

The complete mitochondria genome of *Chrysomya phaonis* (Seguy, 1928) (Diptera: Calliphoridae)

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ABSTRACT

Chrysomya phaonis (Seguy, 1928) is one of the blowflies of great medical and forensic importance. In this paper, we report that the entire genome of mitochondrial DNA of *C. phaonis* is 15,831 bp in length, which consists of 39 genes including 13 protein-coding genes, 24 tRNA genes, 2 mitochondrial ribosomal RNA genes, and a 992 bp non-coding A+T-rich region. The overall base compositions of A, G, C, and T are 38.79%, 9.75%, 14.15%, and 37.31%, respectively. We provide the first complete mitochondrial genome of *C. phaonis*, and should provide useful information for phylogenetic and species identification for *C. phaonis*.

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Blowflies (Diptera: Calliphoridae) are distributed worldwide, some species play important role in the mechanical transfer of disease to human beings and animals (Norris 1965; Zumpt 1965; Greenberg 1971; Greenberg 1973; Kuhlhorn 1983; Ghandour 1988). *Chrysomya phaonis* (Seguy, 1928) is one of the most important oriental species of the blowflies, *C. phaonis* distributes in China, India, Nepal, and Afghanistan (Fan 1992; Fan et al. 1997; Kurahashi et al. 1994). However, there is limited molecular biology information about this species. Viewing it is a potential medical vector and indicator in forensic science, we report here the complete mitochondrial genome of *C. phaonis* for species identification and phylogenetic analysis.

Exampled sample of *C. phaonis* were obtained in Bayi County, Tibet, China ($N29^{\circ}37'31.39''$; $E94^{\circ}23'25.01''$) in July 2013. The studied specimen is stored in the medical vector collections of Zhongshan Entry-Exit Inspection and Quarantine technology center, and the accession number to the specimen is 20130728-232F-Baiyi. We designed 10 pairs of oligo-nucleotide primers according to the conserved regions from reported mitochondria genome sequences of its most related species *C. pinguis*.

The complete mitochondrial genome of *C. phaonis* (GenBank accession KX500359) is 15,831 bp in length, which consists of 39 genes (Table 1) including 13 protein-coding genes, 24 tRNA genes, 2 mitochondrial ribosomal RNA genes (12S rRNA and 16S rRNA), and a 992 bp non-coding A+T-rich region. The 13 protein-coding genes include seven NADH

dehydrogenase (*ND1-6* and *ND4L*), three subunits of cytochrome oxidase (*COI-III*), two subunits of ATP synthase (*ATP4* and *ATP6*), one subunit of cytochrome b (*Cytb*). Twelve of the 13 protein-coding genes were identified with ATN as start codon coding for M except for *COI*, which is similar to the former studies (Yan et al. 2014; Zhong et al. 2016).

Chrysomya phaonis not only could cause harm to human health as medical vectors (Norris 1965; Zumpt 1965; Greenberg 1971; Kuhlhorn 1983; Ghandour 1988), but also are significant in forensic science (Harvey et al. 2008). DNA typing of forensic insect specimens offers a quick and reliable alternative, so more and more researchers have started using the whole sequences of mitochondrial genome or part of the genes to identify species of Blowflies (Wells & Williams 2007; Harvey et al. 2008; Desmyter & Gosselin 2009; DeBry et al. 2013).

Here, we provide the entire genome of mitochondrial DNA of *C. phaonis*. The phylogenetic analysis of *C. phaonis* was performed by comparison with other 39 Diptera species mitochondrial genomes (Figure 1). Phylogenetic tree was generated by a neighbour-joining analysis of MEGA 6.0 program (Tamura et al. 2013) using 1000 bootstrap replicates. The N-J tree revealed that *C. phaonis* was placed mostly close to *C. pinguis*, which could not be distinguished with the common adopted COI DNA barcode sequences. So, it may provide some help for the molecular identification of *C. phaonis*, in particular to distinguish *C. phaonis* from its closely related species *C. pinguis*.

Table 1. Mitochondrial gene profile of *C. phaonis*.

Gene	Direction	Nucleotide number	Size (bp)	OL	Non	Anticodon	Codon Start Stop
<i>tRNA^{Ile}</i>	J	1–64	64		4	GAT	
<i>tRNA^{Gln}</i>	N	69–137	69		8	TTG	
<i>tRNA^{Met}</i>	J	146–214	69			CAT	
<i>ND2</i>	J	215–1229	1015				ATT T
<i>tRNA^{Trp}</i>	J	1230–1297	68	8		TCA	
<i>tRNA^{Cys}</i>	N	1290–1353	64		7	GCA	
<i>tRNA^{Tyr}</i>	N	1361–1426	66	2		GTA	
<i>COI</i>	J	1425–2958	1534				TCG T
<i>tRNA^{Leu(UUR)}</i>	J	2959–3024	66		5	TAA	
<i>COII</i>	J	3030–3717	688				ATG T
<i>tRNA^{Lys}</i>	J	3718–3788	71	1		CTT	
<i>tRNA^{Asp}</i>	J	3788–3854	67			GTC	
<i>ATP8</i>	J	3855–4019	165	7			ATT TAA
<i>ATP6</i>	J	4013–4690	678		4		ATG TAA
<i>COIII</i>	J	4695–5483	789		6		ATG TAA
<i>tRNA^{Gly}</i>	J	5490–5554	65			TCC	
<i>ND3</i>	J	5555–5908	354		2		ATT TAA
<i>tRNA^{Ala}</i>	J	5911–5975	65	1		TGC	
<i>tRNA^{Arg}</i>	J	5975–6038	64		6	TCG	
<i>tRNA^{Asn}</i>	J	6045–6110	66	1		GTT	
<i>tRNA^{Ser(AGN)}</i>	J	6110–6179	70			GCT	
<i>tRNA^{Glu}</i>	J	6180–6247	68		18	TTC	
<i>tRNA^{Phe}</i>	N	6266–6332	67			GAA	
<i>ND5</i>	N	6333–8052	1720		15		ATT T
<i>tRNA^{His}</i>	N	8068–8133	66			GTG	
<i>ND4</i>	N	8134–9472	1339	7			ATG T
<i>ND4L</i>	N	9466–9762	297		2		ATG TAA
<i>tRNA^{Thr}</i>	J	9765–9829	65			TGT	
<i>tRNA^{Pro}</i>	N	9830–9895	66		2	TGG	
<i>ND6</i>	J	9898–10,422	525	1			ATT TAA
<i>Cytb</i>	J	10,422–11,556	1135				ATG T
<i>tRNA^{Ser(UCN)}</i>	J	11,557–11,624	68	16		TGA	
<i>ND1</i>	N	11,641–12,579	939		10		ATA TAA
<i>tRNA^{Leu(CUN)}</i>	N	12,590–12,654	65		1	TAG	
<i>IrRNA</i>	N	12,656–13,983	1328				
<i>tRNA^{Val}</i>	N	13,984–14,055	72			TAC	
<i>srRNA</i>	N	14,056–14,839	784				
A + T rich		14,840–15,831	992	66			
<i>tRNA^{Ile}</i>	J	14,929–14,994	66			GAT	

Disclosure statement

The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

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