Reducing Power Consumption in Huffman Coding

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Keywords:

1. Introduction

In the recent years, application of battery-powered portable devices, e.g., laptop computers, personal digital assistant (PDA), and mobile phones has increased significantly. The reliability and performance of such devices are primarily dependent on their power consumption, i.e., effective battery life. Bigger battery size could help to improve power efficiency, however, in a portable device the size of the battery is restricted by the size and weight of the device itself. The cost of providing power to both portable and non-portable devices has resulted in significant interest in power reduction. CMOS technologies were developed in order to reduce the power consumption both in data processing and transmission. In [1], an approach was proposed to minimise power consumption in CMOS circuits. The method considers optimising the technology used to implement the digital circuits, circuits style and topology, the architecture for implementing the circuits and the algorithms that are being implemented in the devices.

As data transmission is a common operation in both portable and non-portable devices, therefore, reducing power dissipation in transmission systems is important. Authors in [2, 3] have investigated different criteria for low power data transmission. Power consumption for data transmission in IEEE 802.11 multi-hop networks and 3G mobile wireless networks were analysed in [4] and [5] respectively. In order to increase transmission speed and reduce power dissipation, parallel data transmission methods are widely used. However, parallel transmission is limited to short distance communications, e.g. locally connected devices, internal buses. Ruling out the possible availability of parallel transmission links over long distance, we are left with its serial alternative only. If we attempt to transfer big files, e.g. DNA sequences, over a serial transmission link then it would take a significant amount of

time and obviously a significant amount power would be dissipated in the process. However, data compression techniques are widely used to reduce size of data before transmitting over serial medium to reduce transmission time and cost.

Huffman code [6] is an entropy encoding algorithm widely used for lossless data compression. For any given set of symbols and associated occurrence probabilities, Huffman algorithm generates a binary tree (Huffman tree) to encode a message with an optimal number bits. The codeword for a distinct symbol is generated by traversing the tree from the root to the leaf node representing the symbol. In this traversal process, a move towards a left and a right node is represented by 0 and 1 respectively. A major drawback of the classical Huffman code is that the whole stream must be read prior to encoding. To transmit a binary string, power dissipation is proportional to the number of switching, i.e., transition from 0 to 1 and vice versa, present in a transmitted string. Although a number of approaches like [7, 8] have extended the classical Huffman code by treating binary bits differently (with unequal letter cost), a little effort has been made to optimise the switching activities in the Huffman code to develop a power efficient Huffman code. To the knowledge of the authors, the only effort to generate Huffman code considering switching activity is seen in [9]. This approach follows a greedy strategy and works in two steps to reduce total number of switching in the Huffman code. In the first step the switching activity between each pair of symbol (inter-switch) is reduced and then in the second step switching activity in the codeword for each each symbol (intra-switch) is reduced. As the optimisation is done independently in two steps therefore it is highly likely that the gain obtained in the first step may be compromised in the second step and vice versa.

In this paper, a genetic algorithm based approach is proposed to reduce the total number of switching where inter-switch and intra-switch is optimised in a single step, i.e, a balance is made between the two different types of switching. The approach generates a set of trees, each one represents a set of codewords (solution). After that a set of genetic operators such as selection, crossover and mutation is used to evolute (create new trees) the population to minimise number of total switches. The evolution process is controlled by the total number of inter- and intra-switches. Inter-Switch between two adjacent symbols depend on the their codewords, i.e., how the codeword of the preceding symbol finished (with 0 or 1) and how the codeword for the descendent symbol started. If the end bit of the preceding

symbol and the staring bit of the successor symbol differs then their exists a switch and the total number of switch is the number of times the two symbols in consideration occurs one after another. To obtain the total number Inter-Switch between pair of symbols, we used a descendent matrix which is a two dimensional matrix represents the number of occurrence of each symbol after another. This matrix is formed when the message to be transmitted is scanned to obtain the frequency of distinct symbols. The number of intraswitch for a symbol is calculated by multiplying the frequency of that symbol by the number of logic level transition (0 to 1 or 1 to 0) in the codeword of the symbol. The efficiency of the approach is evaluated by applying it to a set of biological datasets. The experiments yield that the proposed approach improves the total number of switches by X% in the best case, by Y% in the average case and by Z% in the worst case.

The rest of the paper is organised as follows: Section 2 presents the background study of the relationship of switching activity with power consumption, the relation of switching with Huffman code and the issues in biological data transmission. The proposed approach is described in Section 3. Experimental results and discussion are presented in Section 4. Finally, concluding remarks are presented in Section 5.

2. Background

- 2.1. Huffman Coding
- 2.2. Power consumption
- 2.3. Issues in Biological data transmission

The size of biological data base increase with an expanding rate and it doesn't stop for increasing. In the last decade the use of biological data history increase which evolve a high request to data from this high volume biological data base. The exponential growth of these databases become a big problem to all biological data processing methods. Data compression methods is the most used methods to evolve the reduce the cost of treatment of high volume data bases. The main objective of data compression methods is minimising the number of bits in the data representation. Authors in [] proposed a new approach to genomic data compression based in suffix and common substring and code the difference with Huffman code. In [16], authors propose a general scheme for coding only the difference between the target genome and the referential one, the coding method used is Huffman.

Recent work on data compression for biological data have been proposed to deal with transmission of genomic data as an email attachment [][][]. Cost of data transmission is based on two point, firstly the size of the data which has been improved considerably in the recent year with the improvement of Huffman code. Secondly, is the number of switches on the generated codes. The power consumption increase linear with the number of switches.

3. GA-SO: Genetic algorithm for Switches Optimising

Genetic algorithm (GA) is one the of the most popular bio-inspired metaheuristic algorithm inspired from the natural evolution of species []. It is a population based algorithm starts with the generation of a random initial population. The population contain a set of feasible solutions called individuals, that are usually far from the optimal. The GA optimisation process uses a set of natural genetic operators such as selection, crossover and mutation to converge to the optimal solution.

3.1. Preparing data and population generation

Huffman algorithm read the whole message to compute frequencies. The proposed algorithm read the whole genome to generate the frequencies of the triplet and by the way to generate the adjacent matrix which represents the frequencies precedence of between triplet. The initial population is generated randomly, each solution on the population represents a tree that generates a set of codes equal to the total triplets number.

3.2. Selection

The first operator of the genetic algorithm is the selection []. The main objective of the selection is to choose the part of the current population to be a candidate for the different genetic operators in order to breed the next generation population. Many selection techniques have been proposed in the literature []. In this approach the process of natural selection is maintained. First we generate a random pair numbers rand between 2 and the population size, this number represent how many parents will be processed. After that, an unbiased random selection of rand individuals (solutions) from the population is maid.

3.3. Crossover

The main operator of the GA is the crossover, allows to construct new solutions from the selected part of the population. The selected solutions are ranked by fitness and crossover two by two from high fitness to low. The main objective of the crossover is to benefits from the two good solutions in order to generate a better solution. An internal node for each tree (solution) is selected (node1,node2), these two nodes must have the same number of leaf nodes (contain the same number of codes). The crossover operation will create two new trees (solutions), the first child (second) contain the nodes that are not children of the node1 (node2) and we replace the children of node1 (node2) by the children of node2 (node1) (see fig.1).

3.4. Mutation

The mutation operator changes the positions of two leaf nodes of the generated children (result of the crossover); this introduce the diversity in the search process, this diversification strategy allow the algorithm do conference to the global optimum. The algorithm for this operator goes through the leaf nodes and change the position of two leaf nodes from a parent to another and selects the ones to change according to a fixed mutation rate (see fig.1).

3.5. Population update

The crossover operation aims to generate new solutions from the current solution to built the second generation of the population. Furthermore, the mutation provide the diversity on the search space solution by changing the position of nodes on the same tree. After these two operations, the next step of the genetic algorithm is two to breed the new population. Firstly the new solutions (children) are added to the current population, after that the population is ranked by fitness. The algorithm remove the worst solutions until the initial size of the population is achieved. The algorithm genetic repeat these operators until the stopped criteria is achieved, in this algorithm the stopped criteria is when the objective function (number of switches) stop decreasing.

4. Experiments and Comparison

The effectiveness of the approach has been evaluated with different real genomic biological data, these genomes were downloaded from a recent version of The National Center for Biotechnology Information (NCBI) available

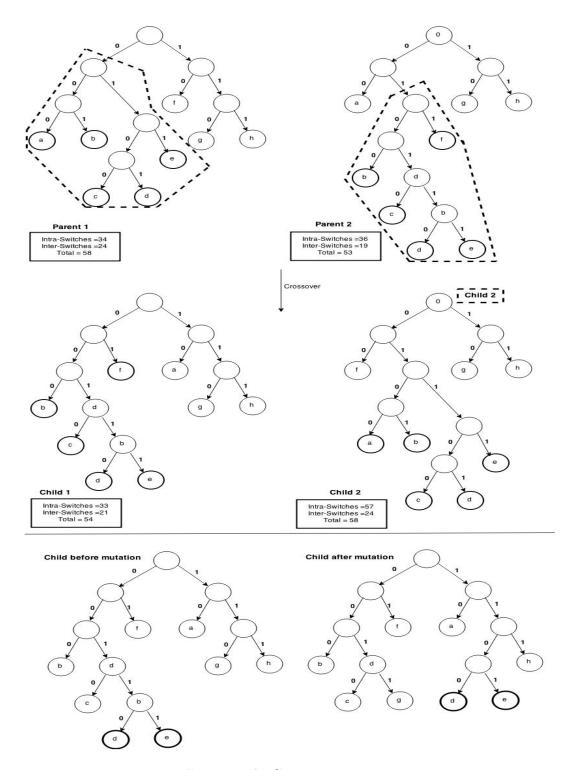


Figure 1: The Crossover operator $\begin{picture}6\end{picture}$

Algorithm 1 Switches optimising Huffman codes

Require: Textual representation of a Genome

Ensure: Low switches Huffman codes

- 1: Generate triplet frequencies and adjacent matrix
- 2: Generate the initial population of trees
- 3: repeat
- 4: Select part of the population
- 5: Generate children candidates via crossover
- 6: Mutate children
- 7: Add children to population
- 8: Rank by fitness
- 9: Remove worst solutions until population limit
- 10: until The switches number stop decreasing

on (http://www.ncbi.nlm.nih.gov) [?]. We focused on the sequences alone, ignoring any header and any other exogenous information. In table 3, the different data sets are described with the size of each of them in megabytes (MB) and the references on the biological data bank.

5. Conclusion

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Table 1: Datasets description

| Data sets | Name | Size (MB) | Reference |
|-----------|---|-----------|-----------------|
| Genome 1 | Mycobacterium smegmatis | 6.66 | CP009496 |
| Genome 2 | Amycolatopsis benzoatilytica | 8.30 | NZ_KB912942 |
| Genome 3 | Mycobacterium rhodesiae NBB3 | 6.11 | CP003169 |
| Genome 4 | Streptomyces bottropensis ATCC 25435 | 8.54 | NZ_KB911581 |
| Genome 5 | Mycobacterium smegmatis str. MC2 155 | 6.66 | CP009494 |
| Genome 6 | Mycobacterium smegmatis MKD8 | 6.76 | NZ_KI421511 |
| Genome 7 | Bradyrhizobium WSM471 | 7.42 | NZ_CM001442 |
| Genome 8 | Amycolatopsis thermoflava N1165 | 8.27 | NZ_CM001442 |
| Genome 9 | Bacillus thuringiensis Bt407 | 5.74 | NZ_CM000747 |
| Genome 10 | Bacillus thuringiensis serovar thuringiensis | 6.03 | NZ_CM000748 |
| Genome 11 | Pseudomonas aeruginosa 9BR | 6.48 | NZ_AFXI01000001 |
| Genome 12 | Bacillus thuringiensis serovar berliner ATCC | 5.97 | NZ_CM000753 |
| Genome 13 | Bacillus thuringiensis serovar pakistani | 5.75 | NZ_CM000750 |
| Genome 14 | Pseudomonas aeruginosa LES400 | 6.28 | CP006982 |
| Genome 15 | Mus musculus chromosome 1 | 25.58 | GL456087 |
| Genome 16 | Danio rerio chromosome 1 | 56.14 | CM002885 |
| Genome 17 | Homo sapiens chromosome 18 | 76.64 | CM000680 |
| Genome 18 | Homo sapiens chromosome 22 | 99.94 | CM000684 |

Table 2: Comparison of performance among classical Huffman code, CCA, and OCCA without penalty

| without penaity | | | |
|-----------------|-------------------|-------|-------|
| | Huffman Algorithm | Р | |
| | | | |
| Genome 1 | 18.16 | 10.05 | 44.65 |
| Genome 2 | 24.66 | 15.63 | 36.61 |
| Genome 3 | 18.46 | 10.66 | 42.25 |
| Genome 4 | 25.18 | 16.26 | 35.42 |
| Genome 5 | 19.24 | 16.08 | 16.42 |
| Genome 6 | 19.61 | 14.96 | 23.71 |
| Genome 7 | 22.40 | 13.21 | 41.02 |
| Genome 8 | 23.87 | 17.70 | 25.84 |
| Genome 9 | 17.66 | 10.04 | 43.14 |
| Genome 10 | 18.14 | 9.89 | 45.47 |
| Genome 11 | 19.63 | 11.48 | 41.51 |
| Genome 12 | 17.59 | 11.86 | 32.57 |
| Genome 13 | | | |
| Genome 14 | | | |
| Genome 15 | | | |
| Genome 16 | | | |
| Genome 17 | | | |
| Genome 18 | | | |
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