



## Larval growth, silk production and economic traits of *Bombyx mori* under the influence of honey-enriched mulberry diet

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**Abstract:** The impact of honey on the silkworm, *Bombyx mori* was demonstrated with reference to the larval growth, silk gland proteins and economic parameters of sericulture. The honey works well at a concentration of 2% in distilled water. At this concentration it promotes growth rates in the silkworm larvae during fifth instar development. It positively reinforces the day-to-day larval growth rate by 4.75 additional percentile points, silk gland growth rate by 4.45 additional percentile points and the gland-body ratio by additional 6.64 percentile points. It stimulates silk protein synthesis in all the three segments of the silk gland, viz., the anterior, middle and posterior parts. Under its influence, the silk gland protein profiles grew significantly by 14.85 additional percentile points in the anterior silk gland (ASG), minimally by 8.68 additional percentile points in the middle silk gland (MSG) and maximally by 15.17 additional percentile points in the posterior silk gland (MSG), compared to their control values. It also stimulates the core shell protein synthesis by 18% and retards floss protein synthesis by ~25% in the three segments of silk gland. In doing so, it contributes to sericulture industry by causing im-provements in profit making economic traits such as gland-body ratio, cocoon weight, shell weight, raw silk weight, denier and renditta and by reducing the production of floss, which contributes to loss in the sericulture industry. Honey is suggested as a profitable supplementary diet for silkworm.

**Keywords:** *Bombyx mori*, Larval growth, Honey, Silk proteins, Economic traits

### INTRODUCTION

The impact studies of exogenous nutrients on the growth, metabolism and development of *B. mori* have been accorded importance in sericultural research (Laskar and Datta, 2000; Kanafi *et al.*, 2007). Such studies largely focused on the enrichment of the silkworm diet (i.e., mulberry leaves) with exogenous nutrients such as proteins, carbohydrates, amino acids, vitamins, minerals, hormones, antibiotics and assessing their influence on desired parameters of *B.mori* (Sanappa *et al.*, 2002; Bhattacharya and Kaliwal, 2004, 2005a, 2005b, 2005c; Chakrabarty and Kaliwal, 2011; Kavitha *et al.*, 2012; Thulasi and Sivaprasad, 2013, 2014).

The honey is a natural sweetener and multi- factorial nutrient produced by the honey bees from the floral nectar (Council of European Union, 2002). It is a rich nutrient and includes a variety of sugars, proteins, enzymes, vitamins and minerals and its chemical composition and characteristics vary depending on its geographical, environmental and botanical origin (Ramirez, 2000; Falco *et al.*, 2003; Anonymous, 2003; Garcia *et al.*, 2005; David Ball, 2007). Further, the honey is an insect product and possess valuable nourishing, healing and prophylactic properties and used as an ingredient of human and veterinary

medicines (Taormina, 2001; Cooper *et al.*, 2002; Anonymous, 2003; Eileen De Mars, 2003; Iglesias *et al.*, 2004). Most of the enrichment studies employed commercial and expensive nutrients, but, no effort has since been made to examine the impact of an insect product and all-in-one nutrient like honey on the silkworm growth, metabolism and economic traits of sericulture.

Keeping in view the high nutritional and medicinal status of honey, the present study explores the possibility of including honey in the silkworm diet by analyzing its impact on the larval growth and protein profiles of *B. mori* and on the economic parameters of sericulture.

### MATERIALS AND METHODS

The present investigation was carried out on PM x CSR<sub>2</sub> hybrid variety of the silkworm (*Bombyx mori*), reared under standard environmental conditions of 28<sup>0</sup>C, 85 % RH as per Krishnaswami (1986). After hatching, the worms were reared on M<sub>5</sub> variety of mulberry leaves by giving 5 feeds per day at 6 AM, 10 AM, 2 PM, 6 PM and 10 PM, under normal 12 hrs light and 12 hrs dark conditions. The experimental design was divided into four phases, namely honey feeding pattern, study of growth patterns, assay of tissue proteins and analysis of economic parameters of

sericulture.

**Honey feeding pattern:** After the third moult, the fourth instars larvae were divided into five batches of 100 worms each. One was treated as the control batch and four others as experimental batches. The control batch was fed with normal feedings 5 times a day as stated earlier and the experimental batches were fed with mulberry leaves soaked in 2% honey in distilled water, the minimum effective concentration of which has been determined earlier by us (Sivaprasad and Thulasi, 2014). Before feeding, the wet mulberry leaves were dried under cool weather conditions. While the honey-enriched leaves were fed to the silkworms at 6 PM, normal feeding pattern was continued at other timings during the day.

**Silkworm growth:** Studies under this head included the impact analysis of honey on the growth of the larval body and silk gland (SG) during fifth instar development. The growth of both the larval body and SG was ascertained in both control and experimental batches and the latter included a single treatment condition i.e.; the larvae fed with fortified mulberry leaves with 2% minimum effective concentration of honey. The body and SG weights were measured in an electronic balance (ELICO; Model BL- 22 OH) on alternative days of fifth instar (i.e.; day1, day-3, day-5 and day-7). The mean weight of 25 randomly selected larvae and 5 pairs of silk glands were taken as the standard weights and the same were expressed in grams.

**Assay of tissue proteins:** Total protein levels of the silk gland (SG), fat body (FB) and haemolymph (HL) were analyzed on day-1, day-4 and day-7 during fifth instar larval development. While the SG and FB were isolated by mid- dorsal dissection of larval body in the Silkworm Ringer (Yamaoka *et al.*, 1971), the HL was extracted by cutting the telson and pro-legs. The total protein content was estimated in 1% homogenates of SG and FB and 1:9 diluted HL (1:9 haemolymph and distilled water) by the method of Lowry *et al.* (1951) and the same was expressed in mg protein / gram wet weight of tissue (or) mg/ml of haemolymph. Similarly, the protein content of the cocoon was estimated in 1 % homogenate in distilled water. Since the silk cocoon is not soluble in distilled water, it was first soaked in diluted sodium hydroxide solution before homogenized in distilled water.

**Analysis of economic parameters:** Nine important economic parameters of sericulture, viz., as the green cocoon weight, shell weight, shell protein, floss weight, floss protein, pupal weight, raw silk weight, denier, renditta, were analyzed as per the methods given by Bohidar *et al.*, 2007; Rahmathulla *et al.*, 2007; Sailaja and Sivaprasad, 2010).

**Statistical analysis:** The data were analyzed by statistical tools such as mean, standard deviation (SD), Percent change and test of significance. While the mean and SD were computed using M.S. Excel, the test of Significance and percent -changes was calculated by

online using the Graph pad and percent change platforms ([www. Graph pad. Com/quick calcs/index cfm/and](http://www.Graphpad.Com/quick_calcs/index_cfm/and) [www. Percent-change com/ index php](http://www.Percent-change.com/index_php)). Further, in order to draw meaningful conclusions, the growth trends in larval body, SG and silk gland proteins were interpreted in terms of an innovative statistical parameter called compound periodical growth rate (CPGR) as given by Sivaprasad (2012).

## RESULTS

The findings of the present investigation on the larval growth, silk gland growth and protein profiles and economic traits profiles of sericulture are presented in Tables 1 to 3 and in Figs. 1 to 3.

**Body versus gland growth:** Under natural conditions, the larval weight of the control batch grew maximally by ~128% on day-3, ~67% on day-5 and minimally by ~15% on day-7 with an overall increase of 334% and a CPGR of 63.18% (Table 1, Fig. 1). At the same time, the SG grew maximally by ~411% on day-3, ~98% on day-5 and ~92% on day-7, representing an overall increase of ~1844% during fifth instar with a CPGR of 227.11% (Table 1). The computed gland- body ratio showed an increase of ~125% on day-3, ~19% on day-5 and ~92% on day-7 with an overall increase of 334% and a CPGR of ~64.82% in the control batch. Under the influence of 2%, minimum effective concentration of honey, the larval body weight recorded an increase of ~155% on day-3, ~58% on day-5 and ~18% on day-7 with an overall increase of 374% and a CPGR of ~68% during fifth instar larval development (Table 1). At the same time, the SG grew by ~483% on day-3, ~167% on day-5 and ~53% on day-7, with an overall increase of 2289% and a CPGR of 250.34%. The gland- body ratio recorded an increase of ~129% on day-3, ~69% on day-5 and ~31% on day-7, representing an overall growth of ~404%, and a CPGR of ~71.47% (Table 1, Fig.1).

**Silk gland proteins (SGP):** Since the honey caused significant improvement in the whole SGP, it is necessary to find as to which region of SG responds more effectively to this exogenous factor. Hence, protein levels were estimated separately in the three regions namely, anterior silk gland (ASG), middle silk gland (MSG) and posterior silk gland (PSG) and the relevant data are presented in table 2 and fig. 2. The protein profiles of ASG in the control and honey treated batch were elevated respectively by ~16% and 51% on day-3, ~3% and 31% on day-5 and just by 17% and 16% on day-7. In the MSG, the corresponding elevations were ~44% and 101% on day-3, ~45% and 9% on day-5 and ~51% and 69% on day-7. While those in PSG were ~41% and 103% on day-3, ~21% and 6% on day-5 and ~51% and 64% on day-7. During the entire period of fifth instar, the overall growth trends in the total protein levels of control and experimental batches were ~39% and 102% in ASG, ~213% and ~272% in MSG and ~158% and 254% in PSG. Similarly, the corresponding CPGRs were 11.64% and 26.49% in ASG, ~46.25%

**Table 1.** Effect of 2% honey on body weight (BW), silk gland (SG) weight and gland- body ratio (GBR) in *B. mori*, during fifth instar larval development.

Day	Statistical tool	Control			Experimental (2% Honey)		
		BW (g)	SG weight (g)	GBR	BW (g)	SG weight (g)	GBR
1	Mean	0.58	0.018	3.10	0.58	0.018	3.10
	S.D	±0.008	±0.0009	±0.009	±0.008	±0.0009	±0.009
3	Mean	1.32	0.092	6.96	1.48	0.105	7.09
	P.C	(127.58)	(411.11)	(124.51)	(155.17)	(483.33)	(128.70)
	S.D	±0.02*	±0.001*	±0.008*	±0.01*	±0.002*	±0.008*
5	Mean	2.20	0.182	8.27	2.34	0.280	11.96
	P.C	(66.66)	(97.82)	(18.82)	(58.10)	(166.66)	(68.68)
	S.D	±0.01*	±0.001*	±0.009*	±0.01*	±0.0009*	±0.01*
7	Mean	2.52	0.350	13.88	2.75	0.43	15.63
	P.C	(14.54)	(92.30)	(67.83)	(17.52)	(53.57)	(30.68)
	S.D	±0.009*	±0.009*	±0.02*	±0.008*	±0.01*	±0.01*
OPC (%)		334.5	1844.4	347.7	374.1	2288.9	404.2
CPGR (%)		63.18	227.11	64.82	68.0	250.34	71.47

\* Statistically significant (P value < 0.001); \*\*Statistically not significant. Each value is a mean, ± standard deviation of four individual observations. The percent changes were calculated taking the previous value as the base value, while the overall percentages (OPCs) and the compound periodical growth rates (CPGRs) were computed on the basis of first and seventh day values as per Sivaprasad, 2012.

**Table 2.** Effect of 2% honey on the total protein profiles (mg/g) of different regions of the silk gland in the fifth instar larva of *B. mori*.

Day	Statistical tools	ASG		MSG		PSG	
		Control	Experimental (2% Honey)	Control	Experimental (2% Honey)	Control	Experimental (2% Honey)
1	Mean	10.88	10.88	20.18	20.18	19.04	19.04
	S.D	±0.83	±0.83	±0.36	±0.36	±0.45	±0.45
3	Mean	12.63	14.56	28.97	40.63	26.88	38.64
	P.C	(16.08)	(50.76)	(43.55)	(101.33)	(41.17)	(102.94)
	S.D	±0.009*	±0.01*	±0.49*	±1.47*	±0.008*	±0.70*
5	Mean	12.98	19.02	41.88	44.27	32.64	41.03
	P.C	(2.77)	(30.63)	(44.56)	(8.95)	(21.42)	(6.18)
	S.D	±0.008*	±0.01*	±0.45*	±1.08*	±0.01*	±0.90*
7	Mean	15.14	22.02	63.13	75.05	49.21	67.40
	P.C	(16.64)	(15.77)	(50.74)	(69.52)	(50.76)	(64.27)
	S.D	±0.20*	±0.14*	±0.22*	±0.55*	±0.02*	±0.64*
OPC (%)		39.2	102.4	212.8	271.9	158.5	254.0
CPGR (%)		11.64	26.49	46.25	54.93	37.23	52.40

\* Statistically significant (P value < 0.001); \*\*Statistically not significant; Each value, expressed in mg/g wet weight of tissue, is a mean, ± standard deviation of four individual observations. The percent changes were calculated taking the previous value as the base value, while the overall percentages (OPCs) and the compound periodical growth rates (CPGRs) were computed on the basis of first and seventh day values as per Sivaprasad, 2012. ASG: Anterior silk gland; MSG: Middle silk gland; PSG: Posterior silk gland

and 54.93% in MSG and 37.23% and 52.40% in PSG (Table 2, Fig. 2).

**Economic traits:** The positive impact of 2% honey has been extended to the economic parameters of sericulture such as the green cocoon weight, shell weight, shell protein, floss weight, floss protein, pupal weight, raw silk weight, denier and renditta (Table 3, Fig. 3). The impact yielded significant gains in profitable traits and

reductions in loss making ones. Under its influence, the green cocoon weight increased by ~23%, shell weight by ~118%, shell protein by ~18%, pupal weight by ~13%, raw silk weight by ~18% and denier by ~18%. The positive impact of honey is further reinforced by reduction in floss weight by ~13% and floss protein by ~25% and renditta by ~2% (Table 3). Further, the raw silk generated from the experimental stock has

**Table. 3.** Effect of 2% honey on the economic traits of *B. mori*.

S. N.	Parameters	Statistical tools	Control	Honey treated
1	Green cocoon Weight (g)	Mean	0.96	1.08
		P.C	-	(22.91)
		S.D	±0.02	±0.0009*
2	Shell Weight (g)	Mean	0.17	0.37
		P.C	-	(117.64)
		S.D	±0.009	±0.01*
3	Shell protein (mg/g)	Mean	36.01	42.6
		P.C	-	18.30
		S.D	1.01	±0.14*
4	Floss Weight (g)	Mean	0.024	0.021
		P.C	-	(-12.5)
		S.D	±0.004	±0.003**
5	Floss protein (mg/g)	Mean	7.27	5.42
		P.C	-	(-25.44)
		S.D	0.17	±0.03*
6	Pupal weight (g)	Mean	0.92	1.04
		P.C	-	(13.04)
		S.D	±0.003	±0.003*
7	Raw silk weight (g)	Mean	14.87	17.48
		P.C	-	(17.55)
		S.D	±0.01	±0.01*
8	Denier	Mean	11.85	13.93
		P.C	-	(17.55)
		S.D	±0.008	±0.07*
9	Renditta	Mean	5.20	5.07
		P.C	-	(-2.5)
		S.D	±0.01	±0.01*

\*Statistically significant ( $P$  value < 0.001): \*\*Statistically not significant. Each value is a mean  $\pm$  standard deviation of four individual observations. The weights of the cocoon, shell and floss represent the mean of 25 individual cocoons, expressed in grams. The values in parentheses represent the percent changes from the control.

light yellow colour compared to the control stock which bear dark yellow colour (Fig. 3).

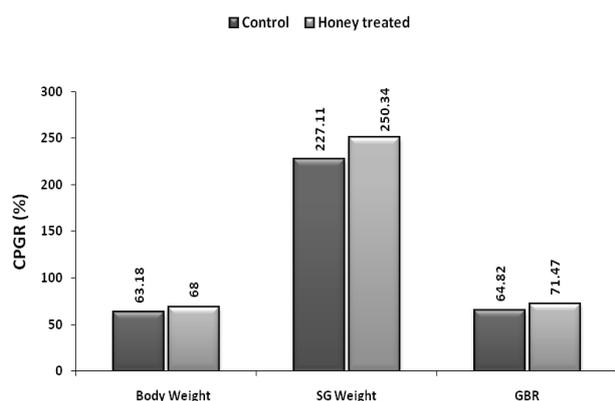
## DISCUSSION

Honey is the richest and natural nutrient comprising chiefly carbohydrates (82%) that meet the energy requirements of metabolism and silk production. It also includes proteins, enzymes (e.g. Diastase, invertase, glucose oxidase, catalase, etc.), free amino acids, trace amounts of B-vitamins and vitamin C, and metals such as Cr, Co, Cu, Fe, Mn and Zn (Falco *et al.*, 2003; Garcia *et al.*, 2005; David Ball, 2007). Due to high nutritive value, it plays a vital role on the growth and metabolism of organism, a fact that has been clearly demonstrated in *B. mori* in its larval growth, protein profiles and economic traits of sericulture.

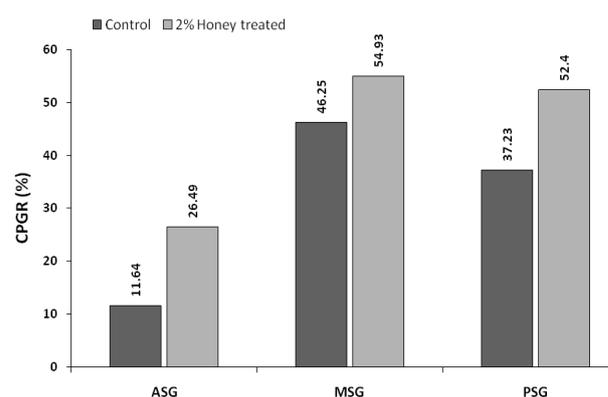
**Honey versus Larval growth:** In the control batch, the larval growth recorded a CPGR of ~25.28% in fifth instar. The larvae fed with 2% honey enriched mulberry leaves the growth was elevated by ~30.03% in fifth

instar. Thus, honey positively reinforced the day-to-day larval growth rates by 4.75 percentile points (30.03 - 25.28) in fifth instar. Another noteworthy feature of silkworm development is that the larvae grow exponentially in early phases of instar and slowly at later phases (Table 1).

**Honey versus gland-body ratio:** The determination of gland- body ratio (GBR) by comparatively analyzing the changes in the weights of the larval body and silk gland during fifth instar development is considered as valuable indicator of silk production (Sailaja and Sivaprasad, 2010). Higher GBR could be achieved by enriching mulberry leaves with potential exogenous modulators and nutrients. The honey is identified as one such modulator. As shown in table 1, this exogenous factor has shown profound effect on GBR in *B. mori*. The data analyzed in terms of CPGR and overall percent change (OPC) clearly demonstrates the positive impact of honey on silkworm. Under the influence of 2% honey, the CPGR of body weight grew by 4.82 percentile points (68.0 - 63.18) while OPC of the SG weight grew



**Fig. 1.** Effect of honey on the silk gland-body ratio during fifth instar larval development in the silkworm, *B. mori*. The values, expressed in percentages, represent the compound periodical growth rates (CPGRs), computed on the basis of initial (Day-1) and final (Day-7) values of fifth instar.



**Fig. 2.** Overall effect of 2% honey on the protein profiles in the different segments of the silk gland in *Bombyx mori* during fifth instar development. The values, expressed in percentages, represent compound periodical growth rates (CPGRs), computed on the basis of initial (Day-1) and final (Day-7) values of fifth instar.

by 4.45 additional percentile points (2289 - 1844). At the same time the GBR grew by 6.64 percentile points (71.47-64.82) in terms of CPGR and by 56.45 percentile points (404.19-347.74) in terms of OPC (Fig.1). As reported earlier, the impact of honey on larval growth is attributable to its constituent vitamins and minerals (Bhattacharya and Kaliwal, 2005a, 2005b; Rajabi *et al.*, 2006a; Kavitha *et al.*, 2012).

**Honey versus silk proteins:** The SGP pool includes two silk proteins (Fibroin and sericin) and 91 other proteins involved in metabolism, immunity, heat-shock mechanism, cytoskeleton formation, protease inhibition transport and transcription (Nirmala *et al.*, 2001; Jin *et al.*, 2004; Takasu *et al.*, 2005; Kyung *et al.*, 2006; Zhang *et al.*, 2006; Hou *et al.*, 2007). Under the impact of 2% honey, the SGP profiles grew by 33.08 percentile points (144.86 - 111.78) during fifth instar development (Table 2, Fig. 2). Within the silk gland, ASG is relatively inert and shows low protein level, while MSG and PSG acts as protein reservoirs (Shimura, 1993). The study yielded interesting results and demonstrated that honey stimulates protein synthesis in all the three regions of silk gland but predominantly in ASG and PSG. The analysis of SGP in terms of overall percent changes (OPC) and compound periodical growth rates (CPGR) reveals this fact. The protein profiles of PSG grew maximally by 99.54 percentile points (OPC: 253.99-154.54) during fifth instar development showing an additional daily growth of 15.17 percentile points (CPGR: 52.40-37.23). The protein profiles of ASG, which were very low in the control batch, grew moderately by 63.23 percentile points (OPC: 102.38-39.15), with a daily growth rates of 14.85 additional percentile points (CPGR: 26.49-11.64) during fifth instar. At the same time the protein base of MSG grew by 59.07 percentile points (OPC: 271.90-212.83), with a daily average of 8.68 additional percentile points (CPGR: 54.93 - 46.25) (Table 2, Fig. 2). Needless to say, the higher protein gain in the three segments of

silk gland is attributable to the high nutritional status of honey in terms of its vitamin and mineral profiles (Hussain and Javed, 2002; Etebari and Matindoost, 2005; Thilsath *et al.*, 2008; Thulasi and Sivaprasad, 2013, 2014).

**Honey versus economic parameters:** The impact of honey on the economic parameters of sericulture is generally positive at its minimum effective concentration of 2% in distilled water. Its effectiveness is attributable to its stimulatory role on growth and silk protein synthesis during larval and pupal stages. Evidently, the positive impact of honey on the larval growth, observed during active fourth (~13%) and fifth instar (~25%) stages, has been extended to the quiescent pupal stage with 13.04% improvement in body weight and its 34% elevation in silk protein synthesis (Table 3). The profitable impact of honey on economic traits appears to be achieved by its two-fold impact on protein synthesis. Firstly, it acts as a potential stimulator of protein synthesis in all the three segments (ASG, MSG and PSG) of the silk gland. Secondly, its stimulatory effect is more specific on shell protein (Fibroin) compared to floss protein (Sericin). Probably, because of this reason, the shell protein levels recorded an increase of over 18% and the same was accompanied by a reduction in the levels of floss protein by ~25%, under the impact of honey. This dual effect in turn caused a rise in the raw silk weight and denier (thickness of silk fibre) by 18% each and a reduction in the renditta (the number of kilograms of cocoons required for production of 1kg of raw silk) by 2.5% (Table 3, Fig.3).

The major impact of honey on SG seems to enhance silk protein synthesis both in qualitative and quantitative terms, leading to improvement in the cocoon economic parameters. More importantly, the prevalence of higher protein levels in the MSG, indicates that silk gland proteins are predominantly synthesized in this region of silk gland. A comparative analysis of SGP vis-à-vis



**Fig.3.** Raw silk reeled from 100 green cocoons produced by *B. mori* in the control and honey treated conditions. Note significant increase in the output of raw silk and its colour under treatment conditions

economic parameters indicates that the honey modulates the tissue protein profiles in the silkworm, either by de novo synthesis or by sequestration from other tissues. Both these mechanisms are likely in *B. mori*.

### Conclusion

Honey is a potent modulator of growth, metabolism and silk production in *Bombyx mori*. Its potential could be realized at the minimum effective concentration of 2% in distilled water. For improvements in the economic parameters of sericulture, the application of 2% honey-enriched mulberry diet is recommended for silkworms. At this concentration, the honey not only stimulates silk protein synthesis in the silk gland, but also mobilizes appropriate protein reserves from fat body and fine tunes the silk output. In doing so, the honey confers two-fold advantage on sericulturists. Firstly, it enhances silk productivity and quality and secondly, it reduces floss output which is treated as the sericultural wastage. However, extensive field trials are recommended before its application in sericulture.

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