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Effects of different cooking and processing methods on the selenium contents of fish species

*Einfluss unterschiedlicher Koch- und Verarbeitungsmethoden
zum Selengehalt in Fischarten*

Nuray Erkan

Summary

The study was carried out to determine the current contents of selenium in different cooked and processed fish products. Mean selenium contents in raw species ranged from 0.361 to 0.563 µg/g. The average selenium contents of cooked samples varied from 0.479 to 0.880 µg/g in grilled fish, from 0.223 to 0.750 µg/g in fried fish and from 0.361 to 0.664 µg/g in steamed fish. Selenium contents from 0.360 to 0.936 µg/g were found in smoked fish and from 0.455 to 1.786 µg/g in salted fish. Measured values are sufficient to meet the daily selenium intake and is far below the upper tolerable level of selenium.

Keywords: Fish, selenium, daily intake, cooking methods, processing methods

Zusammenfassung

Die Studie wurde durchgeführt, um die aktuelle Menge von Selen in verschiedenen gekochten und verarbeiteten Fischprodukten zu bestimmen. Durchschnittliche Selengehalte in den rohen Fischarten wurden von 0.361 bis 0.563 µg/g gefunden. Die mittleren Selengehalte von gekochten Proben variierten von 0.479 bis 0.880 µg/g in gegrilltem Fisch, von 0.223 bis 0.750 µg/g in gebratenem Fisch und von 0.361 bis 0.664 µg/g in gedämpftem Fisch. Die Menge des Selens wurde von 0.360 bis 0.936 µg/g für geräucherten Fisch und von 0.455 bis 1.786 µg/g für gesalzenen Fisch gemessen. Die Messwerte sind ausreichend, um die notwendige tägliche Selenaufnahme zu erfüllen und liegen weit unter dem oberen verträglichen Maß von Selen.

Schlüsselwörter: Fisch, Selen, tägliche Aufnahme, Koch- und Verarbeitungsmethoden

Introduction

Seafood is high nutritional quality food item with rich polyunsaturated fatty acids in n-3 forms, essential amino acids, trace elements and vitamins, and low cholesterol and saturated fatty acids content (Erkan et al., 2010). Its consumption has been widely encouraged to prevent coronary heart disease, hypertension and cancer (Erkan and Çağiltay, 2011). Selenium (Se) is an essential trace element, and foodstuffs such as cereals, meat and fish are the principal source of Se for the general population. The best-known biochemical role of Se is its function as part of the enzyme glutathione peroxidase which protects vital components of cells against oxidative damage. It is also involved in thyroid metabolism, immune system and male fertility (Sirichakwal et al., 2005). Trace concentrations are required for normal growth and development, moderate concentrations can be stored and homeostatic functions maintained and elevated concentrations can result in toxic effects (Hamilton, 2004). Fishing plays an important role in Turkey's economy. Fish consumption per capita of Black Sea, Marmara Sea, Aegean Sea and Mediterranean Sea coastal region were significantly higher than those reported in inland region (TUIK, 2013). Anchovy, sardine, Atlantic mackerel, horse mackerel, small and medium bluefish, red mullet, hake, small and medium Atlantic Bonito, trout and eel are marine resources of major economic and gastronomic importance in Turkey and European countries. Anchovy, sardine, Atlantic mackerel, horse mackerel, red mullet, hake, small and medium Atlantic Bonito and trout is mostly consumed grilled and fried, whereas small-medium bluefish is generally boiled before consumption. Smoked salmon, smoked eel, smoked trout, marinated anchovies, salted anchovies and salted sardines are often consumed in processed sea products. Cooking and processing presents consumer different tastes. Changes of nutrient components in foods occurred due to the cooking and processing must be known since they are important for human health (Ersoy et al., 2006). Recently very valuable data have been published concerning the iodine content assessment of fresh, processed and different cooked fishes (Erkan, 2011). The aim of the study is twofold: first, to determine Se levels in muscle meat of cooked and processed fish of the main species consumed in Turkey, and to compare the results with international levels; second, to estimate the dietary intake of Se derived from this fish product consumption.

Materials and Methods

Samples

Six commercial important fish species were chosen for the study. The fish were purchased from the Istanbul local fish market and included anchovy (*Engraulis encrasicolus*), horse mackerel (*Trachurus trachurus*), small and medium bluefish (*Pomatomus saltatrix*), Atlantic Bonito (*Sarda sarda*), striped red mullet (*Mullus surmuletus*), and whiting (*Merlangius merlangus*). The fishes were prepared and cooked by frying, grilling and steaming according to common household methods. Hot-smoked fish was prepared from small and medium Atlantic Bonito (*Sarda sarda*), Atlantic mackerel (*Scombrus scombrus*), salmon (*Salmo salar*), rainbow trout (*Oncorhynchus mykiss*), and eel (*Anguilla anguilla*). Salted fish (lakerda) was prepared from small and medium Atlantic Bonito (*Sarda sarda*),

salted fish was prepared from sardine (*Sardine pilchardus*). Dried fish was prepared from horse mackerel (*Trachurus trachurus*). Marinated fish was prepared from anchovy (*Engraulis encrasicolus*). Canned fish was prepared from tuna (*Thunnus thynnus*). Fish were obtained from a fish market in Istanbul, Turkey. The fresh fish samples were packed in polystyrene boxes with crushed ice, and then transferred to the laboratory.

Sample preparation

The fish were beheaded, gutted (manually), and then washed with tap water several times to remove adhering blood and slime. The edible muscle was cut into pieces.

Cooking

Regarding frying, fish was previously coated with wheat flour. For frying, samples were immersed in vegetable oil at 180 °C for 10 min. After each treatment the oil was discarded. The grilling process was carried out with an electrically operated grill (1400 Watt, Tefal, France) at 180 °C for 30–40 min. Automatic cooker (2000 Watt, Tefal, France) was used for the steaming process (10–12 min for 250 g fish).

Preparation of processed fish

Smoked fish: The samples were immersed in brine at a ratio of 1:1 (w/w) for 24 h at 2 °C. The brine contained 12 % NaCl. After brining, the samples were removed from the brine, and then briefly dipped in chilled tap water for 1 h. Smoke was produced from oak sawdust. The processing time in the kiln was divided into two stages: (1) a preliminary drying and cooking period (45 min) at 75 °C; (2) a smoking and partial cooking period at 80 °C (60 min). After cooling at room temperature (60 min), the smoked fish were packaged by vacuum and stored at 2 °C.

Salted fish (Lakerda): The Atlantic Bonito were dry salted for 7 days, at 4 °C (fish weight: salt weight (1:3)), and then brined in a 15 % salt solution for 10 days, at 4 °C (fish: brine ratio=1:1, w/w). Salted fish were packaged into glass jars of 500 mL capacity with 200 mL 10 % salt solution.

Salted fish: The sardine were dry salted (fish weight: salt weight (1:3)) for 30 days at 4 °C.

Marinated fish: Anchovy were filleted and washed. Fillets were dipped in 3 % acetic acid/10 % salt marinating solution at a ratio of 1:1.5 (w/w) for 24 h at 4 °C. After marination the fish were drained at ambient temperature (18 ± 1 °C) for 15 min, placed into glass jars of 400 mL capacity with 125 mL sun flower oil.

Dried fish: The horse mackerel were dry salted for 30 days, at 4 °C (fish weight: salt weight (1:3)). Salted samples were dried in a temperature-controlled cabin (12 ± 2 °C) for 20 days.

Canned fish: Tuna were filleted, washed and cooked by steam cooking. They were packaged in water and sunflower oil. After cooking and processing process, the bones and skin were removed. All fish in each lot were homogenized using a laboratory blender and analysed to determine selenium contents. All assays were conducted on duplicate samples of the homogenates.

Analytical procedure of selenium analysis

Determination of total Se was carried out with a Thermo electron X7 inductively coupled plasma mass spectrometry (ICP-MS), (Model X series, Winsford-Cheshire UK). For

each sample, between 0.3 and 0.5 g of tissue (wet weight) was weighed and placed in a Teflon digestion vessel with 7 mL of concentrated (65 %) nitric acid (HNO_3) and 1 mL 30 % hydrogen peroxide (H_2O_2). The sample in the vessel containing concentrated nitric acid was then subjected to a microwave program as follows: Step 1: 25–200 °C for 10 min at 1000 W; Step 2: 200 °C for 10 min at 1000 W. Digests were finally made up with deionized water to 25 mL in acid washed standard flasks (EPA, 1994). ICP-MS operating conditions: Nebulizer gas flow 0.91 L/min, Radio frequency (RF) 1200 W, Lens voltage 1.6 V, Cool Gas 13.0 L/min, Auxiliary Gas 0.70 L/min. Instrument detection limits on the ICP-MS were 2000 ppt for selenium. Three sub-samples of each seafood material were digested using the above method. All measurements were run in triplicate for the sample and standard solutions. Selenium concentrations were calculated in µg/g wet weight. Accuracy of analytical method was monitored by analysing certified reference materials (Mussel tissue (Catalog No: ERMI-CE278, LGC Promochem Middlesex-UK)). Data quality was checked by analysing standard reference materials. Certificated value and observed value were $1.84 \pm 0.10 \mu\text{g/g}$ and $1.82 \pm 0.13 \mu\text{g/g}$, respectively.

Statistical analysis

The descriptive statistics (mean, standard deviation, range) were conducted using the Microsoft Office Excel 2010 software (Seattle, USA).

Results and discussion

Mean selenium levels in raw fish were 0.359 µg/g for anchovy (0.357–0.361 µg/g), 0.488 µg/g for small Atlantic Bonito (0.415–0.622 µg/g), 0.482 µg/g for medium Atlantic Bonito (0.389–0.575 µg/g), 0.545 for small bluefish, 0.428 µg/g for medium bluefish, 0.250 µg/g for eel, 0.893 µg/g for sardine, 0.600 µg/g for horse mackerel (0.563–0.637), 0.258 µg/g for rainbow trout, 0.465 µg/g for red mullet, 0.893 µg/g for sardine, 0.271 for salmon, 0.893 µg/g for sardine, 0.464 µg/g for whiting, 0.847 for tuna, 0.893 µg/g for sardine (Tab. 1). This result is similar to the Se content of Atlantic Bonito (0.021–0.593 µg/g) and of horse mackerel (0.107–0.662 µg/g) described by Özden (2010). Similar result was found by Özden et al. (2011), who studied the effect of season on selenium content. They found that the Se content in red mullet was 0.066–0.618 µg/g, while it was 0.133–0.848 µg/g in whiting. Kadrabova et al. (1997) reported Se levels of sea fish and river fish con-

TABLE 1: The selenium content of cooked and processed fishes.

Raw and cooked fishes	Selenium (µg/g)	The daily Se intake (µg) in cooked fish (about 200 g)	Raw and processed fishes	Selenium (µg/g)	The daily Se intake (µg) in processed fish (about 200 g)
Raw (Anchovy)	$0.361 \pm 0.014\text{a}$	72.2	Raw (Small Atlantic Bonito)	$0.622 \pm 0.021\text{a}$	124.4
Fried	$0.223 \pm 0.014\text{b}$	44.6	Smoked small Atlantic Bonito	$0.936 \pm 0.005\text{b}$	187.2
Grilled	$0.479 \pm 0.005\text{c}$	95.8	Raw medium Atlantic Bonito	$0.575 \pm 0.011\text{a}$	115
Steamed	$0.361 \pm 0.080\text{a}$	72.2	Smoked (Medium Atlantic Bonito)	$0.764 \pm 0.021\text{b}$	152.8
Raw (Horse mackerel)	$0.563 \pm 0.015\text{a}$	112.6	Raw (Atlantic mackerel)	$0.438 \pm 0.013\text{a}$	87.6
Fried	$0.571 \pm 0.011\text{a}$	114.2	Smoked (Atlantic mackerel)	$0.788 \pm 0.012\text{b}$	157.6
Grilled	$0.580 \pm 0.009\text{a}$	116	Raw (Salmon)	$0.271 \pm 0.012\text{a}$	54.2
Steamed	$0.517 \pm 0.016\text{b}$	103.4	Smoked salmon	$0.405 \pm 0.001\text{b}$	81
Raw (Small bluefish)	$0.545 \pm 0.014\text{a}$	109	Raw (Rainbow trout)	$0.258 \pm 0.008\text{a}$	51.6
Fried	$0.623 \pm 0.008\text{b}$	124.6	Smoked trout	$0.360 \pm 0.017\text{b}$	72.0
Grilled	$0.664 \pm 0.017\text{c}$	132.8	Raw (Eel)	$0.250 \pm 0.031\text{a}$	50.0
Steamed	$0.627 \pm 0.011\text{d}$	125.4	Smoked (Eel)	$0.496 \pm 0.022\text{b}$	99.2
Raw (Medium bluefish)	$0.428 \pm 0.002\text{a}$	85.6	Raw (Small Atlantic Bonito)	$0.426 \pm 0.010\text{a}$	85.2
Fried	$0.635 \pm 0.024\text{b}$	127	Salted (Small Atlantic Bonito)	$0.615 \pm 0.016\text{b}$	123
Grilled	$0.880 \pm 0.019\text{c}$	176	Raw (Medium Atlantic Bonito)	$0.389 \pm 0.025\text{a}$	77.8
Steamed	$0.536 \pm 0.020\text{d}$	107.2	Salted (Medium Atlantic Bonito)	$0.455 \pm 0.028\text{b}$	91
Raw (Small Atlantic Bonito)	$0.415 \pm 0.005\text{a}$	83	Raw (Sardine)	$0.893 \pm 0.019\text{a}$	178.6
Fried	$0.582 \pm 0.015\text{b}$	116.4	Saltd (Sardine)	$1.786 \pm 0.260\text{b}$	357.2
Grilled	$0.653 \pm 0.012\text{c}$	130.6	Raw (Horse mackerel)	$0.637 \pm 0.023\text{a}$	127.4
Steamed	$0.545 \pm 0.014\text{d}$	109	Dried horse mackerel	$1.281 \pm 0.024\text{b}$	256.2
Raw (Striped red mullet)	$0.465 \pm 0.012\text{a}$	93	Raw (anchovy)	$0.357 \pm 0.012\text{a}$	71.4
Fried	$0.750 \pm 0.001\text{b}$	150	Marinated anchovy	$0.555 \pm 0.022\text{b}$	111
Grilled	$0.655 \pm 0.017\text{c}$	131	Raw (Tuna)	$0.847 \pm 0.009\text{a}$	169.4
Steamed	$0.562 \pm 0.002\text{d}$	112.4	Canned tuna in water	$0.643 \pm 0.015\text{b}$	128.6
Raw (Whiting)	$0.464 \pm 0.008\text{a}$	92.8	Canned tuna in oil	$0.656 \pm 0.011\text{b}$	131.2
Fried	$0.631 \pm 0.008\text{b}$	126.2	Mean of smoked fish	$0.625 \pm 0.236\text{a}$	125
Grilled	$0.701 \pm 0.005\text{c}$	140.2	Mean of salted fish	$0.952 \pm 0.727\text{a}$	190.4
Steamed	$0.556 \pm 0.007\text{d}$	111.2	Mean of dried fish	$1.281 \pm 0.024\text{a}$	256.2
Mean of raw fish	$0.463 \pm 0.071\text{ab}$	92.6	Mean of marinated fish	$0.555 \pm 0.022\text{a}$	111
Mean of fried fish	$0.574 \pm 0.165\text{a}$	114.8	Mean of canned fish	$0.650 \pm 0.001\text{a}$	130
Mean of grilled fish	$0.659 \pm 0.122\text{a}$	131.8			
Mean of steamed fish	$0.537 \pm 0.083\text{ab}$	107.4			

sumed in Slovak Republic as 0.505–0.196 µg/g. Erkan and Özden (2007) determined selenium content of sea bass (0.28 µg/g) and sea bream catfish (0.24 µg/g). This result is similar to the Se content of sea bass (0.227 µg/g) described by Satovic and Beker (2004). This value was lower than that reported by Önnig (2010) for herring (0.347 µg/g), mackerel (0.498 µg/g), turbot (0.473 µg/g), flounder (0.371 µg/g) and the value reported by Orban et al. (2000) for sea bream (1.37 µg/g). Plessi et al. (2001) reported the selenium content of angler fish (0.173 µg/g), salmon (0.353 µg/g), redfish (0.247 µg/g), and porgy (0.286 µg/g). This is in agreement with findings of Iqbal et al. (2008), who reported Se contents of Pakistani sea fishes between 0.295–0.714 µg/g. Selenium levels in the literature were reported 0.506 µg/g for bogue, 0.117 µg/g for gilthead, 0.297 µg/g for sardine and 0.627 µg/g for trout from Greek market (Pappa et al., 2006) and 0.473 µg/g for catfish, 0.881 µg/g for mackerel, 0.523 µg/g for pomfret from Thai market (Sirichakwal et al., 2005). Se levels for fishes from Slovenian markets (Smrkolj et al., 2005) were reported between 0.153–0.686 µg/g. In the literature, fish-based foods have the highest selenium contents among all food types, with a Se content of 0.23 µg/g; 0.40 µg/g being the average (Amodia-Coccheri et al., 1995; Dashti et al., 2004). Selenium levels in fried fish were found 0.223 µg/g for anchovy, 0.571 µg/g for horse mackerel, 0.623 µg/g for small bluefish, 0.635 µg/g for medium bluefish, 0.582 µg/g for Atlantic mackerel, 0.750 µg/g for red mullet, 0.631 µg/g for whiting. The lowest and the highest Se levels in grilled samples found were 0.479 µg/g in anchovy and 0.880 µg/g in medium bluefish. The minimum and maximum selenium values of steamed samples were found as 0.361 µg/g in anchovy samples and 0.627 µg/g in small bluefish samples (see Tab. 1). The ranges of selenium in µg/g for grilled fishes in literature were reported as 0.97 µg/g for sardine (0.68–1.38 µg/g), 1.85 µg/g for horse mackerel (1.29–2.32 µg/g), 0.38 µg/g for gilthead sea bream (ND–0.61 µg/g) and 1.49 µg/g for silver scabbard fish (1.18–1.87 µg/g) (Martins et al., 2011). Selenium levels in the literature were reported 0.160 µg/g for fried fish and 0.340 µg/g for grilled fish from Australian market (Mc Naughton and Marks, 2002). The ranges of these metals in µg/g for fried fishes in literature (Martins et al., 2011) were reported as 1.28 µg/g for horse mackerel (1.12–1.46 µg/g) and 0.84 µg/g for hake (0.62–1.13 µg/g). Selenium content in the literature (Hussein and Bruggeman, 1999) ranged from 0.308 to 0.387 µg/g for muscles of fried fish from Egyptian. The ranges of Se in µg/g for boiled fish in literature were reported as 1.22 µg/g for hake (0.84–1.41 µg/g) and 0.87 µg/g for octopus (0.77–0.99 µg/g) (Dashti et al., 2004).

Selenium was found at an average level of 0.625 µg/g in smoked fish. Smoked small Atlantic Bonito and smoked Atlantic mackerel again had the highest levels of Se (0.936 and 0.788 µg/g, respectively) followed in decreasing order by smoked medium Atlantic Bonito and smoked eel (0.764 and 0.496 µg/g, respectively). The lowest selenium contents in smoked fish were 0.360 µg/g in rainbow trout (Tab. 1). Selenium levels in the literature were reported 0.632 µg/g for smoked fish from Australian market (Hussein and Bruggeman, 1999). Selenium average levels in salted fish were found 0.952 µg/g. The lowest and highest selenium contents in salted fish were in medium Atlantic Bonito (0.455 µg/g) and in sardine (1.786 µg/g). Selenium contents of dried fish were determined 1.281 µg/g. Selenium values were 0.555 µg/g and 0.643–0.656 µg/g for marinated anchovy and canned tuna, respectively. Selenium levels in the

literature were reported 0.223 µg/g for canned sardine from Egyptian (Mc Naughton and Marks, 2002) and 0.571 µg/g for canned sardine, 0.859 µg/g for canned tuna from Croatia (Klapec et al., 2004), 0.70 µg/g for canned tuna in brine (Murphy and Cashman, 2001), 0.14 µg/g for canned red mullet (Tüzen and Soylak, 2007).

Generally, the increase in selenium content was not significant ($p > 0.05$) for cooked and processed samples when compared with the raw samples. The cause of the rise is in water loss during this cooking and processing. Martins et al. (2011) found an increase of selenium after cooking in grilled sardine, grilled and fried mackerel, fried hake, whereas the same publication reported a considerable decrease in the selenium content after household preparation of boiled hake. Selenium, essential mineral is necessary for good health. Selenium plays a protective role in preventing carcinogenesis and other chronic diseases (Dashti et al., 2004). There is evidence that selenium has an antioxidant role (Hartikainen, 2005). Se deficiency can lead to heart disease, hypothyroidism and a weakened immune system (Ellis and Salt, 2003; Sirichakwal et al., 2005). Selenium is necessary for health, although there may be toxic at high doses. Selenium intake of 900 µg a day for hair loss and the morphological changes in fingernails have been reported in sufficient doses. The U.K Department of Health has recommended the maximum safe Se intake from all sources to be 450 µg per day for adult males (Foster and Sumar, 1997). According to the European Food Safety Authority (2006), tolerable upper intake value is 300 µg per day for adult. The recommended intake value of selenium varies between countries. Recommended daily intake of Se in Turkey has been reported as 30 µg/day (Foster and Sumar, 1997). However, selenium requirement of adults is calculated to be 70 and 55 µg/day for males and females, respectively (RDA, 1989). A joint FAO/IAEA/WHO Expert Consultation (WHO, 1996) gave several modes for the calculation of requirements of the individual and populations. For a 65 kg reference man the average normative requirement of individuals for selenium was estimated to be 26 µg/day, and from this value the lower limit of the need of population mean intakes was estimated to be 40 µg/day. The corresponding values for a 55 kg reference woman were 21 and 30 µg selenium/day, respectively. Reference Intake of 55 µg selenium per day for adults, but also other levels of intakes based on other criteria, was established by the Scientific Committee for Food of the European Commission (1993). Bioavailability of selenium in food samples is affected by its chemical form and also other dietary factors such as total protein, fat and heavy metal contents (Bhattacharya et al., 2003). This is the first study concerning the evaluation of the effects of cooking and processing methods on selenium contents of fish species consumed in Turkey. According to this study results, with a mean fish meal consumption of 200.0 g/person/day, the average daily selenium intake was found 114.8 µg in fried fish, 131.8 µg in grilled fish and 107.4 µg in steamed fish. The highest daily intake were found as 150 µg in fried red mullet samples, 176 µg in grilled medium bluefish samples and 125.4 µg in steamed small bluefish samples. The lowest and highest daily selenium intakes for processed fish products were found as 81 µg in smoked salmon and 357.2 µg in salted sardine samples. Mean daily selenium intake was found of smoked fish and canned fish (125 µg and 130 µg). This value agrees well with the recommended nutrient intake for adults proposed by FAO

(2001) (25–34 µg/day) and by FSA (2003) (60–75 µg/day). It is also far below the upper tolerable nutrient level of 400 µg/day (FAO, 2001).

Conclusions

In this study, the selenium content of consumption preferred and economically important of fishes was found similar to the literature data. The applied different cooking and processing techniques showed positive effects in the selenium content of these fishes. Measured values is more than sufficient to meet the daily selenium intake values and is far below the tolerable upper intake value of selenium. This is the first study concerning the evaluation of the effects of cooking and processing methods on selenium contents of fish species consumed in Turkey.

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Conflict of interest

I certify that there is no potential conflict of interest in relation to this article.

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+++ Nachrichten aus Forschung, Politik und Industrie +++

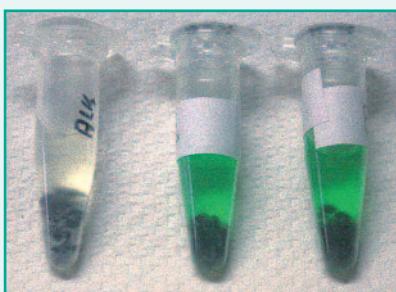
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Falscher Kaviar aus Bulgarien und Rumänien

Ein erheblicher Teil des in Bulgarien und Rumänien verkauften Kaviars wird mit falschem Etikett verkauft oder ist sogar gefälscht. Dies fanden Forscher des Leibniz-Instituts für Zoo- und Wildtierforschung (IZW) und des WWF Österreich durch eine Marktuntersuchung heraus. Die Studie wurde jetzt in dem wissenschaftlichen Fachmagazin *Journal of Applied Ichthyology* veröffentlicht.

Analysen von 27 Kaviarproben aus Rumänien und Bulgarien ergaben, dass die Anzahl an Kaviardosen, die mit falschem oder überhaupt keinem Etikett zum Verkauf angeboten werden, unerwartet hoch ist. Eigentlich müssen alle Kaviargläser und -dosen durch einen universellen Etikettierungscode gekennzeichnet sein, welcher die wichtigsten Informationen über die Herkunft des Kaviars, z. B. Störart, Aquakultur oder Wildfang und Herkunftsland, angibt. In sieben Fällen wurde der Kaviar jedoch illegal ohne Etikett von Straßenverkäufern oder in Geschäften verkauft. Durch genetische Analysen wurde bei allen Kaviarproben die Störart bestimmt. Unter den etikettierten Dosen stimmten lediglich zehn Proben mit der angegeben Art überein. Vier Proben enthielten Kaviar von einer anderen bzw. mehreren, nicht auf dem Etikett genannten Störarten. Bei mindestens einer der falsch etikettierten Kaviardosen wurde der Kaviar von einer preiswerten zu einer teureren Art aufgewertet. Sechs Proben waren gefälscht. Drei dieser Fälschungen enthielten überhaupt keine tierische DNA und wurden wohl gänzlich künstlich erzeugt. Eine Probe stammte vom Seehasen (*Cyclopterus lumpus*), dessen Eier allgemein als Kaviarsatz verkauft werden. Die anderen beiden Fälschungen sind höchstwahrscheinlich aus Störfleisch hergestellt worden.

„Besonders besorgniserregend für den Schutz der Störe waren vier Proben, die im Restaurant oder von Straßenverkäufern explizit als Kaviar von wilden Donau-Stören angeboten wurden. Die Fische wurden also illegal gefangen“, sagt Arne Ludwig vom Leibniz-Institut für Zoo- und Wildtierforschung (IZW). Der Kaviar stammte von dem stark gefährdeten Beluga-Stör (*Huso huso*), dessen Population im Donaudelta wahrscheinlich kurz vor dem Aussterben steht. „Ein wirkungsvoller Schutz der Störe ist nur möglich, wenn Wilderei und illegaler Handel endlich gestoppt werden“, ergänzt Jutta Jahrl vom WWF Österreich.



Aufbereitung von Kaviarproben mit Alkohol für eine genetische Analyse. Bei den gefälschten Proben BG1 und BG3 tritt künstliche grüne Farbe aus. Links sieht man eine Original-Kaviarprobe mit dem gleichen Verfahren behandelt ohne Abfärbung. Foto: Ida Steie

Echter Kaviar ist eines der teuersten Tierprodukte im weltweiten Handel und wird von Stören und Löffelstören entnommen. Der Preis des Kaviars hängt stark von der Herkunftsart ab, wobei der teuerste Kaviar vom Beluga-Stör stammt. Die Wilderei stellt weltweit eine große Bedrohung für das Überleben der Störe dar. Laut Einschätzung der Weltnaturschutzvereinigung (IUCN) von 2009 sind Störe die weltweit am stärksten gefährdete Tiergruppe. Alle 27 Stör- und Löffelstörarten wurden in die Liste des Übereinkommens über den internationalen Handel mit gefährdeten Arten freilebender Tiere und Pflanzen (CITES) aufgenommen, um sie vor illegalem Handel zu schützen. Dadurch soll jeder internationale Handel von Stör exemplaren, -teilen oder -produkten kontrolliert werden, was auch das oben beschriebene Etikettierungssystem beinhaltet. Die relativ hohe Anzahl an falsch etikettierten Kaviardosen wirft jedoch Zweifel an der Effizienz des CITES-Kennzeichnungssystems und dessen Kontrolle auf.

„Rumänien und Bulgarien sind die einzigen Länder der Europäischen Union, in denen noch immer wildlebende Störpopulationen, z. B. im Schwarzen Meer oder in der Donau, vorkommen“, berichtet Harald Rosenthal, Präsident der World Sturgeon Conservation Society (WSCS). Zwar gibt es in beiden Ländern Fang- und Handelsverbote, jedoch hält offensichtlich die illegale Fischerei weiterhin an. „Die Ergebnisse der Studie verdeutlichen die Schwäche des Störschutzes in Rumänien und Bulgarien. Wir empfehlen daher, den Schutz und die Kontrollmaßnahmen zu verstärken“, betont Arne Ludwig. Er und seine Kollegen plädieren dafür, DNA-Kontrollen von Kaviar intensiver durchzuführen und dabei auch Dosen mit scheinbar korrektem Etikett einzubeziehen.

Weitere Informationen (Quelle):

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