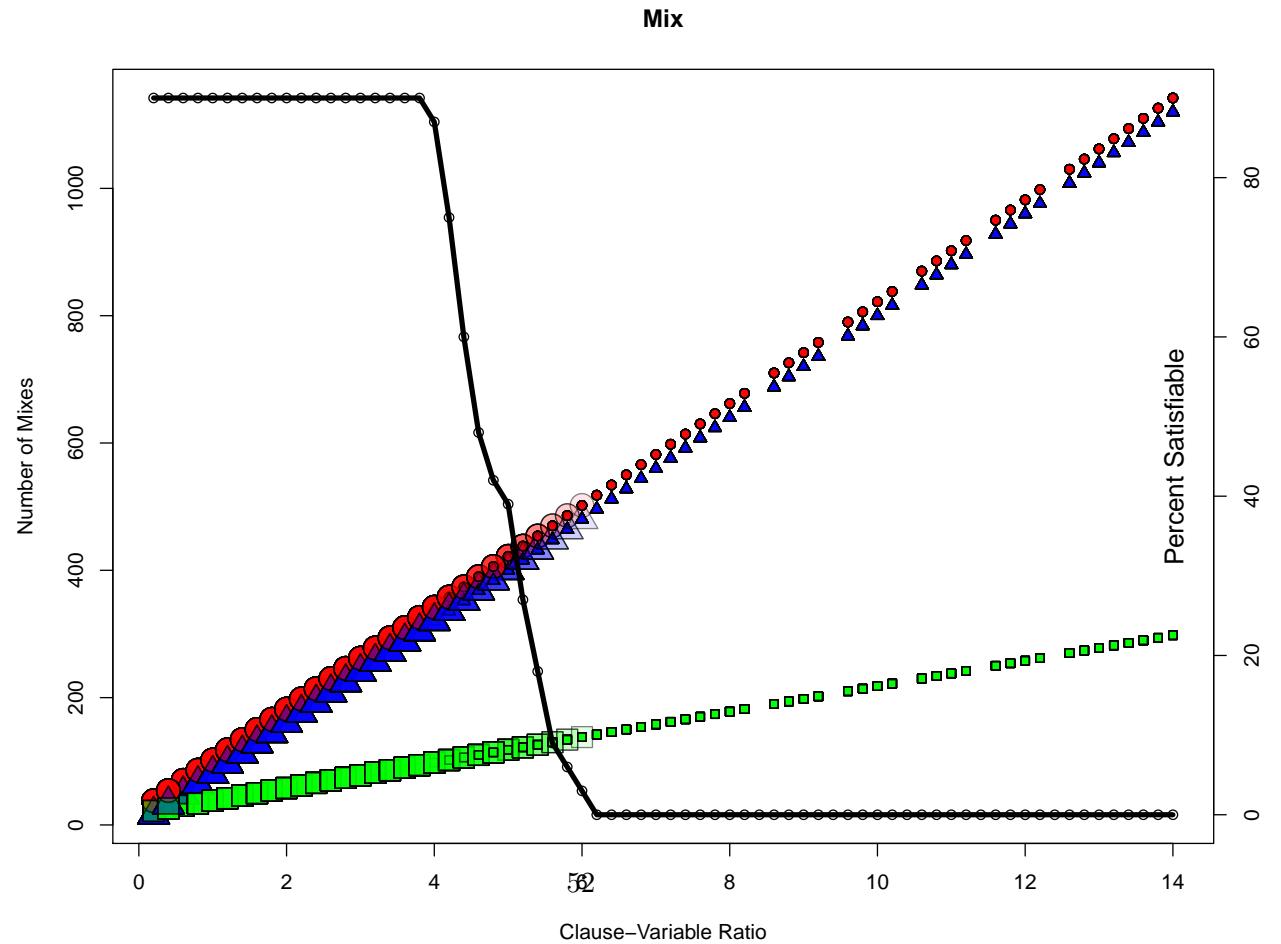


Figure 7.6: Clause to variable ratio α vs. Number of mixes, with $n = 20$.

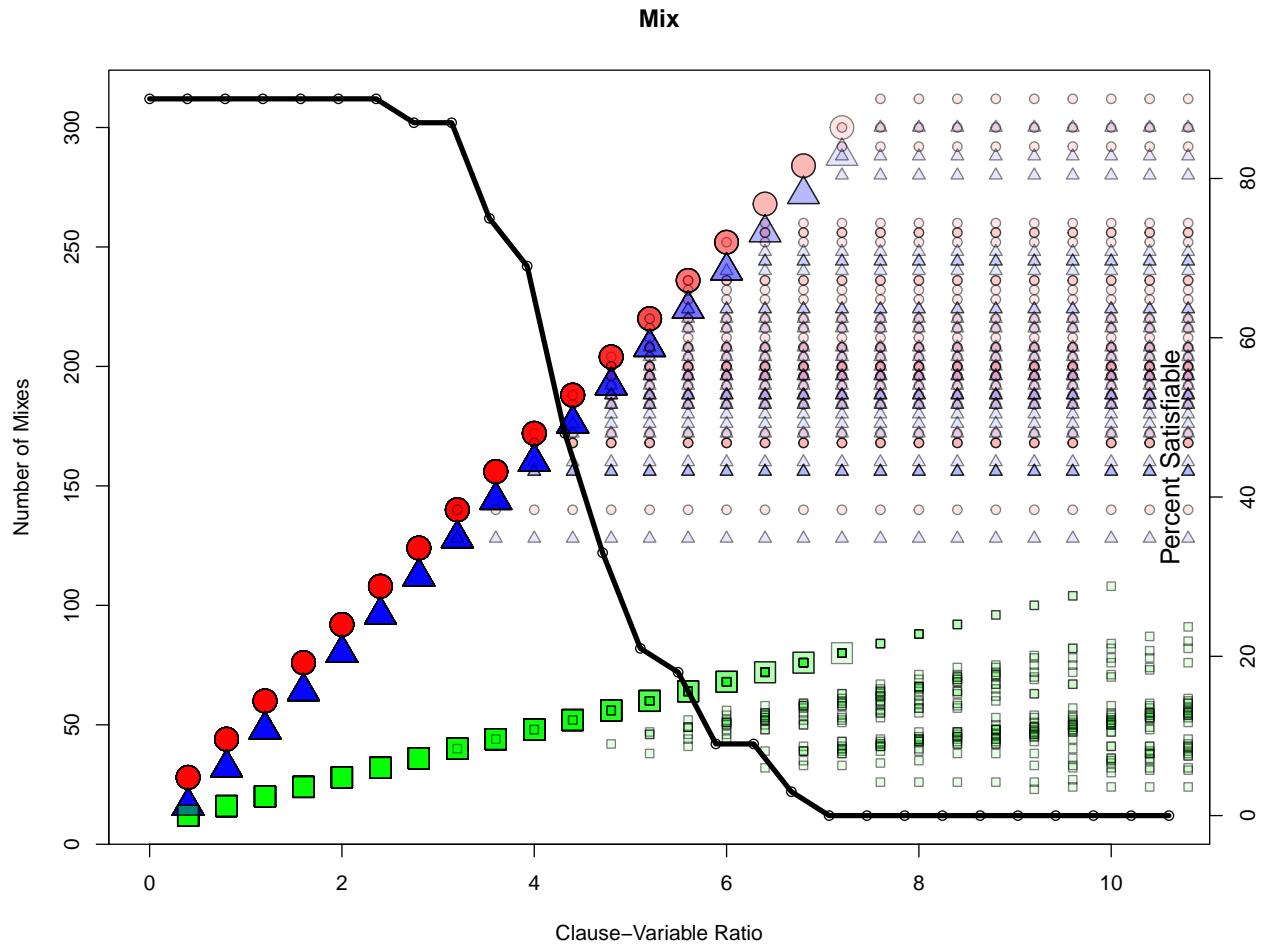


Figure 7.7: Clause to variable ratio α vs. Number of mixes, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Purify ensures a uniform distribution of each independent oligonucleotide in a tube. Figure 7.8 shows the number of purifications used in the naive implementations. Figure 7.9 shows early termination on unsatisfiable 3-CNF instances.

Ogihara and Ray's algorithm requires $O(m + n)$ purifications. The algorithm requires a purification for each of the m clauses, and a purification for each of the n variables.

Lipton's and the Distribution algorithms each use $O(m)$ purifications. Each of these algorithms purify the contents on the iteration of each of the m clauses.

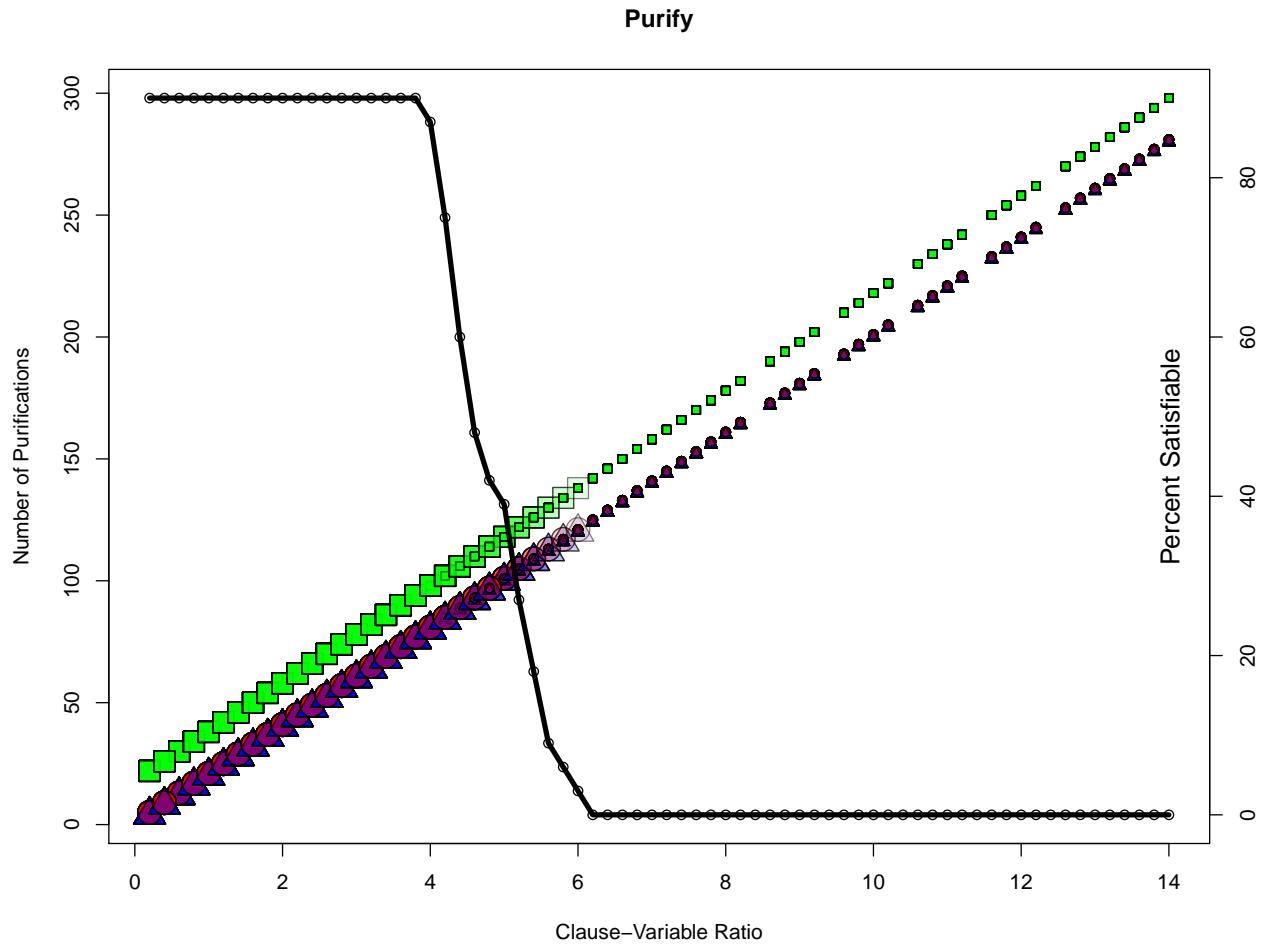


Figure 7.8: Clause to variable ratio α vs. Number of purifications, with $n = 20$.

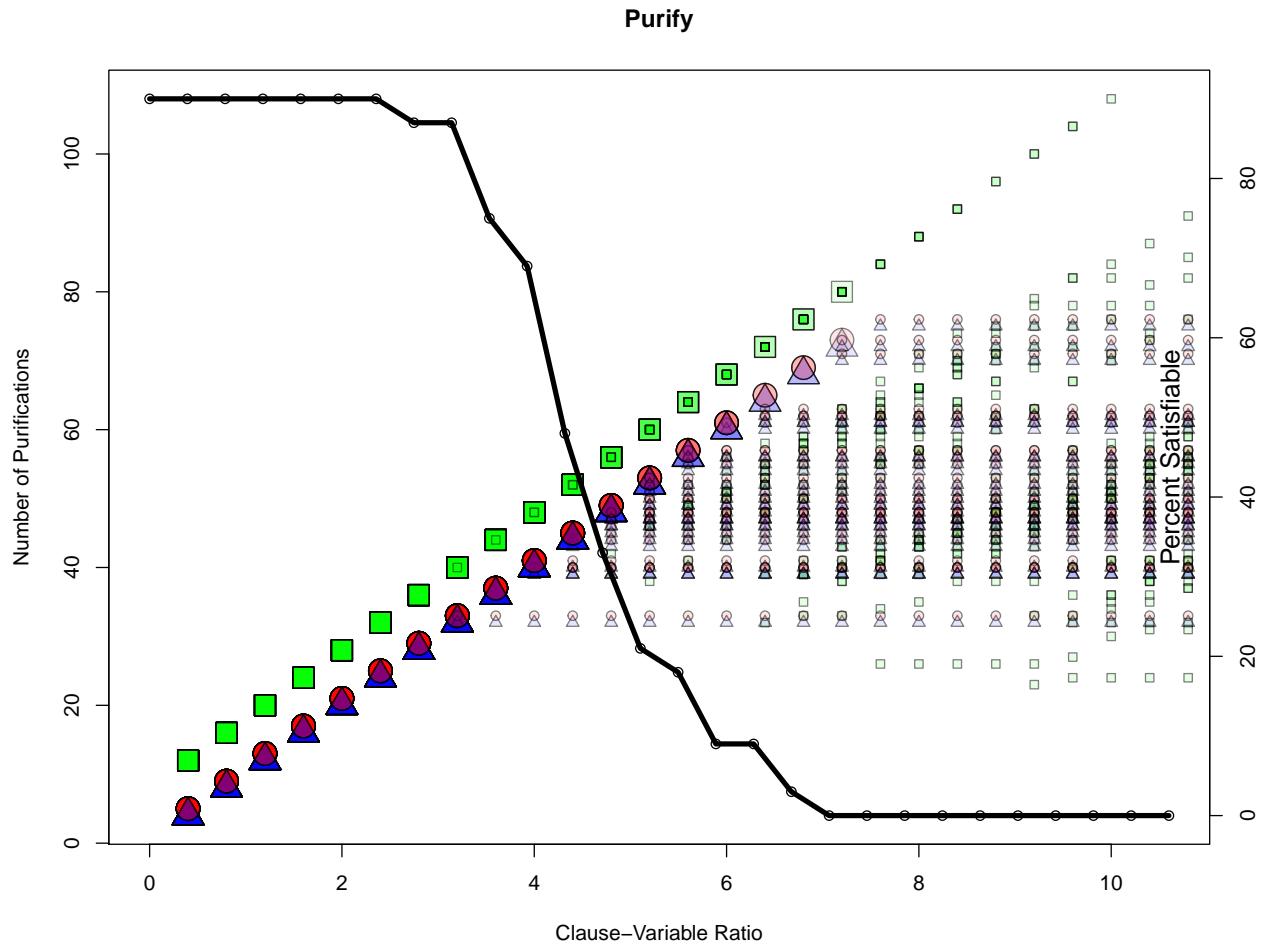


Figure 7.9: Clause to variable ratio α vs. Number of purifications, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Split portions a tube into two exact tubes. Figure 7.10 shows the number of splits used in the naive implementations. Figure 7.11 shows early termination on unsatisfiable 3-CNF instances.

The Distribution algorithm uses $O(k \cdot m)$ splits. The split operation occurs on each of the m clauses of a k -CNF instance.

Lipton's algorithm uses $\Theta(n)$ splits during the creation of a combinatorial space of 2^n witness candidates with COMBINATORIAL GENERATE.

Ogihara and Ray's algorithm uses $O(n)$ splits. The algorithm requires a copy of witness candidates in the tubes (T_P and T_N). The tubes T_P and T_N extend with positive and negative literal assignments.

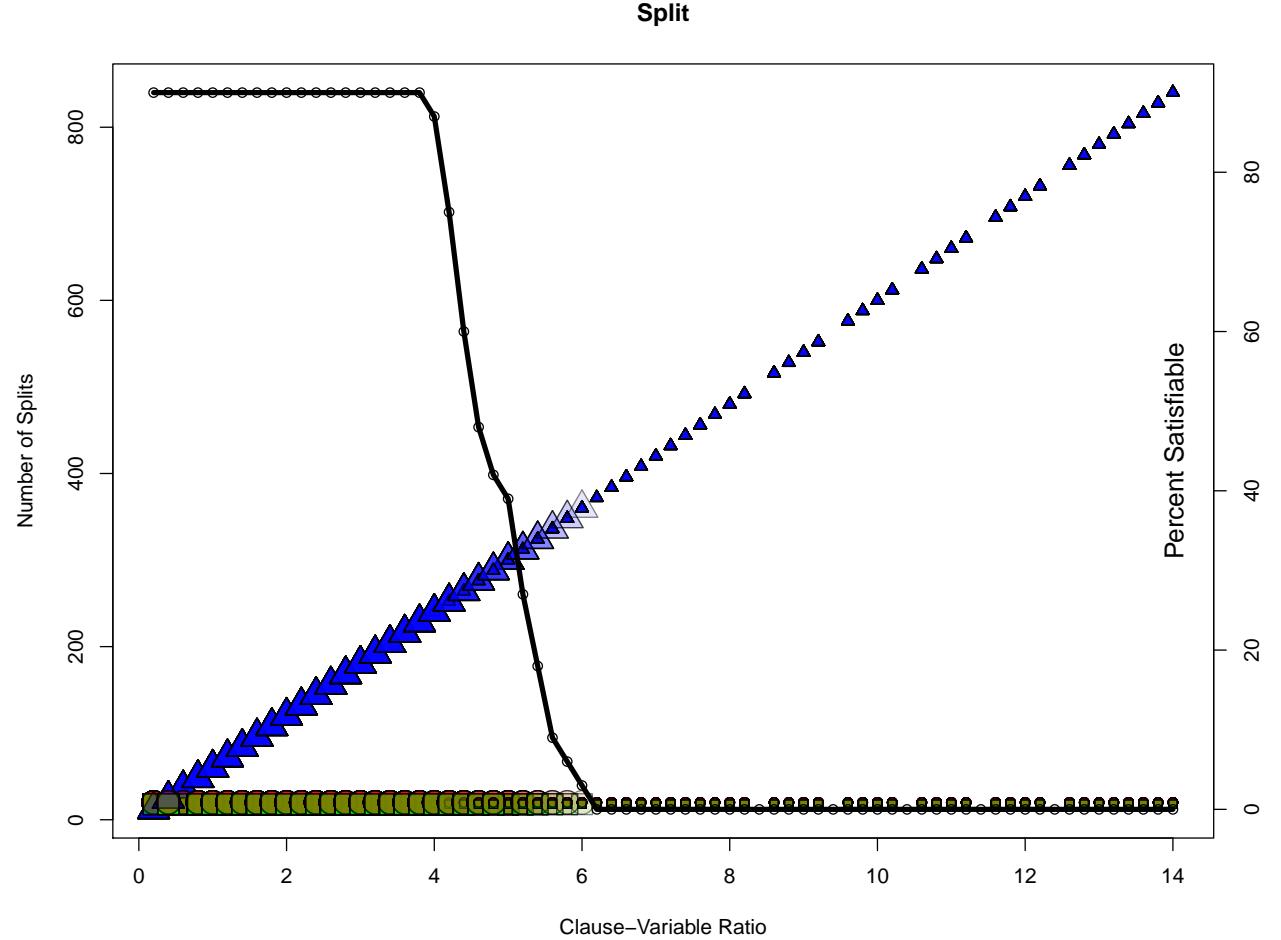


Figure 7.10: Clause to variable ratio α vs. Number of splits, with $n = 20$.

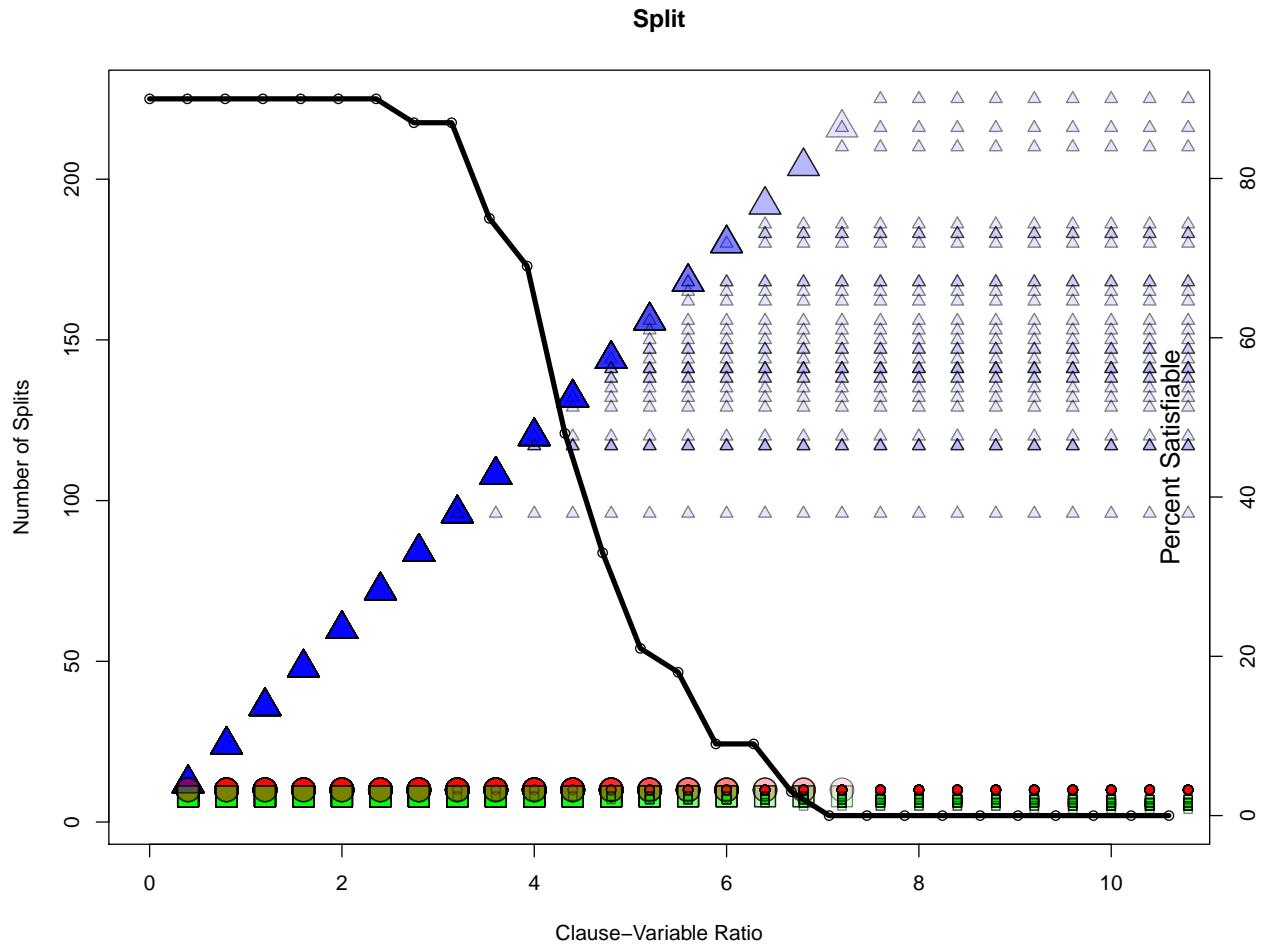


Figure 7.11: Clause to variable ratio α vs. Number of splits, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Time measures algorithm execution time in seconds. Figure 7.12 shows the execution time in seconds used in the naive implementations. Figure 7.13 shows early termination on unsatisfiable 3-CNF instances.

Ogihara and Ray's algorithm requires the least amount of time. The algorithm takes the greatest amount of time for under-constrained instances (causing a large number of witnesses to be generated). Less pruning occurs in over-constrained instances, reducing the execution time of test instances.

Lipton's algorithm executes in exponential time $\alpha \approx [4.0, 6.0]$ (the phase transition region for 3-SAT) taking the longest.

The Distribution algorithm executes in exponential time, and performs better than Lipton's algorithm for low conflict ratios. Instances take the longest from $\alpha \approx [4.0, 6.0]$ (in the 3-SAT phase-transition region).

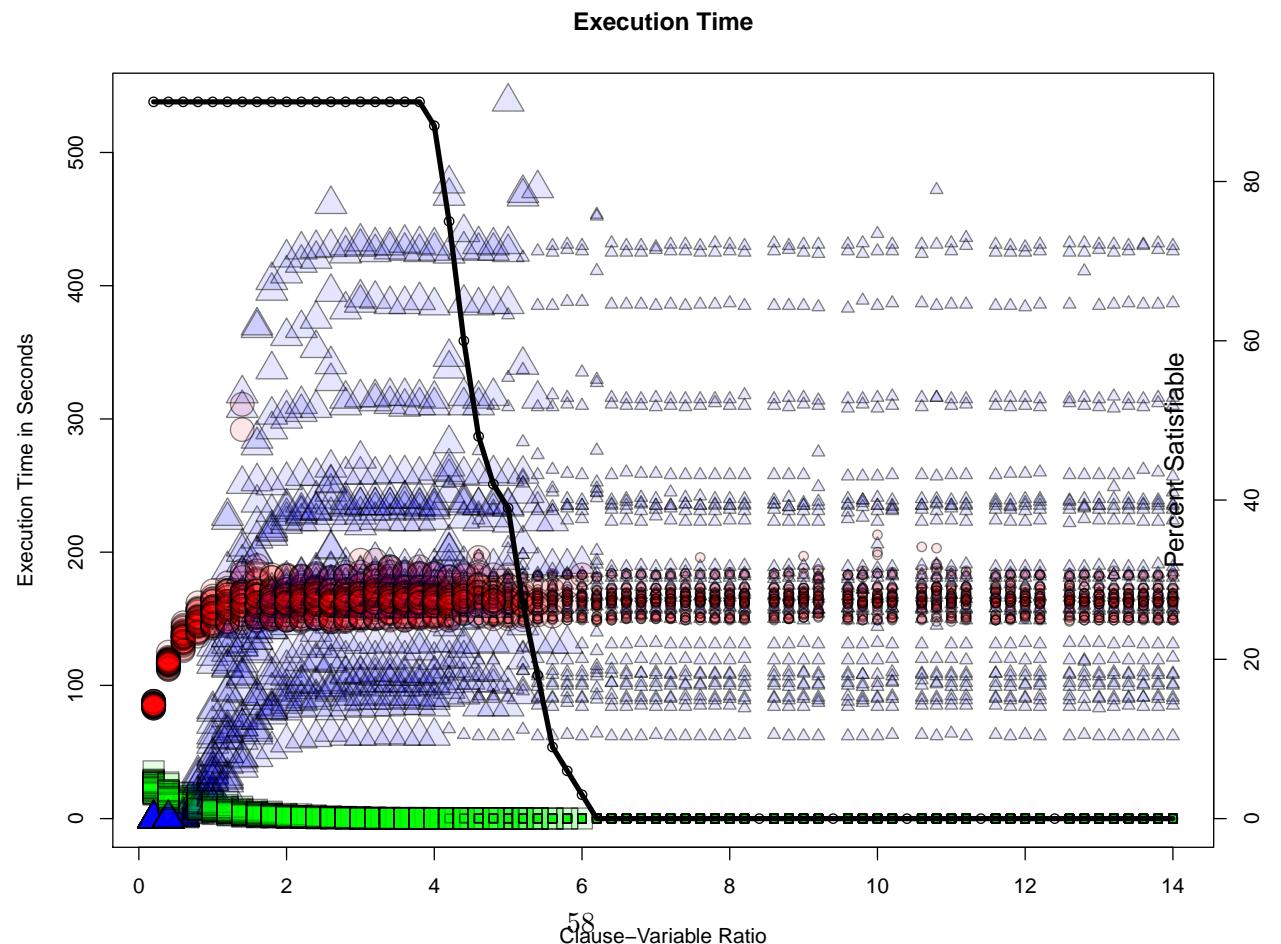


Figure 7.12: Clause to variable ratio α vs. execution time in seconds, with $n = 20$.

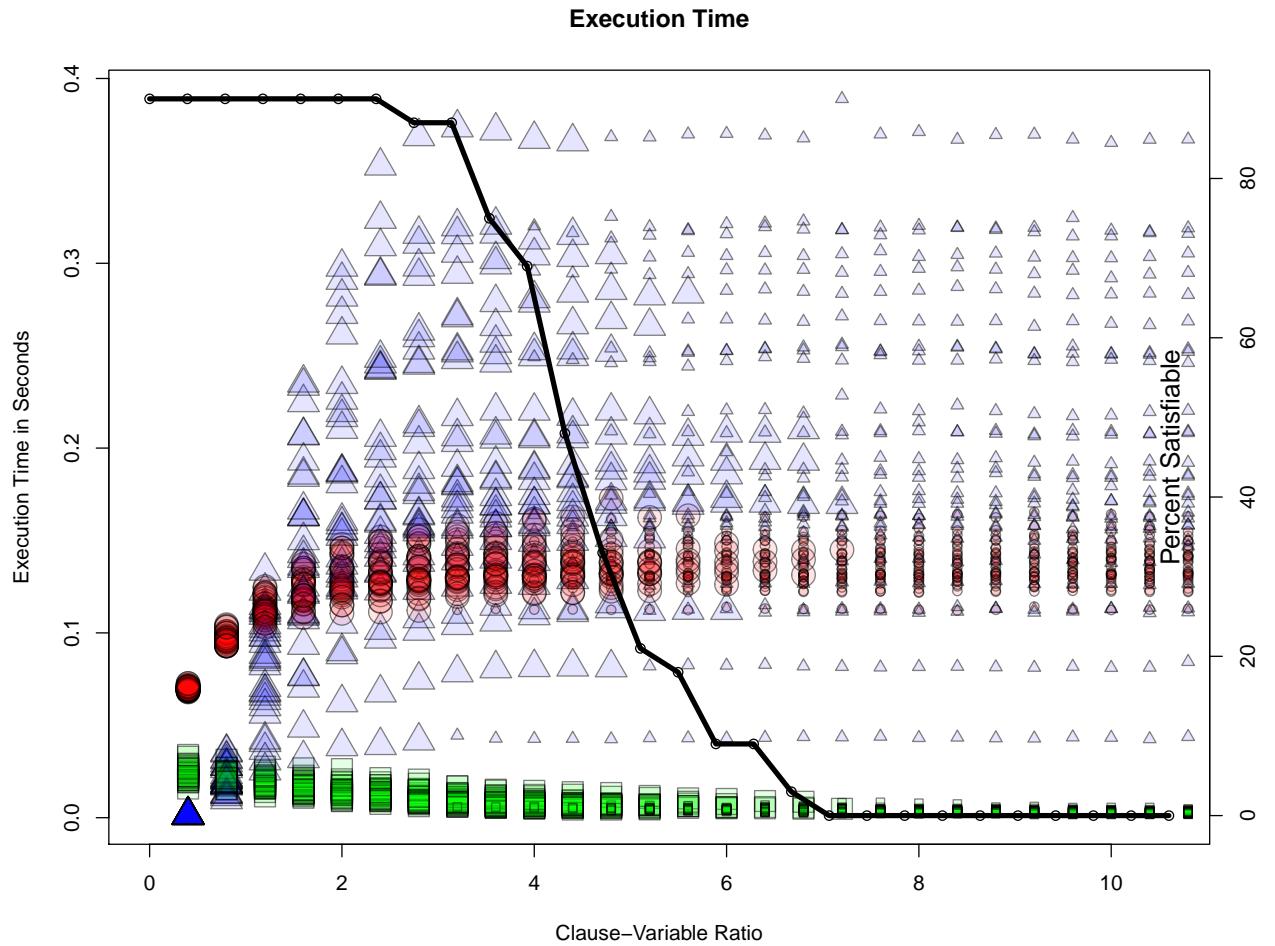


Figure 7.13: Clause to variable ratio α vs. Number of splits, with $n = 10$. Algorithms terminate on detection of unsatisfiable input.

Memory measures witness footprint in Bytes. Figure 7.14 shows the satisfiable instance witness memory. Figure 7.15 shows a detailed view of Figure 7.14 from $\alpha = [3, 6]$.

Lipton's and Ogiwara and Ray's algorithms share the same solution footprint.

The Distribution algorithm contains a larger solution footprint after the trivially satisfiable instances with $\alpha \approx [0.2, 1.0)$. The space contains a set of over-constrained assignments from $\alpha \approx [1.0, 4.0]$.

Each SATISFIABILITY instance has a constrained solution space during the phase-transition region. We scale the axis in Figure 7.14 to observe only satisfiable instances.

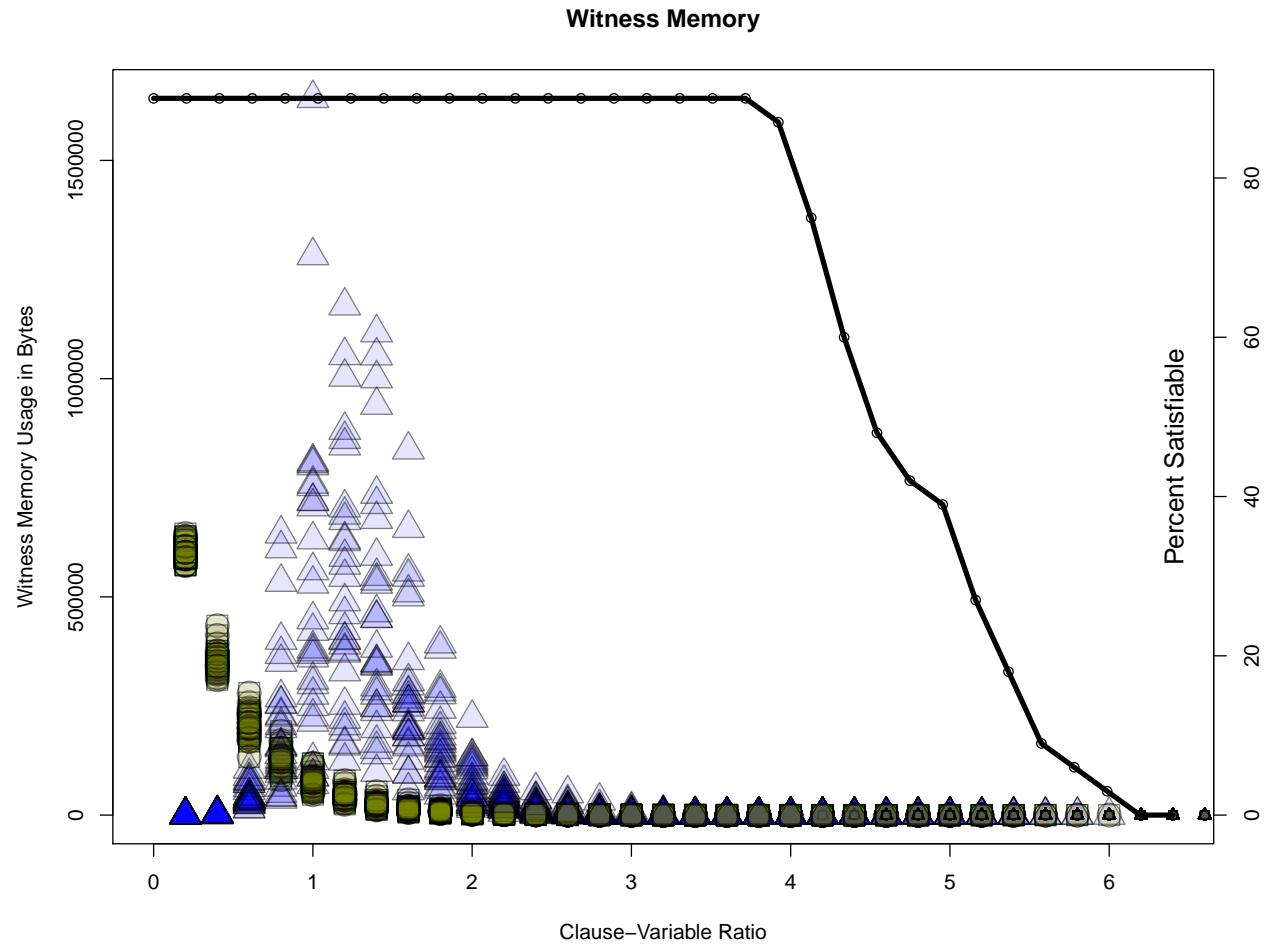


Figure 7.14: Clause to variable ratio α vs. satisfiable solution footprint in Bytes, with $n = 20$.

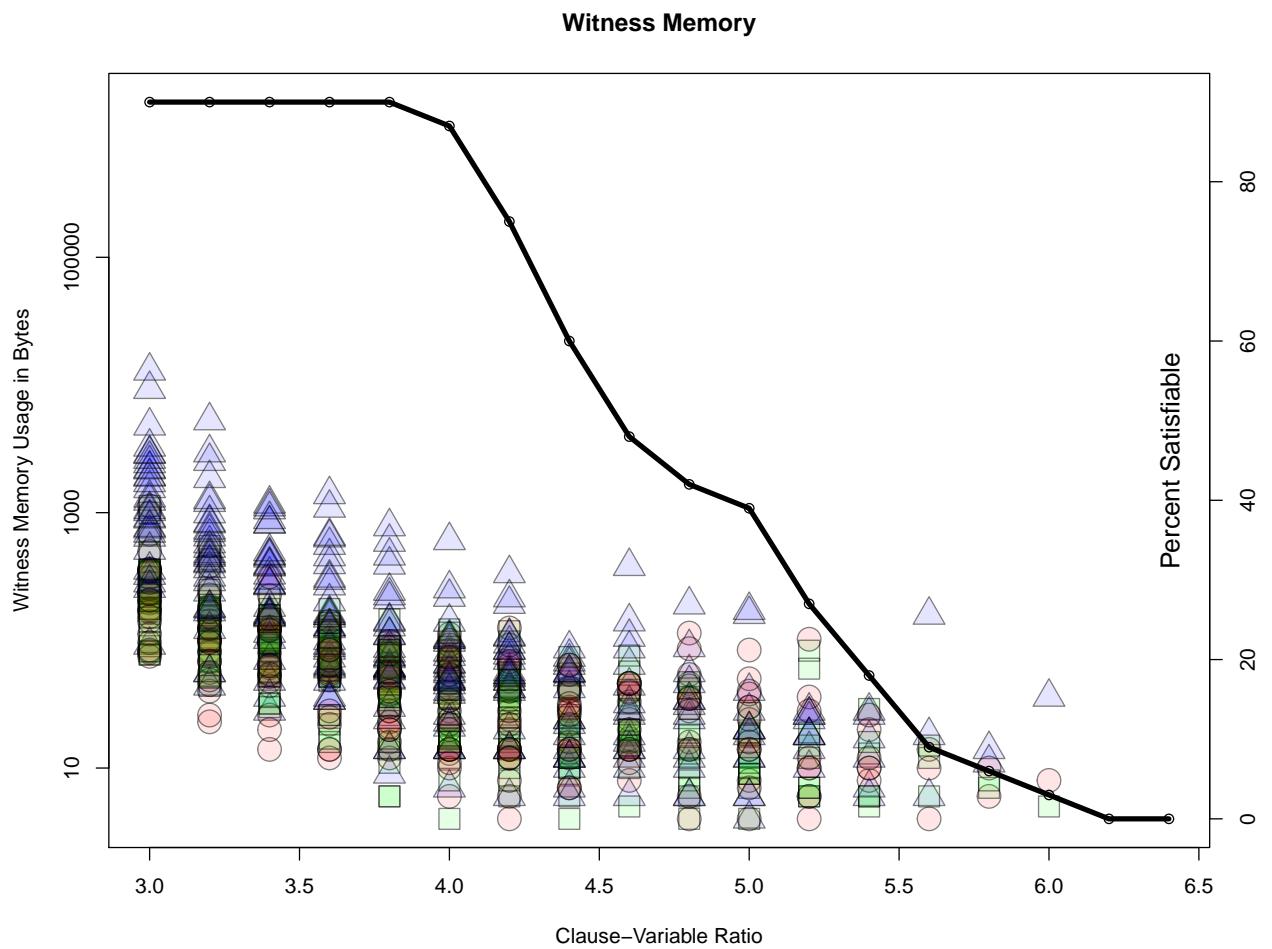


Figure 7.15: Detailed view of Figure 7.14 from $\alpha = [3, 6]$.

7.2 Summary of Molecular Algorithms

Table 7.1 shows a comparison of the molecular operators of the algorithms under test. Once the solution space $T_S = \emptyset$ the algorithms can terminate. This provides the upper bounds for execution of various k -CNF instances.

Table 7.1: Time complexity of molecular operators for each of the molecular algorithms.

Operator	COMBINATORIAL GENERATE	LIPTON	OGIHARA-RAY	DISTRIBUTION
mix	$\Theta(n)$	$O(k \cdot m + m)$	$O(m + n)$	$O(k \cdot m + m)$
extract	—	$O(k \cdot m)$	$O(m)$	$O(k \cdot m)$
split	$\Theta(n)$	—	$O(n)$	$O(k \cdot m)$
purify	$\Theta(1)$	$O(m)$	$O(m + n)$	$O(m)$
append	$\Theta(n)$	—	$O(n)$	$O(k \cdot m)$

Chapter 8

Conclusion

We considered three molecular algorithms for SATISFIABILITY and simulated their execution with a computation framework. Chapter 2 discussed techniques for molecular computation and defined a molecular instruction set (Adleman’s molecular toolbox).

In Chapter 2, we defined SATISFIABILITY along with the standard forms: CNF, k -CNF, and k -SAT. We use random 3-CNF instances for execution of simulated molecular algorithms.

8.1 Contribution

This project introduces the Distribution algorithm for SATISFIABILITY. This algorithm distributes the literals into a growing set of witnesses during the evaluation of a k -CNF instance.

We provide comparisons of molecular operators for each of the molecular algorithms under test. The software framework for this project (Molecular Simulation) was used for simulating and collecting metrics on random 3-CNF instances at fixed $n = 20$ spanning clause-variable ratios $\alpha = m/n = [0.2, 14.0]$ in increments of 0.2.

8.2 Future work

Variations on the Distribution algorithm can decrease the length and quantity of witness candidates. In particular, the Distribution algorithm may select clauses with the maximum number of idempotent literal assignments while regarding all other literals as unassigned. From the remaining clauses, assignments can be made using the standard Distribution algorithm (Chapter 4) or employ a clause selection strategy.

In this project we considered the use of a genetic substrate to store and match witness candidates for SATISFIABILITY. Existing implementation of molecular computers use

laboratory procedures operated by a human or human-like automation. Gene sequencers currently provide an integrated means for reading natural gene expressions.

Practical molecular computation must be integrated into a molecular computing architecture. This architecture must facilitate the storage and transfer of isolated tubes with read/write/extract capabilities. This type of machine provides an environment for observing natural interactions of genes (along with sequencing), and an architecture for molecular computation.

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Appendix A

Source

A.1 Contributed

Download Molecular Simulation:

- <https://github.com/dncarley/MolecularSimulation>

A.2 External

Download David Wilson's *k*-SAT Generator:

- <http://research.microsoft.com/en-us/um/people/dbwilson/ksat/default.htm>

A.3 Dependencies for Molecular Simulation

- GCC, The GNU Compiler Collection (Required)
 - <http://gcc.gnu.org/>
- Perl 5 (Required)
 - <http://www.perl.org/>
- Perl Parallel::ForkManager (Optional for running on multiple cores)
 - <http://search.cpan.org/~dlux/Parallel-ForkManager-0.7.5>
- Doxygen (Optional for generating documentation)
 - <http://www.stack.nl/~dimitri/doxygen/>

Appendix B

Molecular algorithm trace

B.1 Example SATISFIABILITY instance

$$\phi = (x_1 \vee x_2 \vee \neg x_3) \wedge (x_2 \vee x_3 \vee \neg x_4) \wedge (\neg x_1 \vee \neg x_3 \vee x_4)$$

B.2 Lipton's Algorithm

$$T = \text{COMBINATORIAL GENERATE}(4)$$

$$T = \{\text{STTTT}, \text{SFTTT}, \text{STFTT}, \text{SFFT}, \\ \text{STTFT}, \text{SFTFT}, \text{STFFT}, \text{SFFFT}, \\ \text{STTTF}, \text{SFTTF}, \text{STFTF}, \text{SFTTF}, \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF}\}$$

From the tube T , select witnesses for Clause 1:

$$C_1 = (x_1 \vee x_2 \vee \neg x_3)$$

Extract x_1 :

$$T_P = \{\text{STTTT}, \quad , \text{STFTT}, \quad , \\ \text{STTFT}, \quad , \text{STFFT}, \quad , \\ \text{STTTF}, \quad , \text{STFTF}, \quad , \\ \text{STTFF}, \quad , \text{STFFF}, \quad \}$$

Mix T_P into T_C

$$T_C = \{\text{STTTT}, \quad , \text{STFTT}, \quad , \\ \text{STTFT}, \quad , \text{STFFT}, \quad , \\ \text{STTTF}, \quad , \text{STFTF}, \quad , \\ \text{STTFF}, \quad , \text{STFFF}, \quad \}$$

Extract x_2 :

$$T_P = \{\text{STTTT}, \text{SFTTT}, \quad , \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ \text{STTTF}, \text{SFTTF}, \quad , \quad , \\ \text{STTFF}, \text{SFTFF}, \quad , \quad \}$$

Mix T_P into T_C

$$T_C = \{\text{STTTT}, \text{SFTTT}, \text{STFTT}, \quad , \\ \text{STTFT}, \text{SFTFT}, \text{STFFT}, \quad , \\ \text{STTTF}, \text{SFTTF}, \text{STFTF}, \quad , \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \quad \}$$

Extract $\neg x_3$:

$$T_N = \{ \quad , \quad , \quad , \quad , \quad , \\ \text{STTFT}, \text{SFTFT}, \text{STFFT}, \text{SFFT}, \text{SFFT}, \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF} \}$$

Mix contents:

$$T_C = \{\text{STTTT, SFTTT, STFTT, }, \\ \text{STTFT, SFTFT, STFFT, SFFT}, \\ \text{STTTF, SFTTF, STFTF, }, \\ \text{STTFF, SFTFF, STFFF, SFFFF}\}$$

Next select Clause 2:

$$C_2 = (x_2 \vee x_3 \vee \neg x_4)$$

$$T = \{\text{STTTT, SFTTT, STFTT, }, \\ \text{STTFT, SFTFT, STFFT, SFFT}, \\ \text{STTTF, SFTTF, STFTF, }, \\ \text{STTFF, SFTFF, STFFF, SFFFF}\}$$

Extract x_2 :

$$T_P = \{\text{STTTT, SFTTT, }, , , \\ \text{STTFT, SFTFT, }, , , \\ \text{STTTF, SFTTF, }, , , \\ \text{STTFF, SFTFF, }, , \}$$

Mix the contents of T_P into T_C :

$$T_C = \{\text{STTTT, SFTTT, }, , , \\ \text{STTFT, SFTFT, }, , , \\ \text{STTTF, SFTTF, }, , , \\ \text{STTFF, SFTFF, }, , \}$$

Extract x_3 :

$$T_P = \{\text{STTTT, SFTTT, STFTT, }, , , , , , \\ \text{STTTF, SFTTF, STFTF, }, , , , , , \}$$

Mix T_P into T_C

$$T_C = \{\text{STTTT, SFTTT, STFTT, }, , , , , , \\ \text{STTFT, SFTFT, }, , , , , , \\ \text{STTTF, SFTTF, STFTF, }, , , , , , \\ \text{STTFF, SFTFF, }, , , \}$$

Extract $\neg x_4$:

$$T_N = \{ , , , , , , , , , , , , \\ \text{STTTF, SFTTF, STFTF, }, , , , , , \\ \text{STTFF, SFTFF, STFFF, SFFFF} \}$$

Mix T_N into T_C

$$T_C = \{\text{STTTT, SFTTT, STFTT, }, , , , , , \\ \text{STTFT, SFTFT, }, , , , , , \\ \text{STTTF, SFTTF, STFTF, }, , , , , , \\ \text{STTFF, SFTFF, STFFF, SFFFF} \}$$

Finally, select Clause 3:

$$C_3 = (\neg x_1 \vee \neg x_3 \vee x_4)$$

$$T = \{ \text{STTTT}, \text{SFTTT}, \text{STFTT}, \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ \text{STTTF}, \text{SFTTF}, \text{STFTF}, \quad , \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF} \}$$

From the tube T , extract $\neg x_1$:

$$T_N = \{ \quad , \text{SFTTT}, \quad , \quad , \\ \quad , \text{SFTFT}, \quad , \quad , \\ \quad , \text{SFTTF}, \quad , \quad , \\ \quad , \text{SFTFF}, \quad , \text{SFFFF} \}$$

Mix the contents of T_N into T_C

$$T_C = \{ \quad , \text{SFTTT}, \quad , \quad , \\ \quad , \text{SFTFT}, \quad , \quad , \\ \quad , \text{SFTTF}, \quad , \quad , \\ \quad , \text{SFTFF}, \quad , \text{SFFFF} \}$$

Extract $\neg x_3$:

$$T_N = \{ \quad , \quad , \quad , \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ \quad , \quad , \quad , \quad , \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF} \}$$

Mix the contents of T_N into T_C :

$$T_C = \{ \quad , \text{SFTTT}, \quad , \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ \quad , \text{SFTTF}, \quad , \quad , \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF} \}$$

$$T_P = \{\text{STTTT}, \text{SFTTT}, \text{STFTT}, \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ , \quad , \quad , \quad , \quad , \\ , \quad , \quad , \quad , \quad \}$$

Mix the contents of T_P into T_C :

$$T_C = \{\text{STTTT}, \text{SFTTT}, \text{STFTT}, \quad , \\ \text{STTFT}, \text{SFTFT}, \quad , \quad , \\ , \text{SFTTF}, \quad , \quad , \\ \text{STTFF}, \text{SFTFF}, \text{STFFF}, \text{SFFFF}\}$$

B.3 Ogihara and Ray's Algorithm

Initialize the tube T with initial vector assignments for literals x_1 and x_2

$$T = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$$

Iterate variable x_3 :

$$T_P = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$$

$$T_N = \{\text{STT}, \text{STF}, \text{SFT}, \text{SFF}\}$$

Evaluate clause C_1 :

$$C_1 = (x_1 \vee x_2 \vee \neg x_3)$$

The literal $\neg x_3$ matches v_3 from C_1 .

$$\begin{aligned} T_{P1} &= \{\text{STT}, \text{STF}\} \\ T_{N1} &= \{\text{SFT}, \text{SFF}\} \\ T_{P2} &= \{\text{SFT}\} \\ T_P &= \{\text{STT}, \text{STF}, \text{SFT}\} \end{aligned}$$

Evaluate clause C_2 :

$$C_2 = (x_2 \vee x_3 \vee \neg x_4)$$

The literals x_3 or $\neg x_3$ does not match v_3 from C_2 .

Evaluate clause C_3 :

$$C_3 = (\neg x_1 \vee \neg x_3 \vee x_4)$$

The literals x_3 or $\neg x_3$ does not match v_3 from C_3 .

Append assignments for x_3 and $\neg x_3$.

$$\begin{aligned} T_P &= \{\text{STTT}, \text{STFT}, \text{SFTT}\} \\ T_N &= \{\text{STTF}, \text{STFF}, \text{SFTF}, \text{SFFF}\} \end{aligned}$$

Mix T_P and T_N into the tube T .

$$T = \{\text{STTT, STFT, SFTT,}\\ \text{STTF, STFF, SFTF, SFFF}\}$$

Iterate variable x_4 :

$$T_P = \{\text{STTT, STFT, SFTT,}\\ \text{STTF, STFF, SFTF, SFFF}\}$$

$$T_N = \{\text{STTT, STFT, SFTT,}\\ \text{STTF, STFF, SFTF, SFFF}\}$$

Evaluate clause C_1 :

$$C_1 = (x_1 \vee x_2 \vee \neg x_3)$$

The literals x_4 or $\neg x_4$ does not match v_3 in C_1 .

Evaluate clause C_2 :

$$C_2 = (x_2 \vee x_3 \vee \neg x_4)$$

The literal $\neg x_4$ matches v_3 in C_2 .

$$\begin{aligned} T_{P1} &= \{\text{STTT, SFTT, STTF, SFTF}\} \\ T_{N1} &= \{\text{STFT, STFF, SFFF}\} \\ T_{P2} &= \{\text{STFT}\} \\ T_P &= \{\text{STTT, SFTT, STTF, SFTF, STFT}\} \end{aligned}$$

Evaluate clause C_3 :

$$C_3 = (\neg x_1 \vee \neg x_3 \vee x_4)$$

The literal x_4 matches v_3 in C_3 .

$$\begin{aligned} T_{P1} &= \{\text{SFTT, SFTF, SFFF}\} \\ T_{N1} &= \{\text{STTT, STFT, STTF, STFF}\} \\ T_{P2} &= \{\text{STTF, STFF}\} \\ T_N &= \{\text{SFTT, SFTF, SFFF, STTF, STFF}\} \end{aligned}$$

Append assignments for x_4 and $\neg x_4$.

$$T_P = \{\text{STTTT}, \text{SFTTT}, \text{STTFT}, \text{SFTFT}, \text{STFTT}\}$$
$$T_N = \{\text{SFTTF}, \text{SFTFF}, \text{SFFFF}, \text{STTFF}, \text{STFFF}\}$$

Mix T_P and T_N into the tube T , providing the witnesses for ϕ

$$T = \{\text{STTTT}, \text{SFTTT}, \text{STTFT}, \text{SFTFT}, \text{STFTT},$$
$$\text{SFTTF}, \text{SFTFF}, \text{SFFFF}, \text{STTFF}, \text{STFFF}\}$$

B.4 Distribution Algorithm

Initiate the tube T_S with a start medium S .

$$T_S = \{S\}$$

Evaluate clause $C_1 = (x_1 \vee x_2 \vee \neg x_3)$:

$$T_C = \{ \}$$

Select the literal x_1 from C_1 . Split the contents of T_S :

$$T_V = \{S\}$$

$$T_S = \{S\}$$

Extract literals from T_V that contain negative and positive assignments for x_1 .

$$T_N = \{ \}$$

$$T_P = \{ \}$$

$$T_V = \{S\}$$

Append the satisfying literal x_1 to the required variable tube T_V .

$$T_V = \{S+1\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+1\}$$

Select the literal x_2 from C_1 . Split the contents of T_S :

$$T_V = \{S\}$$

$$T_S = \{S\}$$

Extract literals from T_V that contain negative and positive assignments for x_2 .

$$T_N = \{ \}$$

$$T_P = \{ \}$$

$$T_V = \{S\}$$

Append the satisfying literal x_2 to the required variable tube T_V .

$$T_V = \{S+2\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+1, S+2\}$$

Select the literal $\neg x_3$ from C_1 . Split the contents of T_S :

$$T_V = \{S\}$$

$$T_S = \{S\}$$

Extract literals from T_V that contain negative and positive assignments for $\neg x_3$.

$$T_N = \{ \}$$

$$T_P = \{ \}$$

$$T_V = \{S\}$$

Append the satisfying literal $\neg x_3$ to the required variable tube T_V .

$$T_V = \{S-3\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+1, S+2, S-3\}$$

Complete the iteration of the first clause by storing the witnesses in T_C as witnesses to T_S for the evaluated instance ϕ .

$$T_S = \{S+1, S+2, S-3\}$$

Evaluate clause $C_2 = (x_2 \vee x_3 \vee \neg x_4)$:

$$T_C = \{ \}$$

Select the literal x_2 from C_2 . Split the contents of T_S :

$$T_V = \{S+1, S+2, S-3\}$$

$$T_S = \{S+1, S+2, S-3\}$$

Extract literals from T_V that contain negative and positive assignments for x_2 .

$$\begin{aligned} T_N &= \{ \} \\ T_P &= \{S+2\} \\ T_V &= \{S+1, S-3\} \end{aligned}$$

Append the satisfying literal x_2 to the required variable tube T_V .

$$T_V = \{S+1+2, S-3+2\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2, S+1+2, S-3+2\}$$

Select the literal x_3 from C_2 . Split the contents of T_S :

$$T_V = \{S+1, S+2, S-3\}$$

$$T_S = \{S+1, S+2, S-3\}$$

Extract literals from T_V that contain negative and positive assignments for x_3 .

$$T_N = \{S-3\}$$

$$T_P = \{\}$$

$$T_V = \{S+1, S+2\}$$

Append the satisfying literal x_3 to the required variable tube T_V .

$$T_V = \{S+1+3, S+2+3\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2, S+1+2, S-3+2, S+1+3, S+2+3\}$$

Select the literal $\neg x_4$ from C_2 . Split the contents of T_S :

$$T_V = \{S+1, S+2, S-3\}$$

$$T_S = \{S+1, S+2, S-3\}$$

Extract literals from T_V that contain negative and positive assignments for $\neg x_4$.

$$T_N = \{\}$$

$$T_P = \{\}$$

$$T_V = \{S+1, S+2, S-3\}$$

Append the satisfying literal $\neg x_4$ to the required variable tube T_V .

$$T_V = \{S+1-4, S+2-4, S-3-4\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2, S+1+2, S-3+2, S+1+3, S+2+3, S+1-4, S+2-4, S-3-4\}$$

Complete the iteration of the second clause by storing the witnesses in T_C as witnesses to T_S for the evaluated instance ϕ .

$$T_S = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

Evaluate clause $C_3 = (\neg x_1 \vee \neg x_3 \vee x_4)$:

$$T_C = \{ \}$$

Select the literal $\neg x_1$ from C_3 . Split the contents of T_S :

$$T_V = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

$$T_S = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

Extract literals from T_V that contain negative and positive assignments for $\neg x_1$.

$$T_N = \{S+1+2, S+1+3, S+1-4\} \\ T_P = \{ \} \\ T_V = \{S+2, S-3+2, S+2+3, S+2-4, S-3-4\}$$

Append the satisfying literal $\neg x_1$ to the required variable tube T_V .

$$T_V = \{S+2-1, S-3+2-1, S+2+3-1, S+2-4-1, S-3-4-1\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2-1, S-3+2-1, S+2+3-1, S+2-4-1, S-3-4-1\}$$

Select the literal $\neg x_3$ from C_3 . Split the contents of T_S :

$$T_V = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

$$T_S = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

Extract literals from T_V that contain negative and positive assignments for $\neg x_3$.

$$T_N = \{S+1+3, S+2+3\} \\ T_P = \{S-3+2, S-3-4\} \\ T_V = \{S+2, S+1+2, S+1-4, S+2-4\}$$

Append the satisfying literal $\neg x_3$ to the required variable tube T_V .

$$T_V = \{S+2-3, S+1+2-3, S+1-4-3, S+2-4-3\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2-1, S-3+2-1, S+2+3-1, S+2-4-1, S-3-4-1, \\ S-3+2, S-3-4, S+2-3, S+1+2-3, S+1-4-3, S+2-4-3\}$$

Select the literal x_4 from C_3 . Split the contents of T_S :

$$T_V = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

$$T_S = \{S+2, S+1+2, S-3+2, \\ S+1+3, S+2+3, \\ S+1-4, S+2-4, S-3-4\}$$

Extract literals from T_V that contain negative and positive assignments for x_4 .

$$T_N = \{S+1-4, S+2-4, S-3-4\}$$

$$T_P = \{\}$$

$$T_V = \{S+2, S+1+2, S-3+2, S+1+3, S+2+3\}$$

Append the satisfying literal x_4 to the required variable tube T_V .

$$T_V = \{S+2+4, S+1+2+4, S-3+2+4, S+1+3+4, S+2+3+4\}$$

Mix the contents of T_C , T_P , and T_V into the clause witness tube T_C .

$$T_C = \{S+2-1, S-3+2-1, S+2+3-1, S+2-4-1, S-3-4-1, \\ S-3+2, S-3-4, S+2-3, S+1+2-3, S+1-4-3, S+2-4-3, \\ S+2+4, S+1+2+4, S-3+2+4, S+1+3+4, S+2+3+4\}$$

Complete the iteration of the last clause by storing the witnesses in T_C as witnesses to T_S . The tube T_S contains witnesses for ϕ .

$$T_S = \{S+2-1, S-3+2-1, S+2+3-1, S+2-4-1, S-3-4-1, \\ S-3+2, S-3-4, S+2-3, S+1+2-3, S+1-4-3, S+2-4-3, \\ S+2+4, S+1+2+4, S-3+2+4, S+1+3+4, S+2+3+4\}$$