



UNIVERSITI PUTRA MALAYSIA

**HIGH-THROUGHPUT SEQUENCING AND ANALYSIS
OF CHROMOSOME 1 OF EIMERL4 TENELLA**

LING KING HWA

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CHROMOSOME 1 OF *EIMERIA TENELLA***

PERPUSTAKAAN PERUBATAN
UNIVERSITI PUTRA MALAYSIA

By

LING KING HWA

**Thesis Submitted to the School of Graduate Studies, Universiti Putra Malaysia,
in Fulfillment of the Requirements for the Degree of Master of Science**

June 2005



*This thesis is dedicated to my beloved Pike See
for her unwavering support, understanding, strength,
love, constant encouragement and for instilling me
with a belief in my own abilities.*



Abstract of thesis presented to the Senate of Universiti Putra Malaysia in fulfilment of the requirement for the degree of Master of Science

HIGH-THROUGHPUT SEQUENCING AND ANALYSIS OF CHROMOSOME 1 OF *EIMERIA TENELLA*

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Chairman : Associate Professor Rozita Rosli, Ph.D.

Faculty : Medicine and Health Sciences

Eimeria tenella is one of seven *Eimeria* species that causes avian coccidiosis. It is also highly pathogenic and is one of the three most common species occurring in the field. High-throughput sequencing of the 1.05Mb chromosomes one of *Eimeria tenella* (Houghton strain) revealed genomic information which may be useful in the discovery of genes such as those involved in drug resistance, cellular regulation and integration, and mechanisms of invasion.

High-throughput and random chromosomal shotgun sequencing resulted in 61 unfinished contigs representing full shotgun state of chromosome one nucleotide sequences which are ready into the finishing phase. Out of these contigs, 57 of them were arranged into 11 scaffolds whereas 4 contigs remain unordered. All the contigs represent 86.9% or 9.29-fold coverage of chromosome one. The quality of the assembly is assured with 94.1% of consistent paired reads with only 5824.3 (0.5%) errors expected. In addition, contiguity of the assembly vastly improved with the integrated BAC-end sequences and HAPPY map markers. Consensus level assessment showed 99.2% of the unfinished chromosomal sequence has expected error rate less than 1 per 10,000 bases (PHRAP score > 40) and only 7.6% of them need further polishing.

The GC content of chromosome one is 49.35% and long-ranged excursions from its mean are found prominently in three regions whereas chromosomal wide GC fluctuations ranged from 35% to 60% at a 12kb window length analysis. GC skews were found to be correlated with the repeats rich regions of the chromosome. Telomeric sequence at both ends of the chromosome is derived as TTTAGGG / CCCTAAA with undefined real length. A centromeric like region with approximately 1,453bp was found in chromosome one with 81.3% AT composition. Chromosome one is expected to bear at least 25.3% of repetitive elements with the most prominent tandem repeat, TGC, which are distributed throughout the chromosome. The longest minisatellite, *mst1*, is 3,624bp in length and occurs as a single stretch in the chromosome. Besides that, there are a few under-characterized interspersed repeats such as LINE and DNA transposons which were found in the chromosome and preliminary homology-based gene survey demonstrated the



possibility of LTR elements in SC11. Both the GC skews and distribution of repetitive elements divide the chromosome into 7 prominent regions.

Alignments with non-redundant and EST databases during gene survey gave a coarse estimation of coding densities of chromosome one at 1 CDS per 1000bp which also corresponded to 12.6% as coding and 87.4% as non-coding. Careful inspection on the distribution revealed that the coding sequences are centrally arranged within the chromosome. GC composition (53.9%) is higher in coding sequences compared to non-coding sequences (48.6%). The number of genes embedded in chromosome one is unknown until further laboratory investigations are carried out. Some of the significant hits may reflect the presence of the genes in chromosome one such as previously characterized LPMC-61 antigen, elongation factor Tu, proteophosphoglycan, proteases, and AAA ATPase family proteins that are involved in the parasite's mobility, parasite-host interaction and possibly invasion.

However, *in silico* gene prediction using a homology-based technique identified three full length genes, phosphatidylinositol-4-phosphate 5-kinase (PIP5K), glucose-6-phosphate isomerase (PGI) and malate:quinone oxidoreductase (MQO). These genes served as gene models and provided early information regarding the intron, exon and splicing sites. The average exon and intron sizes were predicted as 118.5bp and 535.3bp, respectively. The most commonly utilized splice pairs is AG...GT. Chromosome one nucleotide sequences have been deposited in the data depository of the Interim Laboratory of National Institute for the Genomics and Molecular Biology, BIOVALLEY-UKM, Bangi, Malaysia.



Abstrak tesis yang dikemukakan kepada Senat Universiti Putra Malaysia sebagai memenuhi keperluan untuk ijazah Master Sains

**PENJUJUKAN BERSKALA BESAR DAN ANALISIS
KROMOSOM SATU *EIMERIA TENELLA***

Oleh

LING KING HWA

Jun 2005

Pengerusi : Profesor Madya Dr. Rozita Rosli, Ph.D.

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Eimeria tenella merupakan satu daripada tujuh spesies *Eimeria* yang menyebabkan koksidiosis ayam. Ia juga merupakan spesies yang paling patogenik dan paling kerap ditemui berbanding dengan yang lain. Penjujukan kromosom satu *Eimeria tenella* strain Houghton secara berskala besar akan menghasilkan maklumat genomik untuk penemuan gen-gen yang terlibat dalam kerintangan dadah, integrasi dan pengawalaturan sel serta mekanisme jangkitan.

Penjujukan kromosom berskala besar secara rawak telah menghasilkan 61 kontig yang sedia menjejaki fasa penghabisan. Lima puluh tujuh kontig daripada kontig-kontig ini, telah diaturkan ke dalam 11 kerangka manakala 4 kontig yang lain tidak dapat diaturkan ke dalam mana-mana perancah. Kontig-kontig ini mewakili 86.9% atau 9.29 kali liputan kromosom satu. Pasangan jujukan konsisten yang tinggi (94.1%) dan jangkaan ralat yang rendah (5824.3 atau 0.5%) menjamin kualiti perhimpunan jujukan yang tinggi. Kesenambungan perhimpunan jujukan juga diperbaiki melalui integrasi jujukan-jujukan hujung kromosom buatan bakteria dan penanda peta HAPPY. Sebanyak 99.2% daripada jujukan konsensus kromosom satu dijangkakan mempunyai kadar ralat kurang daripada 1 bes dalam setiap 10,000 bes (Skor PHRAP > 40) dan cuma 7.6% daripada jujukan konsensus ini perlu diperbaiki.

Kandungan GC kromosom satu ialah 49.35% dan penyisihan julat besar daripada nilai min didapati ketara di tiga kawasan manakala julat naik-turun kandungan GC secara keseluruhan adalah dari 35% hingga 60% pada analisis gelangsar tingkap bersaiz 12kpb. Sisihan kandungan GC juga didapati berkaitan dengan kawasan kromosom satu yang kaya dengan jujukan ulangan. Jujukan telomer pada kedua-dua hujung kromosom mengandungi jujukan ulangan TTTAGGG / CCCTAAA dengan saiz sebenar belum dapat ditentukan lagi. Kawasan mirip sentromer dengan saiz 1,453pb dan kandungan AT setinggi 81.3% juga ditemui. Kromosom satu dijangkakan mengandungi sekurang-kurangnya 25.3% elemen ulangan. Jujukan ulangan jenis tandem yang paling kerap ditemui ialah TGC dan tertabur sepanjang kromosom satu. Satelit mini yang paling panjang ialah *mst1*, mempunyai saiz 3,624pb dan wujud cuma dalam satu bentangan dalam kromosom. Selain itu, elemen seperti transposon DNA dan elemen nuklear celahan panjang juga ditemui dalam kromosom satu. Peninjauan gen berdasarkan konsep homologi menunjukkan



kewujudan elemen terminal panjang di SC11. Kedua-dua ciri sisihan kandungan GC dan elemen ulangan telah membahagikan kromosom kepada tujuh kawasan yang menonjol.

Penjajaran jujukan pengkodan kromosom satu dengan jujukan-jujukan penanda terungkap dan jujukan-jujukan di pangkalan data 'non-redundant' semasa peninjauan gen, menganggarkan 1 jujukan pengkodan dalam setiap 1000pb. Ini juga sepadan dengan 12.6% daripada jujukan kromosom satu sebagai jujukan pengkodan dan 87.4% sebagai bukan pengkodan. Pemantauan terperinci menunjukkan bahawa taburan jujukan pengkodan ini diatur ketengah kromosom. Kandungan GC didapati lebih tinggi (53.9%) pada jujukan pengkodan berbanding bukan pengkodan (48.6%). Jumlah sebenar gen yang terkandung dalam kromosom satu masih tidak diketahui sehingga bukti makmal diperolehi. Penjajaran yang ketara menunjukkan kehadiran gen-gen seperti yang mengkodkan antigen LPMC-61, factor pemanjangan Tu, *proteophosphoglycan*, protease dan protein dalam famili AAA ATPase yang kesemuanya terlibat sama ada dalam mobiliti parasit, interaksi hos-parasit dan kemungkinan dalam proses jangkitan.

Ramalan gen secara *in silico* melalui teknik homologi telah menemukan tiga gen bersaiz penuh iaitu, *phosphatidylinositol-4-phosphate 5-kinase* (PIP5K), *glucose-6-phosphate isomerase* (PGI) dan *malate:quinone oxidoreductase* (MQO). Gen-gen ini berfungsi sebagai model gen dan memberi maklumat awal berkaitan intron, ekson dan tapak-tapak penyambatan. Saiz purata ekson dan intron ialah 118.5pb dan 535.3pb masing-masing. Tapak penyambatan yang paling kerap digunakan ialah pasangan AG...GT. Jujukan-jujukan kromosom satu telah disimpan dalam pangkalan data di Makmal Interim untuk Institut Kebangsaan Genomik dan Biologi Molekul BIOVALLEY-UKM, Bangi, Malaysia.



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Ling King Hwa

12th October 2004



I certify that an Examination Committee met on 16th June 2005 to conduct the final examination of Ling King Hwa on his Master of Science thesis entitled "High-throughput sequencing and analysis of chromosome 1 of *Eimeria tenella*" in accordance with Universiti Pertanian Malaysia (Higher Degree) Act 1980 and Universiti Pertanian Malaysia (Higher Degree) Regulations 1981. The Committee recommends that the candidate be awarded the relevant degree. Members of the Examination Committee are as follows:

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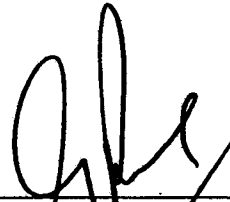
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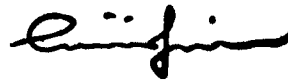
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DECLARATION

I hereby declare that the thesis is based on my original work except for quotations and citations which have been duly acknowledged. I also declare that it has not been previously or concurrently submitted for any other degree at UPM or other institutions.


LING KING HWA

Date: 29th June 2005

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LIST OF ABBREVIATIONS

Ω	Ohm
μ l	Microliter
A	Adenine
ATP	Adenosine triphosphate
BLAST	Basic Local Alignment Search Tool
bp	Base pair
C	Cytosine
cDNA	Complementary deoxyribonucleic acid
ddNTP	Dideoxyribonucleotide triphosphate
DNA	Deoxyribonucleic acid
dNTP	Deoxyribonucleotide triphosphate
EST	Expressed sequence tag
F	Faraday unit
G	Guanine
h	Hour
HSP	High Scoring Segments
IPTG	Isopropyl-beta-D-thiogalactopyranoside
Kb	Kilobase
kV	Kilovolt
LB	Luria-Bertani
LINE	Long interspersed nuclear element
LTR	Long terminal repeat
mA	Milliampere



Mb	Megabase
mg	Milligram
ml	Milliliter
MQO	Malate:quinone oxidoreductase
ng	Nanogram
ORF	Open reading frame
PCR	Polymerase chain reaction
PFGE	Pulse-Field Gel Electrophoresis
PGI	Glucose-6-phosphate isomerase
PIP5K	Phosphatidylinositol-4-phosphate 5-kinase
POP	Performance Optimized Polymer
RNA	Ribonucleic acid
rpm	revolution per minute
T	Thymine
U	Unit enzyme
X-gal	5-bromo-4-cloro-3-indolil-beta-D-galatopiranocid
Hg	Mercury



CHAPTER 1

INTRODUCTION

1.1 *Eimeria* species and coccidiosis

Eimeria species belong to the phylum Apicomplexa and is responsible for the diseases of coccidiosis in intensively reared livestock such as poultry, cattle and sheep. Other members of this phylum include *Plasmodium falciparum*, the causative agent of malaria; *Toxoplasma gondii*, an opportunistic pathogen in immunocompromised individuals; and *Cryptosporidium parvum*, an animal parasite as well as an opportunistic pathogen of humans. These parasites are characterized by the presence of apical complex structures or organelles such as conoid, apical polar ring, micronemes, rhoptries and dense granules (Chobotar and Scholtyseck, 1982). Apicomplexans are obligate intracellular parasites that require host cells in which they can invade and replicate to ensure their survival.

Eimeria species from chickens are the most important parasitic pathogens of poultry. There are seven *Eimeria* species that infect chickens; *Eimeria tenella*, *Eimeria necatrix*, *Eimeria acervulina*, *Eimeria brunetti*, *Eimeria maxima*, *Eimeria mitis* and *Eimeria praecox* (Marquardt *et al.*, 2000). All the seven *Eimeria* species infect the intestinal epithelial lining of chicken at only specific locations. In an intensively reared flock of chickens, coccidiosis is always due to the infections of more than one species of *Eimeria* (Williams *et al.*, 1996).

