

1 **A method for identifying Sound Scattering Layers and extracting key**  
2 **characteristics**

3

4 **Scattering Layer identification and description**

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6 Word count: ~7650

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25 **Abstract**

26

27 1. Mid-trophic level water-column (pelagic) marine communities comprise millions  
28 of tonnes of zooplankton and micronekton that form dense and geographically-  
29 extensive layers, known as Sound Scattering Layers (SSLs) when observed  
30 acoustically. SSLs are ubiquitous in the global ocean and individual layers can  
31 span entire ocean basins. Many SSLs exhibit clear diel vertical migration  
32 behaviour. Vertical migrations contribute substantially to the 'biological pump',  
33 such that SSLs have important global biogeochemical roles: SSLs are important  
34 conduits for vertical energy and nutrient flow. Ship-based remote sensing of SSLs  
35 using acoustic instruments (echosounders) enables their shape and density to be  
36 quantified, but despite SSLs being discovered in the 1940s, there is no consistent  
37 method for identifying or characterizing SSLs. This hampers ecological and  
38 biogeographical studies of SSLs.

39

40 2. We have developed an automated and reproducible method for SSL identification  
41 and characterization, the Sound Scattering Layer Extraction Method (SSLEM). It  
42 functions independently of echosounder frequency and the spatial scale (vertical  
43 and horizontal) of the data. Here we demonstrate the SSLEM through its  
44 application to identify SSLs in data gathered to a depth of 1000 m using 38 kHz  
45 hull-mounted echosounders in the South West Indian Ocean and Tasman Sea.

46

47 3. SSLs were identified in the water-column as horizontally extensive echoes that  
48 were above background noise. For each identified SSL a set of 9 quantitative 'SSL  
49 metrics' (describing their shape, dynamics and acoustic backscattering

50 distribution) were determined, enabling inferences to be made concerning the  
51 spatial arrangement, distribution and heterogeneity of the biological community.  
52 The method was validated by comparing its output to a set of visually-derived SSL  
53 metrics that were evaluated independently by 8 students. The SSLEM  
54 outperformed the by-eye analysis, identifying three times the number of SSLs and  
55 with greater validity; 95% of SSLs identified by the SSLEM were deemed valid,  
56 compared to 75% by the students.

57

58 4. In the same way that data obtained from satellites has enabled the study and  
59 characterization of global phytoplankton distribution and production, we  
60 envisage that the SSLEM will facilitate robust, repeatable and quantitative  
61 analysis of the growing body of SSL observations arising from underway acoustic  
62 observations, enhancing our understanding of global ocean function.

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71 **Key-words:**

72 Sound Scattering Layer Extraction Method; SSLEM; Deep Scattering Layers; biological  
73 layers; marine acoustics; SSL metrics; mid-trophic level; pelagic ecology; biological  
74 communities; diel vertical migration

## 75 **Introduction**

76

77 Sound Scattering Layers (SSLs) or Deep Scattering Layers (DSLs) are vertically discrete  
78 (100s of m or less) water column aggregations of organisms that can extend  
79 horizontally over 1000s of km (Kloser et al. 2009). The layers are comprised of pelagic  
80 organisms (organisms of the water column, as opposed to benthic organisms that live  
81 on or in the seabed), primarily zooplankton and small fish (cm to 10s cm), living  
82 together in distinct communities. When insonified, these organisms produce a distinct  
83 echo that, depending upon depth and incident acoustic frequency, can stand out  
84 prominently as scattering layers above background noise (Simmonds & MacLennan  
85 2005). Such layers have been known since the mid 20<sup>th</sup> century when naval sonars  
86 detected signals thought initially to be echoes from the sea bed: the depths of the echoes  
87 however changed with time of day and it became apparent that these 'false bottoms'  
88 were in fact biological in origin (Brierley 2014).

89

90 The organisms that comprise SSLs are responsible for the transport of vast quantities of  
91 carbon (increasing particle export by up to 40% - Bianchi 2013) from the surface to the  
92 deep sea (the biological pump) via Diel Vertical Migration (DVM), the largest known  
93 daily migration of biomass on the planet (Hays 2003). They thus play an important role  
94 in atmosphere-ocean interactions, and in the global biogeochemical cycle. SSL scales can  
95 vary from the micro, at vertical resolutions of centimetres over timescales of a fraction  
96 of a minute (Holliday et al. 2003; McManus et al. 2003, 2005) to pan-oceanic (Anderson,  
97 Brierley, & Armstrong 2004; Kloser et al. 2009; Irigoien et al. 2014). The former, so  
98 called 'microlayers' provide an explanation for the 'paradox of the plankton' where high  
99 species diversity occurs in what at first glance may appear to be a homogenous volume

100 of water. Vertical structure means that the water column is in fact far from  
101 homogenous. Its physical properties can change dramatically over just a few cm's or  
102 meters enabling the discrete formation of aggregates at specific depth ranges  
103 (Longhurst 1998). In this way, SSLs show the water column divided into multiple  
104 vertically discrete habitats. Despite the importance of SSLs to ocean and earth-system  
105 function, there is no accepted standard method for identifying or classifying them. This  
106 in turn has hampered comparative or integrative studies of SSLs.

107

108 There are parallels between the pelagic and tropical rain forest ecosystems, in as far as  
109 both have vertical (depth) layered structure that with increasing depth is increasingly  
110 light-limited. The sea surface, like the forest canopy, can be studied remotely by  
111 satellites and yield estimates of biomass and primary production (PP) (Longhurst,  
112 2003; Anderson, 2012). The sea bed is a physically-fixed entity and organisms living  
113 there, like organisms and vegetation of the forest floor, are amenable to study because  
114 they are constrained by a two dimensional environment. The ocean interior, however, is  
115 physically dynamic, and its inhabitants – pelagic organisms including zooplankton and  
116 fish – have freedom to move in three dimensions so the dynamics of the sometimes-  
117 dense layers of mid-trophic level communities are poorly understood (Lehodey,  
118 Murtugudde & Senina, 2010) . Developments in forest sampling can perhaps guide  
119 developments in ocean sampling.

120

121 In an effort to improve understanding of rainforest interiors, the 'RAINFOR' project  
122 (Malhi et al., 2002) that was established over a decade ago initialised a network of  
123 sampling sites for Amazonian rainforest ecosystems. As part of the initiative, field  
124 measurements of the forest interior were taken and the data related back to satellite

125 information in order to gain a more complete picture of the ecosystem and to provide  
126 the capability to validate remotely-sensed data. Since the rainforest covers less than 7%  
127 of the earth's surface (Bierregaard et al., 1992; Wilson, 1994), a network of sites can  
128 provide a representative sample. The ocean on the other hand, covers over 71% of the  
129 earth's surface and therefore makes it extremely difficult, logistically and financially to  
130 study the pelagic community by insitu biological sampling alone. The required greater  
131 spatial coverage for the ocean can be achieved by using active acoustic sampling  
132 techniques (scientific echosounding) to rapidly observe large volumes of the ocean,  
133 from ship-based instruments routinely used on both research and fishing vessels. These  
134 acoustic data can help reveal the spatial structure of pelagic communities: many marine  
135 organisms scatter sound waves in a characteristic fashion (dependant on the frequency  
136 of the incident wave and anatomy of the organism) such that 'remote sensing' by  
137 echosounder can provide community insight. These data could then be linked back to  
138 remotely sensed PP data at the surface, acquired from online resources such as the  
139 National Oceanographic Data Centre (NODC, [www.nodc.noaa.gov](http://www.nodc.noaa.gov)) and validated by  
140 biological point samples, available from, for example, the Ocean Biogeographical  
141 Information System (OBIS, [www.iobis.org](http://www.iobis.org)). Acoustic survey data already exist in vast  
142 quantities, with wide geographic coverage, leading to the possibility that a method  
143 capable of identifying and characterising pelagic communities, found within SSLs,  
144 would, akin to the RAINFOR project, potentially enable deep ocean processes to be  
145 inferred from satellite observation of the surface, yielding a more complete  
146 understanding of ocean ecosystem function. To achieve this, first a repeatable technique  
147 to identify and parameterise SSLs is required.

148

149

150 Some quantitative research has already been carried out on SSLs. Regional structure of  
151 SSLs has been identified by comparing total water-column backscattering strength at  
152 sites across the Pacific and Atlantic Oceans, in the Caribbean, Labrador, Norwegian and  
153 Mediterranean Seas and in Baffin Bay (Chapman et al. 1974) and also in the Atlantic and  
154 North Western Pacific using depth-frequency structure of backscattering strength  
155 (Andreeva, Galybin, & Tarasov 2000; Tarasov 2002). Biomass estimates of SSLs have  
156 also been made at the basin scale using both echo counting and echo integration  
157 techniques, in the Tasman Sea for example (Kloser et al. 2009). More recently, a method  
158 was developed to extract SSLs (Cade and Benoit-Bird 2014), that required input  
159 parameters such as an acoustic threshold intensity and minimum separation distances  
160 between SSLs to be defined a priori.

161

162 Research on SSLs has, however, typically been qualitative. The depth structure of SSLs  
163 has been observed to vary over large spatial scales in both longitude and latitude  
164 (Kloser et al. 2009), across ocean basins (Anderson, Brierley, & Armstrong 2004) in the  
165 Irminger Sea and across fronts (Nicol et al. 2000; Kawaguchi et al. 2010) in the  
166 Southern Ocean. Studies at oceanic features have also shown characteristic behaviour,  
167 such as bulges in SSLs at continental shelves (Jarvis et al. 2010) off East Antarctica and  
168 bowl-like SSL features forming under eddy structures (Godø et al. 2012) in the  
169 Norwegian Sea.

170

171 Conversely, discrete, biological aggregations - schools, shoals and swarms - have been  
172 quantitatively defined. The fisheries acoustic community has wrestled for years over  
173 the question of what, as seen in an acoustic record, is a school, and some standard  
174 identification and description protocols have been agreed (Reid and Simmonds 1993;

175 Reid 2000). The importance of particular school metrics for school identification has  
176 differed between studies; for example Coetzee (2000), using the Shoal Analysis and  
177 Patch estimation system algorithm (SHAPES: Barange 1994), identified morphological  
178 aspects to be the chief descriptor; Lawson (2001), identified school energetics and  
179 water-column position as the most important parameters.

180

181 In spite of previous work, no standardised objective approach exists for defining SSLs,  
182 rendering comparisons between studies and between water-column communities and  
183 the environment difficult beyond the merely descriptive. The lack of a consistent  
184 analytical approach was identified by Handegard et al. (2013) as hindering marine  
185 ecosystem monitoring and management.

186

187 Our overarching goal was to develop a standardised analysis method to extract  
188 biological layers consistently from underway-acoustic survey data. We illustrate the  
189 utility of our method here by identifying SSLs from data observed at 38 kHz in the South  
190 West Indian Ocean and Tasman Sea. Application of this method will lead to a better  
191 understanding of mid-trophic level communities, within and across oceanic boundaries  
192 at varying spatial and temporal scales.

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200 **2 Materials and Method**

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202 **2.1 Acoustic data**

203

204 The acoustic data used in this study were all 38 kHz data and were obtained from the  
205 Integrated Marine Observing System data centre (IMOS, [www.imos.au](http://www.imos.au), downloaded on  
206 1<sup>st</sup> June 2013). 38 kHz data were suitable for SSL observations because the moderately  
207 low attenuation rate (5 - 10 dB/km: Ainslie and McColm 1998) enables deep water-  
208 column penetration (up to 1500m) and because the wavelength is appropriate for  
209 detection of many of the fish and plankton species of the order of cm's that inhabit the  
210 layers. Data had been collected by research vessels (RV) *Southern Surveyor* and *Aurora*  
211 *Australis*, as well as several fishing vessels (FVs) including the *Southern Champion*,  
212 *Janas*, *Rehua*, *Austral Leader II* and *Will Watch*. Data were granted by the Marine  
213 National Facility and processed by the Commonwealth Scientific and Industrial  
214 Research Organisation (CSIRO) Oceans and Atmosphere Flagship as part of the IMOS  
215 Bio-Acoustic Ships of Opportunity (BASOOP) Program. The data totalled 24 transects  
216 covering wide areas of the South West Indian Ocean and Tasman Sea (Fig. 1). Transect  
217 length ranged from 200 NM to 1800 NM and included 24hr (day/night) coverage across  
218 all seasons between 2009 and 2012. The spatial coverage of the tracks included 2 of the  
219 4 major global ocean biomes as described by Longhurst (1998) - the Trades and  
220 Westerlies - and also spanned major frontal zones and boundaries including the  
221 Subtropical Convergence Zone (STCZ) and the Polar Front (PF).

222

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225

226

[FIGURE 1]

227

228 We pre-processed IMOS data by removing dropped pings (a ping is a single acoustic  
229 transmit/receive cycle; typical ping rate was 0.5 Hz) and noise spikes (caused for  
230 example by violent ship's motion in rough seas), and partitioned data into separate  
231 day/night segments, bounded by sunrise/sunset. In instances where, for example the  
232 echosounder had been turned off for short periods, or the vertical sampling resolution  
233 of the echosounder was changed, data were further segmented. An acoustic image, or  
234 echogram, was created for each segment. The acoustic image was as a two-dimensional  
235 array of backscatter values on a depth v time (or space) grid. Each cell in the image, an  
236 acoustic pixel, had an associated timestamp, geographical position and depth. For each  
237 image, echo intensity in the form of Mean Volume Backscattering Strength (MVBS -  
238 Simmonds & MacLennan, 2005) was calculated at a cell resolution of 5m in depth (from  
239 the surface down to 1000m) by 1 minute in time.

240

## 241 **2.2 SSL extraction method (SSLEM)**

242

243 Our objective was to provide a method that would function over the range of bio-  
244 acoustical echosounder frequencies in common (and likely future) use, over horizontal  
245 scales from bays to oceans, and on vertical scales that encompass microlayers (cm;  
246 Holliday et al., 2003) upwards to tens and hundreds of m. The common observational  
247 frequency band (18 to 200 kHz) spans the Rayleigh and geometric scattering regions for  
248 most zooplankton and nekton. This means that small changes in frequency can result in  
249 large changes in backscattering intensity and hence in SSL descriptors. Layers only

250 become apparent acoustically when they can be distinguished from background noise  
251 (sufficiently high signal-to-noise ratio – SNR). SNR is a function of organism packing  
252 density, acoustic Target Strength, depth, insonification frequency and power, and  
253 environmental conditions (Simmonds & MacLennan, 2005). SSL appearance may also  
254 be influenced by sampling resolution (Korneliussen et al., 2008): for transect data  
255 resolution is determined by ping rate, beam angle, depth and ship speed. The  
256 geographic scale of data is an important consideration as there are many oceanic  
257 processes that occur over different spatial and temporal scales, from micro-turbulence  
258 to decadal oscillations. A robust general method should be capable of resolving features  
259 of interest at the scale of the study being conducted, and for the organisms of interest in  
260 the environment in which they exist.

261

### 262 **2.2.1 Identification of SSLs**

263

264 The SSL extraction method (SSLEM) is based upon detection of a contrast in MVBS  
265 (Berge et al. 2014) between pixels within SSLs (relatively high MVBS signal) and  
266 background pixels outside (relatively low MVBS noise). For a simple SSL analysis, using  
267 a window of depth range Z and time/space extent X, one could identify a vertically  
268 ‘static’ SSL surrounded by empty water by selecting pixels for which MVBS intensities  
269 were greater than the mean,  $\mu$ , over the entire window. Under such a scheme, for any  
270 acoustic pixel (px) within the analysis window,

271 Equ. 1 
$$ssl = \begin{cases} 1, & px > \mu \\ 0, & px \leq \mu \end{cases}$$

272 where *ssl* is a Boolean variable, taking a value of 1 for pixels that are deemed to belong  
273 to an SSL and 0 for those that are not. This simple process, useful as an introduction to

274 the method, assumes that the SSL is completely contained by the analysis window and  
 275 the surroundings are made up of pixels with low MVBS that is attributable to  
 276 background noise. This may not be the case; for example, a transition between depth  
 277 intervals that exhibit a difference in background noise (inherently caused by time-  
 278 varied gain (TVG) amplification of background noise) would yield a layer-like boundary  
 279 of *ssl* pixels. To ensure that SSLs were surrounded by lower intensity MVBS (both  
 280 towards the surface and the seabed), the depth interval of the analysis window was  
 281 divided into two equal values,  $d_1$  and  $d_2$ , and pixels were only deemed a SSL pixel (*ssl*  
 282 value of 1) when their MVBS value was larger than both the MVBS means,  $\mu_1$  and  $\mu_2$ ,  
 283 over each of the two regions of the split window (Fig. 2a), yielding a new equation for  
 284 *ssl*:

285 Equ. 2 
$$ssl = \begin{cases} 1, & (px > \mu_1) \text{ AND } (px > \mu_2) \\ 0, & \text{otherwise} \end{cases}$$

286 where  $\mu_1$  and  $\mu_2$ , are calculated over the regions X by  $d_1$  and X by  $d_2$  respectively, and  
 287 where  $d_1$  is equal to  $d_2$ .

288

289 [FIGURE 2]

290

291 In practice, using a fixed analysis window does not capture all SSLs, since they are  
 292 rarely vertically static: they may for instance oscillate with internal wave activity or  
 293 migrate vertically. To accommodate this, an analysis column one pixel wide was used  
 294 instead of a window. The column was moved pixel-by-pixel through the image  
 295 evaluating the pixel at the centre of the column at each step, such that  $\mu_1$  and  $\mu_2$  were  
 296 calculated over the specific column (single point in time-series), not the entire window,  
 297 bounded either side of the central pixel by the depth ranges  $d_1$  and  $d_2$ . For constant

298 values of  $d_1$  and  $d_2$  this approach would only work if all SSLs had the same thickness and  
 299 were separated by distances larger than the size of  $d_1$  or  $d_2$ : this is not the case. To  
 300 overcome this problem the depth ranges  $d_1$  and  $d_2$ , for each pixel evaluated, were varied  
 301 in size from 2 pixels in height (this minimum, rather than 1, was used to avoid flooding  
 302 the image with incorrectly assigned *ssl* pixels when analysing highly variable or 'noisy'  
 303 images) up to the vertical extents of the image (Fig. 2b). Then, for any pixel within an  
 304 image, evaluated using a dynamic column,

305

306 Equ. 3 
$$ssl = \begin{cases} 1, & \left( \sum_{d_1=2}^{r-1} \sum_{d_2=2}^{R-r} (px > \mu_1) \text{ AND } (px > \mu_2) \right) > 0 \\ 0, & \text{otherwise} \end{cases}$$

307

308 where  $r$  is equal to the row number of the pixel being evaluated and  $R$  equal to the total  
 309 number of rows within the image; consequently, the first and last two rows of each  
 310 image are not processed. Each pixel thus has multiple opportunities (for varying  $d_1$  and  
 311  $d_2$  values) to be attributed an SSL pixel. This ensured that both vertically static and  
 312 migrant SSLs of varying thicknesses and separation distances would all be identified.

313

### 314 **2.2.2 Phantom SSLs and $SSL_{min}$**

315

316 On occasions when no SSL was present in the analysis column, pixels would sometimes  
 317 erroneously be designated as SSL pixels as a result of the backscatter from individual or  
 318 diffuse arrangements of organisms, tightly packed schools or swarms, variation of the  
 319 physical properties of sea water, or by natural variation inherent within the data. The  
 320 variable,  $SSL_{min}$ , which designated a fixed minimum horizontal extent for SSLs

321 (measured in space/time units), was therefore introduced to enable the identification of  
322 only those SSLs that were relevant to the scale of process being studied e.g. from ocean  
323 basin scale (large  $SSL_{min}$ ) down to krill swarms (small  $SSL_{min}$ : Watkins et al. 1990). In  
324 spite of this precaution, the natural variation within the data still sometimes produced  
325 SSLs. These incorrectly identified SSLs, termed 'phantom SSLs' (Fig. 3) were removed in  
326 post processing (Section 3.1) by analysis of SSL signal-to-noise ratios; SSLs were  
327 removed where the mean SSL MVBS (signal) was smaller than the maximum  
328 background MVBS value (noise) by analysing the pixels immediately surrounding the  
329 layers.

330

331 [FIGURE 3]

332

### 333 **2.2.3 Separation of merged SSLs**

334 The SSL pixels identified in Section 2.2.1 were used to generate an SSL mask (Fig. 4b)  
335 that bounded pixels that were connected, termed SSL features. These features were not  
336 classified as SSLs at this point, as a single feature could consist of several merged SSLs.  
337 Features smaller than  $SSL_{min}$  were removed and internal gaps smaller than  $SSL_{min}$   
338 within accepted features were filled (Fig. 4c).

339

340 [FIGURE 4]

341

342 Each feature was then segmented into individual SSLs which existed over a discrete  
343 depth range at each point along a time-series. This was achieved by implementing a  
344 region-based image segmentation process. A simple growing algorithm moved column  
345 by column through the image, initialising new regions (that would eventually grow into

346 individual SSLs) within features where SSLs merged or split (Fig. 5). SSLs identified in  
347 this process that were smaller than  $SSL_{min}$  were ignored and not analysed further.

348

349 [FIGURE 5]

350

#### 351 **2.2.4 Separation of vertically 'static' and migrant SSLs**

352

353 Vertically static SSLs were separated from upwardly or downwardly-migrating SSLs by  
354 application of a Change Point Analysis (CPA: Page 1954). CPA can detect the existence of  
355 multiple trends within a time series by analysing the cumulative deviation from the  
356 mean over time. The CPA was conducted using a time-series of the mean depth change,  
357  $\Delta Z$ , over a selected time interval,  $CPA_{int}$ , across each SSL (Fig. 6). The choice of  $CPA_{int}$  is  
358 related to the size of  $SSL_{min}$ . For a relatively large value of  $SSL_{min}$  (> 4 hours for example)  
359 a large  $CPA_{int}$  value can be chosen and will reduce the likelihood that undulating SSLs,  
360 caused by internal waves, would be incorrectly segmented. For small values of  $SSL_{min}$ , a  
361  $CPA_{int}$  value should be selected to provide enough samples (>10) for the CPA to be  
362 conducted appropriately. The deviation of  $\Delta Z$  from its mean was calculated (Fig. 6b),  
363 followed by the cumulative sum of this deviation (Fig. 6c), the most significant point of  
364 change was indicated by the largest absolute value (Fig. 6c – black line), and quantified  
365 by the range,  $CPA_{max}$ , of the cumulative sum values. A confidence interval (CI) was  
366 calculated by determining the percentage of 1000 bootstrapped samples of  $\Delta Z$  that  
367 yielded a  $CPA_{max}$  value smaller than the original  $CPA_{max}$  value. Where a significant  
368 change occurred (95% CI), indicating that a SSL changed from simply varying in depth  
369 around a static mean to exhibit migrant behaviour (increasing/decreasing depth), SSLs  
370 were separated into migrant and static components (Fig. 6).

371

372

[FIGURE 6]

373

374 The CPA was conducted iteratively, until no further statistically significant change (95%  
375 CI) within an image segment could be detected: this ensured that multiple migrations,  
376 during a diel cycle for example, would all be separated. Multiple migrations were  
377 unlikely to occur in this study since at an earlier stage we partitioned the data into  
378 separate day and night segments.

379

### 380 **2.2.5 SSL metrics**

381 For each individual SSL identified a set of SSL metrics were evaluated (Table 1). The  
382 depth, duration, MVBS, MVBS standard deviation, MVBS range and layer thickness  
383 described spatial extent and backscatter distribution. The vertical velocity and depth  
384 range were used to identify and describe migratory layers. The Background Noise Level  
385 (BNL) was used to quantify the maximum level of noise surrounding the SSL. The  
386 maximum value was taken to ensure that it would be greater than the mean MVBS value  
387 of SSLs consistent of natural variation within the data (phantom SSLs: Section 2.2.2).

388

389

[TABLE 1]

390

391 The SSL metrics were analysed (Section 3.2) to gain better insight into the nature of  
392 SSLs within the study region, enabling inferences concerning the pelagic community  
393 (spatial arrangement, distribution and heterogeneity) to be made.

394

395



### 396 **2.3 Validation framework**

397 In order to examine the efficacy of our automated SSL identification technique versus  
398 the principal present approach, adopted by most acoustic-trawl surveys when assessing  
399 fish stocks – visual scrutinisation (which may be subjective and prone to between-  
400 operator inconsistencies) - we designed a validation framework to examine potential  
401 differences between SSLs determined by the SSLEM and visually scrutinised acoustic  
402 images. If visual scrutiny gave highly variable results, this would illustrate the difficulty  
403 likely to be encountered in comparative studies and the requirement of an automated  
404 method.

405

406 The validation was conducted using a subset of the IMOS data. Images from this subset  
407 were published online at [www.soundscatteringlayers.com](http://www.soundscatteringlayers.com). Independent visual scrutiny  
408 was performed autonomously by a group of 8 students, all of whom had attended an  
409 acoustic data collection and processing summer school  
410 ([www.depts.washington.edu/fhl/](http://www.depts.washington.edu/fhl/)) and were either PhD candidates at the University of  
411 St Andrews or the Florida International University. Each student estimated 3 SSL  
412 metrics, namely the depth, MVBS and thickness for all SSLs they could identify that  
413 persisted for a time period longer than 1 hour. Each student considered 10 images  
414 selected randomly from a set of 50. The results of the visual scrutiny were compared to  
415 those from SSLEM (see Results 3.1).

416

### 417 **3 Results:**

418 Sound Scattering Layers (SSLs) that persisted for time periods longer than 1 hour  
419 ( $SSL_{\min} = 60$  minutes;  $CPA_{\text{int}} = 6$  minutes: see Sections 2.2.2 and 2.2.4 respectively for  
420 definitions) were identified and extracted from the IMOS dataset (Section 2.1) using the

421 SSL extraction method (SSLEM). SSLs were extracted at these setting in order to  
422 conduct a regional analysis of the study area, ensuring that only persistent and  
423 therefore characteristic SSLs were identified. In total, 2064 SSLs were extracted from  
424 264 images that were on average 3.4 hours in length; an example of the identification of  
425 SSLs for an IMOS image is given in Fig. 4. 108 phantom SSLs (see Section 2.2.2) that  
426 were identified in the validation procedure were removed. For each SSL, SSL metrics  
427 described in Section 2.2.5 were determined and relationships found between select SSL  
428 metrics explored (Section 3.2).

429

### 430 **3.1 SSLEM Validation**

431 SSL Metrics that were estimated by the students from SSLs visually identified were  
432 mapped back on to the original acoustic images for comparison with the SSLEM  
433 identified SSLs. Each visually identified SSL was categorised as being a valid/invalid SSL  
434 identification. This process was mirrored using the output from the SSLEM, where  
435 phantom SSLs (see Section 2.2.2) were visually identified after extraction and deemed  
436 to be invalid SSLs comprised of background noise. The students identified 211 SSLs  
437 from 80 images. For the same SSLs that were identified by more than 6 of the students  
438 from the same image, mean ranges of the estimated metrics were calculated (Table 2).

439 [TABLE 2]

440

441 Of the three SSL identification fields in Table 2 ( $< SSL_{min}$ , Noise & TVG), the Time-  
442 Varied-Gain (TVG) field contained the largest proportion of the students  
443 misclassification of SSLs. The TVG increases the amplitude, as a function of time (or  
444 depth for a fixed sound speed) of the echo return and serves to amplify both signal,  
445 when organisms are present, and the background in an empty pixel, which appears in

446 the acoustic image as depth dependent noise (Simmonds & MacLennan, 2005). This  
447 essentially limits the useful (range over which signal dominates noise) range of the  
448 instrument and causes visible, layer-like bands to form at the far extent of this range.  
449 These layers can resemble SSLs to the untrained eye, but not to the SSLEM; normally  
450 TVG is removed in pre-processing but can just as easily be removed afterwards (e.g.  
451 Watkins & Brierley, 1996).

452

453 The SSLEM identified, on average, over 3 times the number of SSLs per acoustic image  
454 than the group of students. Whereas the SSLEM output included no variance between  
455 repeat identifications and characterisations of SSLs from the same image, the overall  
456 mean standard deviation of the number of SSLs identified per image for the students  
457 was 0.54. This is in fact quite low, and demonstrates that although the students  
458 identified fewer SSLs, the group broadly did agree on the number of SSLs per image.  
459 Metric estimates by the students were, however, notably large, especially the mean  
460 MVBS range (4.8 dB re  $1\text{m}^{-1}$ ) that is equivalent to a factor 3 change in the linear domain.  
461 The SSLs identified by the SSLEM were also much more likely to be valid (94.8%), i.e.  
462 not phantom SSLs, than those identified by the students (only 75.4% valid), who  
463 misidentified SSLs a quarter of the time (Table 2). The SSLEM extracted a total of 108  
464 incorrect SSLs, all of which were considered to be phantom SSLs (Section 2.2.2).

465

466 Phantom SSLs form by the naturally occurring variation in the data. They are an artefact  
467 of the SSLEM, and hence not detected by visual scrutinisation. This variability is of no  
468 consequence within SSLs, but can cause false SSL identification outside (i.e. in empty  
469 water - Fig. 3). In order to remove phantom SSLs, the Background Noise Level (BNL)  
470 metric was used to identify SSLs that had low signal-to-noise ratios or more accurately,

471 low BNL-to-MVBS ratios. SSLs that had a MVBS value that was smaller than the BNL  
472 were identified as being phantom SSLs (black points in Fig. 7). Removing these SSLs  
473 from the results increased the validity of the automated method, for the data analysed  
474 in this study, to 100%.

475

476 [FIGURE 7]

477

478 Phantom SSLs are apparent throughout the entire water-column, except for the region  
479 between 400m and 800m. In this depth region within the study location of the South  
480 West Indian Ocean and Tasman Sea, strong and broad SSLs were persistently present  
481 (Fig. 8), meaning that SSLs dominated, excluding the possibility of phantom SSLs  
482 forming at the same depth.

483

### 484 **3.2 SSL Metrics**

485 The SSLEM output a total of 1956 valid (non-phantom) SSLs from the IMOS data. Each  
486 SSL was summarised by a set of 9 SSL metrics (Table 1).

487

488 [FIGURE 8]

489

490 Physical characteristics of SSLs provide biological/ecological insight into pelagic  
491 community dynamics. For example, the MVBS range increased towards the surface (Fig.  
492 8a) suggesting that the biological community becomes more complex/heterogeneous in  
493 the epi-pelagic region; this could be caused by an increase in species diversity or a  
494 larger range of the orientations of organisms, caused by feeding for example. The  
495 broadest SSLs lie within the central portion of the mesopelagic region of the water-

496 column (Fig. 8b). The highest MVBS values (a proxy for increased biomass/abundance)  
497 occurred within the water-column at depths of around 600m and at the surface (Fig. 8c)  
498 and also geographically towards 40 degrees south (Fig. 8d). This is consistent with the  
499 fact that the zone is the highly productive Subtropical convergence zone (see Fig. 1; as  
500 identified by Longhurst 1998) where previous work has revealed an enhanced prey-  
501 field (Boersch-Supan et al. 2012).

502

#### 503 **4 Discussion:**

504

##### 505 4.1 Sound Scattering Layer Extraction Method (SSLEM)

506

507 The Sound Scattering Layer (SSL) Extraction Method (SSLEM) was demonstrated here  
508 using data observed at a single frequency, 38 kHz. However, the method is independent  
509 of frequency and is entirely appropriate for use with other frequencies. The main  
510 differences that would occur would be subject to the characteristics of the incident  
511 frequency, for example, reduced depth range and increased resolution at higher  
512 frequencies (Simmonds & MacLennan 2005).

513

514 The efficacy of the SSLEM was examined by comparing output from that of visually  
515 scrutinised data (Table 2). The comparison demonstrated that the SSLEM method is  
516 more effective at identifying SSLs (Section 3.1). Visually scrutinised images were subject  
517 to SSL misclassification, under classification and sample variation, whereas the SSLEM  
518 output was perfectly repeatable, with zero variance between repeat extractions of SSLs  
519 and metrics.

520

521 Merged vertically static and migrant SSLs were separated by an application of a change-  
522 point analysis (Section 2.2.4). Analysis of the derived SSL metrics revealed water-  
523 column and geographical trends (Fig. 8). Summarising these data into discrete metrics  
524 offers a method of standardised analysis for assessing variability in biological  
525 communities in the water-column. Importantly, acoustic survey data can now be  
526 condensed down from millions of values to a set of community descriptors that can be  
527 easily stored, shared and analysed.

528

## 529 4.2 SSLEM Applications

530

531 Analysis of the ocean's SSLs will enable the study of the ocean's mid-trophic structure,  
532 providing a global prey field that would be invaluable to predator-prey ecologists. SSL  
533 depths could be used to gauge energy expenditure of diving mammals (Boersch-Supan  
534 et al. 2012; Walters et al. 2014) and spatial arrangements of prey-fields that could be  
535 incorporated into existing biophysical models (for example, SEPODYM: Bertignac,  
536 Lehodey, & Hampton 1998; Lehodey et al. 1998). Monitoring the structure of SSLs over  
537 long time periods could reveal climatic influences and the knock-on effects for SSL  
538 inhabitants (Lehodey et al. 2003). Spatially distinct formations of SSLs made up of  
539 diverse communities are likely to be distinguished and characterized by SSL metrics,  
540 allowing the division of regionally distinct biological communities (Longhurst 1998).  
541 Whilst the SSLEM does not resolve communities at the species level, such as the Species  
542 Identification Methods from Acoustic Multifrequency Information (SIMFAMI: Gajate et  
543 al. 2004) project, the SSLEM offers an alternate and simpler approach for fisheries and  
544 conservation management regimes to assess and monitor open ocean ecosystem health  
545 and stability (Korneliussen et al. 2008; Handegard et al. 2013).

546

547 4.3 Summary

548

549 The SSLEM presented here is directly applicable to all acoustic images, including  
550 echograms output from Acoustic Doppler Current Profilers (ADCPs), because it is  
551 independent of frequency and scale. Furthermore, it naturally lends itself to more  
552 complex multi-frequency analysis (Jarvis et al. 2010). Unlike other methods (e.g. Cade  
553 and Benoit Bird, 2014), the SSLEM was built to facilitate automated processing of data  
554 in a standardised fashion that would vary only with consideration of the resolution and  
555 scale of the study in mind. It is our hope that its introduction will enable the analysis of  
556 a wealth of data that is immediately available, offering insights into the biological  
557 structure of the world's ocean. The derived SSL metrics provide a means to summarise  
558 the extracted layers, making them readily available for a wide range of analysis.  
559 Importantly, the SSLEM offers the opportunity to study the structure of the mid-trophic  
560 communities in the ocean and will aid in improving our understanding of an ocean  
561 ecosystem.

562

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570 Florida International University and Sam Gordine, Mirko Semler, Chris Mueller & Moritz

571 Wiesel from the University of St Andrews, for taking the time to help with the validation  
572 of the method.

573

574 Data Accessibility:

575 Acoustic data are publicly available via the Integrated Marine Observing System (IMOS)  
576 - Bio-Acoustic Ships of Opportunity (BA SOOP) sub-facility found using the IMOS Ocean  
577 data portal: <https://imos.aodn.org.au/imos123/home>

578

579 References:

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<b>SSL metric</b>	<b>Definition</b>	<b>Unit</b>
<b>Depth</b>	mean depth	m
<b>Depth range</b>	Max(depth) – Min(depth)	m
<b>MVBS</b>	MVBS over entire SSL	dB re 1m <sup>-1</sup>
<b>MVBS range</b>	Max(MVBS) - Min(MVBS)	dB re 1m <sup>-1</sup>
<b>MVBS STD</b>	standard deviation of MVBS	dB re 1m <sup>-1</sup>
<b>Thickness</b>	mean SSL thickness	m
<b>Vertical velocity</b>	(change in depth)/time	ms <sup>-1</sup>
<b>Background Noise Level (BNL)</b>	Max (MVBS) of background pixels surrounding SSL	dB re 1m <sup>-1</sup>
<b>Duration</b>	length of SSL duration	(H:M:S)

768

769 Table 1 – Sound Scattering Layer (SSL) metrics: summary metrics for individual SSLs.

770 The unit dB re 1m<sup>-1</sup> represents 10 times the log base 10 value of a variable with units of771 m<sup>-1</sup>, in this case the mean volume backscattering coefficient (Simmonds & MacLennan,772 2005), that is relative to a reference level of 1m<sup>-1</sup>.

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Method	Mean depth range (m)	Mean MVBS range (dB re 1m <sup>-1</sup> )	Mean thickness range (m)	< SSL <sub>min</sub> (%)	Noise (%)	TVG (%)	Valid (%)	Invalid (%)
<b>SSLEM</b>	0	0	0	0	5.2	0	94.8	5.2
<b>VISUAL</b>	26.3	4.8	53	5.7	8.5	10.4	75.4	24.6

783

784 Table 2 – Summary of method output versus visual scrutiny for identification of Sound  
785 Scattering Layers (SSLs). Columns definitions from left to right: Method – method of SSL  
786 identification; Mean depth range, Mean MVBS range & Mean thickness range – mean  
787 ranges of values for the depth, MVBS and thickness metrics for repeat estimations of the  
788 same SSLs; < SSL<sub>min</sub> – percentage of SSLs identified smaller than the pre-set minimum  
789 value; Noise – percentage of SSLs consistent of background noise (including phantom  
790 SSLs); TVG – percentage of SSLs made up of ‘layer-like’ noise bands amplified by time-  
791 varied gain (TVG); Valid – number of correctly identified SSLs; Invalid – number of  
792 incorrectly identified SSLs. Percentages and means are to 1 d.p.

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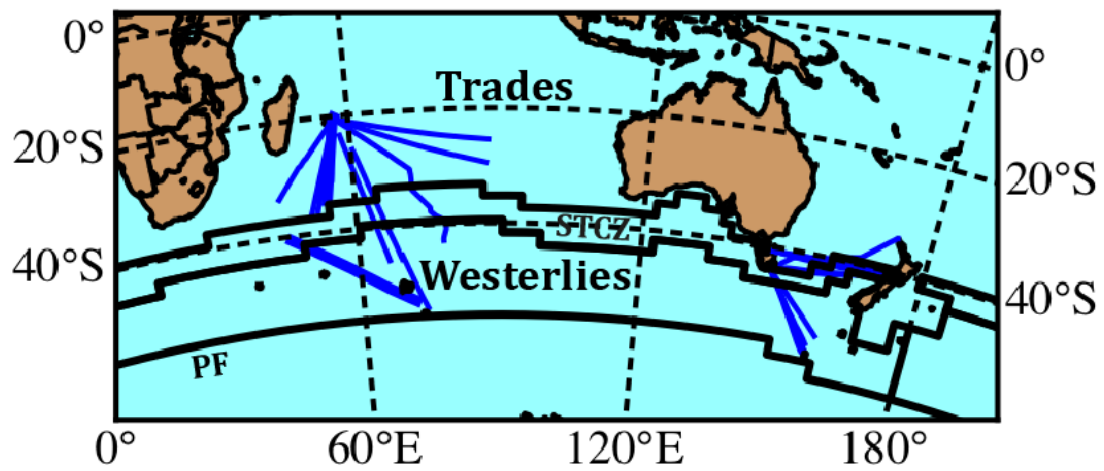
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803 Figure 1 - Map showing ship transect lines (blue) for acoustic data extracted from the

804 Integrated Marine Observing System (IMOS) data centre. Mean positions of the

805 Subtropical Convergence Zone (STCZ) and Polar Front (PF) are marked as well as two

806 Longhurst Biomes, the Trades and the Westerlies, separated by the northern boundary

807 of the STCZ.

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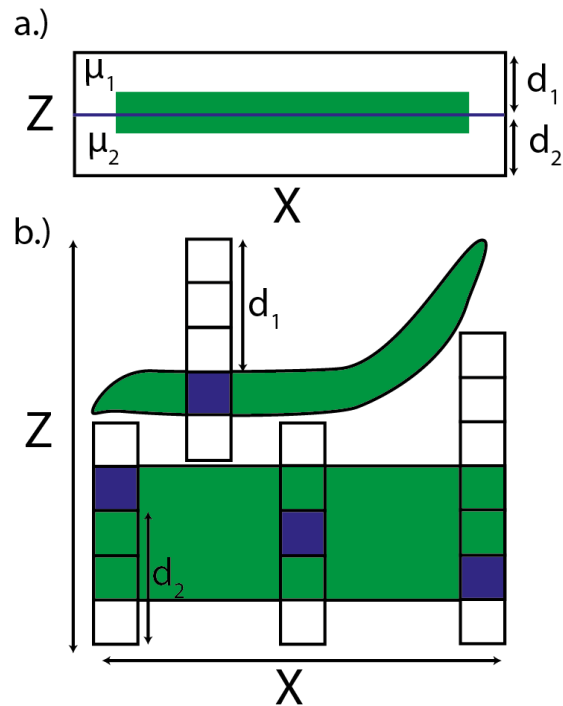
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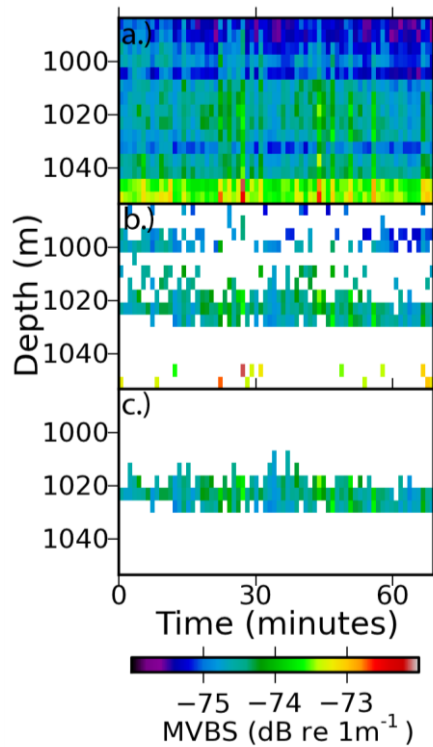
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Figure 2 - Identification of Sound Scattering Layer (SSL) pixels, where green features indicate relatively high intensity SSLs, white background indicates low intensity noise (or empty water) and blue cells represent the SSL pixel being evaluated. a.) Simple SSL analysis window: only vertically static SSLs separated by a distance larger than  $d_1$  or  $d_2$  are detected b.) Dynamic SSL analysis column: a column is moved pixel by pixel through the image, where at each step the column size ranges from the minimum (5 pixels in length; where  $d_1$  and  $d_2$  are both equal to 2 pixels plus the pixel being evaluated) up to the full vertical extent of the Z axis, by stepping through all the possible values for each of the two parameters,  $d_1$  and  $d_2$ ; in doing so, SSLs of varying separation distances and vertical behaviours are captured.



835

836 Figure 3 – Phantom Sound Scattering Layer (SSL) extracted from FV *Austral Leader II*  
 837 transect during May 2012, where  $SSL_{min}$ , the minimum horizontal resolution of SSLs,  
 838 was set to 60 minutes a.) acoustic image segment; b.) SSL pixels identified by the  
 839 method (Section 2.2.1). c.) Extracted phantom SSL: MVBS values of the SSL are similar  
 840 to that of the surrounding background.

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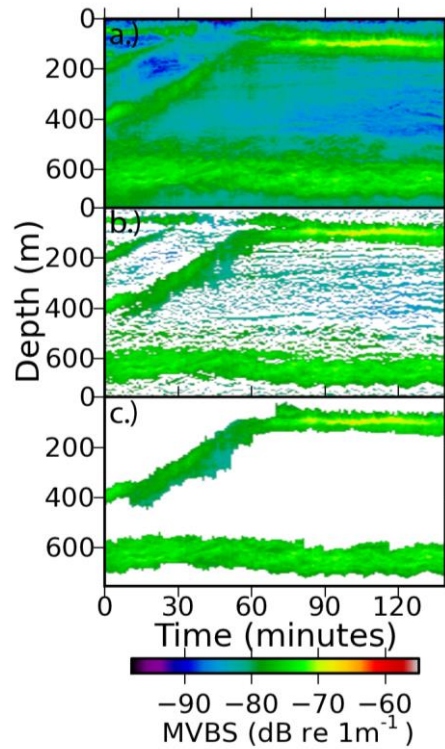
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850 Figure 4 - Processed acoustic image, from FV *Austral Leader II* transect during May  
 851 2012, where  $SSL_{min}$ , the minimum horizontal resolution of Sound Scattering Layers  
 852 (SSLs), was set to 60 minutes. a.) Original acoustic image at a resolution of 5m in depth  
 853 and 1 minute in time. b.) Masked image: only pixels that are deemed to be potential SSL  
 854 pixels are shown. c.) SSLs identified after removing features smaller than  $SSL_{min}$  and  
 855 filling SSL internal gaps (that are smaller than  $SSL_{min}$ ).

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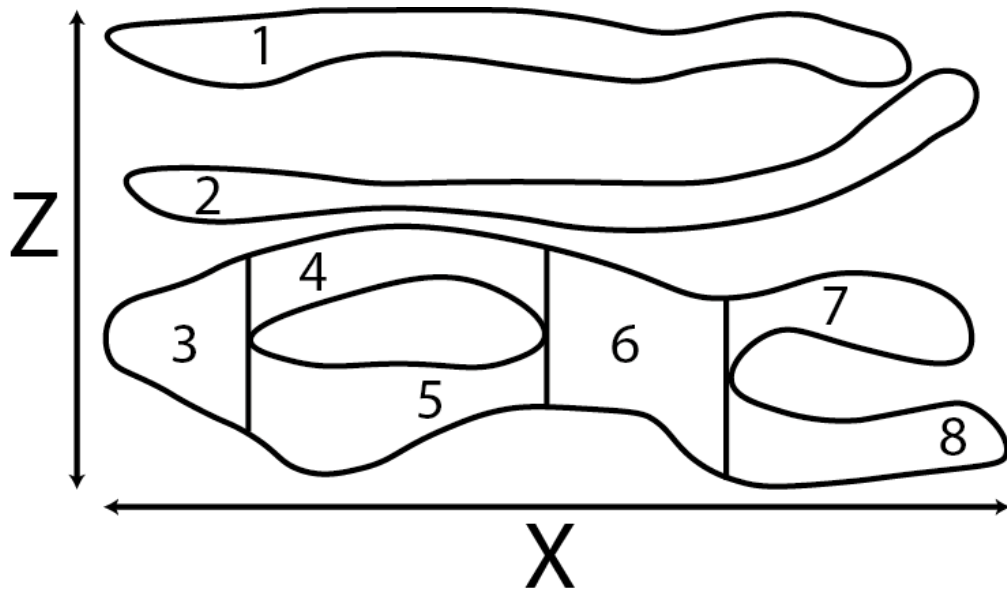
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Figure 5 - Segmentation of Sound Scattering Layers (SSLs) features into individual SSLs. Each SSL was assigned a unique index value.

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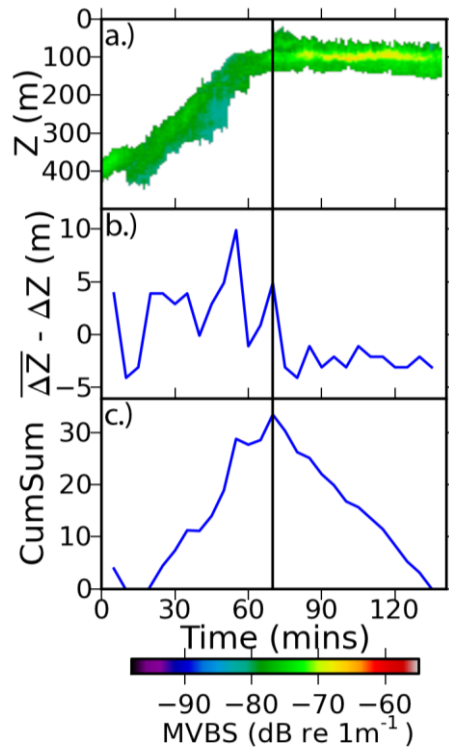
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881 Figure 6 - Change Point Analysis of mean Sound Scattering Layer (SSL) depth – SSL  
 882 taken from the example image in Fig. 4. The vertical black line at 70 minutes - the  
 883 maximum point of the cumulative sum of b.) - indicates the point of separation of a  
 884 static SSL and a migrant SSL. a.) SSL depth b.) The overall mean depth change of SSL  
 885 minus each mean depth change in the time-series (binned at 6 min intervals) plotted in  
 886 time c.) Cumulative sum of b: the maximum value indicates the most significant point of  
 887 change.

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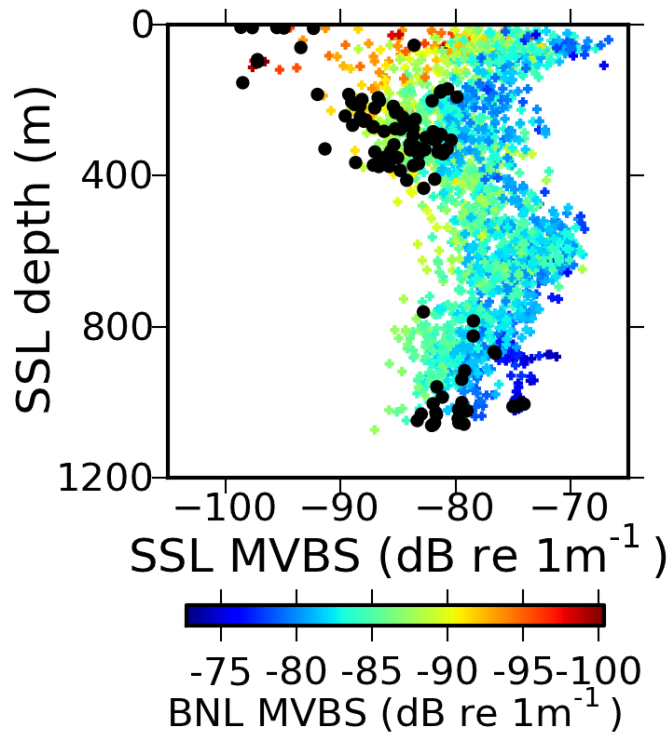
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896 Figure 7 – Background Noise Level (BNL) for Sound Scattering Layers (SSLs) extracted  
 897 from the IMOS dataset by SSL depth and MVBS. Black points represent phantom SSLs  
 898 (incorrectly assigned SSLs) identified where BNL > SSL MVBS.

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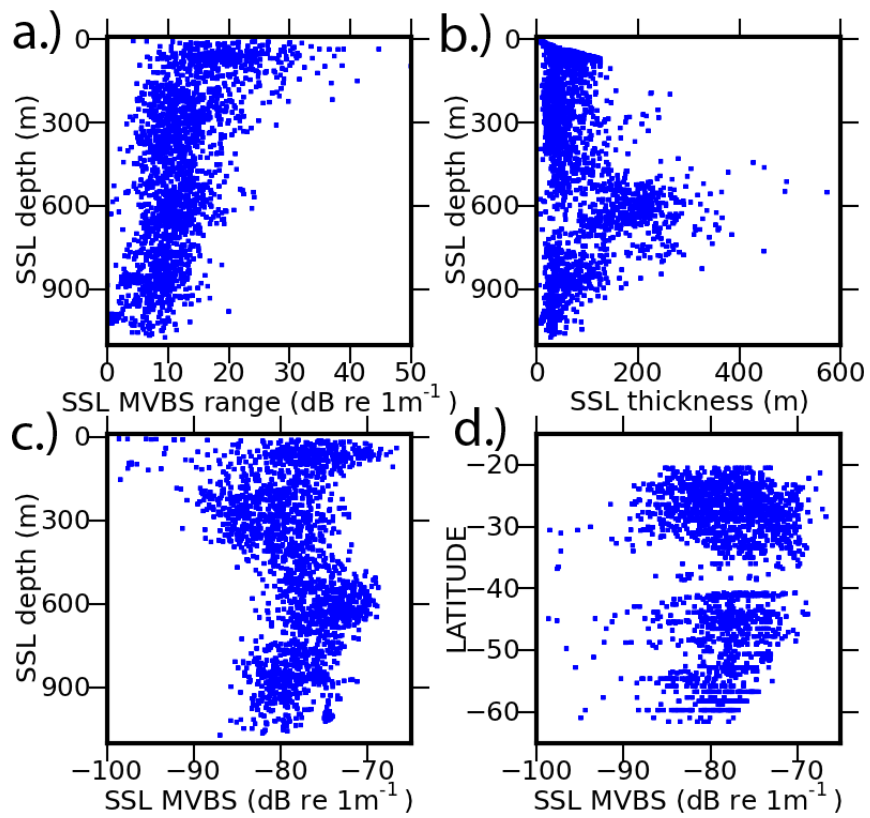
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911 Figure 8 – Sound Scattering Layer (SSL) metrics extracted from the IMOS dataset. a.)

912 Water column heterogeneity: MVBS range serves as a proxy for biological complexity;

913 b.) SSL thickness as a function of depth; c.) Depth distribution of MVBS; d.) Latitudinal

914 distribution of MVBS.

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