Relation of smoking parameters to the yield, colour and sensory quality of smoked Atlantic salmon (Salmo salar)

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Abstract

The relations between smoking parameters and the characteristics of salmon raw material were investigated with respect to yield, colour, flesh content of phenol and salt, and sensory properties. The fish studied were ocean ranched salmon harvested in Iceland in July 1998 and farmed salmon from Norway slaughtered in October 1998 and April 1999. Seven treatments were applied on fresh or frozen raw material combining dry or brine salting with cold smoking at 20 or 30 °C. Electrostatic smoking was tested on dry-salted salmon fillets. The results show a lower yield after filleting and trimming with ocean ranched fish. Although freezing had little effect on yield, total loss was slightly greater, especially for fish with low fat content. Sensory differences were also apparent. The brine salting technique resulted in lower losses. Fish with higher fat content gave a better yield after processing, although careful control of the smoking procedure was required (especially at 30 °C) to avoid a case-hardening effect. With brine salting, salt uptake was higher for smaller, leaner fish. The phenol content of flesh depended on the technique and/or smoking temperature used, regardless of the fish studied. However, for a smoking temperature of 30 °C, the flesh of smaller, leaner fish showed a higher phenol level. Smoking conditions and preliminary treatment such as freezing produced similar differences in sensory characteristics, regardless of the fish studied, although smaller, leaner individuals appeared to be more sensitive to these processes. © 2001 Elsevier Science Ltd. All rights reserved.

Keywords: Atlantic salmon; Smoking; Yield; Salting; Freezing; Quality parameters

1. Introduction

Important changes in food consumption and distribution systems during the second half of the 20th century have been associated with extensive industrialisation of production structures. In this context, smoked salmon, which was initially regarded as a luxury product, has gradually become an element of general consumption. Initially produced in the 1920s in salt-dried, cold-smoked form by simple tradesmen, it has now become an industrial product on a highly developed market (Monfort, 1999). The development of aquaculture of the Atlantic salmon (Salmo salar) has contributed very largely to commercial growth (Hempel, 1998, 1999; Roberts, 1998), allowing producers of smoked salmon to have ready access (less than 6 days) to fresh raw material of uniform quality available year-round.

As the cost of this raw material accounts for a large part of total production costs, firms need to seek supplies of fresh salmon from highly diversified markets. For France, the raw material is mainly of aquacultural origin. Although the main production areas in Europe for Atlantic salmon are in Scandinavian countries, the rearing procedures can vary considerably from one country to another and from one farm to another. These variability factors are numerous and concern rearing conditions as well as slaughtering, packaging and transport (Torrissen, 1995). Moreover, the processing firms now have a large choice of equipment for better
control of production parameters. Thus, the producer must have perfect knowledge of the raw material and the effects of the processes used, so that the characteristics of the finished product will be consistent with market demand and profitability requirements.

Several factors are involved in this notion of profitability and production control (Sigurgisladottir, Torrisen, Lie, Thomassen, & Hafsteinsson, 1997). The most important ones are processing yields and a respect for the characteristics of the finished product. The latter, independently of health considerations, concern chemical (salt, water, lipid and phenol contents), physical (texture and colour) and sensory aspects.

For smoked salmon, yield and quality are related to the operations involved in fillet production, such as head removal, filleting and trimming (Rorá, Kvale, Morkore, Rorvik, Steien, & Thomassen, 1998), the choice of the processes used (salting, drying and smoking techniques), and the parameters relating to these processes, such as brine concentration, length of treatment, temperature or hygrometry (Chan, Toledo, & Morkore, 1995; Sigholt, Erikson, Rustad, Johansen, Knockaert, 1990; Le Gall, 1938), all of which are closely dependent on the characteristics of the raw material.

Various studies have been performed concerning the optimisation of salting, drying and smoking processes (Doe, Sikorski, Haard, Olley, & Sun Pan, 1998; Knockaert, 1990; Le Gall, 1938) and the quality of fresh (Aursand, Bleivik, Rainuzzo, Jorgensen, & Mohr, 1994; Berg, Erikson, & Nordtvedt, 1997; Hillestad, Austreng, & Johnsen, 1995; Sigholt, Erikson, Rustad, Johansen, Nordtvedt, & Seland, 1997) and smoked Atlantic salmon (Indrasena, Hansen, & Gill, 2000; Rorá et al., 1998; Wang, Tang, & Correia, 2000).

Thus, the size and shape of fish have an effect on processing yields (Rorá et al., 1998) and the choice of processing parameters. Likewise, lipids in flesh play an important role as a limiting factor during the salting and drying steps (Bohuon, 1995; Jason, 1965a), either replacing the aqueous phase that serves as a vector for transfers during these steps or acting as a physical barrier. On the contrary, the presence of lipids during the smoking step is a positive factor for the uptake of smoke compounds (Beltran & Moral, 1991; Korhonen, Reagan, Carpenter, Campion, & Stribling, 1978).

The purpose of the present study was to detect and analyse interactions between the characteristics of the raw material and certain processing parameters relative to the yields and quality of smoked salmon. Two salting techniques (brine or dry salt) were associated with two cold smoking temperatures (20 and 30°C). The deposition of smoke compounds by an electrostatic smoking method was also studied, as well as the effect of freezing the raw material. Our intention was to propose technical recommendations for better control of the finished product based on a relevant choice of the characteristics of the raw material with respect to the processes available.

This study is part of a larger undertaking within the scope of European Union FAIR project No. CT 95-1101 (“Interaction between raw material characteristics and smoking process on quality of smoked salmon”) coordinated by the Icelandic Institute (Matra) in association with the Instituto del Frio (Spain), the Institute of Marine Research (Norway), the Institute of Nutrition (Norway), Akvaforsk (Norway) and IFREMER (France).

2. Materials and methods

2.1. Materials

2.1.1. Animal material

Three groups of 105 fish each, from different origins and with different rearing characteristics as described by Sigurgisladottir, Ingvarsdottir, Torrisen, Cardinal, and Hafsteinsson (2000), were compared in this study. The first sample (Group A) was obtained from ocean reared salmon harvested in July 1998 in Iceland. The average weight of these fish was 2.6 kg±300 g (individuals weighing 4 kg were expected, but proved impossible to obtain because of planning problems). In these conditions, no sorting according to weight was performed. The second sample consisted of farmed salmon harvested in October 1998 (Group B) with an average weight of 4 kg±150 g. The third group, also consisting of farmed salmon, was harvested in Norway in April 1999 (Group C) and had an average weight of 3.7 kg±500 g.

All fish were harvested by netting and bled by cutting the gill arches on one side. After a bleeding step in cold seawater, salmon were gutted, cleaned and individually weighed and tagged. Fish were placed on ice in sealed boxes, shipped to AKVAFORSK for fat determination and then transported by refrigerated truck to IFREMER (French Research Institute for the Exploitation of the Sea) in Nantes, France. The delay between slaughtering and the beginning of treatment was 7 days.

To study the effect of freezing, 30 fish within each group were sent to Nantes 1 month earlier and were frozen (1 h) with cryogenic equipment (CO₂ at −60°C) and stored at −20°C. A thawing step in air was performed at 4°C for 24 h.

2.1.2. Processing equipment

Two different techniques were compared for drying and smoking salmon fillets. The traditional method used an HMI Thirode (PC90 Model) apparatus with a capacity of 380 kg mounted on a trolley with 28 grids. The smokehouse measured 150×130×225 cm, and air/smoke circulation was horizontal. The airflow rate and relative humidity were controlled. Smoke was produced with a generator by pyrolysis (450°C) of sawdust from beechwood and sustained airflow (Thirode, France). The electrostatic smoking method (Bardin, Desportes,
Knockaert & Vallet, 1997; Collignan, Knockaert, Raoult-Wuck, & Vallet, 1992, 1993) was performed using an experimental tunnel 3 m long, 35 cm wide and 20 cm high. This allowed continuous smoking at ambient temperature, with a production capacity of 125 kg/h. Tension applied between the positive pole (smoke ionisation) and the cathode (conveyor belt) was 40 kV. Air speed above the fillets was 0.5 m/s, and relative humidity around 70%. Smoke was obtained from a friction generator (Muvero, The Netherlands) using oak beam at a temperature of 350°C.

2.2. Methods

2.2.1. Sample preparation

Gutted fish were weighed and filleted. Each fillet was trimmed by hand and processed according to the preparation scheme in Fig. 1. Each group of 105 fish was processed according to seven different treatments, including different salting and smoking techniques. 15 salmon were used for each procedure, and fillet weight was recorded at each step to calculate yields.

2.2.1.1. Salting. Two salting techniques were studied. The first was a dry salting technique in which fillets salted by hand with refined salt (Salins du Midi, France) were left for 6 h (4 h for ocean-ranched salmon because of their small size) at 12°C. The fillets were then rapidly rinsed with water (15°C) on grids and stored in a cold room at 2°C until smoking.

The brine salting technique used saturated brine (360 g/l) maintained at 12°C in which the fillets were placed [ratio 50/50 (w/v)]. After 6 h (4 h for ocean-ranched salmon), the fillets were removed, rapidly rinsed and stored overnight (12 h) in a cold room at 2°C until smoking. Fillets were weighed before smoking.

2.2.1.2. Drying–smoking. A traditional cold-smoking process was carried out at two different temperatures in accordance with industrial practices based on different materials (Knockaert, 1990). After storage for 12 h at +2°C, the smoking process began with a drying step in the smoking oven for 30 min at 20°C, followed by a smoking step at 20°C±1°C, with a relative hygrometry of 65±3% and air speed of 2 m s⁻¹ above the products and 18 m s⁻¹ at the end of the air channel. Smoking was performed for 2.5 h. The same procedure was carried out for fish smoked at 30°C, with a relative humidity of 50%. With electrostatic smoking, fillets were maintained at +2°C for 12 h and then placed in the smoking tunnel for 15 min.

2.2.1.3. Slicing and packaging. One day before sensory sessions and chemical analyses, fillets were frozen by the cryogenic method (Airgaz) for 20 min at −60°C to an inner temperature of −7°C and mechanically sliced (PNP, France). Skin was removed before slicing. Sliced fillets were immediately vacuum-packed (Boulanger, France). Samples were stored at +2°C until analysis.

2.2.2. Analytic methods

2.2.2.1. Yield measurement. Filleting and trimming loss = 100×(gutted weight of fish − weight after filleting and trimming)/gutted weight of fish.

Salting loss = 100×(fillet weight after salting and trimming − fillet weight after salting)/fillet weight after salting.

Total loss after salting and smoking = 100×(fillet weight after filleting and smoking − fillet weight after smoking)/fillet weight after salting.

Relative loss after salting = (weight loss after salting/weight after salting and smoking)×100.

Relative loss after smoking = (weight loss after smoking/weight after salting and smoking)×100

2.2.2.2. Chemical analyses. Total fat content was determined on gutted fish by non-destructive computer-assisted X-ray tomography, as described by Rye (1991). The correlation equation between data from X-ray tomography and data from chemical analysis allowed fat content values to be expressed as classical chemical results. Dry matter content was analysed by oven drying of 2 g of smoked salmon at 105°C until a constant weight was reached and salt content was measured with Chloride Analyser 926 (Corning, Halstead, UK). Total phenols were quantified by the method described in the French standard for smoked salmon (NF V 45-065, 1995). All these analyses were performed on the front part of each smoked fillet.

2.2.2.3. Sensory evaluation. Descriptive and quantitative analysis (Stone & Sidel, 1985; Stone, Sidel, Oliver, Woolsey, & Singleton, 1974) was performed to evaluate the sensory characteristics of smoked salmon. Twenty panellists belonging to the IFREMER staff, selected for their interest, availability and sensorial capacities, were trained on sensory descriptors chosen during a preliminary step, according to the international procedure, NF ISO 11035 (1994). Three training sessions for profiling were organised to check the panellists’ understanding of the descriptors and analyse performances of each assessor before starting the study. The descriptors chosen related to the appearance, odour, flavour and texture of smoked salmon slices: odour: smoke intensity, wood fire, acrid smell of smoke, toasted bread, fresh salmon; appearance: pink colour, orange colour, homogeneity of colour; darkness of the slice edge, translucent appearance, fatty aspect, white stripes; flavour: smoke intensity, wood fire, salty taste, fresh salmon; and texture: firmness, melting texture, fatty texture, pasty texture.
Whole gutted salmon (*Salmo salar*)
A. Ocean ranched (average weight 2.6 kg), July 1998
B. Farmed salmon from northern Norway (4 kg), October 1998
C. Farmed salmon from western Norway (3.7 kg), April 1999

**Frozen fish**
(storage at −18°C for 1 month)

**Fresh fish**
(in ice)

**Manual Filleting**
Trimming

**Dry salting**

**Salting techniques**

**Brine salting**

**Rinsing**
Storage for 12 hours at 2°C

**Traditional cold smoking**

**Electrostatic smoking**

Ambient Temperature

$20°C$  $30°C$  $20°C$  $30°C$

$2$  $6$  $1$  $5$  $4$

**Smoking procedures**

**Sample code**

$20°C$  $30°C$

$3$  $7$

**Vacuum packing and Storage at +2°C until analysis**

**Mechanical slicing**
After cooling to −7°C

**Analyses**

Fig. 1. Process steps.
Four sessions were proposed to the panel for each raw material. Samples smoked at 20°C and those smoked with the electrostatic process were evaluated twice, after 8 and 9 days of storage at 2°C. Salmon smoked at 30°C were also scored twice, after 12 and 13 days. Samples dry-salted and smoked at 20°C were used as a common reference in these four sessions. Because samples were stored at 2°C, they were considered to keep the same characteristics between 8 and 13 days of storage. Each panellist received two slices taken from the third front part of a fish. Seven or eight fish were used for each session.

Sessions were performed in individual partitioned booths equipped with a computerised system (Fizz system, Biosystèmes, Dijon, France). These conditions were conducive to concentration and avoided communication between assessors and disturbance by external factors (NF V-09-105, 1995). Panellists rated the sensory attributes on a continuous scale presented on a computer screen, from low intensity (0) to high intensity (10). Products were assigned three-digit numbers, randomised and served simultaneously. Twelve sessions were organised to test all the raw material and processes.

2.2.2.4. Colour measurement. Colour was determined by a Hunterlab miniscan XE using the D65 light source and a 10° observer. Reflectance from 400 to 700 nm was recorded as well as $L^*$, $a^*$, $b^*$ according to International Commission on Illumination (CIE) (1976). Colour was measured in the anterior, median and posterior parts of the dorsal and ventral side of raw and smoked fillet. Results are shown as the mean of six measurements per fish fillet.

2.2.2.5. Statistical analysis. The mean, standard deviation, analysis of variance and Duncan’s multiple range test were performed using Statgraphics Plus software (Sigma Plus, Paris, France). The significant statistical level was set at $P<0.05$. Multivariate data processing was performed with Uniwin Plus 3 software (Sigma Plus, Paris, France).

To adjust for variations among assessors in the scoring range, sensory data were standardised using an isotropic scaling factor according to the procedure proposed by Kunert and Qannari (1999). Scores of the dry-salted product smoked at 20°C were used as a reference at each session, thereby allowing data from different sessions to be compared.

The study of differences between samples was done by analysis of variance on each descriptor, and principal component analysis was performed on the means of these data.

3. Results and discussion

3.1. Influence of raw material and processing on yields

3.1.1. Raw material effect on loss after filleting and trimming

Flesh lipid content showed significant differences among the three raw materials: 9.4, 20.0 and 16.8% of wet weight respectively for ocean ranched salmon, farmed salmon slaughtered in October 1998 and farmed salmon slaughtered in April 1999. Analysis of variance for loss after filleting showed a significant difference between the different raw materials. Ocean-ranched salmon from Iceland, which were smaller and leaner than the fish in the other groups, had the highest loss (Fig. 2). According to Rora˚ et al. (1998), this result can be regarded as an effect of weight more than fat content. The difference between groups was as great as 1%. Moreover, wild salmon were not sorted and weight variability in that group was greater than in the other groups. From an industrial point of view, this heterogeneity could raise problems depending on the process used.

3.1.2. Effect of freezing raw material on yields after processing

Losses after dry salting were generally slightly higher with frozen material (Table 1). The difference in losses between fresh and frozen raw material was 0.1% in October 1998, 0.3% in April 1999 and reached 0.6% in July 1998. The difference was only statistically significant in the ocean ranched group. However, losses with frozen material after dry salting were generally low.

![Fig. 2. Loss after filleting and trimming for three raw materials. Group A, ocean ranched salmon (July 1998); Group B, farmed salmon (October 1998); Group C, farmed salmon (April 1999); mean values are given for each group; different letters indicate significant differences ($P<0.05$).](image-url)
compared to total loss. This may seem contradictory with the results of many previous studies indicating that all treatments affecting cell integrity, such as freezing, increase solute intakes at the expense of water (Dussap & Gros, 1980; Ponting, 1973; Saurel, Raoult-Wack, Rios, & Guibert, 1994). However, in our study, the cryogenic method used for freezing, as well as the short storage period (1 month), limited tissue destruction and allowed the raw material to be preserved and to retain its good functional properties, as reported by Sivertsvik (1994) for frozen *Salmo salar*. Nonetheless, if the complete process is considered, including the drying and smoking steps, total losses were greater with frozen raw material, especially for ocean ranched salmon.

After a drying/smoking step at 20°C, the freezing step produced a significant loss of raw material. There was also a relation between the kind of raw material and freezing treatment, which indicated that loss after smoking was higher to the extent that fish were smaller and leaner. For smoking at 30°C, losses were not affected by a freezing step (Table 1).

### 3.1.3. Effect of the salting technique on yields after processing

Our results showed better yields with the brine-salting than the dry-salting technique (Table 1). Solute transfer during the salting step occurs in the water phase between intracellular fluid and the outside medium (Soudan, 1955). However, depending on the kind of salting, transfers do not start in the same way. For dry salting, the water phase (which does not exist initially) has to be created by extraction of intracellular water to the surface of the product (Dieuzaide & Novella, 1951). On the contrary, for brine salting, salmon fillets are already soaked in a liquid solution, so that water diffusion is reduced and losses are lower (Table 1). The effects are greater to the extent that the raw material is lean (i.e. with high water content), as observed in group A for the loss recorded after salting. These results are in accordance with those of Boury (1934) and Jason (1965b), who demonstrated the importance of the size factor and the barrier effect of lipids. Sheehan, O’Connor, Sheehy, Buckley, and FitzGerald (1996) and Wang et al. (2000) also indicated that the transfer of a salted solution decreases with an increase of lipid content.

### 3.1.4. Effect of the smoking technique or drying/smoking temperature on yields after processing

From a physical point of view, drying (which occurs at the surface of the product) is due to a two-step migration of water: first, evaporation of surface water, and then the diffusion of water within the flesh toward the surface of the fillet (Cheftel & Cheftel, 1977). One of the factors affecting the migration mechanism in flesh is the chemical composition, particularly lipid content. The speed of water diffusion in the flesh of lean fish is faster than in fat fish, which leads to more rapid drying and higher losses (Table 1). In the case of wild salmon (group A), the phenomena of surface evaporation remain at a high level during drying because of the greater quantity of water available in the fillet. This is particularly apparent in the case of brine salting for which losses are lower, so that the fillet retains a higher water content.

The drying-smoking step at 20°C leads to greater losses than at 30°C. In fact, at 20°C the hygrometry is easier to control and drying occurs in more favourable conditions. Industrial drying, when well performed, supposes an optimal difference in hygrometry (a maximum of 10%) between the surface of the product and ambient temperature, which can be ensured for a 20°C step, but is difficult to attain with 30°C (Knockaert, 1990). In the latter case, the hygrometric differential is around 15–20%, which accelerates surface evaporation.

<table>
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<tr>
<th>Table 1</th>
<th>Weight loss after each process step (%)</th>
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<tr>
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<td>Dry salting (Frozen material)</td>
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<tr>
<td>Loss after salting</td>
<td>Group A</td>
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<td>Group B</td>
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<td>Group C</td>
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<td>Drying/Smoking process (C)</td>
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<td>Loss after smoking</td>
<td>Group A</td>
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<td>Group C</td>
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<td>Total Loss</td>
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<td>Group B</td>
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<td>Group C</td>
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* a 20°C, traditional cold smoking at 20°C; 30°C, traditional cold smoking at 30°C; E, smoke deposit with the electrostatic method.
* b Group A, ocean ranched salmon (Iceland, July 1998); Group B, farmed salmon (October 1998); Group C, farmed salmon (April 1999).
* c Mean values are given for each group; on each row, different letters indicate significant differences (*P* < 0.05).
In these conditions, the water diffusion rate for fat fish is lower than the rate of surface evaporation, which induces case-hardening and thus a blockage of transfer and a reduction of loss occurs. This phenomenon has been described by Bimbenet (1978), who noted that too low a hygrometry associated with too high a temperature alters drying profiles. The surface of the flesh is modified by a concentration of solutes that block the capillaries and the arrival of water, thus preventing evaporation completely. This case-hardening effect can be detrimental to product quality since it prevents effective drying and can reduce product life.

For smoking at 30°C, the results were the same whether the raw material was frozen or not. As the drying conditions were more favourable at 20°C and water more available in the flesh after the freezing step, the losses were greater.

For ocean ranched salmon, electrostatic smoking led to losses equivalent to those for smoking at 20 or 30°C. Given the very short smoking period and the absence of a drying step, this suggests that losses occurred during storage in the cold room (12 h at 2°C) before the smoking operation. During this storage period, it is likely that dry-salted fillets reached a level of hygrometry similar to that maintained in the smoking unit during processing. As losses in brine after salting were lower, processing at 20 and 30°C still had an effect on weight loss. In the case of fatter farm fish (groups B and C), in which water was less available, overnight storage at +2°C was inadequate to produce a notable drying effect. Thus, drying continued during the drying–smoking step.

3.1.5. Overall effect of salting and smoking conditions on yields after processing

Regardless of the processing technique used, losses were higher with ocean ranched salmon, particularly when frozen (Table 1). This was essentially attributable to the size effect in direct relation with chemical composition. Fig. 3 indicates the relative role of each processing step (salting and smoking) in the losses recorded. For samples smoked by the electrostatic method, losses were preponderant during salting.

For dry-salted ocean ranched salmon, losses were greater during the salting than the smoking step, whereas brine-salted samples showed greater losses during the smoking step. Moreover, when lipid content increased in the raw material, losses were smaller during the salting step.

Because of the organoleptic effect and product preservation, industrial specifications for “smoked finished products” generally recommend a water content in flesh of less than 65% (Red Label standard, Label No. 33-90 and Label No 4-94, France). To obtain this result, producers must take account of the lipid content in raw material and the type of processes used, notably for salting. The choice of smoking parameters is determinant to avoid case-hardening of the product.

3.2. Effect of smoking techniques or smoking temperature on quality parameters

3.2.1. Phenols content

Phenol content allowed samples to be differentiated according to the smoking technique or smoking

![Relative loss (%)](image-url)
temperature (Fig. 4). The lowest levels (0.2–0.55 mg/100 g of flesh) were observed for the electrostatic process. With a 20°C smoking temperature, mean phenol content in the groups ranged from 0.75 to 1.2 mg/100 g, whereas a 30°C temperature involved levels of up to 3 mg/100 g. Various reasons could account for these differences, including the type of smoke production and the type of smoke deposit. Despite the use of oak instead of beechwood with the electrostatic technique, processing time could not exceed 15 min in order to limit surface coloration. Therefore, phenols deposits were reduced. Moreover, the electrostatic field modifies the smoke compound ratio in the vapour phase, mainly by increasing the level of carbonyl compounds (Ruiter, 1979; Sirami, 1981). However, the 30°C temperature used with traditional smoking allows compounds with higher molecular weight, such as phenol compounds, to remain in the vapour phase mainly involved in the smoking effect (Foster, 1961a; b; Girard, 1988; Potthast, 1977, 1978). Therefore, samples can reach high phenol values.

A raw material effect was also noted for the 30°C smoking technique, i.e. the lower the fat content in muscle, the higher was the phenol content in smoked samples. This may have been due to water availability relative to fat content. As noted above, very high surface desiccation leads to a reduction of solute diffusion and prevents interstitial water and lipid compounds from taking up smoke components (Clifford, 1980). The relative thinness of the fillet was another reason for higher phenol values in small fish.

3.2.2. Salt content

Mean salt content in the groups of smoked fish ranged from 2.2 to 4 g/100 g of wet flesh (Fig. 5). Two main factors influenced salt uptake: the nature of the raw material (particularly size and composition) and the kind of salting technique used. As previously reported (Jason, 1965a; Jason & Peters, 1973; Schwartzberg & Chao, 1982; Storey, 1982), high fat content causes greater resistance to the transfer of an aqueous solute such as sodium chloride. This was true for our results, which showed salt uptake of 3.2–4% wet weight for leaner fish and 2.2–3.4% for fatter fish. Moreover, even though salting time had to be reduced for wild salmon (4 instead of 6 h) to produce the same salt level in flesh, the small size of these ocean ranched fish allowed increased salt diffusion. Products salted in saturated brine generally have higher salt content than dry-salted samples because of better contact between surfaces and the salting medium. Brine is recommended for fatty fish because immersion limits exposure to oxidation phenomena in the air (Doe et al., 1998). Salt content in smoked products was slightly higher for frozen raw material than fresh material, especially for leaner fish. This resulted from the freezing step, which modified cell structure slightly, increasing salt diffusivity.

3.2.3. Colour

Colour measurements of raw and smoked fillet surface are given in Figs. 6 and 7. Before processing, ocean ranched fish never fed pigments had lower L*, a* and b* values than the two farmed fish groups. There was no
significant difference in colour parameters between the two reared fish groups. Smoking led to a reduction of \( a^* \) values, as previously reported by Skrede and Storebakken (1986), Choubert, Blanc and Courvalin (1992) and Rø et al. (1998), an increase of \( b^* \) values regardless of the raw material, and a reduction of \( L^* \) values (mainly for farmed fish). Choubert et al. noted that smoking induces a loss of water, with an increase of carotenoid concentration and a decrease of hue and lightness. Ocean ranched salmon showed the lowest \( a^* \) and \( b^* \) values, but there was a colour difference between the two groups of farmed fish. Salmon from group C had lower values than those of group B, possibly because of fat content. Rø et al. found a significant correlation between fat content and colour measurements. In our study, fat content for groups B and C was, respectively, 20.0 and 16.8%. Smoking temperature, however, did not affect parameters \( a^* \) and \( b^* \) in the

![Salt content (g/100 g wet flesh)](image)

Fig. 5. Process effect on smoked salmon salt content (g/100 g). Group A, ocean ranched salmon (July 1998); Group B, farmed salmon (October 1998); Group C, farmed salmon (April 1999); 1: dry salting and smoking at 20°C; 2: frozen raw material, dry salting and smoking at 20°C; 3: brine salting and smoking at 20°C; 4: dry salting and electrostatic smoking; 5: dry salting and smoking at 30°C; 6: frozen raw material, dry salting and smoking at 30°C; 7: brine salting and smoking at 30°C. Different letters indicate significant differences \((P < 0.05)\).

![Colour measurements](image)

Fig. 6. Colour measurements. \( a^* \) and \( b^* \) values of raw and smoked salmon. Colour system \( L^* \), \( a^* \), \( b^* \), CIE 1976 A: ocean ranched salmon (July 1998); B: farmed salmon (October 1998); C: farmed salmon (April 1999); r: fresh raw material, r*: frozen raw material smoking procedures: 1, dry salting and smoking at 20°C; 2, frozen raw material, dry salting and smoking at 20°C; 3, brine salting and smoking at 20°C; 4, dry salting and electrostatic smoking; 5, dry salting and smoking at 30°C; 6, frozen raw material, dry salting and smoking at 30°C; 7, brine salting and smoking at 30°C. Each point is the mean of six readings per fillet for 15 fish.
same way. With a 30°C smoking temperature, \( b^* \) values were greater than \( a^* \) values, i.e. products had a more intense yellowish tone, whereas with smoking at 20°C, \( a^* \) values were higher and the red tone more intense. Regardless of the raw material, \( b^* \) values for raw and smoked fillets were higher when fish had been frozen. This freezing effect is not mentioned by Sheehan, O’Connor, Sheehy, Buckley, and FitzGerald (1998), who noted that colour modification depends on the kind of pigment used in fish diet. Despite the low smoking intensity for samples treated by the electrostatic technique, lightness was low compared to other samples. The level of carbonyl compounds involved in colour and to a lesser extent in flavour is higher with electrostatic smoking (Ruiter, 1979; Sirami, 1981), which could account for the difference of colour.

No colour difference between samples was observed relative to the salting technique.

3.2.4. Sensory properties

Analysis of variance took the raw material effect, the process effect and the interaction between these two factors into account. Table 2 shows the \( F \)-value for each factor. For most sensory criteria, the results show that the process effect was higher than the raw material effect, except for colour descriptors (pink and orange) and firmness (for which the response level depended on the kind of raw material used).

Regardless of smoking temperature, a freezing step before processing affected the quality of smoked salmon, especially if fish were small and lean. Samples whose raw material was frozen had a saltier taste and less translucent slices. Their texture, which was close to that of the electrostatic smoked sample, was less firm and more pasty than that of other products. A texture slightly modified by freezing might actually increase the perception of salt more than the salt content level itself. The results of Sigurgisladottir et al. (2000) also show a significant effect of freezing/thawing on microstructure and texture.

The smoking temperature or technique allowed products to be differentiated according to the intensity of smoke odour and flavour. From the lowest to the highest intensity, the panel ranked samples smoked by the electrostatic method first, followed by those smoked at 20°C and finally those smoked at 30°C. These results are in agreement with those of Simon, Rypinsky, and Tuber (1966) and Girard (1988), who found that a higher temperature increases the deposit of smoke compounds. For products smoked by the electrostatic technique, the panel noted a special toasted bread odour in addition to low flavour intensity. These characteristics could be attributable to various factors, such as the kind of wood used, the smoke production method, pyrolysis temperature, smoke density, or smoking time (Cardinal, Berdague, Dinel, Knocknert, & Vallet, 1997; Holmes, 1991). Moreover, this sample appeared to be fatter, and the texture was described as fatty and melting in the mouth. 20°C smoked samples also showed these characteristics but to a lesser extent. These sensory properties were observed, regardless of the raw material. It is noteworthy that muscle fat content could have modified sensory perception of smoked salmon, as reported by Sheehan et al. (1996). For the same fat level, the kind of smoking procedure may also modify the perception of texture.

Dry salting and brine salting did not appear to have much effect in changing the sensory properties of the

![Fig. 7. Colour measurements on smoked salmon. \( L^* \) and \( a^* \) values. A: ocean ranched salmon (July 1998); B: farmed salmon (October 1998); C: farmed salmon (April 1999); r: fresh raw material; r*: frozen raw material smoking procedures 1, dry salting and smoking at 20°C; 2, frozen raw material, dry salting and smoking at 20°C; 3, brine salting and smoking at 20°C; 4, dry salting and electrostatic smoking, 5 = dry salting and smoking at 30°C; 6, frozen raw material, dry salting and smoking at 30°C; 7, brine salting and smoking at 30°C. Each point is the mean of six readings per fillet for 15 fish.](image-url)
Table 2

<table>
<thead>
<tr>
<th>Descriptors</th>
<th>Raw material effect 2</th>
<th>Process material effect 1</th>
<th>Interaction 1-2</th>
<th>Multiple range tests Duncan for products</th>
<th>Multiple range tests Duncan for raw material</th>
</tr>
</thead>
<tbody>
<tr>
<td>Odour: smoke intensity</td>
<td>3.0NS</td>
<td>116.4***</td>
<td>2.1NS</td>
<td>4a 1b 3b 2b 5c 7c 6c</td>
<td>Ca Aab Bb</td>
</tr>
<tr>
<td>Odour: wood fire</td>
<td>0.94NS</td>
<td>44.8***</td>
<td>1.9NS</td>
<td>4a 2b 6bc 1bc 3cd 7d 5d</td>
<td>–</td>
</tr>
<tr>
<td>Acrid smell of smoke</td>
<td>2.9NS</td>
<td>52.8***</td>
<td>2.6*</td>
<td>4a 1b 3bc 2c 5d 7d 6e</td>
<td>–</td>
</tr>
<tr>
<td>Odour: toasted bread</td>
<td>53.2***</td>
<td>42.1***</td>
<td>19.6***</td>
<td>5a 7a 6ab 3b 1b 2c 4d</td>
<td>Ca Bb Aa</td>
</tr>
<tr>
<td>Odour: salmon/fish</td>
<td>2.9NS</td>
<td>83.8***</td>
<td>7.8***</td>
<td>6a 7a 5b 2c 3d 1d 4e</td>
<td>Aa Cb Bb</td>
</tr>
<tr>
<td>Pink colour</td>
<td>20.3***</td>
<td>7.6***</td>
<td>7.9***</td>
<td>5a 7a 4ab 1bc 3c 2c 6c</td>
<td>Ca Bb Ab</td>
</tr>
<tr>
<td>Orange colour</td>
<td>36.7***</td>
<td>14.2***</td>
<td>15.1***</td>
<td>6a 3b 2b 7c 4c 1c 5c</td>
<td>Aa Bb Cc</td>
</tr>
<tr>
<td>Homogeneity of colour</td>
<td>0.02NS</td>
<td>1.3NS</td>
<td>1.2NS</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Orange-edged slice</td>
<td>22.9***</td>
<td>86.1***</td>
<td>18.5***</td>
<td>3a 1b 7b 2c 5c 6d 4d</td>
<td>Aa Bb Cb</td>
</tr>
<tr>
<td>Translucent aspect</td>
<td>4.3*</td>
<td>15.1***</td>
<td>4.0***</td>
<td>6a 2b 7bc 4bcd 5cd 3cd 1d</td>
<td>Aa Ca Bb</td>
</tr>
<tr>
<td>Fatty aspect</td>
<td>24.7****</td>
<td>21.3***</td>
<td>6.4***</td>
<td>7a 6a 5a 3b 1b 4b 2b</td>
<td>Aa Bb Cc</td>
</tr>
<tr>
<td>White stripes</td>
<td>9.8***</td>
<td>15.2***</td>
<td>2.8*</td>
<td>1a 2a 3a 4a 6b 5bc 7c</td>
<td>Ba Cb Bb</td>
</tr>
<tr>
<td>Flavour: smoke intensity</td>
<td>4.0*</td>
<td>35.3***</td>
<td>2.3*</td>
<td>4a 3b 1b 2b 7c 5cd 6d</td>
<td>Aa Bab Cb</td>
</tr>
<tr>
<td>Flavour: wood fire</td>
<td>0.1NS</td>
<td>63.0***</td>
<td>5.5***</td>
<td>4a 3b 1b 6b 7bc 2bc 5c</td>
<td>–</td>
</tr>
<tr>
<td>Flavour: salmon/fish</td>
<td>1.9NS</td>
<td>8.6***</td>
<td>1.4NS</td>
<td>6a 7ab 5bc 3cd 2de 1de 4e</td>
<td>–</td>
</tr>
<tr>
<td>Firmness</td>
<td>41.8****</td>
<td>29.5***</td>
<td>14.9***</td>
<td>6a 4b 2c 3cd 1de 7e 5c</td>
<td>Aa Cb Bc</td>
</tr>
<tr>
<td>Melting texture</td>
<td>11.5***</td>
<td>95.1***</td>
<td>13.8***</td>
<td>6a 7b 5b 1c 3c 2d 4e</td>
<td>Aa Ba Cb</td>
</tr>
<tr>
<td>Fatty texture</td>
<td>16.1***</td>
<td>88.2***</td>
<td>8.1***</td>
<td>6a 7a 5b 1c 3cd 2d 4e</td>
<td>Aa Bb Cb</td>
</tr>
<tr>
<td>Pasty texture</td>
<td>8.6***</td>
<td>14.4***</td>
<td>3.0*</td>
<td>7a 5a 3a 1a 2b 4bc 6c</td>
<td>Ca Ba Ab</td>
</tr>
</tbody>
</table>

a Value of fisher test and significant level; NS: no significant difference; * significant difference at a level $P < 0.05$; ** significant difference at a level $P < 0.01$; *** significant difference at a level $P < 0.001$.

b 1: dry salting and smoking at 20°C; 2: frozen raw material, dry salting and smoking at 20°C; 3: brine salting and smoking at 20°C; 4: dry salting and electrostatic smoking; 5: dry salting at 30°C; 6: frozen raw material, dry salting and smoking at 30°C; 7: brine salting and smoking at 30°C.

c Raw material A: ocean ranched salmon (July 1998); B: farmed salmon from Norway (October 1998); C: farmed salmon from Norway (April 1999). Different letters (a,b,c) indicate significant differences ($P < 0.05$) between groups A, B and C.

Component 1 (46.3%)
Component 2 (20.2%)

Fig. 8. Projection of variables and samples in the 1-2 plane of principal component analysis on sensory descriptors. A, ocean ranched salmon (July 1998); B, farmed salmon from northern Norway (October 1998); C, farmed salmon from western Norway (April 1999) smoking procedures: 1, dry salting and smoking at 20°C; 2, frozen raw material, dry salting and smoking at 20°C; 3, brine salting and smoking at 20°C; 4, dry salting and electrostatic smoking; 5, dry salting at 30°C; 6, frozen raw material, dry salting and smoking at 30°C; 7, brine salting and smoking at 30°C.  

odour: smoke intensity (osi), wood fire (owf), acrid smell of smoke (oacr), toasted bread (obrea), fresh salmon (osalm); appearance: pink colour (pink), orange colour (oran), homogeneity of colour (homo); darkness of the slice edge (edge), translucent appearance (atran), fatty aspect (afat), white stripes (str); flavour: smoke intensity (fsmi), wood fire (fwf), salty taste (salt), fresh salmon (fsalm); and texture: firmness (firm), melting texture (melt), fatty texture (tfat), pasty texture (tpast). 

M. Cardinal et al./Food Research International 34 (2001) 537–550
products. The only modification observed was a more intense orange colour of slices when the dry-salting technique was used. However, the differences were generally very slight and not detectable by instrumental analysis performed on the fillet surface.

Fig. 8 shows the main characteristics of each product in terms of principal component analysis performed on mean panel scores. The main interactions between the process and raw material were observed with toasted bread odour, for which differences between products were noted mainly for fish from group C. An increase in panel discriminative power for this particular score could account for these results. The process effect on colour (pink and orange colour, dark-edged slice) and translucence was more intense when salmon were leaner (groups A and C). For these fish, slices of previously frozen samples had a more intense pink colour. A storage temperature effect on colour loss and pigment concentration has already been noted by Andersen and Steinsholt (1992) and Sheehan et al. (1998). The electrostatic process gave samples with the most intense dark-orange colour (pink and orange colour, dark-edged slice) and translucence was more intense when salmon were leaner (groups A and C). For these fish, slices of previously frozen samples had a more intense pink colour. A storage temperature effect on colour loss and pigment concentration has already been noted by Andersen and Steinsholt (1992) and Sheehan et al. (1998). The electrostatic process gave samples with the most intense dark-edged slices, followed by samples smoked at 30°C and/or frozen before processing. Similar results were obtained with instrumental measurements in which the L* value of the electrostatically smoked product was low compared to that of other samples (Fig. 7). It is noteworthy that this criterion is considered to be of great interest with respect to consumer demand (Gormley, 1992).

Another process–raw material interaction was observed for the white stripe criterion. With a 30°C smoking temperature, stripes were more visible to the extent that the processed fish had a lower fat content. It would be of interest to determine whether processing parameters have an effect on lipid distribution and/or the connective tissue linking muscle fibres (Bremner, 1992).

In summary, these results show that smoking conditions and preliminary treatment of fish (such as freezing) are major factors determining the sensory characteristics of smoked salmon, even though effects are reduced when fat content is increased in flesh.

4. Conclusion

This study confirms the results of previous work concerning the relation between fat content and yields or the final characteristics of smoked salmon. It also dealt with the nature of the interaction between the process used and the initial raw material characteristics. It would appear that losses after processing increase when fish have reduced lipid content (less than 10%) as compared to fatter fish, especially in the case of frozen fish. A fat fish is less sensitive to a freezing step, and the yield after processing is increased, regardless of the kind of smoking procedure used. Nevertheless, for this type of raw material, smoking temperature needs to be reduced to prevent too high a desiccation of the surface (which generally produces intense surface colouring but little diffusion of smoke compounds into the flesh).

For the three raw materials analysed, the processes studied showed the same kind of effect on smoked salmon properties, but with greater differences between treatments in the case of small lean fish.

A freezing step should be avoided in order to limit losses during the processing of fish with little fat (brine salting and a smoking temperature of 30°C are recommended). However, fatter fish require a lower smoking temperature to prevent possible case-hardening or a colour problem. Analysis of the effect of smoking temperature or the kind of smoke deposit showed a gradient in fat texture perception according to the process used, regardless of the kind of raw material. Samples subjected to electrostatic smoking showed more fat texture, less in those smoked at 20°C and the least in those smoked at 30°C. Smoking conditions would appear to have a greater impact than muscle fat content on texture evaluation. A specialised study is needed to determine the effect of temperature or the electrostatic field on lipid structure.

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References


