A remarkable, precisely timed release of hyperglycemic hormone from endocrine cells in the gut is associated with ecdysis in the crab *Carcinus maenas*

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Molting or ecdysis is the most fundamentally important process in arthropod life history, because shedding of the exoskeleton is an absolute prerequisite for growth and metamorphosis. Although the hormonal mechanisms driving ecdysis in insects have been studied extensively, nothing is known about these processes in crustaceans. During late premolt and during ecdysis in the crab *Carcinus maenas*, we observed a precise and reproducible surge in hemolymph hyperglycemic hormone (CHH) levels, which was over 100-fold greater than levels seen in intermolt animals. The source of this hormone surge was not from the eyestalk neurosecretory tissues but from previously undescribed endocrine cells (paraneurons), in defined areas of the foregut and hindgut. During premolt (the only time when CHH is expressed by these tissues), the gut is the largest endocrine tissue in the crab. The CHH surge, which is a result of an unusual, almost complete discharge of the contents of the gut endocrine cell, regulates water and ion uptake during molting, thus allowing the swelling necessary for successful ecdysis and the subsequent increase in size during postmolt. This study defines an endocrine brain/gut axis in the arthropods. We propose that the ionoregulatory process controlled by CHH may be common to arthropods, in that, for insects, a similar mechanism seems to be involved in antidiuresis. It also seems likely that a cascade of very precisely coordinated release of (neuro) hormones controls ecdysis.

In insects, precydal behavior, ecdysis, and subsequent associated events—such as air swallowing, hemolymph redistribution, and postecdysial behavior—are controlled by a complex, precisely timed interaction of several neuropeptides, including ecdysis-triggering hormone, eclosion hormone, and crustacean cardioactive peptide (1-6). Although analogous ecdysis-related events and behaviors must occur in crustaceans, nothing is known about the hormonal regulation of these processes in these arthropods. In spite of the large size of many decapod crustaceans in relation to other arthropods, which might be deemed a useful feature for physiological research, molting in most large, heavily calcified, adult decapod crustaceans is an annual event. Unlike insects, there are few obvious established behaviors or morphological events (7) useful in predicting ecdysis with the essential precision needed to elucidate the involvement of (neuro) hormones in this process. Nevertheless, during ecdysis in the green shore crab *Carcinus maenas*, we noticed increases in levels of circulating crustacean hyperglycemic hormone (CHH) immunoreactivity during the initial stages of ecdysis, culminating in a dramatic but transient (~10-30 min) hormone surge and declining to basal values on completion of ecdysis. Although this neuropeptide is well known to be involved in energy mobilization in crustaceans (8) and more recently was reported to be involved in inhibition of ecdysteroid and juvenoid biosynthesis (9-11), we reasoned that this pattern of CHH release, which is very precisely temporally correlated with ecdysis (rupture of the epimeral lines and swelling), must be important in some aspect of this process. Herein, we describe the temporal pattern of release of an authentic CHH, which is identical to that expressed by the eyestalk neurosecretory tissues from gut endocrine cells during ecdysis. The source of this hormone (previously undescribed paraneurons) and its action in stimulating water uptake at this time, which is necessary to allow ecdysis and subsequent swelling to postmolt dimensions, are also described.

Materials and Methods

Animals. Specimens of premolt *Carcinus maenas* were collected locally and maintained in a circulating seawater system under ambient conditions of temperature and photoperiod. Preparation for molt was staged by using the appearance and opening of the posterior lateral epimeral line and emergence from the old cuticle as visual markers. Other parameters for molt staging were based on established criteria (12).

Radioimmunoassay (RIA), Blood Chemistry, and Immunohistochemistry. RIA for CHH and CHH precursor-related peptide (CPRP) was performed by using 125I-labeled peptides and polyclonal antisera (raised in rabbits by using HPLC-purified sinus gland peptides or, in the case of CPRP, by using a bovine thyroglu- 

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Abbreviations: CHH, crustacean hyperglycemic hormone; RIA, radioimmunoassay; CPRP, CHH precursor-related peptide; MALDI-TOF, matrix-assisted laser desorption ionization time-of-flight.

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quantities of RNA were reverse transcribed by using AMV reverse transcriptase (Roche Molecular Biochemicals) and cDNA amplified by using the following CHH gene-specific primers: forward 5'-GCCATGCTAGCAATCATCACGAG-3' and reverse 5'-GTTGAGATCTGTTGTTTACTCC. After denaturation at 94°C for 4 min, PCR was done for 30 cycles consisting of denaturation at 94°C for 1 min, annealing at 55°C for 1 min, and extension for 72°C for 2 min. The amplification was terminated by a final extension at 72°C for 7 min. For a positive control, RNA extracted from the medulla terminalis of the eyestalk was used. Products were electrophoresed on 1.2% agarose gels with ethidium bromide visualization. PCR products of expected size (421 bp) were purified (Hybaid PCR purification kit, Middlesex, U.K.) and cloned in Top 10F.

HPLC and Mass Spectrometry. Sep-Pak purified hemolymph and gut extracts were fractionated by HPLC by using the following conditions: 300 × 3.9-mm Phenyl column (Waters), 1 mL min⁻¹ gradient elution, 30–80% solvent B over 1 h [solvent A was 0.1% trifluoroacetic acid; solvent B was 60% (vol/vol) acetonitrile/0.1% trifluoroacetic acid]. Matrix-assisted laser desorption ionization time-of-flight (MALDI-TOF) mass spectrometry was performed on a Micromass TosfEc-E instrument (Micromass, Manchester, U.K.).

Radiography. Molt-staged crabs (beginning of ecdysis: stage E1) were injected with CHH (100 pmol) or saline (controls) and immersed for 30 min in a stirred 20% (vol/vol) barium sulfate suspension in seawater, followed by rinsing and radiography, (X-protector, PLH Medical, Watford, U.K.) for 0.4 s by using Agfa CURIX PR1 x-ray film.

Results

During late premolt (stage D3), levels of circulating CHH rapidly increased from typically low levels observed during intermolt (1–5 fmol/100 µl) to around 25 fmol/100 µl. During ecdysis (stage E1), a consistent pattern of very dramatically increased levels of CHH was observed, with peak levels (150–200 fmol/100 µl) occurring during the latter stages of ecdysis (E30–100). Immediately (within 1 h) after ecdysis, levels of CHH returned to basal values (Fig. 1A). With regard to the general time course of ecdysial, the duration of stage D3 is 4–24 h; that of initial swelling E1–30 is 200–500 min (in experimentally manipulated animals; see Fig. 4); E30 to completion of ecdysis lasts 15–20 min. The source of CHH during this hormone surge was not from the eyestalk neurosecretory tissues, because animals whose stalks had been removed had identical patterns of CHH titer during ecdysis (and molted viably). Screening a variety of tissue extracts (including brain, stomatogastric ganglia, thoracic ganglia, gills, gonads, muscle, hepatopancreas, and gut) for CHH immunoreactivity during premolt revealed that the gut exclusively synthesized large quantities of CHH immunoreactive material during premolt. Immunoreactive content declined rapidly just before the dramatic increase in immunoreactive material in the hemolymph (Fig. 1A).

The immunoreactive material in the hemolymph during ecdysis (in crabs with and without eyestalks) was deemed to be authentic CHH on the basis of cochromatography by HPLC, of parallel dilution characteristics in RIA, and of mass determinations (MALDI-TOF) of HPLC-purified material eluting at the same time as sinus gland-derived CHH, which gave molecular ions corresponding to the Na⁺ (8,545 Da) and K⁺ (8,561 Da) salts of CHH. Additionally, CPRP, a 38-residue polypeptide of unknown function, which is part of the preprocessed CHH molecule (15, 16), was also detected in the hemolymph during the CHH surge.

The immunoreactive material in gut extracts was deemed authentic by cochromatography (HPLC) with sinus gland-derived CHH (Fig. 2A and B) and by MALDI-TOF [molecular ions corresponding to the Na⁺ (8,545 Da) and K⁺ (8,561 Da) salts were identified from HPLC purified premolt gut extract eluting at the retention time of CHH]. Reverse transcription–PCR of premolt gut total RNA with gene-specific primers based on the prepro-CHH nucleotide sequence (15) gave an amplified product identical in size to that obtained from eyestalk tissue (Fig. 2C). Sequencing of the product revealed complete identity with the nucleotide sequence and conceptual amino acid sequence of prepro-CHH from the eyestalk and CPRP (15, 16), which were identical in these respects to one of the three CPRPs produced by the eyestalk neurosecretory system of Carcinus (J.S.C. and S.G.W., unpublished results). During ecdysis, levels of glucose and lactate increased only marginally and fell to intermolt levels within 24 h of molting (Fig. 1B).
Whole-mount immunohistochemistry of Carcinus guts revealed many thousands of endocrine cells, which were immunopositive to antisera raised against both CHH and CPRP (Fig. 3A–D). These cells were found only in the chitin-lined foregut and hindgut; cells were never observed in the midgut or associated diverticulae. Large numbers of cells were associated with the anterioventral and posteriolateral margins of the pyloric ampullae and were most abundant in gut epithelia adjacent to muscle insertions, particularly the external and internal inferior pyloric dilator muscles (17). In the hindgut, the immunoreactive cells were scattered but appeared mainly confined to infoldings of the gut lumen adjacent to the longitudinal muscles. The cells had a very distinctive morphology, including an extensive array of arborizations abutting the hemocoelic spaces and a narrow (<10 μm) tapering cell body terminating at the chitinous surface of the gut. The nuclei were invariably located at the widest part of the cells adjacent to the arborizations.

To determine the function of the CHH hormone surge during ecdysis, crabs at very precisely staged periods of ecdysis were injected with CHH (100 pmol). This procedure resulted in a marked acceleration of ecdysis, the magnitude of which was related to the stage at which animals were injected (Fig. 4).
example, injection at the beginning of ecdysis (stage D_7–E_0) reduced the mean time taken to complete ecdysis from 500 min to 140 min. Injection of CHH during later stages of ecdysis (E_1) reduced the mean time to ecdysis from 60 min to 10 min. If larger doses of CHH were injected, ecdysis was always unsuccessful, judging by the fact that the legs of the crabs became trapped in the exoskeleton because of a very marked and abnormal swelling. Radiography of crabs injected with CHH (Fig. 4) revealed that hormone-induced swelling was associated with accumulation of barium sulfate in the stomach and anterior hindgut, indicative of drinking and water absorption. Radiopaque dark lateral areas represent barium sulfate trapped in gill tissues. This material was difficult to remove in control crabs, whereas, in hormone injected crabs, the impending molt allowed efficient rinsing of the gills.

**Discussion**

In this study, we have shown that ecdysis in *Carcinus maenas* is associated with a precisely timed surge in circulating CHH levels, which is a result of an unusual, almost complete release of this hormone from a previously undescribed endocrine tissue, namely the gut. The endocrine cells in the foregut and hindgut responsible for this hormone surge possess a very similar morphology to previously described endocrine cells of the insect gut (18, 19) or the “closed” type of paraneuron described by Fujita et al. (20), although these cells clearly differ from previously described “putative” endocrine cells from the crustacean midgut (21). Furthermore, no immunopositive cells were seen in this region of the gut. In view of the very distinctive morphology of these cells, reminiscent of the habitus of *Hydra*, we propose that they be called “hydroid cells.” During intermolt, these cells are not visible by immunohistochemical techniques but, during premolt (in accord with the RIA results), the cells become strongly immunoreactive. However, after molting, very little immunoreactivity was observed, except for some cell debris, which suggested that the CHH surge was a result of almost complete secretion of peptide contents, if not cell death. In this context, it is of interest to note that complete secretion of cellular peptide content has been observed in insect (nonneuronal) epitracheal “Inka cells” after the ecdysis-triggering hormone surge before eclosion (1) and the dramatic release of eclosion hormone from proctodeal nerves before eclosion (22). Thus, parallels might perhaps be drawn concerning this unusual mode of complete peptide release for functionally critical hormones during ecdysis in the arthropods.

What is the function of the CHH surge? Because ecdysis is very energy demanding and possibly stressful, an obvious role for CHH at this time might be in regulation of stress-related hyperglycemia (8). However, during ecdysis, blood glucose levels show only a moderate increase, and hypoxia (as evidenced by increased blood lactate levels) is slight (Fig. 1B). Because ecdysis in crustaceans depends on a very impressive uptake of water via the gut (23, 24), it seemed possible that the CHH surge was in some way linked to this phenomenon. When crabs were injected with physiologically relevant (100 pmol) quantities of CHH before the expected endogenous CHH surge, we observed a precocious initiation of significant uptake of water (swelling), which resulted in accelerated ecdysis, relative to saline-injected controls (Fig. 4). When higher levels (300 pmol) of CHH were injected, swelling was very rapid; in addition, molting was unsuccessful, because the crab became trapped in the old exoskeleton of the legs because of the enormous tissue expansion.

At present, we do not know the precise mechanisms by which CHH increases water uptake. Increased drinking is involved, as evidenced from barium meal radiographic experiments (Fig. 4). An ionoregulatory action of CHH is also plausible. Because seawater drinking is inevitably associated with preecdysial swelling, which occurs in an organism essentially isosmotic with its external environment, active uptake of salts across the gut epithelia followed by passive movement of water seems likely. It has long been known, from ultrastructural and physiological studies, that the hindgut and midgut are highly osmotically active (23) and that these tissues are predominantly responsible for water and ion uptake during molting (24). It is known that in locusts, hindgut (ileal) chloride transport is stimulated by an ion transport peptide, which, by activating electrogenic Cl^- pumps, can drive isosmotic fluid reabsorption by the ileum (25–27). Because this peptide (which was the first insect member of the CHH peptide family to be isolated) is highly structurally related to *Carcinus* CHH (39% sequence identity), an attractive hypothesis would be one involving similar ionic mechanisms that drive (isosmotic) water uptake by the hindgut in crustaceans. The CHH-containing cells of the foregut and hindgut seem ideally placed to detect changes in gut tension. In particular, the cells at the anteriointestinal margin of the pyloric ampullae, associated with the external and internal inferior pyloric dilator muscles, may be influenced (mechanically) by changes in the pyloric motor rhythm at ecdysis. In lobsters (*Homarus gammarus*), a deep preecdysial hypoxia modulates motor network expression.
of the stomatogastric ganglion (28). Our preliminary results on changes in oxygen tension in the hemolymph in *Carcinus* during ecdysis (J.S.C. and S.G.W., unpublished results) and the rather moderate increases in blood lactate seen at this time (Fig. 1B) suggest that deep preecdysial hypoxia does not occur in *Carcinus* (as would befit a well adapted intertidal crustacean). However, it is entirely possible that a mechanically stimulated release of CHH from these cells during ecdysis could act locally on the gut and centrally in circulation. Indeed, perfusion of the circulatory system with India ink to reveal the fine structure of the arterial circulatory system has shown that the pyloric ampullae, in particular, have a very impressive arterial microcapillary blood supply (J.S.C. and S.G.W., unpublished results). If the proposed scenario were correct, then, strikingly, a common mechanism would control disparate functions: antidiuresis (insects) and ecdysial swelling (crustaceans). Nevertheless, CHH is certainly not the only neuropeptide that is involved in crustacean molting. Crustacean cardioactive peptide (29) is intimately involved in later stages of ecdysis controls stereotyped ecdysial behavior in crustaceans (31, 32). Additionally, because expression of CHH in gut endocrine cells during premolt closely follows increases in ecdysteroid levels during this period, it is likely that activation of transcription of CHH genes in these cells may be mediated by ecdysteroids. Obviously, the early (swelling) and subsequent (behavioral) processes during ecdysis must be tightly temporally regulated to culminate in successful molting. To paraphrase the statements of Truman (2) and Hesterlee and Morton (31), many more layers of complexity, which remain to be shed, still shroud the remarkable phenomenon of molting in crustaceans.

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