

## The value of host and parasite identification for arripid fish

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**Abstract.** Accurate identification of fishes and their parasites is fundamental to the development, management and sustainability of fisheries and aquaculture worldwide. We examined three commercially and recreationally exploited Australian arripid species (Pisces: Arripidae), namely Australian herring (*Arripis georgianus*), eastern Australian salmon (*A. trutta*) and western Australian salmon (*A. truttaceus*), to determine their metazoan parasite assemblages and infection parameters. We identified 49 parasite species including 35 new parasite–host records and recognised seven ambiguous parasite–host records in the literature, largely a consequence of unsubstantiated host identifications in previous studies. Morphological and molecular methods confirmed a new western extension for the range of *A. trutta*, ~1000 km west of the previous record. Confusion about host identification and the range extension documented here has implications for the management of these economically important arripid species in southern Australian waters. Our examination of an endemic Australian fish family emphasises that accurate identification of fishes and their parasites is a fundamental prerequisite for efficient and sustainable resource management.

**Additional keywords:** Arripidae, Australian herring, Australian salmon, fish parasites, fisheries management, molecular techniques, taxonomy.

### Introduction

Accurate species identification underpins knowledge of biodiversity, biogeography, biosecurity, biology, ecology, conservation, management and animal health. Published examples emphasise this for parasites (e.g. Peters *et al.* 1983; Siddall *et al.* 2007), commercially important fish (e.g. Dulvy and Reynolds 2009) and parasites infecting fish (e.g. Barber *et al.* 2000; MacKenzie 2002; Young *et al.* 2007). Morphological taxonomy remains a fundamental scientific tool and has immediate relevance to applied marine science. There is an increasing need to return to basics, survey fishes and their parasite fauna across broad geographic ranges and accurately identify hosts and parasites to species level. Fundamental knowledge of parasite assemblages is missing for members of the Arripidae, fishes endemic to southern Australian and New Zealand (NZ) waters. The Arripidae contains a single genus *Arripis* of four species (Paulin 1993). The following three species comprise important

commercial and recreational fisheries in Australian waters: Australian herring or tommy rough (*Arripis georgianus*), eastern Australian salmon (*A. trutta*) and western Australian salmon (*A. truttaceus*). About 3000 t of ‘Australian salmon’ (*A. trutta* and *A. truttaceus* combined) are taken commercially in Australian waters per annum, and ~100 t of *A. georgianus* are captured commercially in South Australia (SA) (ABARE 2009).

*Arripis georgianus* and *A. truttaceus* school inshore and share similar distributions and reproductive strategies; however, they are distinct morphologically. They each comprise a single stock distributed from Western Australian (WA) to Victorian waters and waters around Tasmania (Malcolm 1960; Hoedt and Dimmlich 1994; Fairclough *et al.* 2000b). *A. georgianus* also occurs off the eastern coast of New South Wales (NSW) (Gomon *et al.* 2008). *A. georgianus* and *A. truttaceus* share the same single spawning ground off the south-western coast of WA (Fairclough *et al.* 2000b; Jones 2008). *A. trutta* overlaps

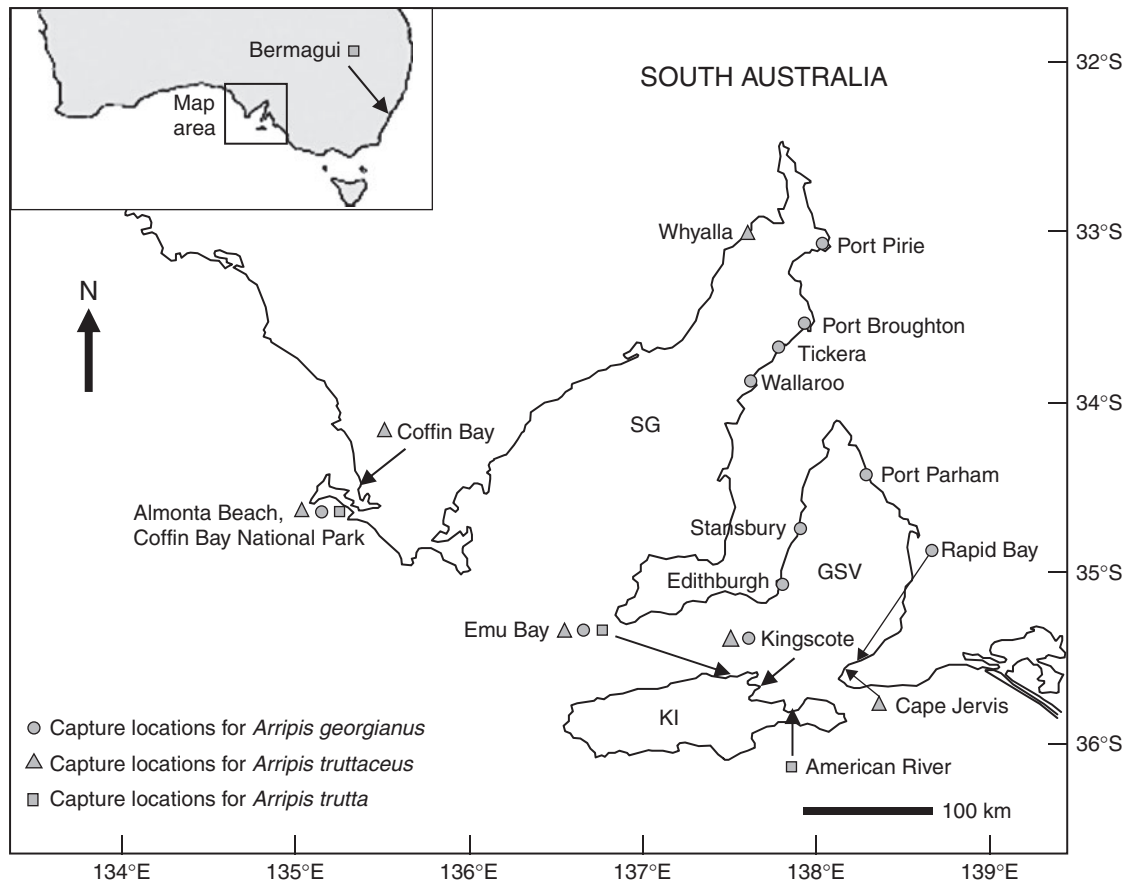


Fig. 1. Sample localities for *Arripis* spp. in southern Australian waters and one site off the coast of New South Wales (inset). GSV, Gulf St Vincent; KI, Kangaroo Island; and SG, Spencer Gulf.

their distribution in Victorian and Tasmanian waters, and is also found off NSW and NZ. Its stock structure is currently unknown (Smith *et al.* 2008).

Morphological similarity among some *Arripis* species has led to an identity crisis (Paulin 1993). *A. trutta* and *A. truttaceus* can be distinguished only by maximum attainable size and the number of gill rakers on the first gill arch (Gomon *et al.* 2008), a character suggested to result from different prey species consumed (Malcolm 1959; Hoedt and Dimmlich 1994). The practical difficulty in distinguishing these species has ultimately confounded scientific research and fishery data (SARDI 2000; ABARE 2009). Indeed, *A. trutta* and *A. truttaceus* are not treated as separate species for current Australian fishery catch statistics (ABARE 2009). Furthermore, the distribution of these species remains unclear. Reports that indicate capture records outside the recognised distribution of these species are available only through 'grey' literature and no corresponding specimens are deposited in museum collections for verification and future study.

Ambiguity in fish identification also mars parasite–host records, with many reports failing to detail methods used to identify hosts. Knowledge of the parasite fauna of arripids is valuable because individuals may aggregate near structure, including fish farms (Dempster and Kingsford 2004; Neira

2005; Catalano and Hutson 2010), a behaviour that presents an opportunity for parasite interactions between farmed and wild fish. Several parasite species are documented from *A. trutta* but few are reported from *A. georgianus* and *A. truttaceus*. The aim of our study was to extend knowledge of the metazoan parasite fauna of the Arripidae in southern Australian waters, provide comprehensive host and parasite identifications and validate host identity via molecular genetics.

## Materials and methods

### Fish collection and identification

In all, 183 *A. georgianus* individuals (mean fork length (FL) = 179 mm, range = 154–220 mm) and 67 *A. truttaceus* individuals (mean FL = 350 mm, range = 185–601 mm) obtained from Spencer Gulf (SG), Gulf St Vincent (GSV), Kangaroo Island (KI) and Coffin Bay (CB) in SA waters (Fig. 1) were examined between February and July 2009 for metazoan parasites (University of Adelaide, Animal Ethics Committee Project S-098-2007). Four *A. trutta* individuals were examined from KI and Almonta Beach, and CB National Park in SA waters (Fig. 1), and 19 were examined from Bermagui in NSW (mean FL = 448 mm, range = 179–545 mm) (Fig. 1, inset). Fishes were collected either via line fishing or from market catches

(Safcol Fish Market, Mile End, Adelaide, SA, and Sydney Fish Market, Pyrmont, NSW). Most fish individuals were examined fresh; others were frozen on capture and processed later. We distinguished *A. trutta* from *A. truttaceus* by using gill raker counts on the first gill arch (Gomon *et al.* 2008) and used molecular genetics to confirm our morphological differentiation.

#### Fish necropsies

Fish were examined for metazoan parasites by standard parasitological methods following Hutson *et al.* (2007). Tissue samples (fin clips) were collected from each fish examined, stored in 95% undenatured ethanol and lodged with the Australian Biological Tissue Collection (ABTC 108509–108777) at the South Australian Museum (SAMA). Four partially dissected *A. trutta* individuals were deposited in the Ichthyology Collection at SAMA ( $n = 2$  from KI, F12179–80;  $n = 1$  from Almonta Beach, CB National Park, F12182) and Museum Victoria (MV) ( $n = 1$  from KI, A 30606-001). One individual of *A. georgianus* and one of *A. truttaceus* captured from Edithburgh and Pt Broughton, respectively, were also deposited (partially dissected specimens) in the Ichthyology Collection at SAMA ( $n = 1$  from Edithburgh, F12378;  $n = 1$  from Pt Broughton, F12379).

#### Parasite preparation

Standard methods following Hutson *et al.* (2007) were used to preserve, stain, mount and examine parasites. Original descriptions, redescrptions (see Tables 1–3), keys and borrowed vouchers assisted identification. Parasite prevalence and intensity are given in whole numbers and follow Bush *et al.* (1997). Mean and maximum intensity for *Ceratomyxa* sp. (Myxozoa) from *A. truttaceus* could not be determined because of the high level of infection observed prohibiting accurate counts of individuals. Parasites are deposited in the Australian Helminthological Collection (AHC) and the Crustacean collection (C) at SAMA, the Parasitology Collection at Queensland Museum (G), the collections at the Natural History Museum, London (BMNH) and the United States National Parasite Collection (USNPC) (Tables 1–3).

#### Molecular methods

Total genomic DNA was extracted, amplified and sequenced from fin clips of *A. truttaceus*, *A. trutta* and *A. georgianus*, sourced from different localities in SA and NSW (Table 4, available as an Accessory Publication to this paper), using the molecular methods in Catalano *et al.* (2010). Three partial gene segments were amplified and sequenced using the following primers: Cox1 fragment – Primers Fish F2 and Fish R2 (Ward *et al.* 2005); 16S fragment – Primers L2510–16S and H3084–16S (Doiuchi and Nakabo 2006); 28S fragment – Primers 28SV and 28SJJ (Smith and Wheeler 2004). Representative sequences are deposited in GenBank (Table 4, available as an Accessory Publication to this paper). The recovered sequences were compared with reference strains (Table 4, available as an Accessory Publication to this paper). All pair-wise similarities

are presented in Table 5, available as an Accessory Publication to this paper.

## Results and discussion

### Range extension for *Arripis trutta* and distribution of arripid species

Our morphological (different gill raker counts) and molecular analyses (0% variance in Cox1 sequences between SA and NSW *A. trutta*) confirmed the presence of *A. trutta* in SA waters and extended its western range by ~1000 km from Port Philip Bay, Victoria (38°19'11"S, 144°42'48"E) to Almonta Beach, CB National Park, SA (Fig. 1). On the basis of the molecular analyses, we have confirmed that gill raker counts morphologically discriminate the species and also noted a consistent difference in the appearance of the gill rakers; those of *A. trutta* are relatively long, thin and closely spaced compared with those of *A. truttaceus*. Primers specific to *A. georgianus* 28S rRNA failed to amplify product from *A. trutta* and *A. truttaceus*, prohibiting sequence comparison for this gene. Furthermore, the 16S rRNA sequence fragment (545–563 bp) from *A. trutta* was identical to that from *A. truttaceus*, also prohibiting the discrimination of these species by using this gene fragment. Given the morphological similarity of these two arripid species, except maximum attainable size and gill raker number on the first gill arch (Gomon *et al.* 2008), it is easy to understand why their coexistence in SA waters has gone undetected. Our results indicated a considerable overlap in the distribution of three *Arripis* spp. in SA waters (Fig. 1). It is likely that *A. trutta* schools with *A. truttaceus* in SA waters because they were always captured together in our study.

Fairclough *et al.* (2000a) indicated that *A. trutta* spawns in south-eastern Australia in summer, when the southward-moving Eastern Australian Current (EAC) and subsequent residual current moving westward are strong (Richardson and Poloczanska 2009). In recent years, the strength of the EAC has intensified, with a predicted increase of >20% by 2100, resulting in changes in the range of several marine species (Ridgway and Hill 2009). This suggests that the occurrence of *A. trutta* in SA waters may increase in the future. We believe our discovery is significant for the Australian salmon fishery in SA waters. However, before any action is taken to modify fishery management, it is crucial to determine whether or not *A. trutta* is common in SA waters and also whether the species contributes significantly to the catch limit. This will require subsampling over an extended time period.

A single fish species that exhibits divergent migration behaviours may exhibit subtle, but demonstrable morphological differences (Secor 1999). It is plausible that *A. truttaceus* and *A. trutta* may represent a single *Arripis* species and that differences in gill raker number and morphology and the maximum body size attained may correlate with different feeding habits along separate migration paths (Malcolm 1959). Fish tissue accessioned for all arripid individuals examined by us may be valuable for future investigation of this hypothesis. Irrespective of the outcome, the distinct life-history differences between *A. truttaceus* and *A. trutta* should be acknowledged by fishery managers and it is recommended that the relative proportions of these species in commercial catches should be assessed.

Table 1. Metazoan parasite fauna from *Arripis georgianus* collected in the present study and from published literature

Under infection parameters, prevalence is expressed as a percentage (%); mean intensity is followed in parentheses by maximum intensity. SA, South Australia; and WA, Western Australia

Parasite	Habitat	Reference	Location	Infection parameters	Accession number
<b>Cestoda</b>					
Lacistorhynchidae					
<i>Callitetrarhynchus gracilis</i> Rudolphi, 1819	Viscera	A	SA <sup>B</sup>	13; 4 (13)	AHC 45416
Order: Tetraphyllidea Carus, 1863					
Type 1	Intestine	A	SA <sup>B</sup>	1; 1 (1)	AHC 29765
Type 2	Intestine	A	SA <sup>B</sup>	1; 1 (1)	AHC 29766
Tentaculariidae					
<i>Nybelinia thyrssites</i> Korotaeva, 1971	Intestine	A	SA <sup>B</sup>	1; 1 (1)	AHC 29788
<b>Digenea</b>					
Acanthocolpidae					
<i>Monostephanostomum georgianum</i> Kruse, 1979	Digestive tract	Bray and Cribb (2002)	SA, SA <sup>B</sup>	11; 2 (7)	AHC 29759–60
<i>Monostephanostomum manteri</i> Kruse, 1979	Digestive tract	Braicovich and Timi (2008)	SA, SA <sup>B</sup>	10; 3 (11)	AHC 29761
Bucephalidae					
<i>Telorhynchus arripidis</i> Crowcroft, 1947	Digestive tract	A	SA <sup>B</sup>	42; 32 (653)	AHC 29764
<b>Hemiuridae</b>					
<i>Elytrophalloides humerus</i> Bray, 1990	Stomach, intestine	A	SA <sup>B</sup>	11; 1 (4)	AHC 29755–58
<i>Erilepturus tiegsi</i> Woolcock, 1935	Digestive tract	Bray (1990)	WA, SA <sup>B</sup>	6; 2 (3)	AHC 29754 AHC 29789
<b>Opcoeloidae</b>					
<i>Pseudopocoeloides arripi</i> Aken'Ova <i>et al.</i> , 2009	Intestine, caeca	Aken'Ova <i>et al.</i> (2009)	WA, SA, SA <sup>B</sup>	30; 3 (21)	AHC 29762–3
<b>Monogenea</b>					
Microcotylidae					
<i>Microcotyle arripis</i> Sandars, 1945	Gills	Sandars (1945)	WA, SA, SA <sup>B</sup>	72; 5 (21)	AHC 29751–53 AHC 29883–84 BMNH 2009.12.28.1–2 USNPC 102673.00–102675.00
<b>Nematoda</b>					
Anisakidae					
<i>Hysterothylacium</i> sp.	Digestive tract	A	SA <sup>B</sup>	3; 11 (51)	AHC 45417–18
Camallanidae					
<i>Procammallanus</i> sp.	Digestive tract	A	SA <sup>B</sup>	1; 40 (40)	AHC 45421
Philometridae					
<i>Philometra</i> sp.	Gonad	A	SA <sup>B</sup>	2; 2 (3)	AHC 45422–23
Trichinellidae					
<i>Capillaria</i> sp.	Digestive tract	A	SA <sup>B</sup>	2; 1 (2)	AHC 45419–20
<b>Branchiura</b>					
Argulidae					
<i>Argulus diversicolor</i> Byrnes, 1985	Skin	Catalano and Hutson (2010)	SA <sup>B</sup>	2; 2 (3)	BMNH 2009.261
<b>Copepoda</b>					
Caligidae					
<i>Caligus punctatus</i> Shiino, 1955	Gills	Catalano and Hutson (2010)	SA <sup>B</sup>	1; 1 (1)	C6814

<sup>A</sup>New host record; <sup>B</sup>From the current study.

Table 2. Metazoan parasite fauna from *Aripis truttae* collected in the present study and from published literature

Parasite	Habitat	Reference	Location	Infection parameters	Accession number
Myxozoa					
Ceratomyxidae					
<i>Ceratomyxa</i> sp.	Gall bladder	A	SA <sup>B</sup>	33; NR	G465430–31
Cestoda					
Lacistorhynchidae					
<i>Callitetrarhynchus gracilis</i> Rudolphi, 1819	Viscera	Beveridge and Campbell (1996)	Vic., SA <sup>B</sup>	16; 5 (25)	AHC 45424
Order: Tetrathyllidea Camus, 1863					
Type 1	Intestine, caeca	A	SA <sup>B</sup>	3; 1 (1)	AHC 29777–78
Type 3	Caeca	A	SA <sup>B</sup>	12; 3 (12)	AHC 29779
Tentaculariidae					
<i>Nybelinia thyrssites</i> Korotaeva, 1971	Stomach	Beveridge and Campbell (1996)	SA, SA <sup>B</sup>	6; 1 (1)	AHC 29780 AHC 45425
Digenea					
Acanthocolpidae					
<i>Monostephanostomum georgianum</i> Kruse, 1979	Digestive tract	A	SA <sup>B</sup>	16; 6 (21)	AHC 29773–74
<i>Monostephanostomum manteri</i> Kruse, 1979	Digestive tract	A	SA <sup>B</sup>	4; 1 (2)	AHC 29912–14
Bucephalidae					
<i>Telorhynchus arripidis</i> Crowcroft, 1947	Digestive tract	A	SA <sup>B</sup>	40; 169 (2351)	AHC 29775 AHC 29791
Hemiuridae					
<i>Eritepturus tiegsi</i> Woolcock, 1935	Digestive tract	A	SA <sup>B</sup>	25; 2 (5)	AHC 29770–72 AHC 29790
<i>Elytrophalloides humerus</i> Bray, 1990	Stomach	A	SA <sup>B</sup>	1; 1 (1)	AHC 29792
<i>Elytrophallus</i> sp. Manter, 1940	Gills – artefact	A	SA <sup>B</sup>	1; 1 (1)	AHC 29793
Lepocreadiidae					
<i>Opechona kahawai</i> Bray & Cribb, 2003	Intestine, caeca	Bray and Cribb (2003)	Tas., SA <sup>B</sup>	12; 4 (12)	AHC 29776
Monogenea					
Microcotylidae					
<i>Kahawaita truttae</i> Lebedev, 1969	Gills	A	SA <sup>B</sup>	39; 3 (14)	AHC 29767–69, 29882, 29885–86 BMNH 2009.12.28.3–4 USNPC 102676.00–102677.00
Nematoda					
Anisakidae					
<i>Contracaecum</i> sp.	Stomach, caeca	A	SA <sup>B</sup>	6; 1 (1)	AHC 45428–29
<i>Hysterothylacium</i> sp.	Digestive tract	A	SA <sup>B</sup>	33; 2 (14)	AHC 45426–27
Philometridae					
<i>Philometra</i> sp.	Stomach	A	SA <sup>B</sup>	3; 1 (1)	AHC 45430
Branchiura					
Argulidae					
<i>Argulus diversicolor</i> Byrnes, 1985	Skin	Catalano and Hutson (2010)	SA <sup>B</sup>	1; 1 (1)	C6823
Copepoda					
Calligidae					
<i>Calligus bonito</i> Wilson, 1905	Gills	Catalano and Hutson (2010)	SA <sup>B</sup>	13; 2 (3)	C6819
<i>Calligus longipedis</i> Bassett-Smith, 1898	Gills	Catalano and Hutson (2010)	SA <sup>B</sup>	3; 3 (4)	C6815–6
<i>Calligus punctatus</i> Shino, 1955	Skin	Catalano and Hutson (2010)	SA <sup>B</sup>	3; 7 (9)	C6817–8

<sup>A</sup>New host record; <sup>B</sup>From the current study.

Table 3. Metazoan parasite fauna from *Arripis truttia* collected in the present study and from published literature

Under infection parameters, prevalence is expressed as a percentage (%); mean intensity is followed in parentheses by maximum intensity. GAB, Great Australian Bight; NA, not applicable; NR, not recorded; NSW, New South Wales; NZ, New Zealand; PR, previously recorded from the literature but not found in the current study; SA, South Australia; Tas., Tasmania; TS, Tasman Sea; and Vic., Victoria

Parasite	Habitat	Reference	Location	Infection parameters	Accession number
<b>Myxozoa</b>					
<b>Ceratomyxidae</b>					
<i>Ceratomyxa arripica</i> Su & White, 1994	Gall bladder	Su and White (1994)	Tas.	NA	PR
<i>Leptotheca annulata</i> Meglitsch, 1960	Gall bladder	Meglitsch (1960)	NZ	NA	PR
<i>Leptotheca minima</i> Meglitsch, 1960	Gall bladder	Meglitsch (1960)	NZ	NA	PR
<b>Cestoda</b>					
<b>Lacistorhynchidae</b>					
<i>Callitetrarhynchus gracilis</i> Rudolphi, 1819	NR	Beveridge and Campbell (1996)	Vic.	NA	PR
Order: Tetraphyllidea Carus, 1863					
Type 3	Intestine	A	NSW <sup>B</sup>	11; 19(19)	AHC 29803
<b>Tentaculariidae</b>					
<i>Nybelinia</i> sp. larva Baker, 1971	Stomach	Baker (1971)	NZ	NA	PR
<b>Digenea</b>					
<b>Acanthocolpidae</b>					
<i>Monostephanostomum georgianum</i> Kruse, 1979	Digestive tract	A	SA <sup>B</sup>	22; 3(4)	AHC 29796-97
<i>Monostephanostomum manteri</i> Kruse, 1979	Intestine, caeca	Kruse (1979) <sup>C</sup> ; Bray and Cribb (2002)	SA <sup>C</sup> , Tas., SA <sup>B</sup>	22; 3(3)	AHC 29798
<b>Bucephalidae</b>					
<i>Prosorhynchus</i> sp. Manter, 1954	NR	Manter (1954)	NZ	NA	PR
<i>Telorhynchoides longicollis</i> Lebedev, 1968	NR	Lebedev (1968) <sup>C</sup>	GAB <sup>C</sup>	NA	PR
<i>Telorhynchus arripidis</i> Crowcroft, 1947	Digestive tract	Crowcroft (1948); Manter (1954); Lebedev (1968) <sup>C</sup>	Tas., NZ, GAB <sup>C</sup> , TS, NSW <sup>B</sup>	33; 4(6)	AHC 29800-01
<i>Telorhynchus kahawai</i> Lebedev, 1968	NR	Lebedev (1968) <sup>C</sup>	GAB <sup>C</sup>	NA	PR
<i>Telorhynchus peachei</i> Lebedev, 1968	NR	Lebedev (1968) <sup>C</sup>	GAB <sup>C</sup> , TS	NA	PR
<b>Hemiuridae</b>					
<i>Eriopterurus tiegsi</i> Woolcock, 1935	Stomach, caeca	Yamaguti (1958)	Vic., SA <sup>B</sup>	33; 1(2)	AHC 29794-95
<i>Hemiurus (Anahemiurus)</i> sp. Manter, 1947	Digestive tract	Baker (1971)	NZ	NA	PR
<i>Parahemiurus arripidis</i> Lebedev, 1971	NR	Bray and Cribb (2002)	NZ	NA	PR
<i>Parahemiurus</i> sp. Lebedev, 1971	NR	Lebedev (1971)	NZ	NA	PR
<b>Lepoecreadiidae</b>					
<i>Opechona kahawai</i> Bray & Cribb, 2003	Caeca	Bray and Cribb (2003)	Tas., NSW <sup>B</sup>	11; 1(1)	AHC 29802

(Continued)

Table 3. (Continued)

Parasite	Habitat	Reference	Location	Infection parameters	Accession number
Opcoelidae					
<i>Pseudopocoeloides arripji</i> Aken'Ova <i>et al.</i> , 2009	Stomach	<sup>A</sup>	NSW <sup>B</sup>	11; 1(1)	AHC 29799
Syncoelidae					
<i>Syncoelium filiferum</i> Sars, 1885	Gills	Rohde <i>et al.</i> (1980)	NZ	NA	PR
Monogenea					
Microcotylidae					
<i>Kahawata truttae</i> Lebedev, 1969	Gills	Dillon and Hargis (1965); Lebedev (1969) <sup>C</sup>	NZ, GAB <sup>C</sup> , SA <sup>B</sup> , NSW <sup>B</sup>	48; 2(5)	AHC 29781–82
Nematoda					
Anisakidae					
<i>Anisakis</i> sp. larva	Viscera	Hewitt and Hine (1972)	NZ	NA	PR
<i>Contracaecum aduncum</i> Rudolphi, 1802	Digestive tract	Baker (1971)	NZ	NA	PR
<i>Contracaecum</i> sp.	Stomach, intestine, body cavity	Hewitt and Hine (1972)	NZ	NA	PR
<i>Hysterothylacium</i> sp.	Digestive tract	<sup>A</sup>	NSW <sup>B</sup>	11; 2(2)	AHC 45431
Annelida					
Piscoliolidae					
<i>Austrobdella translucens</i> Badham, 1916	Tail fin	Bolton <i>et al.</i> (2005) <sup>C</sup>	SA <sup>C</sup>	NA	PR
Copepoda					
Calligidae					
<i>Caligus bonito</i> Wilson, 1905	Gills	Catalano and Hutson (2010)	NSW <sup>B</sup>	9; 2(2)	C6821
<i>Caligus kahalawai</i> Jones, 1988	Body surface	Jones (1988)	NZ	NA	PR
<i>Caligus pelamydis</i> Hewitt, 1963	Gills	Jones (1988)	NZ, NSW <sup>B</sup>	4; 1(1)	C6822
<i>Caligus punctatus</i> Shiino, 1955	Gills	Catalano and Hutson (2010)	NSW <sup>B</sup>	4; 1(1)	C6820
Chondracanthidae					
<i>Chondracanthus australis</i> Ho, 1991	Gills	Ho (1991)	NZ	NA	PR
Ergasilidae					
<i>Abergasilus amplexus</i> Hewitt, 1978	Gills	Hewitt (1978)	NZ	NA	PR
Isopoda					
Cymothoidae					
<i>Codonophilus imbricatus</i> Fabricius, 1787	Tongue	Baker (1971)	NZ	NA	PR
<i>Nerocila orbigny</i> Guérin-Méneville, 1832	Body	Hewitt and Hine (1972)	NZ	NA	PR

<sup>A</sup>New host record; <sup>B</sup>From current study; <sup>C</sup>ambiguous parasite-host record (see Results and discussion).

### Metazoan parasite fauna of the Arripidae

We recovered a total of 8455 parasites, 4959 from *A. truttaceus*, 3417 from *A. georgianus* and 79 from *A. trutta*. A diverse community of metazoan parasites infects arripids, with representation across a wide spectrum of parasite taxa in five phyla (Tables 1–3). In all, 17 parasite species were recorded and identified from *A. georgianus*, 20 from *A. truttaceus* and 12 from *A. trutta* (Tables 1–3). Including previous accounts of parasite species from *A. trutta*, together with those recorded here, a total of 34 species is now documented from this species (Table 3). Of the parasites recorded in our study from all three arripids, 71% represent new host records. Poulin and Morand (2000) stated that a central limitation in the current parasite-diversity estimates is that not all living host species are identified and/or described. The three arripid species examined here were described in the early 1800s, so discovery of such a high percentage of new host records is surprising. Host species that are well known but not examined carefully in sufficient numbers also contribute significantly to underestimates of parasite diversity.

We also recorded four parasitic crustacean species that have been associated with mass mortality of cultured fishes overseas. The importance of this discovery for southern Australian finfish sea-cage aquaculture is reviewed elsewhere (Catalano and Hutson 2010). Larval tetraphyllidean metacestodes, which are difficult to classify using basic morphology (e.g. Chambers *et al.* 2000), were the only parasites not identified to genus or below. They were designated as Types 1–3 on the basis of differences in length, width and position of the four trilobulate bothridia and apical sucker.

### Similarities and differences in parasites among *Arripis* spp.

Our results showed similarities as well as clear differences in the parasite assemblages and parasite species richness among *A. georgianus*, *A. truttaceus* and *A. trutta* (Tables 1–3, respectively), which can be explained by a range of host attributes. Similarities and differences in parasite assemblages could be due to host phylogeny because shared parasite species may be inherited from a common ancestor (Poulin and Rohde 1997), whereas distinct parasite species may reflect host speciation over time (Guegan *et al.* 1992). *A. trutta* is more closely related to *A. truttaceus* than to *A. georgianus* (see Ward and Holmes 2007), a fact mirrored by *A. trutta* and *A. truttaceus* having greater similarities in parasite fauna than *A. trutta* and *A. georgianus* (compare Tables 1–3). For example, *Kahawaia truttae* (Monogenea), *Contraecum* sp. (Nematoda) and Tetraphyllidea Type 3 (Cestoda) are shared by *A. truttaceus* and *A. trutta* (Tables 2, 3) whereas *A. georgianus* has its own monogenean species (*Microcotyle arripis*) and different nematode and cestode species (Table 1).

Nonetheless, differences observed may simply be attributed to spatial scale restrictions. For instance, Marcogliese (2002) and Poulin (2007) stated that parasite distributions are superimposed on distributional patterns of free-living animals that participate as hosts in the parasite's lifecycle. Price and Clancy (1983) asserted that the geographical range of a host comprises multiple habitats, with each contributing additional parasite

species to the overall fauna. Our study, together with published literature, sampled *A. georgianus*, *A. truttaceus* and *A. trutta* from only a small part of their total geographical and size range. Therefore, there is no information on the contribution of additional geographical habitats and fish ontogeny to the overall parasite fauna of each arripid species.

Parasite species richness was greatest for *A. trutta* (Table 3). This may be because *A. trutta* is the most widely distributed arripid species compared with *A. georgianus* and *A. truttaceus* and, therefore, has the potential to encounter greater numbers of parasite species across its broad range (e.g. Guegan *et al.* 1992). Nonetheless, numerous studies have also shown that larger fish hosts have greater parasite abundance and parasite species richness than do small hosts (e.g. Kearns 1967; Ho 1991; Guegan *et al.* 1992; Poulin 1997). Larger host bodies provide more space and a greater diversity of niches for parasites to colonise (Guegan *et al.* 1992; Lo *et al.* 1998). *A. trutta* and *A. truttaceus* reach a larger maximum size than *A. georgianus* (see Gomon *et al.* 2008), possibly explaining the observed result that the first two host species have a greater parasite species richness compared with the latter. Nevertheless, *A. trutta* is historically the more studied arripid species for parasites and therefore sampling bias may also explain this observation.

Food web structure is hypothesised to affect transmission of parasite species (Marcogliese and Cone 1997; Marcogliese 2002). Therefore, diet may influence the parasite assemblage associated with an individual (Braicovich and Timi 2008). Diet and feeding studies for all three arripid species are limited. Information focuses on the different prey items consumed by *A. truttaceus* and *A. trutta* corresponding to differences in gill raker anatomy, with little information on the diet for *A. georgianus* (K. Jones, pers. comm.). In our study, the highest maximum infection intensities were for the endoparasitic digenetic *Telorhynchus arripidis* from *A. georgianus* and *A. truttaceus* (Tables 1, 2). Infection by *T. arripidis* occurs by ingesting intermediate hosts harbouring encysted metacercariae (Lo *et al.* 1998), suggesting prey containing these infective stages comprise a major part of the diet of *A. georgianus* and *A. truttaceus*.

### Ambiguous parasite–host records

Several ambiguous parasite–host records for *A. trutta* in SA waters were noted (Table 3). These may have resulted from confusion surrounding discrimination and identification of *A. trutta* and *A. truttaceus* in previous literature (see Paulin 1993 for a review), but are also a result of authors failing to acknowledge methods used to identify hosts (e.g. Lebedev 1968, 1969; Kruse 1979; Bolton *et al.* 2005; Table 3). Before our study, the known western limit for *A. trutta* was Port Phillip Bay (Hutson *et al.* 2004), rendering previous parasite–host identifications for *A. trutta* in SA ambiguous. It is clear from our study and from previous reports (e.g. Hoedt and Dimmlich 1994; Jones and Westlake 2003) that *A. truttaceus* is the common Australian salmon species in SA waters and that most previous ambiguous records are likely to be *A. truttaceus* misidentified as *A. trutta*. These doubtful reports plainly demonstrate the critical importance of stating the methods used to identify hosts (e.g. keys and/or literature consulted, authorities asked for advice) as



well as the value of lodging host specimens and/or host-tissue samples in established permanent collections for verification and future study.

## Conclusions

The present study has provided new information on the distribution of *A. trutta*, with validation of morphological identification (based on gill raker number and morphology on gill arch one) via molecular genetics. The extension of the western range by ~1000 km, from Port Philip Bay, Victoria, to Almonta Beach, CB National Park, SA, has implications for the management of arripids in southern Australian waters. We have here reported 35 new metazoan parasite–host records from three arripid species in southern Australian waters. Our thorough documentation of parasite assemblages for *A. georgianus*, *A. truttaceus* and *A. trutta* provides valuable baseline data that will permit the detection of changes in parasite fauna over time.

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