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Abstract.—Analysis of surface and subsurface plankton collections in the Middle Atlantic Bight (MAB) yielded larvae and juveniles of Phycis chesteri and five species of Urophycis. Identification was based on numbers of epibranchial gill rakers, abdominal vertebrae, and fin rays (dorsal, caudal, pelvic), patterns of pterygiophore interdigitation, and morphometric characters including body depth at the vent and a ratio between height of the pelvic-fin base and length of the mandible. Urophycis tenuis accounted for 99% of the Urophycis larvae and pelagic juveniles collected during spring off Virginia and New Jersey and was most abundant offshore. Urophycis tenuis larvae were smallest at offshore stations and increased in size as collections proceeded shoreward. Urophycis chuss was found in summer and fall collections off the coasts of New Jersey and Virginia, with abundances highest at midshelf stations. Urophycis chuss was the only species of hake found during August and early September, and it dominated summer ichthyoplankton collections. Urophycis regia was found primarily in midshelf areas off Virginia during fall, but was also collected offshore from both Virginia and New Jersey during winter. Phycis chesteri, also found in fall and winter collections, was restricted to offshore stations. Southern species, found exclusively in offshore winter collections, included U. floridana and U. cirrata.

Identification and distribution of Urophycis and Phycis (Pisces, Gadidae) larvae and pelagic juveniles in the U.S. Middle Atlantic Bight*

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Species of the gadid genera *Urophycis* (Gill) and Phycis (Artedi), collectively referred to as 'hakes', are abundant on the continental shelf and slope of the northwest Atlantic Ocean. Six species of *Urophycis* and one species of Phycis are found in this area (Svetovidov 1948, Wenner 1983): U. tenuis (Mitchill), U. chuss (Walbaum), U. regia (Walbaum), U. floridana (Bean and Dresel), U. earlli (Bean), U. cirrata (Goode and Bean), and P. chesteri (Goode and Bean). Larval hake are present at all times of the year in the Middle Atlantic Bight (MAB) and dominate summer plankton collections (Comyns 1987), but persistent taxonomic problems (Dunn & Matarese 1984) have hindered the accumulation of ecological data on these important components of offshore ichthyoplankton communities (Kendall & Naplin 1981, Hermes 1985).

Newly hatched *U. chuss* and *U.* regia of known parentage were described by Hildebrand & Cable (1938), Miller & Marak (1959), Barans & Barans (1972), and Serebryakov (1978). Although these sources describe pigmentation differences between the two species, this information alone is insufficient to positively identify field-caught larvae.

Older larvae and juveniles of U. chuss, U. regia, U. floridana, and a single iuvenile specimen of *U. earlli* were described by Hildebrand & Cable (1938). Larvae and juveniles of U. regia were collected off Beaufort NC and were identified by the presence of relatively few second dorsal-fin rays and lack of dark ventral-fin pigment. A second larval morph collected off Beaufort was tentatively identified as U. floridana because adult *U. floridana* was the only other species of Urophycis commonly found in the collection area. These specimens differed from U. regia in having darkly-pigmented ventral fins and more second dorsal-fin rays. A single juvenile specimen (37 mm) collected off Beaufort was identified as *U. earlli* because this specimen was darker than U. regia and U. floridana, and possessed smaller scales. Specimens of a fourth morph, collected off Cape Henry VA were identified as U. chuss because they possessed dark ventral-fin pigment, were relatively slenderbodied, and it was assumed that

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U. floridana would not be found as far north as Cape Henry.

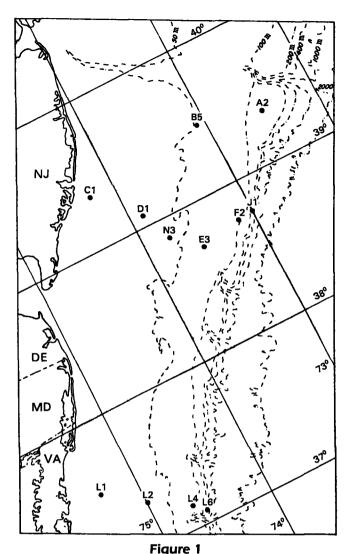
Methven (1985) presented a size-dependent key to the identification of young *U. chuss*, *U. tenuis*, and *P. chesteri* from the Northwest Atlantic. Identifications were based on body depth, numbers of epibranchial gill rakers (Musick 1973, Wenner 1983), and numbers of caudal-fin rays. Material for Methven's study came primarily from Canadian waters, and he did not encounter *U. cirrata*, *U. earlli*, *U. floridana*, or *U. regia*. As a result, Methven's key is of limited use in more southerly locations where these species occur.

The objective of this paper is to describe additional morphometric and meristic characters that aid in the identification of *Urophycis* and *Phycis* larvae, and to describe the spatial and temporal distribution of these larvae collected 1975–77 off Virginia and New Jersey in the Middle Atlantic Bight.

Materials and methods

Sampling locations and shipboard procedures

Sampling extended from October 1975 until August 1977 and was conducted quarterly at 12 stations off Virginia and New Jersey (Fig. 1, Table 1). Neuston samples were collected with a floating sampler developed at Woods Hole Oceanographic Institution (Bartlett & Haedrich 1968, Craddock 1969). The net, constructed with 505 um mesh Nitex, was 1 m wide and fished to a depth of 12cm in calm seas. Tows were of 20 min duration at a ship speed of ~2 kn. The net was deployed from a boom and the towing course followed a widely-circular track to prevent sampling in the ship's wake. A single neuston tow was made at 3h intervals over a 24h period at each station, resulting in eight samples per station during each cruise. Two oblique tows between nearsurface and bottom were made at all stations with 60 cm openingclosing bongo systems (McGowan & Brown 1966), the first with paired 202 µm Nitex nets and the second with paired 505 µm nets. To prevent surface contamination, all nets were closed during passage through the surface layer (upper meter). Both bongo and neuston nets were equipped with flowmeters (General Oceanics, Inc.). The flowmeter attached to the underside of the neuston frame provided an estimate of horizontal distance relative to sea surface fished by the net. Estimates of volume filtered by the neuston sampler were determined by multiplying distance fished by net area fished (1 m×12 cm). In calm seas the neuston sampler consistently fished to a depth of 12 cm, but in rough seas the net opening would occa-



Ichthyoplankton sampling locations off New Jersey and Virginia. Stns. F2, J1, A2, L4, and L6 are considered offshore stations.

sionally be almost completely filled or empty (the flowmeter was always submerged). This variability decreased precision of neuston volume estimates but was not expected to bias volume estimates. Comparisons between neuston and bongo collections were performed only to emphasize the relative importance of the surface layer to larval hakes. Patterns of the spatial and temporal distribution of larvae were based only on comparisons among neuston collections because most specimens were collected with this gear type.

Laboratory procedures

Fish larvae were sorted from whole collections. All specimens of *Urophycis* and *Phycis* were cleared and stained (Dingerkus & Uhler 1977, Potthoff 1984, Tay-

Table 1
Sampling schedule for twelve stations occupied off New Jersey (NJ) and Virginia (VA), 1975–77.

	1975		19'	76	1977				
Stn.	Oct	Feb	Jun	Aug Sep	Nov	Feb Mar	May	Aug	
C1 (NJ)	X	X	X	X	X	X	X	х	
D1	X	X	X	X	X	X	X	X	
N3	X	X	X	X	X	X	X	X	
E3	X	X	X	X	X	X	X	X	
F2	X	X	X	X	X	X	X	X	
J1	X	X	X	X	X	X	X	X	
B5 (NJ)					X	Х	X	X	
A2					X	X	X	X	
L1 (VA)					X	X	X	X	
L2					X	X	X	X	
L4					X	X	X	X	
L6					X	X	X	X	

lor & Van Dyke 1985), except those occurring in collections taken during August–September 1976 (*n*>16,000) and August 1977 (*n*>4000). During these periods of high abundance, subsamples of over 2000 larvae from August–September 1976 and >900 larvae from August 1977 were randomly selected and similarly processed. Herein specimens <18 mmSL are arbitrarily termed larvae, whereas fish ≥18 mm are termed juveniles (Markle et al. 1982). Fish <~12 mm were measured with an ocular micrometer, while lengths of larger specimens were measured with a dial caliper ruler. The largest pelagic juveniles found were ~40 mmSL.

The following morphometric criteria were used in the analysis: (1) height of pelvic fin/vertical distance from base of pelvic fin to ventral margin of body; (2) mandible length/distance from anterior tip of the dentary to posteroventral tip of the angular; and (3) body depth at anus/vertical distance from anterior end of anal-fin base to dorsal surface immediately above. Morphometric measurements were made with an ocular micrometer. The first interneural space was defined as the space anterior to the first neural spine.

Hake larvae and juveniles possessing the adult meristic complement were initially identified using published and unpublished meristic data (Table 2). Meristic observations included epibranchial gill rakers (left side examined), abdominal vertebrae, and fin rays (dorsal, caudal, pelvic). Observations were taken from both cleared and stained material and from radio-

graphs of juvenile and adult museum specimens (App. Table 1).

Identification of smaller larvae was facilitated by using morphometric criteria, patterns of interdigitation between pterygiophores supporting the median fins and the neural or haemal spines, and by defining the size at which larvae attained various stages of fin-ray, vertebral, and gill-raker development. Unfortunately, faded pigmentation caused by specimen storage in formalin and subsequent clearing and staining prevented use and further description of larval pigmentation in the present study.

Table 2

Ranges of meristic characters in *Phycis chesteri* and six species of *Urophycis*. Numbers in parentheses indicate meristic ranges that were extended by the present study. Data sources are (1) Svetovidov 1948, (2) Hildebrand & Cable 1938, (3) Bigelow & Schroeder 1953, (4) Leim & Scott 1966, (5) Miller & Jorgenson 1973, (6) Musick 1973, (7) Hoese & Moore 1977, (8) Markle 1982, (9) Fahay 1983, (10) Wenner 1983, (11) Methven 1985, (12) J.A. Musick, pers. commun., VIMS, Gloucester Point VA 23062.

	U. tenuis	U. chuss	U. regia	U. floridana	U. earlli	U. cirrata	P. chesteri
Caudal-fin rays	33–38(40)	28–34	28-32(34)	28-32(34)	27-30(31)	28–33	28–35(36)
1st dorsal-fin rays	9-10(12)	9-11(12)	8–10	10-13	8–11	9-11(12)	8-11(12)
2nd dorsal-fin rays	50-59(62)	(52)53-64	43-51(52)	54-63	57-63(68)	54–68	50-63
Anal-fin rays	41-52(53)	45-57	41-50(52)	45-54(55)	49-56(60)	46-58	43-54
Vertebrae (total)	47-50(51)	45-50(51)	(44)45-47(48)	44-50(51)	45-47(48)	47-53	45-52
Caudal vertebrae	(32)34-35	33-36	(30)31-33(34)	30-34(35)	31-33(34)	32-37	31–37
Abdominal vertebrae	*13-17	14-17	13–15	14–17	14–15	15-17	13–16
Pelvic-fin raysb	3	3	3	3	3	3	3
Epibranchial gill	°2	3	3	2	2	3	4–5
Rakers (1st arch)							
Data source	1,3,6,8,11	1,2,4,5,6,8,11	1,2,3,4,5,8	1,7,12	1,5,12	1,12	1,10,11,12

^a In our material (n=205) *U. tenuis* never possessed <15 abdominal vertebrae.

^b The third pelvic-fin ray in adult *Urophycis* and *Phycis* is rudimentary.

^{&#}x27; Urophycis tenuis occasionally has three epibranchial gill rakers.

Results

Identification of *Urophycis* and *Phycis* larvae and juveniles

Meristics

Epibranchial gill rakers (Table 3) A complete sizeseries of all species was not available, but size (mm) at which U. regia, U. chuss, and U. tenuis larvae attain the adult complement of epibranchial gill rakers and other meristic elements is shown in App. Table 2. The following sections are abbreviated to avoid repeating in the text what the tables and figures succinctly show. Phycis chesteri does not attain the adult complement of epibranchial gill rakers until 16-18 mm (Methven 1985), but by 13 mm the third gill raker has developed and serves to separate larvae of this species from U. tenuis, U. earlli, and U. floridana. Occasionally U. chuss and *U. regia* possess two or four epibranchial gill rakers on one side, but most of these specimens have the normal complement of three gill rakers on the other side. Urophycis tenuis occasionally possesses a 3rd epibranchial gill raker, but only very rarely is this third gill raker found on both sides of a specimen.

Caudal-fin rays (Table 3) All but two specimens of *U. tenuis* (n=195) had more caudal-fin rays than all other species of *Urophycis*. Numbers of caudal-fin rays of *U. tenuis* overlapped those of *P. chesteri*, but more than half of our *U. tenuis* were distinct in having >36 rays, and over 40% of *P. chesteri* (n=56) differed from *U. tenuis* in having <34 rays. As few as 28 caudal-fin rays have been reported in *P. chesteri* (Wenner 1983) and *U. cirrata* (J.A. Musick, VA Inst. Mar. Sci., pers. commun.), but this may be because some of the small procurrent rays are not easily seen in radiographs of larger fish.

No U. earlli specimens (n=31) possessed >31 caudal-fin rays, while all other hake commonly have >31 rays.

Dorsal-fin rays (Table 3) Despite overlapping extremes, numbers of first dorsal-fin rays helped distinguish *U. floridana* from other species of hake. In our material, *U. regia* and *U. earlli* never possessed >10 and 11 first dorsal-fin rays, respectively, while over 80% of *U. floridana* (n=45) had >11 rays. One-third of *U. floridana* specimens examined possessed 13 first dorsal-fin rays, delimiting these from all other hake taxa.

The relatively low number of second dorsal-fin rays in *U. regia* separated this species from *P. chesteri* and other *Urophycis* species with little overlap. *Urophycis chuss* and *U. regia* with incomplete development of second dorsal-fin rays were delimited by numbers of pterygiophores supporting these rays at sizes as small as 6 mm (Fig. 2). Although extremes in numbers of

second dorsal-fin rays overlapped in all other taxa, many of the available specimens of *U. earlli* and *U. cirrata* were distinct in possessing >63 rays.

Abdominal vertebrae (Table 3) Numbers of abdominal vertebrae cannot be used alone to identify individual specimens because of overlapping extremes, but this meristic character is useful when identifying collections comprised entirely of *U. tenuis* or *U. chuss*. Urophycis tenuis larvae, identified by numbers of epibranchial gill rakers and caudal-fin rays, were found in the MAB only in spring and accounted for 99% of the *Urophycis* collected at this time (*U. regia* juveniles accounted for the other 1%). Urophycis larvae <10 mm (n=154) that were present in spring collections had not yet developed the adult complement of caudal-fin rays, but these larvae (>4 mm) had developed the adult complement of abdominal vertebrae and were identified as *U. tenuis* because their frequency-distribution of numbers of abdominal vertebrae was identical to that found in larger U. tenuis; 88% of the larvae had 16 abdominal vertebrae, and no specimens were found with <15. It is unlikely that any of these small larvae were U. floridana or U. cirrata, two other species with similar numbers of abdominal vertebrae, because these two southern species were found only in offshore winter collections and most specimens were pelagic juveniles.

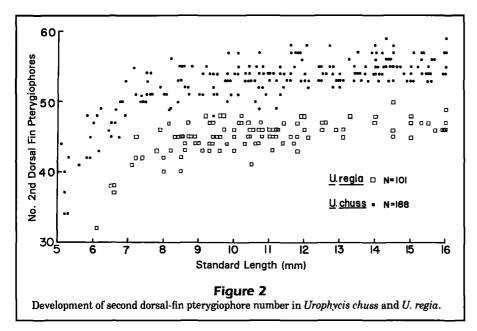
Urophycis chuss >4 mm (n=448) possessed 14–16 abdominal vertebrae, but the majority of specimens (n=391) had 15. In all other species of *Urophycis* the count of 15 occurred in <20% of the specimens; and although extremes of *U. chuss* and *P. chesteri* were similar, P. chesteri commonly had 14 or 16 abdominal vertebrae (17%). Consequently, in the MAB during late summer when U. chuss larvae are extremely abundant (and, in this study, were the only species of hake found at this time), complete meristic counts to check for species other than *U. chuss* need be performed only on those specimens that do not have 15 abdominal vertebrae. If species other than *U. chuss* are found in late-summer collections, numbers of abdominal vertebrae are no longer taxonomically useful and complete meristic counts are necessary to identify larvae.

Urophycis regia (n=698) had 13–15 abdominal vertebrae, but only eight specimens had 15, and seven of these specimens had an anomalous 15th abdominal vertebra. This anomalous vertebra possessed one short transverse process characteristic of abdominal vertebrae and one long transverse process typical of caudal vertebrae. Because 99.9% of U. regia examined had <15 normal abdominal vertebrae, it was assumed that specimens with \geq 15 abdominal vertebrae were not U. regia. This meristic character aided in the separation of small (<6 mm) U. chuss and U. regia in fall collec-

Table 3

		Epi	branc	hial	gill r	aker	s on	left fi	irst g	gill ar	ch. S	lash	sepa	rate	s nu	mber	s on	left a	nd r	ight	sides				
		•		2	-		3/2			3			4/3			4			5	-					
P. chesteri U. chuss U. regia U. cirrata U. tenuis				16			8 4 6			596 631 19			8 4			24 2 2	***		8						
I. floridana I. earlli				3:	4	_																			
									Nu	mber	of ca	audal	-fin	rays											
	29		3	0	3	1		32		33		34		35		36		37		3		3			10 1
J. tenuis P. chesteri J. regia J. chuss J. cirrata J. floridana J. earlli	1		1 1 5 13		1 2 1	1 9 2 3 3]	8 34 13 8 21		15 16 10 2 14		2 19 1 3		28 10		56 3		65	1	3	4	•	,		1
							-	1	Vum	ber of	firs	t dor	sal-fi	n ray	/s					_				-	
			_	8			9	_		10			11			12			13						
U. floridana U. cirrata U. chuss U. tenuis P. chesteri U. earlli U. regia				1 14			2 10 19 6			2 33 24 38 21 7			8 51 26 13 5			22 3 9 3 2			15						
								N	umb	er of a	secor	nd do	rsal-	fin ra	ıys	-									
U. cirrata U. earlli U. floridana P. chesteri U. chuss U. tenuis U. regia	2	45 1	46 15	22	30	49 36	50 29	51 1 16	1 3 2	3 3 9	8 4 7	55 6 5 6 8	6 14 16 3	57 1 7 13 16 12	58 9 5 16 5	59 1 6 5 20 4	2 5 3 9 1	61 8 3 1 6 1	62 1 8 3 6 2	1 3 1 3	64 2 7	2 1	5 1	67	1 1
								N	umb	er of	abdo	mina	al ve	rtebr	ae										
						13		1	4		1	5		10	6		17								
U. tenuis U. cirrata U. floridana U. chuss P. chesteri U. earlli U. regia Jote: Although ormally develo	ı 8 sp	ecin	nens	of <i>U</i>		66 ia h	ad 18	5 62	8 7 4	nal v	39 5	2 7 1 7 4 8	in (0 8 7 4	of th	5 1 3	speci	men	в (О.	1%)	was :	the 1	լ5 ^{ւհ} ՝	verte
					Nui	nbei	of a	nal-fi	n pte	erygio	pho	res a	nteri	or to	first	hae	mal s	pine						-	
		_		3			4			5			6				7			8			9		_
U. earlli U. floridana U. regia U. chuss U. tenuis U. cirrata P. chesteri				3			3 1 28			8 18 36 8 34			150 173 23	3		2	14 23 02 52 1			8 7 14 7			1		
	Inte	rneı	ıral s	pace	into	whi	h pro	jects	the	ptery	/giop	hore	supj	porti	ng th	e fir	st ray	of t	he se	conc	dors	sal fir	ı		
					7				8				9				10								
U. tenuis U. floridana P. chesteri U. chuss U. regia					1 3 4 141			2	8 10 31 203 41				52 25 20 20				6 6 1 4								

Note: If pterygiophore was aligned with the tip of a neural spine, it was arbitrarily recorded as pointing into the space posterior to the spine in question.



tions when numbers of second dorsal-fin pterygiophores were not yet taxonomically useful.

Numbers of abdominal vertebrae may help separate *U. earlli* from *U. floridana* and *U. cirrata*, the other two southern species of *Urophycis*. Over 80% of *U. floridana* and *U. cirrata* possessed 16 or 17 abdominal vertebrae, but *U. earlli* has never been recorded with this many.

Anal-fin pterygiophores (Table 3) The number of anal-fin pterygiophores positioned anterior to the first haemal spine helps distinguish *P. chesteri*, *U. cirrata*, and *U. tenuis* from *U. earlli*, *U. floridana*, *U. regia*, and

U. chuss. Only one specimen of U. tenuis (n=65) and no U. cirrata (n=11) or P. chesteri (n=70) were found with >7 analfin pterygiophores positioned anterior to the first haemal spine. hut. 45% U. chuss (n=350) and over half of U. earlli (n=27), U. floridana (n=42), and *U. regia* (n=380) had at least 7 of these pterygiophores. More than 60% of U. tenuis. U. cirrata, and P. chesteri had <6 anterior anal-fin pterygiophores, whereas <2% of *U. regia* and no U. earlli or U. floridana had this few.

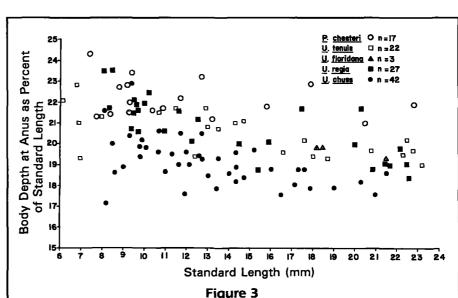
Second dorsal-fin pterygiophores (Table 3) The interneural space into which points the first pterygiophore of the second dorsal fin helped separate U. chuss from U. regia, and U. floridana from U. earlli. In over half of U. chuss examined (n=431) the first pterygiophore of the second dorsal fin pointed into the 9th or 10th interneural space, whereas in all U. regia examined (n=182) this pterygiophore pointed into the 7th or 8th interneural space. In >75% of U. regia this pterygiophore pointed into the 7th interneural space, whereas <1% of U. chuss showed this pattern.

In >70% of U. floridana examined (n=42) the first pterygiophore of the second dorsal fin pointed into the 9th or 10th interneural space, whereas in all juvenile and adult U. earlli examined (n=27) this pterygiophore pointed into the 7th or 8th interneural space. In

over half of *U. earlli* examined, the first pterygiophore of the second dorsal fin pointed into the 7th interneural space, but in only 2% of *U. floridana* did this pterygiophore project this far forward.

Morphometrics

Body depth at anus (Fig. 3) Body depth at the anus separated some species of hake larvae at sizes >12-13 mm. Extremes of body depth as percent of standard length for cleared and stained *P. chesteri*, *U. tenuis*, and *U. chuss* were 21.0-23.4, 19.0-21.1,



Body depth at anus as a percent of standard length plotted against standard length for larvae and juveniles of *Phycis chesteri* and four species of *Urophycis*.

Ranges of pelvic-fin-base he Urophycis. Ranges of values		rcent of ma					e species of
			Size	-interval (mm)		
	5–9	10–14	1519	20–24	25–29	30–34	35–45

				Size	:-interval (mm)		
		5–9	10–14	1519	20–24	25–29	30–34	35–45
U. regia	(n=31)	21–30	19–33	19–25	16–28	12-17	ND	ND
U. floridana	(n=19)	ND	ND	29-36	23-37	23-28	24-29	ND
U. chuss	(n=38)	20-39	23-33	24-36	19-22	15-16	16	16
U. cirrata	(n=4)	ND	ND	ND	39	31	ND	19-31
P. chesteri	(n=29)	44-74	52-61	54-61	46-61	52-64	50-59	26-57
U. tenuis	(n=39)	28–42	24–42	33–40	29–37	26~30	32	ND

and 17.6-19.7, respectively. Body depth of *U. floridana*, however, was found to overlap extremes of *U. tenuis* and *U. regia*, while *U. regia* exhibited the greatest variation in this character, overlapping the extremes of *P. chesteri* and all other species of *Urophycis* studied.

Mandible length and height of pelvic fin (Table 4, Fig. 4) Height of the pelvic fin plotted against mandible length separated larval *P. chesteri* from other hake at sizes between ~6 and 35 mm. At sizes > 3 5 m m m. P. chesteri was similar to Urophycis with respect to this character because *P. chesteri* became more slender-bodied and the pelvic-fin origin moved closer to the ventral margin of the body. Ranges of pelvic-fin height as percent of mandible length in cleared and stained larvae ranging in length from 6 to 35 mm varied from 44 to 74% in *P. chesteri* (n=29), but the highest value of this ratio in five species of Urophycis (n=131) was only 42%.

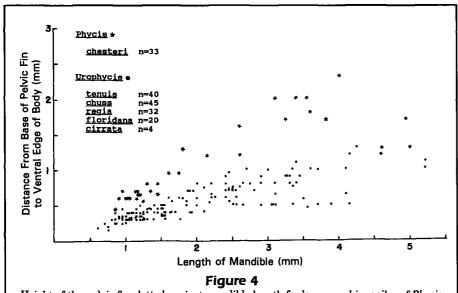
Distribution and abundance of hake larvae

Urophycis chuss Urophycis chuss was found only in summer and fall plankton collections from the MAB, and was the only species of larval hake found in August and early September. Densities of *U. chuss* in summer collections off the coast of New Jersey were up to two orders of magnitude greater than densities found off Virginia (Fig. 5). In October 1975 and November 1976, *U. chuss* were still present off both Virginia and New Jersey, but were far less abundant than during summer.

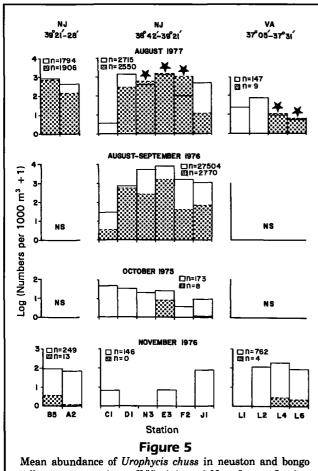
Densities of larval *U. chuss* also varied with distance from shore (Fig. 5), particularly during summer when lowest densities occurred inshore and highest densities were found in midshelf regions in water depths of 40–120 m. Variations in larval density with both latitude and water depth were not well defined in f a l collections.

An increase in mean size of U. chuss was evident in fall collections (Fig. 6). As larval size increased in fall collections, the number of larvae collected with bongo nets decreased greatly. More than 1300 specimens were collected in October 1975 and November 1976, but only 25 of the larvae were collected with bongo gear. Onshore-offshore variation in size of *U. chuss* was most evident in fall collections off both Virginia and New Jersey; size tended to increase with decreasing water depth.

Urophycis regia Urophycis regia was collected in the MAB from October until May, with highest densities of larvae occurring in fall collections off the Vir-



Height of the pelvic fin plotted against mandible length for larvae and juveniles of Phycis chesteri and five species of Urophycis.



Mean abundance of *Urophycis chuss* in neuston and bongo collections at stations off Virginia and New Jersey, October 1975—August 1977. n= number of larvae collected. Neuston catches are denoted by clear histograms; bongo catches by stippled histograms. NS = no samples taken. Star denotes bongo catches exceed neuston catches.

ginia coast at midshelf station L2 (Fig. 7). Densities of *U. regia* were much lower in collections taken in February and March, and most specimens were pelagic juveniles found at offshore stations off both Virginia and New Jersey. By May, *U. regia* was scarce; only seven neustonic juveniles were found at offshore stations.

Urophycis tenuis Apart from an occasional *U. regia* juvenile found at offshore stations, *U. tenuis* was the only species of hake present in spring plankton collections off Virginia and New Jersey. Abundance of *U. tenuis* in May 1977 was up to one order of magnitude greater than abundances in June 1976 (Fig. 8). Larvae were collected at all but inshore stations off both Virginia and New Jersey, but were most abundant at offshore stations. Larvae were smallest at offshore stations and increased in size as collections proceeded inshore (Fig. 9, page 220).

Urophycis floridana and U. cirrata Urophycis floridana $(n=41, 13-32 \,\mathrm{mmSL})$ and U. cirrata $(n=5, 20-42 \,\mathrm{mmSL})$ were found exclusively in offshore winter collections (Fig. 10, page 221). With the exception of a single juvenile U. floridana (23.0 mmSL) captured in a bongo tow, all specimens were found in neuston samples.

Phycis chesteri Phycis chesteri larvae first appeared in fall neuston collections from the Middle Atlantic Bight; 16 larvae 6–13 mm in length were collected in November 1976 at offshore stations off Virginia (Fig. 11, page 221). Phycis chesteri larvae and pelagic juveniles remained in surface waters during winter and were found in water deeper than ~ 100 m off both Virginia and New Jersey (n=41). All specimens were collected with the neuston net.

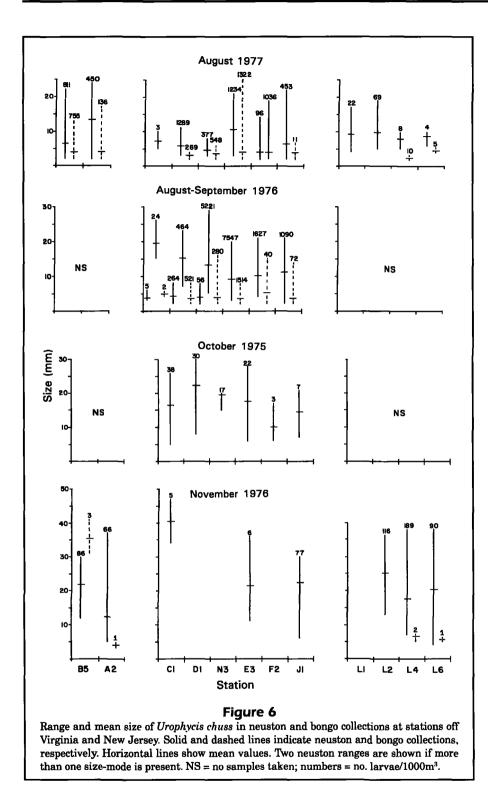
Discussion

Because of similarities between larvae of the seven hake species found in the MAB, a dichotomous key is not a practical tool with which to identify hake larvae in this area. However, the specific identification of larval and pelagic juvenile hake is feasible using a suite of diagnostic characters (App. Table 2). Identifications in this study were based on comparison of larval meristics with adult meristics. Further examination of larvae revealed diagnostic characters comprised not only of meristic information, but also morphometric and pterygiophore interdigitation data. Spawning season and capture location were not used as 'characters' to identify larvae in this study.

Methven (1985) had limited success using pigment characters to separate *U. chuss* and *U. tenuis* >7–8 mm. Problems will persist with the identification of small hake larvae until ontogenetic pigment patterns of all species have been described. These ontogenetic pigment patterns, when used in concert with meristic characters, will hopefully enable relatively routine identifications of these taxa.

The only species of *Urophycis* not found in the present study was *U. earlli*. Adult *U. earlli* are rare and larvae remain undescribed, but they are expected to co-occur with *U. floridana* (Hildebrand & Cable 1938). Both species are similar in having two epibranchial gill rakers, but numbers of first dorsal-fin rays, abdominal vertebrae, and caudal-fin rays delimit most specimens of these two species.

Larval and juvenile *Urophycis* or *Phycis* were present in the MAB throughout the year, and patterns of spatial and temporal distribution of larvae were consistent during both years of this study. *Urophycis chuss* larvae were found in summer and fall collections, with greatest abundances occurring during summer in the



central and northern MAB where water depth was 40-60 m. *Urophycis chuss* was the only species of larval hake found in summer collections, and accounted for 80% of all hakes collected during this 2-year study.

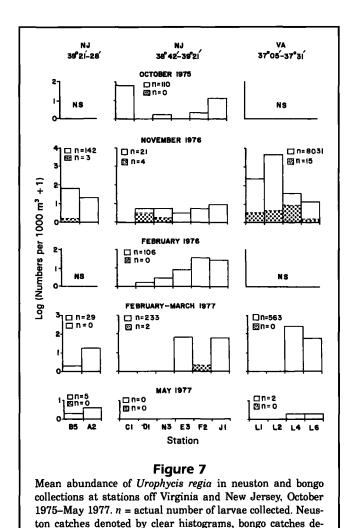
Most *U. regia* in the present study were collected in November, but some larvae or neustonic juveniles were

collected from October to May. Urophycis regia in the MAB is reported to spawn from late September through November, and possibly to February, with peak activity in October (Barans & Barans 1972). Urophycis regia was most abundant during fall in the southern MAB in the relatively shallow (41-43 m) midshelf area. Size range of Urophycis regia collected in this area was 2-34 mm, and although some of the larger specimens may have drifted from deeper water, small larvae were most likely spawned on the shallower central shelf. Evidence of *U. regia* spawning in shallow water was also found in October 1975 off New Jersey where larvae as small as 4mm were found in water as shallow as 12 m. However, not all specimens off New Jersey originated in shallow water; a second group of larvae 6-23 mm in length was found offshore.

The offshore distribution of U. regia became quite distinct in winter collections, with abundances being greatest at offshore stations in February 1976, February-March 1977, and May 1977. These U. regia were probably spawned in offshore waters of the MAB or in offshore waters of the South Atlantic Bight and transported northward. Larval U. regia have been found in abundance in winter collections from offshore waters off North Carolina in the South Atlantic Bight (Fahay 1975, Powles & Stender 1976).

Late-summer spawning by *U. tenuis* occurs in shallow water of the southern Gulf of St. Lawrence and the Scotian Shelf

(Markle et al. 1982). Fahay & Able (1989) suggest the existence of a second stock of *U. tenuis* that spawns in deep water during early spring on the slope of Georges Bank, and probably also along the slopes of the Scotian Shelf, southern New England, and the MAB. The present study found direct evidence of spring spawn-

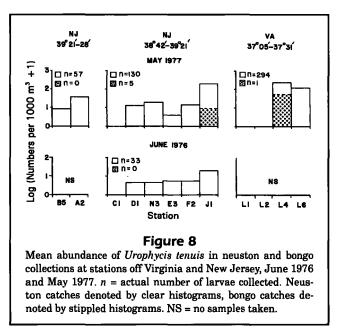


ing by *U. tenuis* in deep water of the MAB; in May 1977 *U. tenuis* larvae as small as 3–4 mm were found over the continental break and slope off both New Jersey and Virginia. In June 1976 *U. tenuis* found at offshore stations were 16–38 mm in length. Based on estimated larval and pelagic juvenile growth rates of 10–22 mm/mo (Markle et al. 1982) and demersal juvenile growth rates of ≈30 mm/mo (Fahay & Able 1989), these fish were probably spawned in late April and May.

noted by stippled histograms. NS = no samples taken.

Fahay & Able (1989), studying young *U. tenuis* in the Georges Bank area, found a shoreward migration with growth. Recruitment to nearshore areas was also indicated in the present study by the increasing size of *U. tenuis* as collections proceeded shoreward. Neustonic juveniles (35–53 mm) were captured in water as shallow as 32 m off the coast of New Jersey.

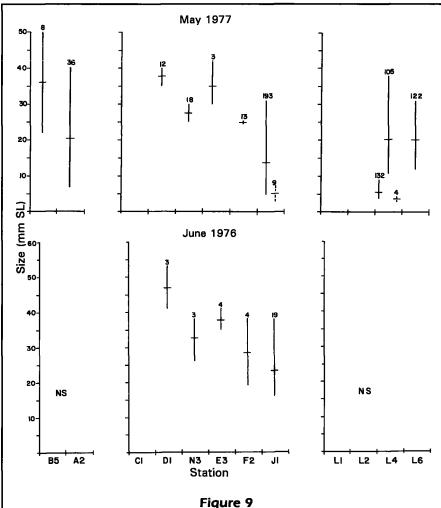
Urophycis floridana and U. cirrata, two southern species of hake, were found off New Jersey and Virginia only in offshore winter collections. The large size



and offshore distribution observed for both species suggest that these pelagic juveniles may have been transported northward into the study area. Larvae of U. earlli, another species found south of the MAB, are rare and remain undescribed, but this species may also occur occasionally in offshore waters of the MAB during winter. Hildebrand & Cable (1938) expected U. earlli to be a winter spawner after collecting three juveniles (37, 75, 103 mm) in March and April off North Carolina, and Fahay (1975) collected a few neustonic U. earlli in winter in the South Atlantic Bight.

Phycis chesteri larvae and pelagic juveniles appeared at offshore stations in fall and winter off Virginia and New Jersey. This larval distribution concurs with Wenner (1983) who found adult P. chesteri generally at depths >183 m on the continental slope from 36°N to 47°N in the western North Atlantic, and noted that spawning off Virginia took place between late September and April, with peak spawning occurring in December and January. Methven & McKelvie (1986) collected 51 P. chesteri larvae and pelagic juveniles along the edge of the continental shelf in the MAB, Grand Bank, and Labrador Shelf, and based on estimated growth rates suggested that most spawning occurs in October.

This study has shown the spatial and temporal distribution of hake larvae in the MAB to be more complex than previously thought. Additional taxonomic characters, particularly ontogenetic pigment patterns, are still needed in order to routinely identify small hake larvae, and more research is needed to explain the observed patterns of larval distribution. Of particular interest is an understanding of the processes that result in the northward transport of larvae and



Range and mean size of $Urophycis\ tenuis$ in neuston and bongo collections at stations off Virginia and New Jersey. Solid and dashed lines indicate neuston and bongo collections, respectively. Horizontal bars show mean values. Two neuston ranges are shown if more than one size-class is present. NS = no samples taken; numbers = no. larvae/1000m³.

pelagic juveniles of southern species into the MAB. Assuming that these individuals are transported northward by the Gulf Stream, it remains to be shown how they leave the influence of this current and move shoreward.

Acknowledgments

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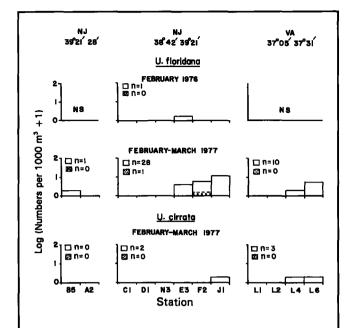


Figure 10

Mean abundance of Urophycis floridana and U. cirrata in neuston and bongo collections at stations off Virginia and New Jersey, February 1976 and February–March 1977. n = actual number of larvae collected. Neuston catches denoted by clear histograms, bongo catches denoted by stippled histograms. NS = no samples taken.

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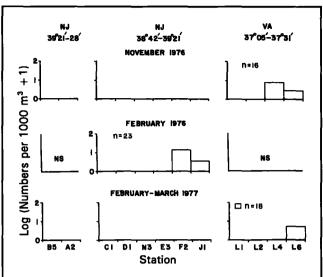


Figure 11

Mean abundance of *Phycis chesteri* in neuston collections at stations off Virginia and New Jersey, during February 1976, November 1976, and February–March 1977. n = actual number of larvae collected. Neuston catches denoted by clear histograms, bongo catches denoted by stippled histograms. NS = no samples taken.

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Species	Collection #	Location	No. specimens	SL(mm)
U. earlli	USNM 025295	N. Carolina	1	124
	USNM 155746	32°34'N,79°05'W	ī	55
	USNM 155747	Wilmington, NC	$ar{f 2}$	50-60
	USNM 226521	32°29'N,79°42'W	3	88-129
	USNM 226522	32°29'N.79°41'W	3	91-122
	USNM 226523	32°29'N,79°41'W	ĭ	113
	USNM 226524	33°14'N,78°24'W	ī	130
	USNM 226525	34°14'N,78°24'W	ī	82
	USNM 226526	32°28'N,79°42'W	$\overline{4}$	91–132
	USNM 226530	32°29'N,79°40'W	4	96–157
	USNM 226531	32°29'N,79°41'W	5	138-166
	USNM 226543	28°48'N,80°38'W	1	74
	VIMS 06557	Gulf of Mexico	î	195
TI A:-I			1	63
U. floridana	USNM 073010 USNM 116729	Key West, FL Beaufort, NC	16	35-49
	USNM 131586	26°18'N,83°09'W	1	59 59
	USNM 151386 USNM 155738	Zo 1811,65 09 W Texas	1	77
	USNM 155782	Cape Canaveral, FL	1	86
	USNM 155783	St. Augustine, FL	1	67
	USNM 156146	Pelican-Stn. 120-5	1	94
	USNM 214118	Brickhill Creek, GA	· 5	64–75
	VIMS 03756	Silver Bay	1	165
	VIMS 03730 VIMS 04142	Brunswick Sound, GA	5	78–113
	VIMS 04142 VIMS 04152	Silver Bay	4	52-85
	VIMS 04192 VIMS 04192	Pensacola, FL	2	81–109
	VIMS 04192 VIMS 04193	Cumberland Id., GA	2	129-184
	VIMS 04193 VIMS 04194	N.Cumberland R., GA	$\frac{2}{2}$	133–157
	VIMS 04194 VIMS 04195	Santa Rosa Sound, FL	ī	66
	VIMS 04196	Oregon S646	1	185
U. cirrata	GCRL 433	29°09'N,88°33'W	1	281
	GCRL 436	29°22'N,87°30'W	1	290
	GCRL 525	29°11'N,88°07'W	3	318-343
	GCRL 2783	Louisiana	4	109-130
	GCRL 17534	28°27′N,90°38′W	1 1	145
	USNM 115686	22°23'N,91°45'W	_	141
	USNM 116929	Tortugas, FL	1	140
	USNM 155642	29°04'N,88°44'W	1	114
	USNM 218169	29°18'N,88°51'W	1	108
	USNM 218192	28°58'N,84°44'W	1	109
	USNM uncat.	24°32'N,83°36'W	1	197
	USNM uncat.	28°59'N,88°48'W	1 2	198
	USNM uncat.	28°35'N,91°12'W		186-220
P. chesteri	USNM 025903	Newport, RI	17	73–98
	USNM 026081	Martha's Vineyard, MA	5	68–83
	USNM 026097	No data	1	79
	USNM 028732	No data	9	58–76
	USNM 083821	GA, SC	12	54-65
	USNM uncat.	Atlantic Arctus Exped.	6	105–147
	USNM 092695	No data	1	63
	VIMS 05238	36°43'N,74°39'W	4	67–150

Appendix Table 2

A summary of key meristic, morphometric, and pterygiophore interdigitation characters used to separate *Phycis chesteri* and six species of *Urophycis*. Given for each species: length (mm) at which characters are attained (upper), morphometric or meristic value (middle), and explanatory notes (**bold** letters in parentheses). Percentages indicate proportion of a species that possesses a particular character.

	Epibranchial gill rakers	Caudal- fin rays	2nd dorsal- fin rays	1st dorsal- fin rays	Abdominal vertebrae	Anal-fin pterygiophores anterior to first haemal spine	Interdigitation of second dorsal-fin pterygiophore	Body depth at vent as %SL	fin base a	of pelvic- as percent ble length
U. tenuis	11–13 mm 2 (95%) (A)	11 mm ≥35 (99%)	14 mm ≥53 (93%)	15 mm ≤12	4 mm ≥15 (H)	8–9 mm >6 (2%) <6 (60%)	≥9 (88%) (L)	12 mm 19.0–21.1 (M)	6–19 mm 24–42 (N)	20-34 mm 26-37
U. chuss	11-13 mm 3 (97%)	8–9 mm <35	14 mm ≥53 (E)	14 mm ≤12	4 mm ≥15 (87%) (I)	8–9 mm >6 (45%) <6 (5%)	12 mm ≥9 (52%)	12 mm 17.6–19.7	6–19 mm 20–39	20-34 mm 15-22
U. regia	11-13 mm 3 (97%)	8–9 mm <35	14 mm ≤52 (E)	13 mm <12 (G)	4 mm ≤14 (J)	8–9 mm >6 (57%) <6 (2%)	12mm <9	12mm 18.4–21.7	6–19 mm 19–33	20-34 mm 12-28
U. floridana	2	<35	≥53	13 (33%)	≥15 (98%) (K)	>6 (74%) <6 (0%)	≥9 (73%)	12 mm 17.7–19.8	15–19 mm 29–36	20–34 mm 23–37
U. earlli	2	<35 (C)	>53	<12 (G)	≤14 (87%) (K)	>6 (85%) <6 (0%)	<9			
U. cirrata	3	<35	>53 (F)	≤12	≥15 (K)	>6 (0%) <6 (82%)				20-34 mm 31-39
P. chesteri	16–18 mm 4–5 (B)	<35 (77%) (D)	≥53	≤12	≥15 (88%)	>6 (0%)	≥9 (38%)	12 mm 21.0-23.4	6–19 mm 44–74	20-34 mm 46-61 (O)

Notes

- (A) U. tenuis occasionally possessed a third gill raker, but in only
- one specimen (n=167) were three gill rakers found on both left and right sides.
- (B) Three epibranchial gill rakers had developed by 13 mmSL.
- (C) U. earlli has never been recorded with >31 caudal-fin rays, while all other hake commonly have >31 rays.
- (D) Although ranges in *U. tenuis* and *P. chesteri* overlapped, numbers
- of caudal-fin rays separated over 40% of P. chesteri (<34 rays) from more than half of U. tenuis (>36 rays).
- (E) Numbers of second dorsal-fin pterygiophores separated U. regia
- from U. chuss at sizes as small as 6 mmSL. Although not shown here, one specimen of U. chuss (n=106) possessed 52 second dorsal-fin rays.
- (F) Although ranges overlap, almost 70% of U. cirrata had at least 64
- second dorsal-fin rays, while P. chesteri and other species of Urophycis (except U. earlli) had <64 rays.
- (G) U. regia and U. earlli have never been found with >10 and 11 first dorsal rays, respectively, while over 80% of U. floridana (n=45) possessed >11 rays.
- (H) Numbers of abdominal vertebrae helped identify *U. tenuis* larvae
- <10 mm, the size below which numbers of caudal-fin rays no longer afford confident identifications. See Results.
- (I) Numbers of abdominal vertebrae aided in the identification of U.
- chuss in summer collections from the MAB when this was the only species of hake found. See Results.
- (J) Only eight specimens of *U. regia* from the Middle Atlantic Bight
- (n=698) had 15 abdominal vertebrae, and seven of these specimens had an anomalous 15th vertebra.
- (K) U. floridana and U. cirrata commonly possessed 16 or 17 abdominal vertebrae, but U. earlli has never been recorded with this many.
- (L) Numbers refer to the interneural space into which points the
- pterygiophore supporting the first ray of the second dorsal fin. The first interneural space was defined as the space anterior to the first haemal spine.
- (M) Specimens of all species were cleared and stained.
- (N) Size ranges do not define size when character first became useful, but bracket the size-range over which particular morphometric values were found.
- (O) At sizes >35 mmSL, P. chesteri was similar to Urophycis with respect to this character.