

Allozyme analysis reveals a complex population structure in the southern calamary *Sepioteuthis australis* from Australia and New Zealand

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ABSTRACT: Allozyme electrophoresis was used to investigate species boundaries and population genetic structure within the southern calamary *Sepioteuthis australis* Quoy and Gaimard. Samples collected from 17 localities around southern Australia and northern New Zealand were examined for allozyme variation at 49 loci. Of 13 polymorphic loci detected, 7 were sufficiently variable to be useful as routine genetic markers of population structure. There was little or no genetic differentiation across the entire range sampled at 5 of these 7 loci. In marked contrast, the allozyme data at 2 loci (*Fdp* and *PepD*) unequivocally sorted all individuals into 1 of 3 genetic types, the geographic distributions of which exhibited a markedly non-random pattern. One type was mainly found near the western and eastern limits of the sampled area, the other type predominantly in the intervening region. Where these 2 types overlapped, a third hybrid-type was found at frequencies predicted under Hardy-Weinberg expectations. The 2 most-likely explanations for these data are: (1) there are 2 taxa within *S. australis* which produce only F₁ hybrids wherever they overlap, or (2) the 2 loci *Fdp* and *PepD* are tightly linked and thus are not independent measures of population structure. Preliminary morphological and reproductive data support the hypothesis of 2 taxa, while mitochondrial DNA-sequence data are inconclusive. It is argued that some combination of the 2 explanations may be operating. Regardless of the final outcome, the data indicate that there are a number of discrete stocks of *S. australis* in this region, a result at variance with current management perspectives on this important fishery.

KEY WORDS: Population structure · Allozyme · Southern calamary

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INTRODUCTION

The southern calamary, *Sepioteuthis australis* Quoy and Gaimard, is a neritic species, endemic to southern Australia and northern New Zealand (Lu & Tait 1983). It is an important component of coastal ecosystems, not only as a primary consumer of crustaceans and fishes, but also as a significant food source for numerous marine animals (Coleman 1984, Gales et al. 1993). The fishing effort directed at this species by both the commercial and recreational sectors has increased substantially over the last decade (Triantafillos 1997), re-

sulting in localised depletions of stocks (Triantafillos 1998). This has raised considerable concern about the status of this resource, suggesting that continued exploitation at current levels might place the stocks at high risk of collapse. Consequently, there is an urgent need to determine the status of southern calamary stocks, and to utilise this information to help implement management strategies that will ensure the long-term sustainability of this fishery.

Like most squid species, *Sepioteuthis australis* is a strong swimmer and therefore has the potential for long-distance dispersal. Despite this potential, tagging studies have revealed only limited migration (Smith 1983, Triantafillos 1998). Nevertheless, such results have to be treated with caution, as recent aquarium

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trials have found that the average life expectancy of southern calamary, once tagged, is less than a few weeks (L.T. unpubl. data). No other documented information on population structure is available for the species. Clearly, more information is required for any informed management of the fishery, be it at the local or global level.

Molecular genetic studies have demonstrated a high incidence of cryptic species in squid (Augustyn & Grant 1988, Brierley et al. 1995, Izuka et al. 1996), revealing taxa that are not routinely detectable using traditional approaches. For example, Yeatman & Benzie (1994) identified 4 species of *Photololigo* using allozyme electrophoresis after a detailed morphological study had revealed only 2. Any ecological studies applied to cephalopods prior to the availability of molecular systematic evidence are suspect, since they possibly involved sample sets that were composites of 2 or more species.

Clearly, there is a need for molecular genetic data at the outset of any detailed study of stock structure. Of the many molecular techniques available, allozyme electrophoresis is still one of the most appropriate techniques for an initial assessment of both species boundaries and population structure (Awise 1994, Hillis et al. 1996). The ability to provide a rapid and comparatively inexpensive survey of a large number of individuals for a wide range of independent nuclear genetic markers makes allozyme data ideal for an examination of species boundaries. In addition, the same data can also provide some insight into the genetic structure of stocks (Ihssen et al. 1981, Ryman & Utter 1987). Allozyme data have already proved useful for assessing intraspecific differentiation in a number of economically important cephalopod species (e.g. Katugin 1995).

With reference to *Sepioteuthis australis*, another systematic issue emerges. As well as the more general consideration of cryptic species, there is a need to address the status of the broadfin squid *S. bilineata* from New Zealand. A morphological study by Lu & Tait (1983) concluded that the 2 named forms were conspecific. The validity of this conclusion can be independently assessed with allozyme data.

The present study uses a combination of molecular and morphological techniques to provide insight into 2 overlapping areas of systematic interest for the southern calamary. The first objective was to clarify the taxonomic status of *Sepioteuthis* throughout southern Australia and northern New Zealand. The second and more important objective was to identify the population genetic structure of *Sepioteuthis* in southern Australasia. Both these objectives were achieved using a variety of techniques including allozyme electrophoresis, mitochondrial DNA (mtDNA)-sequencing, and

analyses of morphological characters and reproductive condition.

MATERIALS AND METHODS

Sample collection. Samples of *Sepioteuthis australis* Quoy and Gaimard were collected from 17 sites along the coast of southern Australia and northern New Zealand between October 1995 and January 1998 using a variety of techniques (Fig. 1, Table 1). Distances between sites ranged from 30 to 5500 km. In addition, 3 sites were represented by replicate sample sets, 2 temporal (Pearson Island and Albany) and 1 spatial (Coles Bay; Table 1). Both the temporal replicates were collected within 16 mo of the original collection date. The spatial replicate was collected on the same day, from a location <5 km from the initial site.

Soon after collection, a piece of tentacle tissue was removed from each individual and frozen in liquid nitrogen. Tissues samples were returned to the laboratory and stored at -80°C , pending genetic analysis.

With the exception of the Perth, Coles Bay, Hazards Beach, and Eden sample sets as well as a few individuals from Albany, Pearson Island, and New Zealand, all calamary ($n = 456$) were characterized for 3 basic parameters, dorsal mantle length, gender, and sexual maturity stage. Dorsal mantle length (DML) was measured, to the nearest millimetre, from the anteriormost point of the mantle to the posterior tip. Sexual maturity and gender were assigned in accordance with the universal scale of Lipinski (1979), with Stages IV, V or VI considered mature.

The bodies of 133 calamary from Esperance, Albany, Franklin Island, Kangaroo Island and Myponga Beach were opportunistically kept for a more detailed morphological examination (Table 1). These calamary were placed in individually labelled plastic bags and frozen at -30°C .

Allozyme electrophoresis. The allozyme study was carried out in 2 different stages. Initially 58 calamary, comprising a small number (3 to 6) of *Sepioteuthis australis* from 10 sites across southern Australia (Table 1), were examined for allozyme variation at a large number of enzyme loci. An additional 10 calamary from the New Zealand sample set were also screened for all loci, once they became available near the completion of the study. The aims of this overview study were 4-fold: (1) to assess the evidence for cryptic taxa/species; (2) to identify polymorphic loci suitable for determining population structure; (3) to find electrophoretic conditions under which allozyme genotypes at these polymorphic loci could be readily assigned to individuals; (4) to set sample sizes for the second stage of analysis based on preliminary allozyme frequencies.

Table 1. *Sepioteuthis australis*. Details of sample sets used in the electrophoretic study. n = sample size. Three replicate sample sets are presented: WA2a, WA2b and SA4a, SA4b = same place, different time; TAsa, TAsb = same time, ~5 km apart. *Sites included in overview study. Numbers in parentheses = number of specimens opportunistically frozen for detailed morphological analysis. WA: Western Australia; SA: South Australia; TAS: Tasmania; NSW: New South Wales

Sample set	Code	Date caught	n	Peripheral (P)	Central (C)	Hybrid (H)
Perth, WA	WA1*	1 Jan 96	50	50 ^a	–	–
Albany 96, WA	WA2a	29 Sep 96	40	14 (7)	8 ^a (6)	18 (12)
Albany 98, WA	WA2b	26 Jan 98	42	27 (27)	1 (1)	14 (14)
Esperance, WA	WA3	24 Jan 98	34	11 (11)	2 (2)	21 (21)
St Francis Island, SA	SA1*	6 May 96	33	–	26	7
Franklin Island, SA	SA2*	7 May 96	31	–	25 (7)	6
Flinders Island, SA	SA3*	5 May 96	18	2	14	2
Pearson Island 96, SA	SA4a*	4 May 96	27	2	14 ^a	11
Pearson Island 97, SA	SA4b	14 May 97	30	–	25	5
Port Lincoln, SA	SA5	2 May 96	52	–	52 ^a	–
Wedge Island, SA	SA6*	1 May 96	51	1	50	–
Kangaroo Island, SA	SA7	18 Apr 97	35	–	35 (15)	–
Glenelg, SA	SA8*	30 Jan 96	50	–	49 ^a	1
Myponga Beach, SA	SA9*	6 Feb 96	50	–	50 (10)	–
Coles Bay, TAS	TASa*	9 Jan 96	50	–	49 ^a	1
Hazards Beach, TAS	TASb	9 Jan 96	50	–	49	1
Eden, NSW	NSW1	4 Sep 96	30	5	15 ^a	10
Newcastle, NSW	NSW2*	14 Oct 95	50	48 ^a	–	2
New Zealand	NZ	4 Nov 98	34	21	1	12
			757	181 (45)	465 (41)	111 (47)

^a1 individual included in mtDNA study

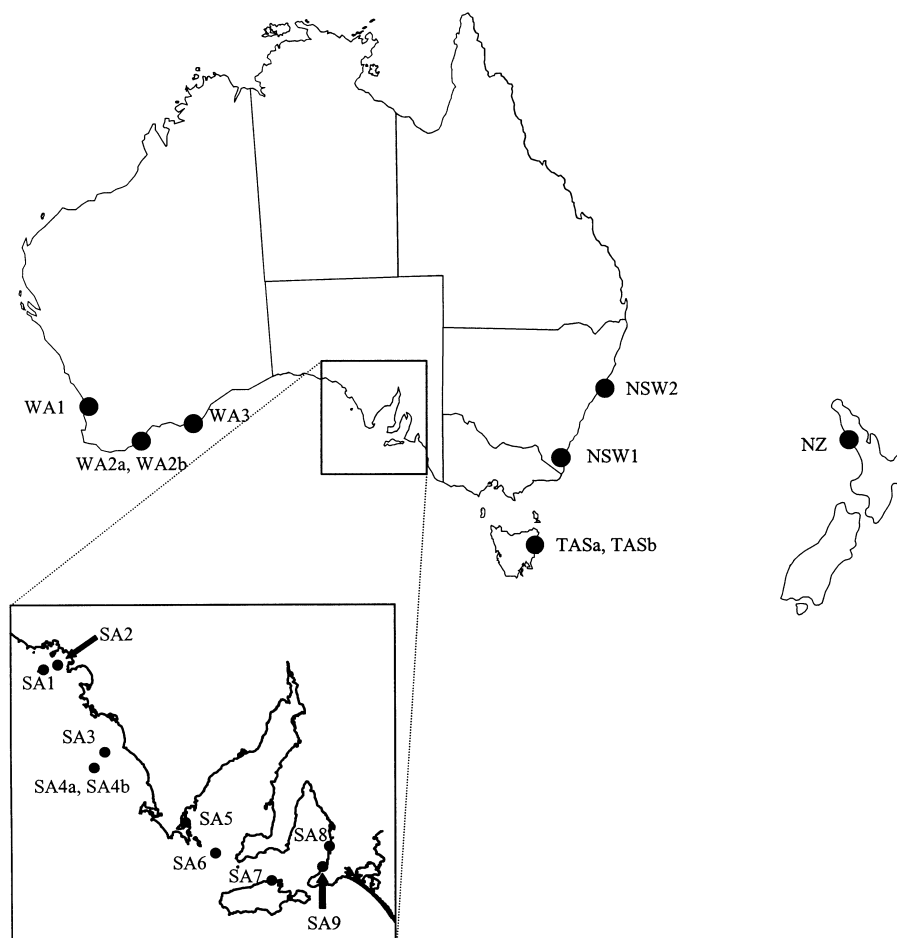


Fig. 1. Map showing collection of *Sepioteuthis australis* in southern Australia and New Zealand. WA: Western Australia; SA: South Australia; NSW: New South Wales; TAS: Tasmania; NZ: New Zealand

Given that enough polymorphic loci were found in the overview study to warrant an assessment of population structure (Richardson et al. 1986), a second stage of allozyme analysis was undertaken. Here, a large number of southern calamary from all 16 sites were genotyped at the polymorphic loci (Table 1, Fig. 1). Based on the overview study, the optimum sample size was set at 50 per sample set, although this was not always achieved for all sample sets. Allozyme electrophoresis was conducted according to the principles and procedures outlined in Richardson et al. Tissues were homogenized by sonication in 2 volumes of homogenizing solution (deionised water containing 0.2% 2-mercaptoethanol and 0.2 mg ml⁻¹ NADP). The following enzymes or non-enzymatic proteins were interpretable in the overview study: ACON (EC 4.2.1.3), ACYC (EC 3.5.1.14), ADA (EC 3.5.4.4), ADH (EC 1.1.1.1), ALD (EC 4.1.2.13), ALDH (EC 1.2.1.5), AP (EC 3.1.3.1), ARGK (EC 2.7.3.3), CA (EC 4.2.1.1), DIA (EC 1.6.99), ENOL (EC 4.2.1.11), EST (EC 3.1.1), FDP (EC 3.1.3.11), GAPD (EC 1.2.1.12), GDA (EC 3.5.4.3), GDH (EC 1.4.1.3), GLO (EC 4.4.1.5), GOT (EC 2.6.1.1), general protein, G6PD (EC 1.1.1.49), GPD (EC 1.1.1.8), GPI (EC 5.3.1.9), GPT (EC 2.6.1.2), IDH (EC 1.1.1.42), LAP (EC 3.4.11.1), LDH (EC 1.1.1.27), MDH (EC 1.1.1.37), ME (EC 1.1.1.40), MPI (EC 5.3.1.8), NDPK (EC 2.7.4.6), NP (EC 2.4.2.1), PEPA (EC 3.4.13; substrate val-leu), PEPB (EC 3.4.11; substrate leu-gly-gly), PEPD (EC 3.4.13; substrate phe-pro), PGAM (EC 5.4.2.1), 6PGD (EC 1.1.1.44), PGK (EC 2.7.2.3), PGM (EC 5.4.2.2), PK (EC 2.7.1.40), SORDH (EC 1.1.1.14) and TPI (EC 5.3.1.1). The nomenclature for referring to loci and allozymes follows Adams et al. (1987).

The allozyme data were analysed for population structure using the computer program GENEPOP Version 3.1b (Raymond & Rousset 1995). All p values were adjusted to allow for multiple tests using the sequential Bonferroni technique (Rice 1989), applied separately for each locus and/or population combination, with an initial α -level of 0.05. *F*-statistics were calculated using the program DIPLOID (Weir 1990). The overall genetic differentiation between sample sets was calculated as Nei's unbiased measure of genetic distance (Nei's *D*; Nei 1978), under the assumption that the loci monomorphic in the overview study were invariant in all sample sets. These genetic distances were depicted diagrammatically as a dendrogram in which clustering was determined by the unweighted pair-group method algorithm (UPGMA; Sokal & Sneath 1963). The results of the allozyme study indicated the need for additional biological characterisation of selected individuals and stocks of southern calamary. This was achieved through an assessment of variability in mtDNA sequence data, external morphology, and reproductive condition. As 2 of the key genetic groups could not be

re-sampled locally, the morphological characterisations were only applied *a posteriori* to the 133 individuals from the allozyme study that were preserved by freezing. Unfortunately, no further *a posteriori* examination of fertility was possible on these individuals, as the reproductive tissues deteriorate after freezing.

Mitochondrial DNA. Eight individuals of *Sepioteuthis australis* from 7 sites were used in a pilot study of sequence variation in portions of the mitochondrial genes COII and COIII (Table 1). These 2 genes were chosen because of the ready availability of primers shown to be successful on a wide range of squid genera (Bonnaud et al. 1997). No specimens from New Zealand were included because this sample set did not become available until near the end of the study, after the mtDNA-sequencing had been undertaken. Sequence data for both genes were obtained from the same individual in all cases, except for 1 sample from Newcastle which resisted all attempts to sequence COII; an additional individual from this site was therefore used for COII analysis. A standard phenol/chloroform extraction procedure was used to isolate DNA (Sambrook et al. 1989). Polymerase chain reaction (PCR) amplification and sequencing was carried out for an ~400 bp portion of COII and an ~700 bp portion of COIII using 2 sets of oligonucleotide primers, namely COII1 (5'ATTGCTCTGCCTTCACTACG3') and COII2 (5'CAAATTTCTGAGCATTGACC3'), plus COIII1 (5'AGCCCATGACCTTTA-ACAGG3') and COIII2 (5'GACTACATCAACAAAATGTCAGTATCA3') (Bonnaud et al. 1997). PCR reactions were carried out in 50 μ l reaction volumes consisting of 1X *Taq* reaction buffer (Promega), 4 mM MgCl₂, 0.8 mM dNTP's, 0.2 mM of each primer, 0.75 units of *Taq* DNA polymerase (Promega) and 50 to 100 ng of template DNA. Amplifications were carried out on a Corbett FTS 320 Thermal Sequencer under the following conditions: 1 cycle at 94°C for 120 s, 48°C for 45 s, 72°C for 80 s; 34 cycles at 94°C for 45 s, 48°C for 45 s, 72°C for 60 s; 1 cycle at 72°C for 360 s. PCR products were purified using Bresaclean, and then cycle-sequenced in 20 μ l reaction volumes on a Corbett FTS1 Thermal Sequencer using ABI Prism[®] (Perkin Elmer) and procedures specified by the manufacturer. DNA sequences were determined using an ABI 373A auto-sequencer. DNA sequences were edited using SeqEd[™] (Version 1.0.3, Applied Biosystem Inc., Madison, WI). Multiple sequence alignments were performed using Clustal V (Higgins et al. 1992). Aligned sequences of 912 bp of combined COII (312 bp) and COIII (600 bp) for each individual (the Newcastle site being represented by the COII of one individual and the COIII of a second) were analysed in the phylogenetic analysis program PAUP (Version 3.1.1; Swofford 1993). Trees were generated using maximum parsimony with a heuristic search option, with all sites weighted equally.

A COIII partial sequence of *S. lessoniana* was used as an outgroup in order to root the tree. Clade robustness was evaluated by bootstrapping over 500 pseudoreplicates. The program MEGA (Kumar et al. 1993) was used to determine the resulting amino acid sequence for each individual using the *Drosophila* mitochondrial genetic code, to ensure that there were no stop codons present which might indicate that a pseudogene had been amplified.

A posteriori morphological characterisation. A range of external morphological characters was initially surveyed on a handful of key specimens for the ability to distinguish the genetic groups evident from the allozyme study. Only 1 character, the number of denticles (teeth) on the largest suckers of the largest tentacles, proved likely to be informative. Consequently, this character alone was measured on all the 133 calamary available for characterization. The 3 largest sucker rings from the 2 largest tentacles were removed from each calamary and viewed under 20× magnification. Sucker teeth were counted, and the internal diameter of the suckers were measured using a graticule eyepiece and calibration slide. A general linear model (GLM) was used to test for significant differences in sucker counts between sexes, sites and genetic types using the SPSS (Subprogram of the Statistical Package for the Social Sciences).

RESULTS

Allozyme variation in *Sepioteuthis australis*

A total of 49 presumptive allozyme loci displayed interpretable banding patterns after histochemical staining. Of these, 13 loci displayed electrophoretic variation consistent with the existence of ≥ 2 alleles at a Mendelian gene. Of these loci, all except 4 (*Enol*, *Got2*, *PepB*, and *6Pgd*) exhibited $q > 10\%$ in at least 1 sample set, and 5 (*Dia*, *Fdp*, *Np1*, *PepD*, *Pgk*) were polymorphic across the range, displaying an average of 4 alleles per locus (range 2 to 8 alleles).

The comprehensive allozyme screen comprised 757 individuals from 19 sample sets (including the 2 tem-

poral and 1 spatial replicates), genotyped at 13 loci. A most striking outcome of this screen was complete linkage disequilibrium between certain genotypic combinations at 2 loci, *Fdp* and *PepD* (Table 2). This occurred despite there being a large number of alleles ($n = 8$) and therefore a very large number of observed genotypes ($n = 25$) at *PepD*.

With respect to individual alleles at the 2 loci, the outcome can be explained by postulating 2 'linkage' associations; the *Fdp*^a allele with any of the 5 most cathodal alleles at *PepD* (a to e), and the *Fdp*^b allele with any of the 3 more anodal *PepD* alleles (f, g, h). No 'recombinant' genotypic pairs were found amongst the 757 individuals examined, and no other evidence of linkage disequilibrium was found for genotypes at any other pairwise combination of the 13 loci. Most importantly, there was also no evidence of deviation from Hardy-Weinberg expectations within any sample set for either *Fdp* or *PepD*, ruling out a common cause of apparent linkage between loci, the Wahlund effect.

The above outcome is consistent with either of 2 alternative biological scenarios. Firstly, it may be that the 2 loci are very tightly linked, such that recombination occurs at less than, on average, $\sim 1/750$ occasions. Alternatively, the results may reflect the presence of 2 taxa, characterized by fixed allelic differences at *Fdp* and *PepD*, plus their F₁ hybrids. For convenience, we hereafter refer to these 2 putative taxa as the 'peripheral' form (*Fdp*^a, *PepD*^{a,b,c,d,e}) and the 'central' form (*Fdp*^b, *PepD*^{f,g,h}), based on their geographic distributions across southern Australasia (Table 1, Fig. 1). The allele frequencies at the 13 polymorphic loci are presented separately for each of the 3 forms (Tables 3 to 5).

How can we distinguish between these 2 very different situations, and consequently their very different implications on stock structure within southern calamary? If the former situation is valid, then no other biological differences should exist between these 3 groups except for those influenced by genes that also lie within the *Fdp-PepD* linkage group. Conversely, if the latter were true, one would expect to find that the 2 putative taxa display genetic differences for other characters/biological traits. Before proceeding to analyse the allozyme data, it is therefore necessary to consider the results of these other investigations.

Table 2. *Sepioteuthis australis*. Association between genotypes at *Fdp* and *PepD* ($p < 0.00001$). Letters in **bold** indicate a 'central' allele

<i>Fdp</i>	<i>PepD</i>		
	aa, ab, ad, ae, bb, bc, bd, be, cc, cd, ce, dd, de, ee	af, bf, cf, df, ef, eg, eh	ff, fg, fh
aa	181	–	–
ab	–	111	–
bb	–	–	465

MtDNA sequence data

Two sequences each from individuals representing 2 'peripheral' and 5 'central' sample sets, identified from their allozyme profiles, were generated in the mtDNA pilot study. Low

		COII Position	COIII Position
		-----	-----
		133	1133455555
Animal's		1700	460505212367
profile	Site	670518	212661398111

Central	WA2	CTTCTC	TAGAAGCGCCGG
Central	SA4a	TCC.C.	CG...ATAT.C.
Central	SA5	.CC..A	CGA.GAT.T..A
Central	SA8	.CCT..	CG.G.AT.TT.A
Central	TAS1	TCC.C.	CG...ATAT.C.

Peripheral	WA1	.CC...	.G....T.T...
Peripheral	NSW2*	.CC...	.G....T.T...

Fig. 2. *Sepioteuthis australis*. Summary of the variable positions found amongst 2 peripheral and 5 central calamary for the COII and COIII sequence data. COII and COIII sequences for Newcastle were derived from separate individual. Dots (.) indicate identical with the reference individual on the first line; *additional individual used in the COII analysis

levels of nucleotide diversity were detected at both protein-encoding genes. For COIII, only 12 of 600 nucleotide positions displayed variation (ignoring the outgroup *Sepioteuthis lessoniana*) and 7 of these involved a unique substitution in a single individual (Fig. 2). Similarly, all but 6 of the 312 nucleotide positions sequenced for COII were invariant, and only 2 of these did not involve a unique substitution in a single individual (Fig. 2). Most variants involved synonymous codon changes, with only 1 in COII (Position 308), and 2 in COIII (Positions 61 and 571), resulting in amino acid substitutions. The COIII nucleotide sequences of *S. lessoniana* and *S. australis* differed at 128 to 131/600 sites, equating to ~6.5% (13 to 14/200) difference in amino-acid sequence.

A phylogenetic analysis was undertaken on the combined 912 bp COII + COIII sequence data, using the COIII sequence from *Sepioteuthis lessoniana* as an outgroup to root the tree. One of the 2 most parsimonious trees is shown in Fig. 3 (the other displayed the same overall topology but a minor difference in 2

Table 3. *Sepioteuthis australis*. Allozyme frequencies in 'peripheral' calamary at the 13 polymorphic loci. All individuals were Genotype *aa* at *Fdp*. Frequencies as % (max. 2n in parentheses)

Locus	Allele (100)	WA1 (28)	WA2a (54)	WA2b (22)	WA3 (4)	SA3 (4)	SA4a (2)	SA6 (10)	NSW1 (96)	NSW2 (42)	NZ
<i>Dia</i>	<i>c</i>	70	71	74	82	50	75		80	69	
	<i>b</i>	26	29	22	18	50		100	20	28	100
	<i>a</i>	4		4			25			3	
<i>Enol</i>	<i>c</i>			2					10	1	
	<i>b</i>	100	100	98	100	100	100	100	90	99	100
<i>Got1</i>	<i>c</i>	1	4	2	9				10		
	<i>b</i>	90	89	94	91	100	100	100	90	91	100
	<i>a</i>	9	7	4						9	
<i>Got2</i>	<i>b</i>	100	100	100	100	100	100	100	100	100	100
<i>Idh</i>	<i>b</i>	100	100	100	100	100	100	100	100	100	100
<i>Np1</i>	<i>e</i>	10	7	8						2	12
	<i>d</i>	7		11	10	25				6	
	<i>c</i>	62	62	72	70	50		100	100	79	13
	<i>b</i>	21	31	9	20	25	75			13	75
<i>Np2</i>	<i>c</i>	1	7	2						1	
	<i>b</i>	99	93	98	100	100	100	100	100	99	100
	<i>a</i>									1	
<i>PepB</i>	<i>b</i>	99	100	100	100	100	100	100	100	99	100
	<i>a</i>	1								1	
<i>PepD</i>	<i>e</i>	33	32	26	32	25	50	50	100	99	100
	<i>d</i>	23	11	22	27		25				
	<i>c</i>	22	24	35	36			50			
	<i>b</i>	17	29	13	5	25	25				
	<i>a</i>	5	4	4		50					
<i>6Pgd</i>	<i>b</i>	99	100	100	100	100	100	100	100	100	100
	<i>a</i>	1									
<i>Pgm</i>	<i>e</i>	1		2							
	<i>d</i>	97	96	96	100	100	100	100	80	97	100
	<i>c</i>	1							10	2	
	<i>b</i>	1	4								
	<i>a</i>			2					10	1	
<i>Pgk</i>	<i>b</i>	73	64	74	77	75	50	100	60	82	100
	<i>a</i>	27	36	26	23	25	50		40	18	

Table 4. *Sepioteuthis australis*. Allozyme frequencies in 'central' calamary at the 13 polymorphic loci. All individuals were Genotype *bb* at *Fdp*. Frequencies as % (max. 2n in parentheses)

Locus	Allele	WA2a (16)	WA2b (2)	WA3 (4)	SA1 (52)	SA2 (50)	SA3 (28)	SA4a (28)	SA4b (50)	SA5 (104)	SA6 (100)	SA7 (70)	SA9 (100)	SA8 (98)	TASa (98)	TASb (98)	NSW1 (30)	NZ (2)
<i>Dia</i>	<i>c</i>	68	50	100	61	56	61	64	68	63	58	54	67	60	57	63	73	
	<i>b</i>	13	50		35	42	39	36	28	31	37	37	27	28	38	33	27	100
	<i>a</i>	19			4	2			4	6	5	9	6	12	5	4		
<i>Enol</i>	<i>c</i>				4	2	4		2	3	4	3	2	1	4	1		
	<i>b</i>	100	100	100	96	98	96	100	98	96	96	97	95	99	96	99	100	100
	<i>a</i>								1				3					
<i>Got1</i>	<i>c</i>								1				2	2		2		
	<i>b</i>	94	100	100	98	94	96	96	98	95	94	91	94	91	89	88	93	100
	<i>a</i>	6			2	6	4	4	2	4	6	9	4	7	11	10	7	
<i>Got2</i>	<i>d</i>														1			
	<i>c</i>														1			
	<i>b</i>	100	100	100	100	100	100	96	100	100	100	97	100	100	98	100	100	100
<i>Idh</i>	<i>a</i>							4				3						
	<i>b</i>	100	100	100	100	100	100	100	100	99	100	100	100	100	100	100	100	100
	<i>a</i>									1								
<i>Np1</i>	<i>e</i>				4		11	7	8	8	11	5	7	7	6	7		
	<i>d</i>	7			6	6	7	4	4	5	7	4	7	5	8	7		
	<i>c</i>	43	50	100	71	75	71	71	69	73	60	73	62	77	60	72	100	50
<i>Np2</i>	<i>b</i>	43	50		19	19	11	18	19	13	21	18	24	11	26	14		50
	<i>a</i>	7								1	1							
	<i>c</i>										1		1					
<i>PepB</i>	<i>b</i>	100	100	100	100	100	96	100	100	100	99	100	99	99	100	99	100	100
	<i>a</i>						4							1	1	1		
	<i>c</i>	100	100	100	100	100	100	96	100	100	99	100	97	100	98	100	100	100
<i>PepD</i>	<i>a</i>									1		2		1	1			
	<i>h</i>		50	25					8				1	2				1
	<i>g</i>					4	4			1	1	1	1	2	1	1		1
<i>6Pgd</i>	<i>f</i>	100	50	75	100	96	96	100	92	99	99	99	99	96	99	98	100	100
	<i>c</i>								2	1				1	1	1		
	<i>b</i>	100	100	100	100	100	100	100	98	99	100	100	100	99	97	98	100	100
<i>Pgm</i>	<i>a</i>													3	3	1		
	<i>d</i>	100	100	100	96	96	96	100	94	98	100	96	99	98	99	97	100	100
	<i>c</i>				4		4					3			1	1		
<i>Pgk</i>	<i>b</i>					4			6	1		1		2	1	1		
	<i>a</i>	81	100	75	73	74	79	61	70	68	67	63	70	67	70	73	77	100
	<i>a</i>	19		25	27	26	21	39	30	32	33	37	30	33	30	27	23	

branch lengths). Of the 4 clades identified, 3 have bootstrap values of $\geq 60\%$, but neither the 'peripheral' nor 'central' haplotypes are strictly monophyletic with respect to each other. While the 2 'peripheral' specimens display identical sequences and form a monophyletic cluster, they also cluster with one of the 'central' specimens (from Albany, Western Australia, a region of pronounced overlap between the 2 putative taxa). As there is no clear support for the genetic distinctiveness of the 2 taxa, and given the low levels of nucleotide diversity and the additional expense involved, no further sequencing was undertaken.

Field examination of specimens *a priori*

Significant differences between the 3 genetic groups were evident for all 3 parameters (gender, stage of sexual maturity, and age-adjusted DML: see Table 6) rou-

tinely determined on most calamary. The proportion of females amongst hybrid calamary (18.2%) was significantly less than in peripheral (38.3%) or central (25.7%) forms ($p = 0.0155$ for 3×2 contingency table; $p = 0.0555$ for central vs peripheral only). Simple comparisons of DML between groups are not appropriate without first correcting for age-related differences between individuals. This analysis will be presented in detail elsewhere (L.T. unpubl. data). Considering only sexually mature individuals, both female and male hybrids were significantly larger than their peripheral or central counterparts (Table 6; $p < 0.001$), a trend that exists with all hybrid age classes (L.T. unpubl. data).

One of the critical pieces of evidence needed to distinguish between the competing hypotheses of simple linkage and 2 hybridizing taxa is the reproductive potential of hybrids. In the absence of any linkage, the allozyme data indicate hybrids would need to be

Table 5. *Sepioteuthis australis*. Allozyme frequencies in 'hybrid' calamary at the 13 polymorphic loci. All individuals were Genotype *ab* at *Fdp*. Frequencies as % (max. 2n in parentheses)

Locus	Allele (36)	WA2a (28)	WA2b (42)	WA3 (14)	SA1 (12)	SA2 (4)	SA3 (22)	SA4a (10)	SA4b (2)	SA8 (2)	TASa (2)	TASb (20)	NSW1 (4)	NSW2 (24)	NZ
<i>Dia</i>	<i>c</i>	67	71	69	79	50	50	64	40			50	50	50	
	<i>b</i>	19	25	31	14	50	50	36	50	100	100	45	50	100	
	<i>a</i>	14	4		7				10			50	5		
<i>Enol</i>	<i>c</i>							9					5		
	<i>b</i>	100	100	100	100	100	100	91	100	100	100	100	95	100	100
<i>Got1</i>	<i>c</i>							5				50			
	<i>b</i>	97	100	98	93	92	75	86	80	100	100	50	95	100	100
	<i>a</i>	3		2	7	8	25	9	20			50	5		
<i>Got2</i>	<i>c</i>				7										
	<i>b</i>	100	100	100	93	100	100	100	100	100	100	100	100	100	100
<i>Idh</i>	<i>c</i>							5							
	<i>b</i>	100	100	100	100	100	100	95	100	100	100	100	100	100	100
<i>Np1</i>	<i>e</i>	12	8	10	7			5			50	50	?		
	<i>d</i>					17		5	20		50		?	25	
	<i>c</i>	69	88	77	86	75	50	75	80	100		50	?	75	12
<i>Np2</i>	<i>b</i>	19	4	13	7	8	50	15					?		88
	<i>c</i>		7												4
	<i>a</i>	100	93	98	100	100	100	100	100	100	100	100	100	100	96
<i>PepB</i>	<i>b</i>	100	100	100	100	100	100	100	100	100	100	100	100	100	100
<i>PepD</i>	<i>h</i>	3													
	<i>g</i>	3			7										
	<i>f</i>	44	50	50	43	50	50	50	50	50	50	50	50	50	50
	<i>e</i>	28	15	9	15	8		9	10	50	50	50	45	50	50
	<i>d</i>	8	14	10	14	25		4	20						
	<i>c</i>	3	21	17	7	17	25		20				5		
	<i>b</i>	11		2	7			32							
<i>6Pgd</i>	<i>a</i>			12	7		25	5							
	<i>c</i>													25	
	<i>b</i>	97	100	100	100	100	100	100	100	100	100	100	95	75	100
<i>Pgm</i>	<i>a</i>	3											5		
	<i>e</i>			2											
<i>Pgk</i>	<i>d</i>	100	100	98	100	100	100	100	100	100	100	100	100	100	100
	<i>b</i>	61	75	83	57	67	50	77	70	100	50	100	65	75	100
	<i>a</i>	39	25	17	43	33	50	23	30		50		35	25	

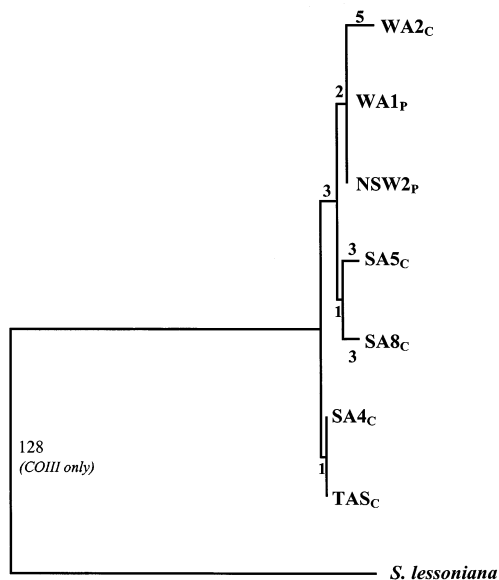


Fig. 3. *Sepioteuthis australis*. Maximum parsimony tree for 2 peripheral (p) and 5 central (c) calamary for the COII and COIII sequence data. Sample abbreviations as in Fig. 1

totally infertile to maintain the integrity of the 2 diagnostic loci. Our assessment here is limited to 1 simplistic measure of reproductive potential, namely the number of individuals that display the overall morphological features of sexual maturity. Given that an individual that appears sexually mature may still nevertheless be infertile, such a measure would be expected to underestimate the extent of any hybrid infertility if it existed. An examination of the reproductive status of hybrids indicates that they are indeed significantly less sexually mature ($p < 0.0001$) than the other 2 types ($p > 0.05$ for peripheral vs central), although a moderate proportion (39 of 88) still showed signs of maturity (Table 6). This is despite hybrids being, on average, much larger than either parental taxon.

Morphological distinctiveness *a posteriori*

Because teeth counts on different suckers from the same calamary showed some minor variation, an aver-

Table 6. *Sepioteuthis australis*. Results of a *a priori* examination of 456 southern calamary in relation to genetic type. DML: dorsal mantle length

Parameter	Southern calamary genetic type		
	Peripheral	Central	Hybrid
No. of immature females (Stages I–III)	4	22	5
No. of immature males (Stages I–III)	5	53	44
No. of mature females (Stages IV–VI)	22	55	11
No. of mature males (Stages IV–VI)	37	170	28
Proportion of females	26/68 (38.3%)	77/300 (25.7%)	16/88 (18.2%)
Proportion immature	9/68 (13.3%)	75/300 (25.0%)	49/88 (55.7%)
DML at which 50% of females mature	148 mm	150.4 mm	206 mm
DML at which 50% of males mature	148 mm	103.7 mm	340.8 mm

age of both teeth counts and sucker diameter was obtained for each individual calamary. A clear relationship was found between DML and average sucker diameter (Fig. 4A). In marked contrast, no discernible relationship was found between average teeth counts and DML, indicating that teeth counts were not size-dependent (Fig. 4B). No significant differences were found in average teeth counts between sexes or sites within genetic type ($p = 0.86$). There was, however, significant variability in the counts between genetic types ($p < 0.001$). The peripherals generally had fewer teeth than the centrals, while the hybrid counts were mostly intermediate. This relationship between average teeth counts and genetic type for the 133 calamary is shown in Fig. 5. The minor variation displayed amongst teeth counts from the same individual is insignificant against the backdrop of the differences between the 3 genetic types, and a single count per calamary provides the same level of discrimination.

Distribution of the 3 forms

An examination of the relative abundance of peripheral, central and F_1 individuals in regions of overlap provides additional insight into the 2 competing hypotheses. On the one hand, the proportion of hybrids in any one sample set does not differ significantly from that predicted for random mating under Hardy-Weinberg expectations (raw data from Table 1), a result entirely consistent with linkage (but not inconsistent with 2 taxa hybridizing at random). In contrast, the proportions of peripheral, central and F_1 individuals differ significantly from each other in different years when the 2 sets of temporal replicates are compared (Samples WA2a vs WA2b, $p = 0.001$; Samples SA4a vs SA4b, $p = 0.013$; raw data from Table 1). This is an unlikely outcome for simple linkage within a stable subpopulation in the absence of strong selection, and implies the existence of 2 taxa whose relative abundance

in regions of overlap can vary markedly between catch efforts. In conclusion, the available evidence supports the notion that the southern calamary comprises 2 taxa with partially overlapping distributions and which hybridize at random in zones of overlap. Given this, the allozyme data have been analysed separately for each taxon (Tables 3 & 4).

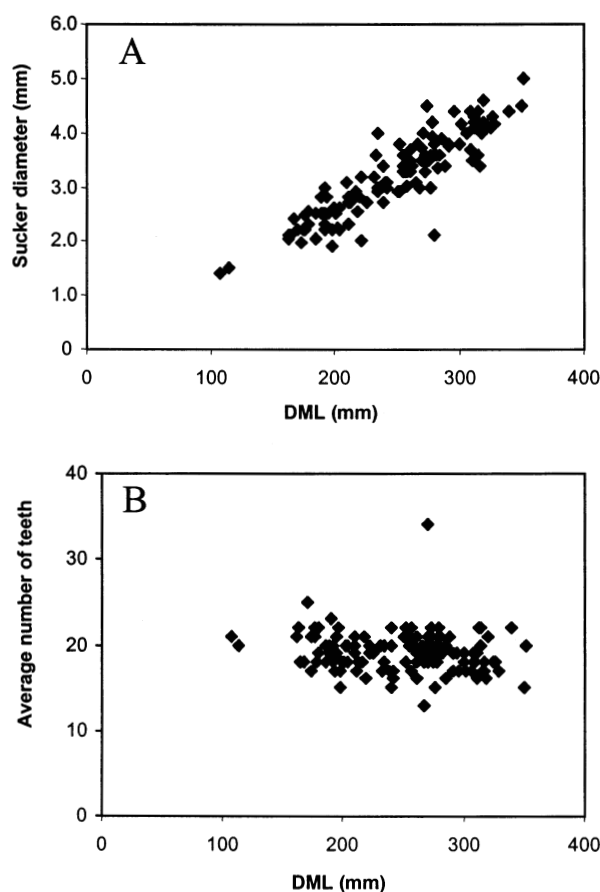


Fig. 4. *Sepioteuthis australis*. Sucker growth. Relationship between dorsal mantle length (DML) and (A) average sucker diameter and (B) average teeth counts

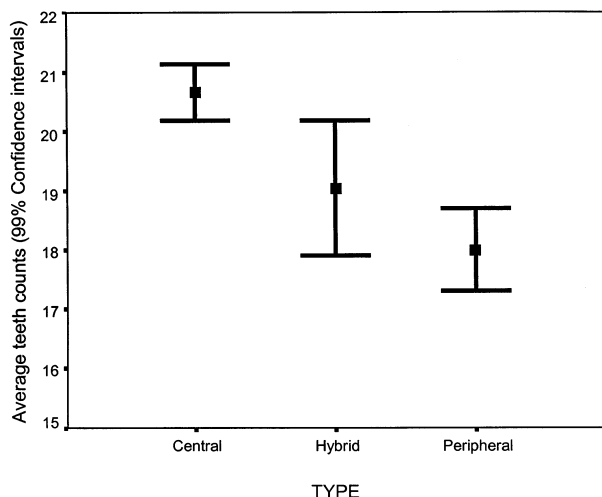


Fig. 5. *Sepioteuthis australis*. Mean \pm 99% confidence intervals of teeth counts for the 3 genetic types of southern calamary from Western Australia and South Australia

Population structure within the peripheral form

The allozyme data (summarised in Table 3) for the peripheral form of *Sepioteuthis australis* were analysed for a suite of population genetic measures available in the program GENEPOP. The measures examined were (1) deviation from Hardy-Weinberg expectations at each locus in each sample set (testing for both a deficit and an excess of heterozygotes), (2) linkage disequilibrium between each locus in each sample set, and (3) genic differentiation between pairwise comparisons of sample sets at each locus. In all cases the 5 individuals from South Australia (at Sites SA3, SA4, and SA6) were pooled into a single sample set ('SA_p') The replicate sample sets at Albany (WA2_a_p and WA2_b_p) were initially treated as separate, and subsequently pooled after failing to show significant differences in any measure.

There were no significant deviations from Hardy-Weinberg expectations at any locus when testing for heterozygote excess, and only one significant outcome testing for heterozygote deficiency (*Np1* in sample set NSW2_p, $0.01 < p < 0.05$). No evidence of linkage between any 2 loci was found in any sample set. Pairwise comparisons of allele frequency indicate the presence of 3 discrete stocks in the peripheral form, characterised by significant differences at ≥ 1 of 4 loci (Table 7). These are referred to hereafter as the New Zealand (NZ), eastern and western stocks. The most distinctive sample set is that from New Zealand, distinguishable at 2 to 4 loci from all other sample sets, even including those with small sample sizes ('SA_p' and Site NSW1; both $n = 5$). Amongst the Australian sample sets, the eastern 2 from Eden (NSW1_p) and Newcastle (NSW2_p) are distinguishable from most others at *PepD*. The single exception involves the small 'SA_p' sample set; a qualitative examination of the allozyme frequency data (Table 3) reveals that the SA individuals appear to belong with the western stock.

Estimates of genetic divergence (Nei's genetic distance, *D*) between sample sets range from 0.000 to 0.001 within the western stock, and 0.001 within the eastern stock. The eastern and western stocks differ on average at 0.008 (Nei's *D*; Fig. 6), whilst both differ from the New Zealand stock at much greater levels (eastern vs New Zealand, average *D* = 0.027; western vs New Zealand, average *D* = 0.026). There were insufficient sample sets within any stock to attempt an isolation-by-distance analysis.

The *F*-statistics for all 8 sample sets support the assertion that there is significant genetic divergence between New Zealand and Australia (Wright's fixation index, $F_{ST} = 0.155$; 95% confidence intervals 0.0307 to 0.245), but no deviation from panmixia within sample sets (inbreeding coefficient, $F_{IS} = 0.031$; 95% confidence intervals = 0.047 to 0.137). Both values are non-significant when the New Zealand sample set is removed.

Table 7. *Sepioteuthis australis*. Summary of pairwise comparisons of allele frequency between sample sets for peripheral calamary. Below diagonal: number of significant differences where $p < 0.05$; above diagonal: loci involved. * $0.01 < p < 0.05$; ** $0.001 < p < 0.01$; *** $p < 0.001$; number of haploid genomes for each sample set in parentheses; all *p* values are corrected

Sample set	WA1 _p (100)	WA2 _p (82)	WA3 _p (22)	'SA _p ' (10)	NSW1 _p (10)	NSW2 _p (96)	NZ _p (42)
WA1 _p	–				<i>PepD</i> *	<i>PepD</i> ***	<i>Dia</i> ***, <i>PepD</i> ***, <i>Pgk</i> ***
WA2 _p	0	–			<i>PepD</i> *	<i>PepD</i> ***	<i>Dia</i> ***, <i>Np1</i> *, <i>PepD</i> ***, <i>Pgk</i> ***
WA3 _p	0	0	–		<i>PepD</i> *	<i>PepD</i> ***	<i>Dia</i> ***, <i>PepD</i> ***
'SA _p '	0	0	0	–		<i>PepD</i> ***	<i>Dia</i> ***, <i>PepD</i> ***
NSW1 _p	1	1	1	0	–		<i>Dia</i> ***, <i>Pgk</i> *
NSW2 _p	1	1	1	1	0	–	<i>Dia</i> ***, <i>Np1</i> ***, <i>Pgk</i> *
NZ _p	3	4	2	2	2	3	–

Population structure within the central form

The allozyme data for the central form of *Sepioteuthis australis* (summarised in Table 4) were analysed for the same array of population genetic parameters, plus the additional calculation of F_{ST} values between sample sets for an isolation-by-distance analysis. In all cases, the 11 Western Australian calamary (WA2a, WA2b, and WA3) were pooled into a single sample set (WA_C). The replicate samples from Pearson Island (SA4a_C and SA4b_C) and Tasmania (TASa_C and TASb_C) were initially treated as separate sample sets, and subsequently pooled after failing to reveal significant differences at any parameter. Consisting as it did of only a single individual, the New Zealand sample set could not be used 'as is' in any analysis. To partially circumvent this problem, the assessment of genic differentiation was carried out using a composite New Zealand gene pool (n = 14 haploid genomes) consisting of the single pure central calamary plus one allozyme from each of the 12 hybrids (only where there was no ambiguity as to what allozyme would have been derived from any putative central parent).

There was no evidence of deviation from Hardy-Weinberg expectations (involving either heterozygote deficiency or excess) at any locus, and no indication of linkage disequilibrium between loci within any sample set. Pairwise comparisons of allele frequencies revealed minor evidence of stock structure within the central form. The composite NZ_C sample set appeared to represent a distinct stock, displaying significant allelic differences from all Australian sample sets at *Dia* (all $p < 0.001$ except for SA2_C, SA3_C, SA5_C, and SA6_C, where $p < 0.01$, and SA7_C, where $p < 0.05$). There was no indication of further substructuring within the Australian stock.

The Australian and New Zealand stocks (as represented by a single specimen) differed at an average Nei's D of 0.007 (range 0.005 to 0.013; Fig. 6). Sample sets within the Australian stock exhibited levels of genetic divergence ranging between 0.000 and 0.003 (Nei's D) from each other. An analysis comparing pairwise F_{ST} values against geographic distance (Fig. 7) revealed no compelling evidence that an isolation-by-distance model is appropriate for these data. The F -statistics for all 13 sample sets also provide weak support for the existence of a separate stock in New Zealand. A marginally significant positive value for F_{ST} ($F_{ST} = 0.0087$; 95% confidence intervals 0.0188 to 0.0001) becomes non-significant after removing the NZ sample set ($F_{ST} = 0$; 95% confidence intervals 0.0014 to -0.0041). F_{IS} values for both analyses do not differ significantly from zero, supporting the assumption of panmixia within sample sets.

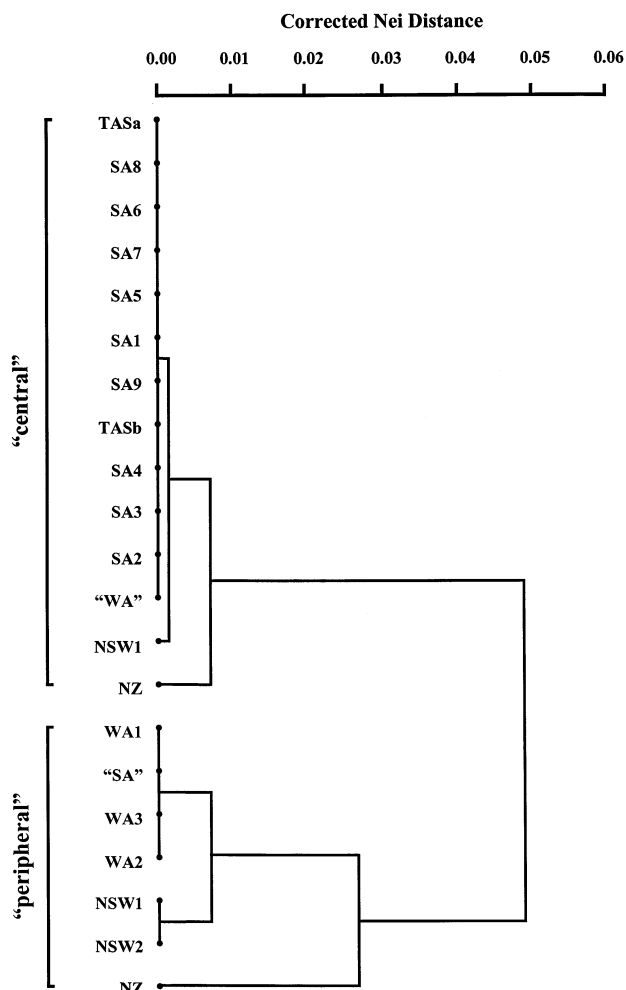


Fig. 6. UPGMA Dendrogram based on Nei's genetic distance for all peripheral and central sample sets

Comparisons of peripheral, central, and hybrid forms in overlap zones

An examination of Tables 3 to 5 reveals that peripheral, central and hybrid sample sets display similar allozyme frequencies in zones of overlap at all loci except the 2 diagnostic markers. Although sample sizes were not always adequate for rigorous statistical analysis, pairwise comparisons of peripheral, central, and hybrid allele frequencies within initial sample sets did not reveal any significant differences at the non-diagnostic loci. Such an outcome is most notable for the New Zealand sample sets (using the composite central/hybrid data to represent the central form). For both the peripheral and central forms, the New Zealand stock is distinguishable from some or all Australian sample sets by correlated differences in allele frequency at the markers *Dia*, *Np1*, and *Pgk*, and yet NZ_p and NZ_C do not differ significantly from each other at

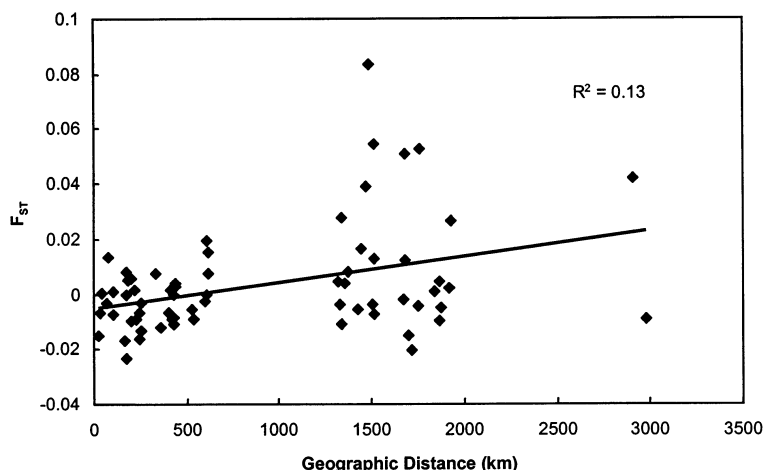


Fig. 7. *Sepioteuthis australis*. Isolation-by-distance analysis for the Australian sample sets of the central form (plot of Wright's fixation index F_{ST} vs geographic distance)

any locus other than *PepD* and *Fdp*, the markers diagnostic for the 2 taxa. Given that 2 Australian stocks of the peripheral form are distinguishable (based on major differences in the frequency of alleles $a+b+c+d$ at *PepD*; Tables 3 & 7), it is worth noting that hybrid calamary from most sample sets (Table 5) bear the genetic signature of the co-existing peripheral stock rather than of the alternate peripheral stock. Thus, distinctive western and eastern 'stocks' of hybrids can be recognized (Sites SA8, TAS, and NSW2 were not able to be assigned because of small sample sizes) based on their *PepD* profile.

DISCUSSION

Systematic status of the 3 genetic forms in *Sepioteuthis australis*

The allozyme data presented herein demonstrate complete linkage disequilibrium between *Fdp* and *PepD* amongst 757 southern calamary examined across a geographic range of more than 5500 km. This despite no indication of heterozygote deficiency in any sample set at either locus, nor, with one exception, at any other locus. Two obvious hypotheses can be envisaged to account for this extraordinary pattern: (1) the 2 loci are very tightly linked, or (2) there are 2 taxa within *Sepioteuthis australis*, diagnosable by fixed differences at *Fdp* and *PepD*, which mate at random wherever they co-exist, but where the hybrids are infertile.

The available morphological and reproductive data largely support the second hypothesis, although they do not specifically exclude the first. Southern calamary

belonging to the peripheral and central forms were distinguishable from each other using the simple morphological feature of the number of teeth on the largest sucker(s) of the largest tentacle(s). They also show significant differences in growth rates after adjusting for sex and site (L.T. unpubl. data). These findings are only consistent with the hypothesis of linkage if genetic control of both these characters also resides within or adjacent to the *Fdp/PepD* linkage group. Furthermore, hybrid calamary differed significantly from both putative parental forms in 4 biological parameters; sex ratio, proportion of sexually-mature individuals, dorsal mantle length at maturity (present study), and growth rates (L.T. unpubl. data). If linkage alone is involved, these characters too must presumably in some way be associated with the *Fdp/PepD* linkage group.

The existence of 3 diagnostic characters and other biological correlates in sympatry would ordinarily be sufficient to warrant the recognition of 2 species within *Sepioteuthis australis*. However, the common occurrence of calamary with the genetic profiles of F_1 hybrids, whose abundance can be predicted under expectations of random mating between the species, creates some uncertainty about such a conclusion. Thus, it is important to explore in more detail the alternative scenarios under which such a result could be obtained within a single species. Pronounced linkage disequilibrium in natural populations between unrelated genes (i.e. not members of a gene family) can occur when the genes concerned are (1) very tightly linked, (2) associated with a polymorphism for some structural chromosomal alteration (usually an inversion or reciprocal translocation, both of which suppress recombination during meiosis in a heterozygote), or (3) not subjected to recombination due to some other mechanism which either suppresses crossing-over or does not allow recombinants to survive (Hartl & Jones 1998). Can any of these situations be operating in the southern calamary?

Explanation 1 is highly implausible in this instance. Firstly, 2 of 13 randomly selected polymorphic markers would have to be so tightly linked that recombinants have never been recovered over successive generations. Loci showing as little as ~1% recombination are usually close to linkage equilibrium in natural populations, and even adjacent loci usually show some evidence of recombination over moderate periods of evolutionary time (Maynard Smith 1994). In addition, a minimum of 2 other genetic makers (i.e. increasing the proportion of 'linked' markers to ~25%), one influencing sucker teeth and the other

growth rates, would also have to be part of the same tight linkage group.

Explanation 2 is also unlikely given the available information on squid karyotypes. Although the karyotype of the southern calamary is not known, it can be reasonably inferred from studies of other species of squid across 5 genera and 2 families. Two sepioids and 3 other loliginids, including the northern calamary *Sepioteuthis lessoniana*, all possess a karyotype of $2N \sim 92$, with no individual chromosome larger than 4% nor smaller than 1.1% of total karyotype length (Gao & Natsukari 1990). Given such evolutionary conservatism, *S. australis* is unlikely to differ greatly from this general pattern. Thus, even if 2 entire chromosomes were involved in a reciprocal translocation, linkage disequilibrium would still only be evident for at most 8% of loci examined. Polymorphism for multiple chromosomal rearrangements, such as is found associated with semi-sterility in some plants (Hartl & Jones 1998) would need to be operating to generate the results obtained here within a single species. Moreover, even where inversion or translocation polymorphisms are evident within a species, there is rarely complete linkage disequilibrium between included loci, due to phenomena such as double cross-overs, gene conversion and 'middle-gene' effects (Loukas et al. 1979, Krimbas & Powell 1992).

For similar reasons, Explanation 3 is also unlikely to be a factor for these data. Assuming a karyotype of $2N \sim 92$, any suppression of crossing-over or recombination would need to operate simultaneously and concertedly on a number of different chromosomes to generate pronounced linkage disequilibrium amongst at least 25% of loci sampled. Some species are known in which recombination does not occur within individuals, but these instances only occur in one sex (Maynard Smith 1994). Absence of recombination would need to feature in both sexes of *Sepioteuthis australis* for this explanation to be tenable and, if present, might also be expected to lead to many other instances of linkage disequilibrium.

Thus we argue that the results obtained here for southern calamary are inconsistent with the hypothesis of a single species, and instead suggest the existence of 2 taxa, capable of hybridizing freely in areas of overlap. Nevertheless, such a biological scenario does not preclude the existence of chromosomal differences between these 2 taxa. Closely-related taxa differing in structural chromosomal features such as translocations or inversions often exhibit partial or total reproductive isolation (White 1978). Any hybrids produced may experience difficulties in chromosomal pairing and proper disjunction during meiosis, resulting in partial or complete sterility (Avisé 1994). And, while total sterility would maintain the genetic integrity of both

taxa (in this case species), partial sterility would result in some degree of genetic exchange between the 2 gene pools for all loci not included within the structural rearrangement distinguishing the 2 taxa. As a consequence, allele frequencies at most loci would tend to converge on intermediate levels in both taxa, the extent of introgression depending on the degree of sterility encountered in hybrids and the effects of selection.

Such an explanation would be consistent with 2 of the major outcomes of this study, namely (1) only F_1 hybrids were found, and (2) allele frequencies at loci other than *Fdp* and *PepD* did not differ significantly between central and peripheral individuals in any zone of overlap, including New Zealand (where both the NZ_P and by inference the NZ_C stocks were diagnosable from their eastern counterparts but not from each other). Hybrids could still be partially fertile, but would only produce gametes with parental genotypes at loci located within the chromosomal rearrangement (*Fdp* and *PepD* in this instance). Loci not linked to the rearrangement would participate in some level of genetic exchange such that in any one site the 2 'parental' taxa would converge on similar allelic profiles at all non-linked loci. As a consequence, each of the 'peripheral', 'central' and 'hybrid' genetic types would actually harbour a small percentage of calamary of hybrid ancestry ($F_1 \times$ peripheral, $F_1 \times$ central, F_2 etc.), depending upon the fertility of each type of second-generation cross. Nevertheless, partial sterility in the hybrids would still ensure that other genetic differences would be maintained at loci under the direct influence of selection.

Unfortunately, the data presented herein cannot resolve whether there is introgression between the 2 taxa. Our limited data plus those of Triantafillos (unpubl. data) suggest that hybrids, whilst more vigorous than either parental form, are at best semi-sterile. Hybrid calamary were, in general, substantially larger than mature peripheral and central calamary, yet a majority were sexually immature, as assessed by a macroscopic examination of the gonads. Interestingly, the only site where mature hybrids were commonly found was New Zealand. Despite having fully formed reproductive organs, these hybrids may still not be able to produce reproductively competent offspring. Galbreath & Thorgaad (1995) found that seemingly mature hybrid salmon produced offspring that never progressed past 30 d, and Lincoln (1981) found that male triploid plaice crossed with flounder produced hybrids that seemed sexually mature, but that only produced sterile gametes.

We hesitate to apply the term 'species' to these 2 taxa until further investigation has addressed the reproductive potential of hybrids. As such, our data do not allow

us to draw conclusions about the nomenclature of *Sepioteuthis* spp. in Australia and New Zealand. The allozyme data do offer general support for the conclusions of Lu & Tait (1983) that the same species of *Sepioteuthis* occurs in both Australia and New Zealand. However, if hybrids turn out to be largely infertile (and therefore both peripheral and central forms are good biological species), the name '*S. bilineata*' may nevertheless still apply to 1 of the 2 species in New Zealand. Such ambiguity could then only be resolved by a re-examination of holotypes for sucker-teeth counts and any other diagnostic markers that may emerge in the future.

Unlike the morphological and the growth data, the pilot mtDNA data did not fully support the hypothesis of 2 taxa. Several explanations are possible here. As closely related taxa often do not display reciprocal monophyly for their mitochondrial gene trees (Avice 1994), the results of our pilot analysis may reflect the true mitochondrial phylogeny. Alternatively, the protein-encoding genes chosen may be too conservative to reveal any fine-scale phylogenetic structure. Low levels of genetic diversity have been found in other loliginids such as *Loligo pealei* (Garthwaite et al. 1989), and may be characteristic of squid in general (Ally & Keck 1978, Brierley et al. 1993). A third possibility is that the mtDNA data are informative in that they identify a clade comprising the 2 'peripheral' calmary (identical sequences despite being from opposite ends of the continent) plus a 'central' calmary from Albany, a region where hybrids are commonly encountered but where the central form is rare. Such a finding may actually reflect the existence of some introgression, with the Albany individual displaying a peripheral mtDNA haplotype originally derived from its original maternal parent. However, any discussion of introgression must remain pure speculation given the low levels of diversity and the absence of further sequence data.

Population structure within each genetic group

Regardless of the extent of introgression occurring between peripheral and central forms of the southern calamary, the existence of these 2 taxa and their hybrids requires that all 3 genetic groups be assessed separately for any evidence of population structure. Unfortunately, all 3 analyses suffer from the same deficiency, namely inadequate sample size. Few sample sets conform to our *a priori* goal of $n = 50$ (2 of 10 peripheral, 6 of 17 central, 0 of 14 hybrid) and this severely limits the ability of the allozyme data to probe too deeply into population structure. This limitation is further compounded by the high probability of a Type II error accompanying all allozyme analyses of popula-

tion structure (Richardson et al. 1986). Nevertheless, it is possible to make some statistically rigorous observations regarding the minimum number of stocks present within each genetic form.

Peripheral form

Three subpopulations were shown to exist within the peripheral form, one each in Western Australia, south-east Australia and New Zealand. Multi-locus heterogeneity between the Australian and New Zealand stocks rules out the possibility of a locus-specific rather than population-structure effect (Richardson et al. 1986). It is not surprising to find genetic differences between mainland Australia and New Zealand samples, because these 2 regions are separated by >2000 km of open ocean, much of which is of considerable depth (>2000 m). Regular migrations and subsequent genetic exchange across such a geographic barrier by a neritic squid are unlikely (see Brierley et al. 1995). The eastern and western stocks of Australia are diagnosable only at 1 locus (*PepD*), albeit involving major clear differences in allele frequency. These 2 stocks are also, to a certain extent, oceanographically isolated. The eastern stock is kept more or less isolated from the western stock by the East Australian Current (Boland & Church 1981).

Central form

In stark contrast to the population heterogeneity observed within the peripheral type, very little population substructuring was detected within the central form. Allele frequencies from Tasmania were similar at all loci to those found in Western Australia, a distance of nearly 3000 km. These data are consistent with a single, randomly-mating stock across Australia. As with the peripheral form, there were some significant allelic differences between Australia and New Zealand, suggesting that the New Zealand sample was a separate stock.

Prior to this study, the only information concerning the stock structure in southern calamary was gained from 2 short-term tagging studies (Smith 1983, Triantafillos 1998). Both these studies showed that although most individuals are predominantly sedentary, the species is capable of long and fairly rapid migrations. Such mobility would potentially facilitate gene flow over considerable distances. In the absence of strong selection pressures, the effective number of migrants between regions that is necessary to prevent genetic drift from producing significantly different allele frequencies is small, i.e. 1 to 10 individuals per generation (Allendorf et al. 1987). Given that large

stretches of the southern Australian coast are suitable habitat for calamary, it is feasible that there is sufficient movement of individuals from one region to the next, in a stepping-stone fashion, to prevent subpopulations emerging within the entire Australian central stock.

Hybrid form

Three stocks of the hybrid form of *Sepioteuthis australis* are broadly identifiable from the allozyme data, based on the distribution of parental stocks and from an examination of allozyme frequencies (Table 5). The western and eastern hybrid stocks differ only at one marker (*PepD*), whilst the New Zealand hybrid stock is diagnosable from the 2 Australian stocks using the same loci that distinguish both New Zealand central and New Zealand peripheral stocks from their respective Australian counterparts (*Dia*, *Pgk* and/or *PepD*). Although the rarest overall, hybrid calamary were the most abundant of the 3 genetic groups in 2 of the 19 sample sets (Albany 96 and Esperance), demonstrating that they are likely to constitute a major part of the catch in the major zones of overlap, i.e. (1) the south coast of WA through to the west coast of SA, (2) the south-east coast of Australia, and (3) at least 1 region in New Zealand. As such, they need to be considered as part of any management plan for the southern calamary fishery.

Biogeographic regions

The geographic distributions of the 2 taxa are closely related to the biogeographical regions proposed by Knox (1963). The distribution of the central type coincides with the Flindersian and Maugean Provinces postulated by Womersley & Edmonds (1958), which have affinities intermediate between cool- and warm-temperate regions. In comparison, the peripheral type prefers the warmer waters of the warm temperate regions of the Western Australian, Peronian and Auporian Provinces found in Western Australia, New South Wales, and New Zealand, respectively. Most of the hybrids were found in the transitional zones between these water masses. Such affinities to different oceanographic masses could account for the genetic divergence between the 2 taxa. For example, an allozyme study of the ommastrephid squid *Nototodarus gouldii* around New Zealand also revealed an allopatric sibling species, *N. sloanii* (Smith et al. 1981). The distributions of these 2 species were divided by the Substructuring Tropical Convergence Zone (Smith et al. 1981). It may be that under normal conditions the peripheral

and central forms of *Sepioteuthis australis* may be largely allopatric, coming together only occasionally when prevailing oceanographic conditions allow (Brierley et al. 1993). For example, the assorted peripherals around the islands of southwestern South Australia may be attributable to the Leeuwin Current (e.g. Maxwell & Cresswell 1981). This current carries warm water from northwestern Australia, where the peripherals dominate, through Albany, and then across the Great Australian Bight (Cresswell 1991). At its strongest, the Leeuwin Current eventually dissipates after it reaches the islands of southwestern South Australia (Cresswell 1991). This may explain why there are no peripherals east of these islands. Similarly, the East Australian Current probably controls the distribution of the peripherals on the eastern side of Australia and New Zealand. Gill (1997) suggested that many Australian marine reptiles were assisted or carried to New Zealand by the East Australian Current.

Implications for management of the southern calamary fishery

The results of our study have clear implications for the management of southern calamary. The existence of sibling species, and their unknown contribution to respective fisheries, would complicate markedly the construction of stock-assessment models and provisions of subsequent fishery advice (Beddington et al. 1990), especially as part of the observed variance in reproductive, size and age-related parameters are species-specific. At present, southern calamary in Australia are managed as if all individuals were members of a single, interbreeding stock. The results presented here indicate a minimum of 5 stocks, namely (1) western peripheral, (2) eastern peripheral, (3) central, (4) western hybrids, and (5) eastern hybrids. Three additional stocks occur in New Zealand: NZ peripheral, NZ central, and NZ hybrids. With the exception of Perth, Port Lincoln, Myponga Beach and Kangaroo Island, all sites had more than 1 genetic type present. At 6 localities, all 3 genetic types were present at various frequencies (Albany, Esperance, Flinders Island, Pearson Island, Eden, and New Zealand). Moreover, both temporal replicates revealed significant differences in the proportions of the 3 stocks taken at random. This result may reflect genuine temporal differences in abundance, or alternatively the differential sampling of taxon-specific differences in the microhabitat. Regardless of the explanation, where stocks differ in their relative strengths in space and time, none can be harvested at an optimal level, since either the weaker stocks are over-exploited or the stronger ones remain under-exploited (Ryman & Utter 1987).

Concluding remarks

The results of this study and the occurrence of cryptic speciation in yet another species of loliginid squid reinforces the need for a systematic molecular assessment of the target 'species' as a first step in any study on the biology of squid. Ambiguity of this sort reiterates the relevance of systematics and the importance of unequivocally identifying whether the populations under study are conspecific before undertaking any assessment of population structure. In the case of *Sepioteuthis australis*, the present study casts doubts over the validity of all previous ecological work, the results of which were not interpreted according to the true systematic complexity present.

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