Cuticle Protein Homologue of the Chinese Oak Silkworm, Antheraea pernyi: cDNA Cloning and Expression

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We have cloned a cDNA encoding the cuticle protein homologue from the Chinese oak silkworm, Antheraea pernyi. In this paper, the cloning, sequencing and expression of a cDNA of A. pernyi cuticle protein homologue are described. The cDNA sequences were 447 bp in length, encoding 149 amino acid residues. The predicted molecular masses for A. pernyi cuticle proteins were approximately 16.4 kDa (ApCP16.4). The deduced amino acid sequences of the A. pernyi cuticle protein cDNA showed protein sequence identity to insect cuticle proteins known. Northern blot analysis revealed that the A. pernyi cuticle protein showed epidermis-specific expression. The expression profile of A. pernyi cuticle proteins revealed by Northern blot analysis that the high-level mRNA expression of A. pernyi cuticle proteins was detected on the first day of larval ecdysis and on the first day after larval-pupal metamorphosis. The cDNA encoding the cuticle protein homologue of A. pernyi was expressed as a 16.4 kDa polypeptide in the baculovirus-infected insect cells.