FINAL REPORT

A SURVEY OF CADMIUM IN BROWN CRABMEAT AND BROWN CRABMEAT PRODUCTS: FOLLOW-ON STUDY ON CADMIUM IN CRAB HEPATOPANCREAS AND OTHER EDIBLE ORGANS FS102010

08 April 2013

Centre for Environment, Fisheries & Aquaculture Sciences (Cefas)



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Submitted to:	Richard Burden, The Food Standards Agency
Date submitted:	08 April 2013
Project Manager:	Thi Bolam
Report compiled by:	Thi Bolam and Philippe Bersuder
Quality control by:	Robin Law
Approved by & date:	Steve Millward, 04 th April 2013
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Final Report



Cefas Project C5700B –A Survey of Cadmium in Brown Crabmeat and Brown Crabmeat Products: Follow-on Study on Cadmium in Hepatopancreas and Other Edible Organs.

Authors: Thi Bolam and Philippe Bersuder

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Area 3B Aviation House,

125 Kingsway, London, WC2B6NH

Contact person: Richard Burden

e-mail: Richard.Burden@foodstandards.gsi.gov.uk



Head office

Centre for Environment, Fisheries & Aquaculture Science Pakefield Road, Lowestoft, Suffolk NR33 0HT, UK Tel +44 (0) 1502 56 2244 Fax +44 (0) 1502 51 3865 www.cefas.co.uk

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Executive Summary

The European Commission has recommended that Member States should provide advice on brown crabmeat consumption to their consumers as the brown meat from crabs (brown crabmeat) has the potential to contain elevated levels of cadmium (Cd). Following a recent survey on Cd levels in brown crabmeat and its products on sale in the United Kingdom (UK) commissioned by the Food Standards Agency (FSA) in 2012, additional information with regards to Cd levels in the hepatopancreas and other edible tissues from the crab's cephalothorax was required to provide this advice.

A total of fifty six live brown crab (*Cancer pagurus*), representing four geographical locations (Fraserburgh, Aberdeen, Dorset, Newlyn) were procured by the FSA from Billingsgate. The crabs were transported to the Cefas Lowestoft laboratory where they were tagged for traceability and stored under appropriate conditions. The crabs were subsequently killed humanely and immediately dissected, all hepatopancreases being homogenised and bulked into one sample, the remaining edible tissues from the cephalothorax making up a second bulked sample.

The analysis of Cd and a suite of other trace metals were carried out on six replicates of each sample using a fully validated and accredited methodology based on the acid digestion of crab tissues using an enclosed vessel microwave. The determination of Cd was performed by Inductively Coupled Plasma-Mass Spectrometry.

For the hepatopancreas sample, the mean Cd concentration [±standard deviation] was found to be 4.0 [±0.18] mg/kg wet weight (w.w.), while the mean Cd concentration in other edible tissues from the cephalothorax was found to be 0.27 [±0.02] mg/kg w.w. The concentration data reported here will support the FSA's risk assessment and management processes regarding the safety of consuming brown crab meat products that are available for purchase in the UK.

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Glossary

Cd Cadmium

CRM Certified Reference Material

EC European Commission

FSA United Kingdom's Food Standards Agency

ICP-MS Inductively Coupled Plasma-Mass Spectrometry

LIMS Laboratory Information Management System

LSN Laboratory Sample Number

RSPCA The Royal Society for the Prevention of Cruelty to Animals

SOP Standard Operating Procedure

w.w. Wet weight

1 Introduction

In 2011, the EC released an information note to raise awareness of national authorities regarding the consumption of brown crabmeat as it has the potential to contain elevated levels of cadmium (Cd) (EC, 2011), with the anticipation that individual Member States would produce consumption advice relevant to their consumers. However, the available information on Cd concentration in brown crabmeat and its products on sale in the United Kingdom (UK) has not been sufficient to allow the Food Standards Agency (FSA) to provide this advice. Consequently, the FSA commissioned a survey to provide a data base which describes the range of Cd concentrations in brown crabmeat and food products containing brown crabmeat sold in UK retail outlets that were readily available to the UK consumer (Bolam and Bersuder, 2013). As a result of this initial survey, additional information with regards to Cd levels in various edible tissues present in the cephalothorax of freshly killed crab is required. This survey provides quality assured evidence with regards to mean levels of Cd in the hepatopancreas and other edible tissues composing the brown crabmeat. The concentration data reported here will support the FSA's risk assessment and management processes regarding the safety of consuming brown crab meat products that are available for purchase in the UK

2 Methods

The study was based on two distinct phases. The first phase involved the collection of fifty six live crabs from at least four different landing locations and their delivery to the Cefas Lowestoft laboratory, their tagging and storage. Subsequently, crabs were killed humanely, dissected for their hepatopancreases and other edible organs, bulked into two distinct samples and analysed for cadmium and other trace elements using reliable, quality assured analytical methodologies.

2.1 Sampling strategy and plan

Live brown crabs (*Cancer pagurus*) were purchased by the FSA from Billingsgate Fish Market. A total of fifty six crabs were required, with the objective of obtaining as even a distribution as possible between the geographical areas that are sold at Billingsgate and the sex of crab e.g. if crabs were able to be purchased from four geographical areas, the ideal split for the fifty six crabs would be as follows:

- Area 1: fourteen live brown crabs (seven male and seven female)
- Area 2: fourteen live brown crabs (seven male and seven female)

- Area 3: fourteen live brown crabs (seven male and seven female)
- o Area 4: fourteen live brown crabs (seven male and seven female)

Additionally, the sampling plan also stipulated the following:

- no more than fifteen crabs were to be purchased from a single geographical area;
- the minimum number of geographical areas to be covered was to be four;
- crabs were to be packaged by catch area for transport and information (i.e. location, sex, sample number) was to be clearly recorded and handed over to Cefas staff;
- all crabs were to be nicked prior to delivery at the Cefas laboratory as they were to be kept alive in tanks at the Cefas aquarium facilities before dissection.

2.2 Sample receipt and storage

2.2.1 Sample receipt

Upon sample receipt, Cefas checked for sample numbers, recorded any mortality or other observations (e.g. claw shedding). All dead crab were stored separately from live crab and, depending on the conditions of the dissected organs from the dead crabs, a decision on whether these should be bulked with the remaining samples was taken. Subsequently, crab sex and length were recorded and an identification tag was fixed to each animal (both dead and alive).

Crabs were tagged using Super Heavy Duty monofilament Double T anchor tags (SHD FD-94 anchor tags) manufactured by Floy Tag and Manufacturing inc. (Seattle, USA). The tag had two 10mm long T-bars that were 10mm apart. The external monofilament of the tag was 55mm long surrounded by coloured polyolefin tubing which contained the numbering and legend, protected by a shrink lock covering. Each tag was serially inscribed with the tag batch and series numbers, e.g.:

E04 0001 WWW.CEFAS.CO.UK PLEASE RECORD

TAG NO, DATE, LAT & LONG, WIDTH, SEX

The last 4 digits of the tag number being sequentially incremented.

Avery Dennison Mark II SHD tagging guns, with a 2.3mm external diameter needle, were used to insert the tags. The guns were modified to reduce possible damage to the crabs' internal organs when tagged. The long point of the needle was ground off and a guard attached, limiting the maximum penetration of the needle to around 1cm. The needle was inserted into the branchial

chamber through a hole made in the epimeral line at the back of the carapace of the crab and, when triggered, the head of the tag was driven inside the carapace and anchored by the terminal T-bar. The second T-bar remained outside the carapace, preventing the trailing tag section from being drawn inside the crab.

2.2.2 Live crab storage

Live crabs were transferred into three storage tanks (volume 1.35 m³) containing saline water (31 ppt salinity) refreshed at a rate of 3 L/min. The water was aerated with compressed air through a medium pore air stone, and the water temperature was maintained at 4.5°C. Crab mortality was checked daily and recorded.

2.2.3 Killing of crabs:

Cefas followed the guidelines of handling and storage of live crabs provided in the RSPCA protocol for the humane stunning/killing of Crustacea (RSPCA, 2012). Electrical stun/killing is the most humane and effective method for killing crabs, as it renders them immediately insensible with death ensuing (whilst insensible) within seconds, and this method was therefore applied in this study. For this, a CrustaStun™ single stunner device (Crustastun, Studham, Beds, UK), designed to administer a lethal shock (230 volt, 1-8 amp electrical charge) to crab was used to kill the live crab humanely. The CrustaStun™ device was operated by initially dissolving 60g of Analar grade sodium chloride (VWR, Leicestershire, UK) in purified water (reverse osmosis) to fill the base of the device. The electrode sponge located in the lid was dampened with the saline solution, and one crab at a time was placed abdomen down onto the sprung tray in the unit. The lid was closed and secured before the stun button specific to "crab" was depressed. Within half a second, the nerve functions of the crab are interrupted and the animal cannot feel pain, the animal being killed within ten seconds of application.

2.3 Sample preparation

Each dead crab was dissected following Cefas Standard Operating Procedure 2150: "Whole crab handling, storage (for pre-killing), killing and dissection" (provided as Supplementary Information separately to this report). The hepatopancreas and other edible tissues from the cephalothorax (including the reproductive organs, soft skin tissue etc.) were manually separated from the whole crab. The hepatopancreas of fifty six crabs were combined in batches of 5-6, homogenised and bulked into one single sample. The same bulking method was applied to the remaining edible tissues, producing a second bulked sample. Additionally, the sex, weight and size of the whole crab,

as well as the weight of the hepatopancreases and other edible organs of each crab, were also

recorded during the sample preparation stage.

2.4 Sample analysis

Six sub-samples of each bulked tissue (hepatopancreases and other edible organs) underwent an

acid digestion using an enclosed vessel microwave (Multiwave 3000, Anton Paar, Hertford, UK).

Typically, approximately 3 g of homogenised sample was weighed out and pre-digested overnight in

6mL of nitric acid (Aristar grade 69%, VWR, Leicestershire, UK). The digestion was performed using a

temperature-controlled microwave programme specific for the sample matrix. The digest was then

further diluted prior to analysis by inductively coupled plasma-mass spectrometry (ICP-MS) using an

Agilent 7500ce (Agilent Technologies, Waldbronn, Germany). Quantification of Cd was performed

by external calibration, using eight levels (0, 0.5, 1, 5, 10, 20, 100 and 500µg/L) of working standard

solutions, which were prepared from a customised mixed metal standard solution of 100mg/L (SPEX

Certiprep Ltd, Middlesex, UK).

A reagent blank and a certified reference material (CRM TORT-2-Lobster hepatopancreas; National

Research Council Canada, Halifax, Nova Scotia, Canada) were included with the batch of 12 samples.

Concentration data derived from the analysis of the CRM were then added to existing quality control

Shewhart charts (using North West Analytical Quality Analyst™, Northwest Analytical Inc., USA) for

the assessment of the day-to-day method performance and validity of Cd concentration data from

the batch analysis of real samples. The validity of results was established using the warning and

control limits of the Shewhart chart, which are defined as 2σ and 3σ , respectively.

3 Results

3.1 Sampling survey

A retailer from Billingsgate was commissioned by the FSA to source the live crabs for this study. A

total of fifty six live crabs, all of the Cancer pagurus species, were delivered at the Cefas laboratory in

Lowestoft. Six crabs were found to be dead, probably as a result of longer distance transport as all

six originated from Scotland. The crabs were from four different landing locations:

Fraserburgh: 2x12 samples;

Aberdeen: 10 samples;

Dorset: 10 samples;

Newlyn: 12 samples.

All crabs, whether alive or dead, were tagged with a unique number to cross-reference their landing location (See Figure 1). Details of the crab landing information and condition upon receipt are tabulated in Appendix 1.



Figure 1: Example of crab tagging

All six dead crabs were put aside for dissection on the same day. The remaining crabs were put into three tanks filled with aerated seawater (Figure 2), with tank 3 containing only one crab from Fraserburgh (tag number EO04 2630) as it was not nicked. The crabs were not fed whilst they were waiting to be killed.



Figure 2: Tank facilities used for storing live crab

Regular monitoring was carried out to check for any mortality: during the whole study, only one crab died at the end of day 1 (tag number EO04 2636), it was removed, frozen and dissected the following day.

3.3 Crab humane killing method

The RSPCA believes that electrical stun/killing is the most humane and effective method for killing crabs, lobsters, crayfish and langoustines. The electrical stun/killing method renders them immediately insensible with death ensuing (whilst insensible) within seconds. Figure 3 shows the CrustaStun™ single stunner device, in use with a live crab.



Figure 3: The Crustastun™ single stunner device

3.4 Sample preparation

The crabs were dissected as per SOP 2015 to remove the hepatopancreas and other edible organs in batches of five or six samples (Figures 4 and 5). The tissues were bulked and homogenised respectively for hepatopancreases and other edible organs. As the dissection was carried out over 2 days, the bulked and homogenised samples were stored frozen at the end of each day and were combined on the last day of dissection into a final container (one for hepatopancreas, one for the other edible organs) and homogenised once more.

All fifty six crabs were dissected and their hepatopancreas and other edible organs (including the tissues from the dead crabs) were bulked and homogenised respectively.

The information on sex, weight (hepatopancreas, other edible organs and total weight of whole crab) and size of each individual crab is summarised in Appendix 2.

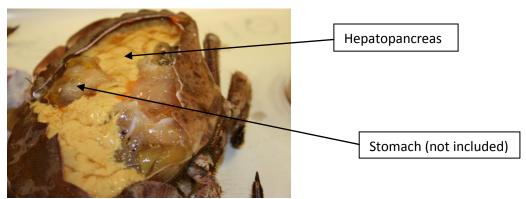


Figure 4: Hepatopancreas (and stomach)

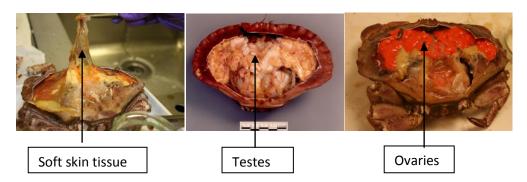
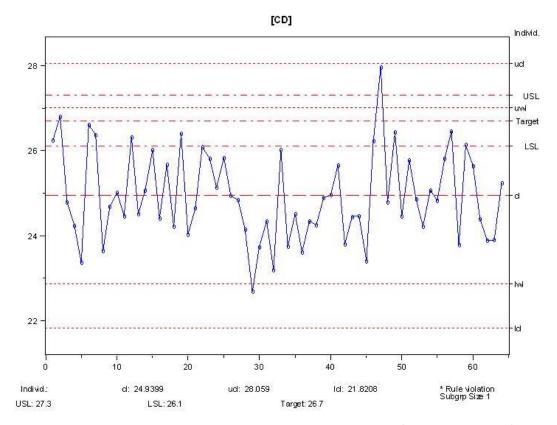


Figure 5: Other edible organs: including the soft skin tissue (left) and reproductive organs: testes in male crab (middle) or ovaries in female crab (right)

3.5 Internal quality assurance and control

The mean recovery of Cd from the extraction and analysis of certified reference material (CRM Tort-2, certified Cd value of 26.7mg/kg) was 93.4% (n=64; RSD of 4.2%). The inter-batch variations of Cd concentrations from the CRM are presented in Figure 6.

The method limit of detection and limit of quantification for Cd are 8 $\mu g/kg$ w.w. and 27 $\mu g/kg$ w.w. respectively.



3.6 Cadmium levels hepatopancreas and other edible organs tissues

Six sub-samples from each bulked tissue were analysed for Cd and other trace elements (arsenic, chromium, copper, iron, lead, manganese, mercury, nickel, selenium and zinc). The individual results and summary statistics are tabulated in Table 1.

Table 1: Cadmium concentrations in bulked crab hepatopancreas

Laboratory	Sample Description		Trace and heavy metal concentration (mg/kg w.w.)											
Sample Number	Sample Description	Cr	Ni	Cu	Zn	As	Cd	Pb	Se	Mn	Fe	Hg		
2013/00475/01	Heppatopancreas 1	0.10	0.46	46	17	15	4.2	0.02	2.2	4.6	47	0.10		
2013/00475/02	Heppatopancreas 2	0.10	0.46	46	19	15	4.1	0.02	2.2	5.0	51	0.10		
2013/00475/03	Heppatopancreas 3	0.11	0.46	45	18	15	4.1	0.02	2.1	4.8	48	0.11		
2013/00475/04	Heppatopancreas 4	0.09	0.41	41	17	13	3.7	0.02	1.9	4.5	45	0.09		
2013/00475/05	Heppatopancreas 5	0.10	0.43	42	19	14	3.9	0.02	2.0	5.0	51	0.10		
2013/00475/06	Heppatopancreas 6	0.12	0.45	44	18	14	4.0	0.02	2.1	4.8	49	0.10		
	Mean	0.10	0.44	44	18	14	4.0	0.02	2.1	4.8	48.4	0.10		
	SD	0.01	0.02	2.16	0.73	0.76	0.18	0.002	0.11	0.22	2.30	0.00		
	%RSD	7.8	4.9	4.9	4.1	5.3	4.5	6.7	5.4	4.5	4.8	4.4		

Table 2: Cadmium concentrations in bulked edible organs from the cephalothorax, not including the hepatopancreas

Laboratory	Samula Description			Trac	e and he	avy met	al concer	ntration (m	g/kg w.v	v.)		
Sample Number	Sample Description	Cr	Ni	Cu	Zn	As	Cd	Pb	Se	Mn	Fe	Hg
2013/00476/01	Other Edible Organs 1	0.03	0.05	16	30	12	0.28	0.02	1.4	1.0	8.2	0.03
2013/00476/02	Other Edible Organs 2	0.03	0.06	17	31	13	0.29	0.02	1.5	1.1	8.8	0.03
2013/00476/03	Other Edible Organs 3	0.02	0.05	16	30	12	0.28	0.02	1.4	1.0	8.4	0.02
2013/00476/04	Other Edible Organs 4	0.10	0.05	16	29	12	0.27	0.02	1.4	1.0	8.5	0.02
2013/00476/05	Other Edible Organs 5	0.02	0.05	16	29	12	0.27	0.02	1.4	1.0	8.1	0.02
2013/00476/06	Other Edible Organs 6	0.03	0.05	14	27	11	0.24	0.02	1.2	1.0	7.6	0.02
	Mean	0.03*	0.05	15.7	29.4	11.9	0.27	0.02	1.37	1.02	8.27	0.03
	SD	0.005*	0.002	0.90	1.35	0.77	0.02	0.001	0.07	0.04	0.43	0.004
	%RSD	16*	4.1	5.7	4.6	6.5	6.3	4.5	5.4	3.8	5.1	16.9

Key =*not including outlier of 2013/00476/04 due to possible Cr contamination

4 Conclusions

A total of fifty six crabs were successfully dissected for the extraction of the hepatopancreas and all other edible organs from the cephalothorax. The respective tissue groups from the fifty six crabs were bulked and homogenised. Six sub-samples from each bulked tissue group were analysed for cadmium and a suite of other trace elements.

Levels of Cd were found to be significantly higher in the crab hepatopancreas than in other edible organs, with a Cd mean concentrations of 4.0mg/kg w.w. and 0.27mg/kg w.w., respectively.

5 Acknowledgments

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6 References

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- EC (2011). Information note: Consumption of brown crabmeat. Accessed March 2012 on http://ec.europa.eu/food/food/chemicalsafety/contaminants/information_note_cons_brown_crab_en.pdf.
- 3. RSPCA (2012). Humane electrical stun/killing of Crustacea, March 2012. Accessed 15/06/2012 on

http://www.rspca.org.uk/ImageLocator/LocateAsset?asset=document&assetId=123271630198 8&mode=prd

7 Appendices

Appendix 1 Crab details as receipted at the Cefas Lowestoft laboratory

Appendix 2 Crab individual measurements

Appendix 1. Crab details as receipted at the Cefas Lowestoft laboratory

Crab number	Tag Number	Location	Date in Tank	·		Tank Number	Sex	Length (mm)
				dissection				()
1	EO04 2583	Fraserburgh	27/02/2013	28/02/2013	2 legs missing	2	F	167
2	EO04 2584	Fraserburgh	n/a	27/02/2013	recently deceased	n/a	М	162
3	EO04 2586	Fraserburgh	n/a	27/02/2013	recently deceased	n/a	М	156
4	EO04 2588	Fraserburgh	n/a	27/02/2013	recently deceased	n/a	F	170
5	EO04 2589	Fraserburgh	27/02/2013	28/02/2013	reactive	2	F	177
6	EO04 2590	Fraserburgh	27/02/2013	28/02/2013	reactive	2	F	175
7	EO04 2591	Fraserburgh	27/02/2013	27/02/2013	reactive	2	М	159
8	EO04 2592	Fraserburgh	27/02/2013	28/02/2013	reactive	2	М	167
9	EO04 2593	Fraserburgh	n/a	27/02/2013	recently deceased, 4 legs missing	n/a	М	150
10	EO04 2594	Fraserburgh	27/02/2013	28/02/2013	reactive, 1 leg missing	2	F	168
11	EO04 2595	Fraserburgh	27/02/2013	28/02/2013	reactive, 1 leg missing	2	М	185
12	EO04 2596	Fraserburgh	27/02/2013	28/02/2013	reactive	2	F	204
13	EO04 2597	Dorset	27/02/2013	28/02/2013	reactive	2	F	147
14	EO04 2598	Dorset	27/02/2013	28/02/2013	reactive	2	М	140
15	EO04 2599	Dorset	27/02/2013	28/02/2013	reactive	2	F	153
16	EO04 2600	Dorset	27/02/2013	28/02/2013	reactive	2	М	145
17	EO04 2601	Dorset	27/02/2013	28/02/2013	reactive	2	М	156
18	EO04 2602	Dorset	27/02/2013	27/02/2013	reactive	2	М	156
19	EO04 2603	Dorset	27/02/2013	28/02/2013	reactive	2	F	157
20	EO04 2604	Dorset	27/02/2013	28/02/2013	reactive	2	F	147
21	EO04 2605	Dorset	27/02/2013	27/02/2013	reactive	2	М	148
22	EO04 2606	Dorset	27/02/2013	28/02/2013	reactive	2	F	144
23	EO04 2607	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	2	F	179
24	EO04 2608	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	F	168
25	EO04 2609	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	166
26	EO04 2610	Cornwall Newlyn	27/02/2013	28/02/2013	reactive, 2 legs missing	3	F	153
27	EO04 2611	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	F	164
28	EO04 2612	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	161
29	EO04 2613	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	F	152
30	EO04 2614	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	168
31	EO04 2615	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	F	150
32	EO04 2616	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	161
33	EO04 2617	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	168
34	EO04 2618	Cornwall Newlyn	27/02/2013	28/02/2013	reactive	3	М	159
35	EO04 2619	Aberdeen	n/a	27/02/2013	recently deceased	n/a	М	170

Crab number	Tag Number	Location	Date in Tank	Date of removal for dissection	Observation on receipt	Tank Number	Sex	Length (mm)
36	EO04 2620	Aberdeen	27/02/2013	27/02/2013	reactive	3	М	161
37	EO04 2621	Aberdeen	27/02/2013	28/02/2013	reactive	3	F	168
38	EO04 2622	Aberdeen	27/02/2013	28/02/2013	reactive, black hole= disease ?	3	М	144
39	EO04 2623	Aberdeen	27/02/2013	28/02/2013	reactive	3	М	148
40	EO04 2624	Aberdeen	27/02/2013	28/02/2013	reactive, 1 leg missing	3	F	170
41	EO04 2625	Aberdeen	27/02/2013	28/02/2013	reactive	3	F	177
42	EO04 2626	Aberdeen	27/02/2013	28/02/2013	reactive	3	М	170
43	EO04 2627	Aberdeen	27/02/2013	28/02/2013	reactive, 2 legs missing	3	М	169
44	EO04 2628	Aberdeen	27/02/2013	28/02/2013	reactive	3	М	174
45	EO04 2629	Fraserburgh	27/02/2013	28/02/2013	reactive	3	М	159
46	EO04 2630	Fraserburgh	27/02/2013	28/02/2013	reactive, not nicked	7	F	153
47	EO04 2631	Fraserburgh	27/02/2013	28/02/2013	reactive, 1 leg missing	3	F	154
48	EO04 2632	Fraserburgh	n/a	27/02/2013	recently deceased, 3 legs missing	n/a	М	150
49	EO04 2633	Fraserburgh	27/02/2013	28/02/2013	reactive	3	М	137
50	EO04 2634	Fraserburgh	27/02/2013	28/02/2013	reactive	3	М	151
51	EO04 2635	Fraserburgh	27/02/2013	28/02/2013	reactive, 1 leg missing	2	F	154
52	EO04 2636	Fraserburgh	27/02/2013	27/02/2013	reactive but dead at 16h30, frozen	2	F	144
53	EO04 2637	Fraserburgh	27/02/2013	28/02/2013	reactive	3	М	154
54	EO04 2638	Fraserburgh	27/02/2013	28/02/2013	reactive	3	М	157
55	EO04 2639	Fraserburgh	27/02/2013	28/02/2013	reactive	2	F	138
56	EO04 2640	Fraserburgh	27/02/2013	28/02/2013	reactive	2	F	153

Appendix 2. Crab individual measurements

Crab number	Tag Number	Location	Sex	Length (mm)	Whole Weight (g)	Hepatopancreas Weight (g)	Other Edible Organs Weight (g)	% brown meat
1	EO04 2583	Fraserburgh	F	167	661.5	70.7	29.7	15.2
2	EO04 2584	Fraserburgh	М	162	589.1	50.8	17.6	11.6
3	EO04 2586	Fraserburgh	М	156	723	40.1	24.1	8.9
4	EO04 2588	Fraserburgh	F	170	619.8	63.8	25.3	14.4
5	EO04 2589	Fraserburgh	F	177	753	81.5	35	15.5
6	EO04 2590	Fraserburgh	F	175	806.5	99.4	43.5	17.7
7	EO04 2591	Fraserburgh	М	159	725.9	47.4	24.9	10.0
8	EO04 2592	Fraserburgh	М	167	778.4	33.1	31.9	8.4
9	EO04 2593	Fraserburgh	М	150	617.7	48.5	18.6	10.9
10	EO04 2594	Fraserburgh	F	168	644.5	76.3	38.1	17.8
11	EO04 2595	Fraserburgh	М	185	1280.9	97.2	29	9.9
12	EO04 2596	Fraserburgh	F	204	1225	124.2	60.1	15.0
13	EO04 2597	Dorset	F	147	493.9	55.8	27.4	16.8
14	EO04 2598	Dorset	М	140	470	46.1	20.4	14.1
15	EO04 2599	Dorset	F	153	549.2	65.1	32.3	17.7
16	EO04 2600	Dorset	М	145	483.4	59.7	25.3	17.6
17	EO04 2601	Dorset	М	156	614.8	71.3	30.3	16.5
18	EO04 2602	Dorset	М	156	684.6	78.7	35.2	16.6
19	EO04 2603	Dorset	F	157	603.7	70.2	42.1	18.6
20	EO04 2604	Dorset	F	147	515.5	93.4	54.1	28.6
21	EO04 2605	Dorset	М	148	500.8	48.7	22	14.1
22	EO04 2606	Dorset	F	144	465.9	49.5	33.3	17.8
23	EO04 2607	Cornwall Newlyn	F	179	921.1	72.2	160.7	25.3
24	EO04 2608	Cornwall Newlyn	F	168	755.8	68.3	58.8	16.8
25	EO04 2609	Cornwall Newlyn	М	166	858.9	47	45.5	10.8
26	EO04 2610	Cornwall Newlyn	F	153	498.8	58.6	29.3	17.6
27	EO04 2611	Cornwall Newlyn	F	164	675.6	75.7	46	18.0
28	EO04 2612	Cornwall Newlyn	М	161	761.2	91.3	35.9	16.7
29	EO04 2613	Cornwall Newlyn	F	152	506.3	66.5	29	18.9
30	EO04 2614	Cornwall Newlyn	М	168	861.5	59.5	54.7	13.3
31	EO04 2615	Cornwall Newlyn	F	150	575.2	45	74.5	20.8
32	EO04 2616	Cornwall Newlyn	М	161	730	87.2	38.4	17.2
33	EO04 2617	Cornwall Newlyn	М	168	833.8	76.9	42.3	14.3
34	EO04 2618	Cornwall Newlyn	М	159	668.1	84.5	31.6	17.4
35	EO04 2619	Aberdeen	М	170	711.2	37.8	25	8.8
36	EO04 2620	Aberdeen	М	161	725.9	54	17.3	9.8

Crab number	Tag Number	Location	Sex	Length (mm)	Whole Weight (g)	Hepatopancreas Weight (g)	Other Edible Organs Weight (g)	Brown Meat (%)
37	EO04 2621	Aberdeen	F	168	708.4	56.7	96.4	21.6
38	EO04 2622	Aberdeen	М	144	492.3	49.7	16.8	13.5
39	EO04 2623	Aberdeen	М	148	487.1	30.2	17	9.7
40	EO04 2624	Aberdeen	F	170	616.4	58.1	31	14.5
41	EO04 2625	Aberdeen	F	177	776.4	68.5	38.4	13.8
42	EO04 2626	Aberdeen	М	170	819	60.6	35.9	11.8
43	EO04 2627	Aberdeen	М	169	792.9	33.2	30.5	8.0
44	EO04 2628	Aberdeen	М	174	925.4	67.3	32.4	10.8
45	EO04 2629	Fraserburgh	М	159	730	62.4	26.7	12.2
46	EO04 2630	Fraserburgh	F	153	554.1	57.5	34.5	16.6
47	EO04 2631	Fraserburgh	F	154	498.2	54.7	35.6	18.1
48	EO04 2632	Fraserburgh	М	150	498.6	43	16	11.8
49	EO04 2633	Fraserburgh	М	137	500.5	39.2	17.6	11.3
50	EO04 2634	Fraserburgh	М	151	556.5	35.6	17.8	9.6
51	EO04 2635	Fraserburgh	F	154	549.4	61.7	28.9	16.5
52	EO04 2636	Fraserburgh	F	144	450.2	49.7	24.2	16.4
53	EO04 2637	Fraserburgh	М	154	695.8	66.4	37	14.9
54	EO04 2638	Fraserburgh	М	157	668.2	45.6	24.1	10.4
55	EO04 2639	Fraserburgh	F	138	412.9	55.5	22.3	18.8
56	EO04 2640	Fraserburgh	F	153	572.8	69.4	25.8	16.6



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Head office Centre for Environment, Fisheries & Aquaculture Science Pakefield Road, Lowestoft, Suffolk NR33 0HT UK

Tel +44 (0) 1502 56 2244 Fax +44 (0) 1502 51 3865 Web www.cefas.co.uk

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