

1 **Essential oils for antimicrobial and antioxidant applications in fish and**
2 **other seafood products**

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21 **Abstract**

22 *Background:* Fish and other seafoods are highly perishable food products due basically to
23 microbiological growth and lipid oxidation, which are known to be the principal causes of
24 quality deterioration of such products. Therefore, offering safe and high quality seafoods
25 combined with consumers' desire for natural products free from chemical preservatives
26 creates real challenging problems. In the recent past, there has been extensive focus on
27 antioxidant and antimicrobial effects of natural preservatives such as essential oils (EOs), as
28 effective alternative to synthetic additives, in order to enhance oxidative and microbial
29 stability of foods and extend their shelf life.

30 *Scope and approach:* In this review, the main spoilage mechanisms of fish and seafood
31 products and the most common techniques used to preserve quality and extend shelf life of
32 such products are first discussed. The chemistry and modes of action of some selected EOs
33 are then briefly presented. The antioxidative and antimicrobial activities of some common
34 EOs, either alone or in combination with other preservative systems, in fish and other
35 seafoods are reviewed. Finally the limitations and the future trends are shown.

36 *Key findings and conclusions:* Several EOs have shown i) great antimicrobial activities versus
37 many spoilage and pathogenic microorganisms, and ii) remarkable antioxidant powers against
38 lipid oxidation in fish and other seafoods during processing or storage. However, much more
39 works are still required in order to better understand the exact mechanism of action of EOs or
40 their main components, the effective dose, and the best combination strategy.

41 *Keywords:* Fish, Preservation, Oxidation, Quality, Shelf life, Microbial spoilage, Natural
42 additives

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44 **1. Introduction**

45 In recent years, food quality and safety have become a major concern to consumers,
46 producers, food industries, and regulatory agencies worldwide. Such recent trends may be due
47 to the globalization of the food trade and changes in eating habits and consumer behavior,
48 such as increasing demand for natural, fresh, minimally processed, easily prepared, and ready-
49 to-eat products (Jayasena & Jo, 2013; Lucera, Costa, Conte, & Del Nobile, 2012). Therefore,
50 production of safe and high quality food products in general and fish and other seafoods in
51 particular has gained more and more attention around the world in the recent past. Due to their
52 high nutritional value, fish and other seafoods are considered among the most important
53 commodity for human diet, and hence their consumption has risen substantially over the past
54 few decades (Ghanbari, Jami, Domig, & Kneifel, 2013; Sampels, 2015a). Indeed, according to
55 the Food and Agriculture Organization of the United Nations, fish consumption increased
56 from an average of 9.9 kg in the 1960s to around 20 kg in 2015 (FAO, 2016).

57 Fish and other seafoods are extremely perishable food products and are especially
58 susceptible to both chemical and microbiological spoilage during processing or storage. For
59 this reason, one or more adequate preservation methods are required in order to maintain the
60 safety and quality and extend the shelf life of such products (Ghanbari et al., 2013; Hassoun
61 & Karoui, 2017; Nosedá, Vermeulen, Ragaert, & Devlieghere, 2014). Various traditional
62 processing methods including drying, salting, smoking, marinating, fermentation and so on,
63 have been widely used since ancient times to preserve fish quality or add more value to the
64 product (Sampels, 2015a). Moreover, low temperature storage and chemical preservatives
65 used for controlling water activity, enzymatic, oxidative, and microbial spoilage are
66 extensively used in food industry (Ghaly, Dave, Budge, & Brooks, 2010). However, due to
67 the growing concerns regarding the safety of chemical and synthetic preservatives, alternative
68 mechanisms based on the use of natural compounds have been increasingly tested over the

69 last years (Amorati, Foti, & Valgimigli, 2013; Calo, Crandall, O'Bryan, & Ricke, 2015;
70 Hyldgaard, Mygind, & Meyer, 2012; Lucera et al., 2012). In this context, essential oils (EOs)
71 could represent a promising option since numerous reports have confirmed their antioxidant
72 (Amorati et al., 2013; Jayasena & Jo, 2014) and antimicrobial (Burt, 2004; Jayasena & Jo,
73 2013; Swamy, Akhtar, & Sinniah, 2016) effects. Thus, these natural preservatives could meet
74 perfectly the increasing consumer demands for clean-label products that are fresh and free of
75 chemical additives.

76 Although there have been several prior reviews on the use of EOs in food applications
77 (Calo et al., 2015; De Souza, da Cruz Almeida, & de Sousa Guedes, 2016; Jayasena & Jo,
78 2013, 2014), the antimicrobial and antioxidant properties of EOs for application in fish and
79 other seafoods have not yet been reviewed. Therefore, this review provides up-to-date
80 information about the most recent published data regarding antimicrobial and antioxidant
81 mechanisms of common EOs or their main components as well as their potential applications
82 in fish and other seafood products.

83 **2. Fish spoilage mechanisms**

84 Although fish flesh is generally regarded as sterile when fish is alive, fish spoilage can
85 occur very rapidly after catch or harvest and during the different stages of the production
86 chain, processing, and subsequent storage conditions. Although the importance of the
87 enzymatic autolysis, occurring mainly after capture or harvest, the following section will
88 focus only on microbial and chemical (oxidation) spoilage occurring during processing and
89 storage of fish.

90 *2.1. Microbial spoilage*

91 Fish and other seafoods have high contents of free amino acids, a high *post mortem* pH,
92 high water contents, and many fish species contain trimethylamine oxide (TMAO) (Chaillou

93 et al., 2015; Gram & Dalgaard, 2002). Such characteristics promote growth of bacteria,
94 including both the Gram-positive and Gram-negative types which survive well in a wide
95 range of temperatures. That is why the microbial growth is considered to be the major cause
96 of quality deterioration of fish and other seafood products, causing up to 25-30% loss of such
97 products (Ghaly et al., 2010; Gram & Dalgaard, 2002). There is a general agreement that each
98 food product has its own unique flora, which is determined by the raw materials, the
99 processing parameters and subsequent storage conditions, and the abilities of microorganisms
100 to tolerate the preservation conditions. For example, it was reported that psychrotolerant
101 Gram-negative bacteria such as species within the genera *Pseudomonas* and *Shewanella* are
102 the most commonly spoilage bacteria of aerobically stored chilled fish, while CO₂-tolerant
103 microorganisms, including *Photobacterium phosphoreum* and lactic acid bacteria, may
104 dominate the microflora and become responsible for spoilage of packed fish products
105 (Chaillou et al., 2015; Giuffrida, Valenti, Giarratana, Ziino, & Panebianco, 2013; Gram &
106 Dalgaard, 2002; Gram & Huss, 2000).

107 Although freshly caught fish is contaminated naturally with various microbiota, only a
108 small fraction of these microorganisms, called specific spoilage organisms (SSOs), are
109 responsible for seafood spoilage (Gram & Dalgaard, 2002). In particular, the seafood SSOs
110 have the ability to convert TMAO to TMA-N, produce ammonia, biogenic amines, organic
111 acids and sulphur compounds from amino acids, hypoxanthine from ATP degradation
112 products, and acetate from lactate. Microorganisms capable of converting TMAO to TMA
113 include *Aeromonas spp.*, *Enterobacteriaceae*, *Photo-bacterium phosphoreum*, *Shewanella*
114 *putrefaciens*, and *Vibrio spp.* (Gram & Dalgaard, 2002). Research studies demonstrated that
115 *Pseudomonas* was the dominant bacteria for Atlantic salmon (*Salmo salar*) packed in a
116 modified atmosphere (Milne & Powell, 2014) and for bighead carp (*Aristichthys nobilis*)

117 fillets sprinkled with 2% salt, whereas *Aeromonas* was the SSOs of unsalted fillets during
118 storage at 4 °C (Liu, Zhang, Li, & Luo, 2017).

119 Several microbial growth parameters such as total viable counts (TVC), mesophilic aerobic
120 counts (MAC), and aerobic plate count (APC) have been used to gives a quantitative idea
121 about the presence of microorganisms in the investigated sample (Cheng & Sun, 2015;
122 Rodrigues et al., 2016). For example, when the TVC of bacteria exceeds a microbial load of
123 10^7 colony-forming units (CFU) per gram or cm^2 , it means that the fish muscle becomes
124 dangerous for consumption and can cause very severe health problems due to the possibility
125 of toxic substances produced (Ellis, 2002). Additionally, counts of SSOs obtained on Lyngby
126 Iron Agar plates (Oxoid LTD., Basingstoke, Hampshire, England) after 3 days incubation at
127 20 °C, have been used as microbial growth parameters for number of Gram-negative and non-
128 fermentative bacteria (Gram., Trolle, & Huss, 1987). Moreover, various other parameters
129 have been widely measured to reveal microbiological quality of fish, such as the nucleotide
130 degradation, the formation of biogenic amines, the production of total volatile basic nitrogen
131 (TVB-N), trimethylamine nitrogen (TMA-N), among others (Rodrigues et al., 2016; Zhu, Ma,
132 Yang, Xiao, & Xiong, 2016).

133 2.2. *Oxidative spoilage*

134 Spoilage caused by oxidation is another prevalent problem, especially for fish species
135 containing high amounts of polyunsaturated fatty acids, resulting in several problems such as
136 off-flavor formation, changes in colour and texture, and altered nutrient value (Maqsood,
137 Benjakul, Abushelaibi, & Alam, 2014; Secci & Parisi, 2016). Although lipid oxidation could
138 undergo several types of oxidation, such as photo-oxidation, thermal oxidation, enzymatic
139 oxidation, and auto-oxidation; this latter, defined as the spontaneous reaction of atmospheric
140 oxygen with lipids, is the most common process causing oxidative deterioration (Shahidi &

141 Zhong, 2005). This process occurs via a free radical chain reaction, and proceeds through
142 three phases: initiation, propagation, and termination. Initiation phase starts with the
143 abstraction of a hydrogen atom adjacent to a double bond in a fatty acid, and this may be
144 catalyzed by light, heat, or metal ions to form a free radical. The resultant free radicals react
145 with oxygen to form peroxy radicals, which in turn react with other lipid molecules to form
146 hydroperoxides and a new free radical during the propagation phase. Termination phase
147 occurs when a build up of these free radicals interact to form non-radical products. Lipid
148 hydroperoxides have been identified as primary products of autoxidation; being unstable,
149 decomposition of hydroperoxides results in a complex mixture of products including
150 aldehydes, ketones, alcohols, hydrocarbons, volatile organic acids, and epoxy compounds,
151 which are known as secondary oxidation products (Ghaly et al., 2010; Shahidi &
152 Wanasundara, 2002; Xu, Riccioli, & Sun, 2015).

153 **3. Fish preservation methods**

154 Several traditional preservation techniques can be applied in order to retard deterioration of
155 seafood products and extend their shelf life as much as possible. Preservation techniques are
156 usually based on the control of temperature, available oxygen, water activity, microbial loads,
157 or several of these parameters at the same time.

158 *3.1. Temperature-based techniques*

159 It is well-known that temperature has a marked effect on the microbial growth and
160 oxidation process occurring during *post mortem* storage or processing and handling of fish
161 and other seafood products. Indeed, on one hand, temperatures have a direct physical impact
162 on microbial growth and may lead to retardation of the growth and spoilage activity of
163 microorganisms. On the other hand, according to the Arrhenius relation, the rates of

164 undesirable biochemical and chemical reactions decrease as temperature is lowered (Hall,
165 2010; Jessen, Nielsen, & Larsen, 2014).

166 The cooling (or chilling) of fish with normal ice flakes, chilled seawater, or ice slurries has
167 been considered as simple and efficient preservation method, keeping the fish in a cool
168 condition with a temperature ranging between 0 and 4 °C. However, it is important to ice the
169 fish as quickly as possible after catch or harvest in order to minimize biochemical and
170 microbiological reactions (Ghaly et al., 2010; Sampels, 2015b). Although the importance of
171 the chilling in keeping fish freshness, it must be emphasized that this technique cannot
172 prevent enzyme activities or microbial spoilage (Sampels, 2015b). So, chilling process should
173 be completed with other preservation method. Another low temperature-based technique is
174 superchilling. This term has been used to describe the decrease in temperature of a food
175 product to 1–2 °C below the freezing point, so that only a minor part of the product's water
176 content is frozen (Kaale, Eikevik, Rustad, & Kolsaker, 2011; Stonehouse & Evans, 2015). In
177 fish sector, the superchilling has been applied successfully and shown to extend shelf life of
178 many seafood products as a result of inhibition of most autolytic and microbial reactions in
179 fish compared with normal cooling (Duun & Rustad, 2008; Kaale et al., 2011; Sampels,
180 2015b).

181 Freezing has been considered the most popular method of conservation and successfully
182 employed to retain the quality of food products, especially fish and other seafoods, over long
183 storage periods (Hall, 2010; Jessen et al., 2014). Although freezing (-18 to -30 °C) inhibits the
184 rate of chemical reactions and microbial growth, enzymatic and non enzymatic reactions
185 persist but at lower rate. An important consideration to be in mind when using freezing
186 technique is the formation of ice crystals during the process, being a critical point, since the
187 formation of large ice crystals may increase the risk of texture damage, loss of water holding
188 capacity, and oxidation (Alizadeh, Chapleau, de Lamballerie, & Le-Bail, 2007; Ghaly et al.,

189 2010; Karoui, Hassoun, & Ethuin, 2017). That is why a fast freezing should be conducted in
190 order to provide small and regular ice crystal formation.

191 3.2. *Modified atmosphere packaging*

192 Modified atmosphere packaging (MAP) has received increasing attention, becoming a
193 popular preservation technique in a wide range of application in food products to meet
194 consumer demands for fresh and natural foods with an extended shelf life (Mastromatteo,
195 Conte, & Del Nobile, 2010a; Santos et al., 2013). This technique is based on the modification
196 of percentage of the three principal gases (i.e., % CO₂, %O₂, and %N₂) inside the package
197 containing food product to provide an optimal condition for effective retardation of
198 microbiological and chemical processes. Generally speaking, the modification of the
199 atmosphere within the package can be achieved by reducing the oxygen content while
200 increasing the levels of carbon dioxide and/or nitrogen (Mastromatteo et al., 2010a; Nosedo et
201 al., 2014). The effect of MAP on the shelf life of foods in general and fish in particular has
202 been reviewed by several authors (Bouletis, Arvanitoyannis, & Hadjichristodoulou, 2017;
203 Sivertsvik, Jeksrud, & Rosnes, 2002). By using different CO₂ and N₂ levels, Provincial and
204 co-workers obtained the best results in term of shelf life of sea bass (*Dicentrarchus labrax*)
205 for MAP samples stored with high CO₂ levels (Provincial et al., 2010). These results were
206 then confirmed by other research study which was conducted on turbot (*Psetta maxima*)
207 fillets, indicating the protective effect of the different MAP studied, especially those with a
208 higher percentage of CO₂ (Santos et al., 2013). Recently, our result obtained on whiting
209 (*Merlangius merlangus*) fillets allowed recommending the use of MAP with 50% CO₂ and
210 50% N₂ to maintain quality and extend the shelf life of fish samples (Hassoun & Karoui,
211 2016).

212 3.3. *High pressure processing*

213 High pressure processing (HPP) has attracted widespread attention in recent years due to
214 its potential of inactivating microorganisms and autolytic enzymes at low temperature, thus
215 extending the shelf life of fish products (Rastogi, Raghavarao, Balasubramaniam, Niranjana, &
216 Knorr, 2007; Truong, Buckow, Stathopoulos, & Nguyen, 2014). As it is performed at room
217 temperature, this technique holds the characteristics of low energy consumption, making it an
218 environmentally friendly processing technology compared with traditional thermal processing
219 methods (Huang, Wu, Lu, Shyu, & Wang, 2017; Rastogi et al., 2007; Truong et al., 2014).

220 The HPP has shown to be effective in inhibiting microbial growth and maintaining the quality
221 in raw octopus (*Octopus vulgaris*) (Hsu, Huang, & Wang, 2014), reducing microbial loads in
222 shrimp (*Penaeus monodon*) (Kaur, Srinivasa Rao, & Nema, 2016), and extending the shelf
223 life of fresh salmon (*Salmo salar*), cod (*Gadus morhua*), and mackerel (*Scorpaenidae*)
224 fillets (Rode & Hovda, 2016). However, this technique may cause some undesirable effects
225 on flesh color and texture, on lipid and protein oxidation, as well as on protein denaturation in
226 the fish (Guyon, Meynier, & de Lamballerie, 2016; Truong et al., 2014)

227 3.4. Chemical preservatives and natural alternative solutions

228 Several chemical preservatives have been used to control microbial, oxidative, and
229 autolytic enzymatic spoilage of fish and fish products (Ghaly et al., 2010). For example, the
230 effects of salts of organic acids, such as sodium acetate, sodium lactate, and sodium citrate, on
231 the quality and shelf life of sliced salmon (*Onchorhynchus nerka*) were investigated during
232 refrigerated storage (Sallam, 2007). The author reported that the use of these preservatives
233 extended the shelf life of the fish by 5 - 8 days compared with control samples. Additionally,
234 synthetic phenolic compounds such as butylated hydroxyanisole (BHA), butylated
235 hydroxytoluene (BHT), and dodecyl gallate (DG) have been widely used as antioxidants and
236 antimicrobial agents for fish and other seafoods (Brewer, 2011).

237 However, the increasing consumers' concern regarding the safety of such compounds has
238 encouraged food industry to develop new natural alternative food preservation strategies
239 (Amorati et al., 2013; Brewer, 2011; Lucera et al., 2012). Among alternative preservation
240 methods, the use of lactic acid bacteria and their metabolites as biopreservation techniques to
241 extend the shelf life and enhance the hygienic quality of fish and other seafood, has received
242 much attention by the scientific community in the last two decades (Ghanbari et al., 2013).
243 Moreover, the use of natural compounds, such as tea polyphenols, rosemary, and sage extracts
244 has become very popular for food preservation (Emir Çoban & Özpolat, 2013; Kenar, Özogul,
245 & Kuley, 2010; Li et al., 2012; Pezeshk, Ojagh, & Alishahi, 2015). Additionally, application
246 of chitosan has widespread in the last years in several applications in the seafood industry,
247 due to its useful biological activities, including among other the antibacterial and antioxidant
248 characteristics (Alishahi & Aider, 2012; Yuan, Chen, & Li, 2016).

249 *3.5. Hurdle technology*

250 The combination of two or more preservation methods, referred as "hurdle technology"
251 may lead to synergistic or additive interactions, offering a greater inhibitory effect against the
252 targeted microorganisms than any single treatment (De Souza et al., 2016; Khan, Tango,
253 Miskeen, Lee, & Oh, 2017). Examples for the application of combined preservation methods
254 are given by Duun and Rustad (2008) for Atlantic salmon (*Salmo salar*) and Fernández et al.
255 (2009) for the same fish species, as well as Zhu et al. (2016) for catfish (*Clarias gariepinus*).
256 In details, the use of vacuum packaging combined with superchilling storage at two
257 temperature levels (-1.4 or -3.6 °C) was evaluated in salmon fillets by using several quality
258 parameters (Duun & Rustad, 2008). The findings revealed that the storage time of vacuum
259 packed samples can be doubled by superchilled storage, maintaining good quality of fish up to
260 17–21 days compared to ice chilled storage. In another study, superchilling storage (-1.5 °C)
261 was combined with MAP at different gas concentrations, and the combined effects of these

262 technologies on salmon fillets were monitored by sensory, chemical, and microbiological
263 analysis (Fernández, Aspe, & Roeckel, 2009). The authors noticed an important increase of
264 shelf life from 11 days for control sample to 22 days in superchilled fish stored in the
265 presence of MAP at high CO₂ (90% CO₂: 10% N₂). These findings were confirmed in a recent
266 study (Zhu et al., 2016) conducted on catfish fillets stored at superchilling temperature (−0.7
267 °C) combined with MAP at high levels of CO₂ (60% CO₂: 40% N₂). Compared to the other
268 storage conditions, the authors reported that this combination maintained effectively the
269 quality of fish fillets and prolonged significantly their shelf life. Other combination method
270 was proposed by Rodrigues and others using MAP (80% CO₂: 20% N₂) and short-wave
271 ultraviolet radiation in order to extend shelf life of rainbow trout (*Oncorhynchus mykiss*)
272 fillets. The findings demonstrated that this combination was effective in reducing the total
273 microbial count and delaying the chemical changes and, consequently, enhancing the shelf
274 life of the fish fillets at least twice (Rodrigues et al., 2016). Recently, a research study was
275 conducted to determine the impact of combination of two treatments using chitosan and
276 pomegranate peel extract on the quality of Pacific white shrimp (*Litopenaeus vannamei*)
277 during 10 days of iced storage (Yuan, Lv, Tang, Zhang, & Sun, 2016). The authors observed a
278 synergistic effect between these treatments since the efficacy of chitosan coating to inhibit the
279 microbial growth, melanosis, changes in color and texture, and other sensory parameters was
280 increased when it was applied in combination with pomegranate peel extract.

281 **4. Essential oils**

282 Essential oils (EOs) are produced by different part of plants as defence mechanisms against
283 microorganisms. These naturally occurring antimicrobial and antioxidant agents are highly
284 complex mixtures of often hundreds of individual aromatic volatile oily compounds, which
285 are extracted from different plant materials, such as leaves, barks, stems, roots, flowers, and
286 fruits (Calo et al., 2015; Jayasena & Jo, 2013). In total, more than 3000 types of EOs are

287 known, of which only 300 are of commercial interest for applications in the food or other
288 industries (Bakkali, Averbeck, Averbeck, & Idaomar, 2008; Burt, 2004)

289 4.1. Main chemical components and principal sources

290 It has been well documented that the biological properties of EOs are primarily due to the
291 presence of major compounds, accounting up to 85% of the oil, while minor compounds,
292 present only in trace quantities, may have synergistic impact with other compounds (Bakkali
293 et al., 2008; Burt, 2004). Chemically, the EOs consist of a diverse family of organic
294 compounds with low molecular weight, which could be divided into several groups according
295 to their chemical structure: terpenes, terpenoids, aromatic (phenylpropanoids) and other
296 compounds (Bakkali et al., 2008; Hyldgaard et al., 2012). Terpenes are hydrocarbons
297 consisting of several isoprene units, which could be classified by the number of isoprene units
298 in the molecule (mono-, sesqui- and diterpenes). Terpenoids are terpenes containing oxygen,
299 and could be classified into alcohols, esters, aldehydes, ketones, ethers, and phenols.
300 Examples of well-known terpenoids found in EOs are thymol, carvacrol, linalool, linalyl
301 acetate, citronellal, piperitone, menthol, and geraniol, while eugenol and cinnamaldehyde are
302 the best known phenylpropanoids (Hyldgaard et al., 2012; Jayasena & Jo, 2013). It should be
303 stressed that phenolic compounds such as thymol, carvacrol, and eugenol are the main group
304 responsible for the preservative effects of EOs (Burt, 2004; Jayasena & Jo, 2014).

305 According to our literature review, EOs from oregano, rosemary, thyme, laurel, sage,
306 cinnamon, clove, and basil have been the most used antimicrobial and antioxidant agents in
307 fish and seafood products.

308 Oregano (*Origanum vulgare*) leaves are a characteristic spice of the Mediterranean cuisine
309 and have been widely used in raw or cooked food due to their distinct pleasant aroma and
310 taste. Besides, the oregano EO has been studied for its antimicrobial and antioxidant activity

311 in various commercial or model foods (Goulas & Kontominas, 2007; Vatavali, Karakosta,
312 Nathanailides, Georgantelis, & Kontominas, 2013). The carvacrol and thymol are reported to
313 be the main compounds responsible for the antimicrobial and antioxidant activity of oregano
314 EO (Rodriguez-Garcia et al., 2016). EO extracted from thyme (*Thymus vulgaris*) has received
315 much attention from researchers and food processors as a potential natural antimicrobial and
316 antioxidant agent as a result of its high content of phenolic compounds (Hyldgaard et al.,
317 2012; Kostaki, Giatrakou, Savvaidis, & Kontominas, 2009).

318 Due to its antimicrobial activity against a wide range of microorganisms, basil (*Ocimum*
319 *basilicum*) EO has been used extensively for many years in flavouring food (Suppakul, Miltz,
320 Sonneveld, Bigger, & Qd, 2003). This activity has been attributed to the major active volatile
321 components, including linalool, methylchavicol, eugenol, methyl eugenol, methyl cinnamate,
322 1,8-cineole, and caryophyllene (Kuorwel, Cran, Sonneveld, Miltz, & Bigger, 2011; Perricone,
323 Arace, Corbo, Sinigaglia, & Bevilacqua, 2015). Rosemary (*Rosmarinus officinalis*) EO has
324 been reported to exhibit an effective antioxidant and antimicrobial activity, which is mainly
325 related to phenolic diterpenes compounds such as carnosol and carnosic acid (Bozin, Mimica-
326 Dukic, Samojlik, & Jovin, 2007; Kenar et al., 2010; Makri, 2013).

327 Many recent studies have investigated the preservative effects of EOs obtained from other
328 sources such as clove (*Eugenia caryophyllata*) (Emir Çoban & Patir, 2013), sage (*Salvia*
329 *officinalis* L.) (Emir Çoban, Patir, Özpolat, & Kuzgun, 2016), *Zataria multiflora* Boiss (Emir
330 Çoban & Kelestemur, 2016), turmeric and lemongrass (Masniyom, Benjama, & Maneesri,
331 2012), and lemon (Alfonzo et al., 2017). The results of these studies demonstrated that the use
332 of these EOs applied to the fish or other seafoods alone or in combination with other
333 preservation methods, was effective in improving the quality and extending the shelf life of
334 the treated products.

335 4.2. Methods of application

336 EOs can be applied using various methods in the fish industry: the direct treatment of fish
337 and seafood products with EOs during manufacturing and processing is the most commonly
338 employed approach, followed by the use of EOs as edible films and coatings and the addition
339 of EOs to animal feed.

340 Although the direct addition of EOs (Emir Çoban & Patir, 2013; Karoui & Hassoun, 2017)
341 or their compounds (Giarratana et al., 2016; Mahmoud et al., 2004) to fish and other seafoods
342 has been the most common method of application, this technique has some disadvantages and
343 criticisms that limit its application to such products. Indeed, it has been generally observed
344 that a greater concentration of EOs is needed to achieve the same effect in food compared to
345 *in vitro* assays. Moreover, even at low doses some EOs could have a negative impact on the
346 sensory attributes (Lv, Liang, Yuan, & Li, 2011; Sánchez-González, Vargas, González-
347 Martínez, Chiralt, & Cháfer, 2011). Thus, some authors suggested the use of edible coating
348 films enriched with EOs as alternative and interesting option in order to reduce the required
349 doses (Doğan & İzci, 2017; Ojagh, Rezaei, Razavi, & Hosseini, 2010; Sánchez-González et
350 al., 2011; Yuan, Chen, et al., 2016). Additionally, some authors reported that fish sedated with
351 EOs during transport before slaughter could delay the loss of fish freshness and increase the
352 shelf life. In this regard, Daniel and co-workers demonstrated that silver catfish (*Rhamdia*
353 *quelen*) exposed to 40 µL/L of *Aloysia triphylla* (L'Her.) Britton EO during *in vivo* transport
354 delayed the nucleotide degradation and loss of quality compared to the control (Daniel et al.,
355 2014).

356 Recently, another technique has been presented in the literature to minimize the
357 organoleptic effects of EOs using the preparation of micro- and nanoemulsions, which
358 improves not only the antimicrobial and antioxidant stability, but also the functional

359 properties and organoleptic quality of the product (Acevedo-Fani, Soliva-Fortuny, & Martín-
360 Belloso, 2016; Alfonzo et al., 2017; Calo et al., 2015; Ozogul et al., 2017; Perricone et al.,
361 2015). Indeed, the encapsulation of EOs into such emulsions may increase the stability of
362 volatile components, protecting them from interacting with the food matrix, thereby
363 increasing the antimicrobial activity due to increased passive cellular uptake (Sugumar,
364 Ghosh, Mukherjee, & Chandrasekaran, 2016). More recently, numerous reviews have just
365 been published reporting an emerging application of EOs in yet more sophisticated approach
366 as active food packaging, which could extend food shelf life and maintain nutritional and
367 sensory quality (Atarés & Chiralt, 2016; Kapetanakou & Skandamis, 2016; Maisanaba et al.,
368 2017; Ribeiro-Santos, Andrade, Melo, & Sanches- Silva, 2017). Active food packaging
369 includes the incorporation of EOs, among other natural compounds, into the food package in
370 such a way that allows these compounds to be released in a controlled way to maintain or
371 enhance the organoleptic properties and microbiological integrity of food (Atarés & Chiralt,
372 2016; Ribeiro-Santos et al., 2017).

373 The use of EOs as fish dietary additives is considered to be an effective method to
374 incorporate natural antioxidant and antioxidant agents into flesh of fish products. For
375 example, one study examined the capacity of rosemary, thymol, carvacrol, and BHT
376 incorporated in the diet of gilthead seabream (*Sparus aurata*) in order to delay lipid oxidation
377 and microbial growth (Alvarez, Garcia Garcia, Jordan, Martinez-Conesa, & Hernandez,
378 2012). Compared to the control group, the results revealed that fillets from fish fed diet with
379 carvacrol (500 mg/kg) during 18 weeks had the lowest thiobarbituric acid (TBA) content (0.2
380 mg MDA/kg fillet), while BHT and thymol groups achieved the lowest bacteria counts. These
381 results were confirmed later in another study, where the addition of thyme EO as a feed
382 supplement at different concentrations (500, 1000, 1500 and 2000 mg kg⁻¹) revealed
383 inhibitory effects on microbial growth and lipid oxidation in gilthead seabream fillets during

384 storage at 4 °C for 21 days. Interestingly, the authors reported that high doses of thyme EO
385 resulted in both lower microbiological counts of *Enterobacteriaceae* and coliforms, and
386 higher oxidative stabilities measures as TBA (Hernández, García García, Jordán, &
387 Hernández, 2015).

388 4.3. Mechanisms of action

389 4.3.1. Antimicrobial activities

390 The antimicrobial properties of EOs have been known since antiquity. Most studies
391 investigating the use of EOs as an antimicrobial agent have been performed on bacteria, while
392 less is known about their action on yeast and molds (Hylgaard et al., 2012). EOs can be
393 applied either to inhibit the bacterial growth (bacteriostatic), which means that the microbial
394 cells will recover their reproductive capacity after neutralization of the agent, or to kill
395 bacterial cells (bactericide), if EOs are used at high concentrations (Swamy et al., 2016). It
396 was reported that lipoteichoic acids in cell membrane of gram positive bacteria may facilitate
397 the penetration of hydrophobic compounds of EOs, while the presence of an extrinsic
398 membrane, surrounding the cell wall of gram negative bacteria limits the diffusion rate of
399 hydrophobic compounds through the lipopolysaccharide layer. That is why gram positive
400 bacteria are slightly more susceptible to EOs than gram negative ones (Rodriguez-Garcia et
401 al., 2016; Tongnuanchan & Benjakul, 2014).

402 Even though that the possible modes of action for EOs as antimicrobial agents have been
403 widely reviewed, their exact mechanism of action is not yet clear (Calo et al., 2015; Maqsood,
404 Benjakul, & Shahidi, 2013; Tajkarimi, Ibrahim, & Cliver, 2010). Several studies have
405 reported that the antimicrobial activity of EOs can be attributed to their major constituents
406 mainly the phenolic constituents, as well as their interaction with minor constituents present
407 in oils (Burt, 2004; Hylgaard et al., 2012; Jayasena & Jo, 2013; Perricone et al., 2015). Due

408 to the complexity of the chemical composition of EOs, it was reported that the antimicrobial
409 activity of EOs may not be attributable to a unique mechanism (Burt, 2004). Nonetheless,
410 there is almost a universal agreement on the fact that the hydrophobicity of compounds
411 present in EOs enables them to pass through the cell wall and cytoplasmic membrane, disrupt
412 the structure of their different layers of polysaccharides, fatty acids and phospholipids and
413 permeabilize them. Additionally, EOs can inhibit several enzyme systems including the
414 enzymes responsible for regulation of energy and synthesis of structural components (Bakkali
415 et al., 2008; Burt, 2004; Jayasena & Jo, 2013).

416 *4.3.2. Antioxidant activities*

417 Recently, synthetic antioxidants, such as BHA and BHT have been suspected of causing
418 potentially harmful consequences on human health. On the other side, the use of EOs has been
419 considered as a good alternative since the majority of EOs are classified as generally
420 recognized as safe (GRAS) (Kapetanakou & Skandamis, 2016; Maqsood et al., 2013; Ribeiro-
421 Santos et al., 2017). The application of EOs as natural antioxidants is a field of growing
422 interest due to the inherent ability of some of their components to stop or delay the oxidation
423 of lipids and extend the shelf life of the food products (Amorati et al., 2013; Patel, 2015).
424 Numerous studies reported that the EOs, as antioxidants, have several modes of direct or
425 indirect actions including, among other mechanisms, prevention of chain initiation and free-
426 radical scavenging activity (Maqsood et al., 2013; Rodriguez-Garcia et al., 2016). Again, it
427 has been reported that phenolic compounds such as carvacrol, eugenol, and thymol are the
428 main group responsible for the antioxidant activity of EOs (Amorati et al., 2013; Jayasena &
429 Jo, 2014). The role of phenolic compounds in the retardation of lipid oxidation in fish muscle
430 is mainly due to their redox properties, allowing them to act as hydrogen donors, reducing
431 agents, singlet oxygen quenchers as well as metal chelators (Maqsood et al., 2014;
432 Tongnuanchan & Benjakul, 2014).

433 Several methods have been used to assess the antioxidant performance of EOs. Although the
434 peroxide value (PV) and TBA are the most commonly used methods for measuring
435 respectively the primary and secondary products of oxidation, other methods, such as the
436 DPPH (2,2-diphenyl-1-picrylhydrazil) radical scavenging method, the absorption capacity of
437 oxygen radicals, and the total phenolic compounds could be used (Amorati et al., 2013; Bozin
438 et al., 2007; Maqsood et al., 2013).

439 **5. Application of EOs to fish preservation**

440 In recent years, the effectiveness of a wide range of EOs against lipid oxidation and
441 microbial growth has been extensively demonstrated by many authors. It has been reported
442 that oregano EO is the most frequently used for applications as fish preservatives, followed by
443 rosemary and thyme EOs (Patel, 2015). Different effects have been observed depending on
444 the EO used, its concentration, as well as the characteristics of the raw material. An overview
445 of the literature reporting studies on the antioxidant and antimicrobial activity of some EOs in
446 fish and fish products are presented in **Table 1**.

447 A typical example of lipid oxidation inhibition induced by addition of EOs is presented in
448 **Figure 1**, where the effect of *Zataria multiflora* Boiss EO on quality of catfish (*Silurus glanis*
449 Linnaeus, 1758) burgers stored at 4 °C was studied (Emir Çoban & Kelestemur 2016).
450 Among other results, the authors showed that, at both the concentrations tested (0.2% and
451 0.4%), the PV (**Figure 1A**) and the TBA (**Figure 1B**) were significantly ($P < 0.05$) reduced
452 by the addition of this EO compared with untreated samples, which was attributed to the
453 presence of phenolic compounds such as carnosol, carnosic acid, and rosmarinic acid.
454 However, treatment of the catfish burgers at the concentration of 0.4% *Zataria multiflora*
455 Boiss EO exhibited a greater inhibitory impact on lipid oxidation and microbial growth
456 compared with that obtained for the samples treated with 0.2%. This dose-dependent

457 inhibitory activities of EOs confirmed our previous results, where the higher concentration of
458 clove EO was found to be more effective to inhibit microbial growth and lipid oxidation
459 occurring in sliced smoked *Oncorhynchus mykiss* (Emir Çoban & Patir, 2013).

460 However, it should be considered that EOs used as natural food additive at high
461 concentrations may lead to undesirable sensory properties on treated fish and may even cause
462 allergic reactions. Indeed, some EOs are characterized by a strong odor and flavor which
463 could leave a bad aftertaste, thus minimizing the acceptance or liking degree for fish and
464 seafood product (Atarés & Chiralt, 2016; Ribeiro-Santos et al., 2017). That is why the
465 antimicrobial effectiveness of EOs is often described using the concept of "minimum
466 inhibitory concentration" which is the lowest concentration capable of inhibiting the growth
467 of challenging organisms (Burt, 2004; Hyldgaard et al., 2012; Mann & Markham, 1998).

468 One method that has been proposed in the literature in order to reduce organoleptic effects of
469 EOs added to fish and other seafoods is to use coatings enriched with EOs (Lucera et al.,
470 2012; Sánchez-González et al., 2011). For instance, a gelatin coating enriched with cinnamon
471 (*Cinnamomum zeylanicum*) EO at different concentrations (1%, 1.5%, and 2%) was tested as
472 antioxidant and antimicrobial agent on refrigerated rainbow trout (Andevvari & Rezaei, 2011).
473 The findings showed that this treatment decreased the lipid oxidation rate, measured by means
474 of TBA and free fatty acids (FFA), and the microbial growth, determined by TVC, APC, and
475 psychrotrophic count. From the obtained results the authors concluded that the gelatin coating
476 enriched with cinnamon EO was suitable for the preservation of quality attributes of rainbow
477 trout fillets to an acceptable level during storage.

478 In more recent years, micro- and nanoemulsions have been suggested, instead of direct
479 addition of EOs to fish products, as interesting area of research in order to transport active
480 compounds of EOs to food and even enhance functional properties of treated products

481 (Acevedo-Fani et al., 2016). For instance, one recent study has investigated the effects of a
482 microemulsion containing 0.3% or 1% lemon EO on the quality of salted sardines during 150
483 days of ripening. The finding revealed a reduction in the concentrations of all examined
484 microbial groups, including *Enterobacteriaceae*, *Staphylococci* and rod *Lactic acid bacteria*.
485 Besides, the addition of this EO, in particular at concentration of 1%, showed a lower
486 accumulation of histamine in the treated sardines compared to those of the control. The
487 authors ascribed the preservative effect of lemon EO to several volatile organic compounds
488 belonging to monoterpene hydrocarbons, oxygenated monoterpenes, and sesquiterpene
489 hydrocarbons (Alfonzo et al., 2017).

490 The scientific literature seems to indicate that the impact of EOs or their compounds as
491 antimicrobial and antioxidant agents depend on the source of these natural food additives. In
492 more details, Karoui and Hassoun reported that basil and rosemary EOs used at the same
493 concentration (1%) resulted in different preservative activities since the former was found to
494 be more effective at retarding fish spoilage than the latter (Karoui & Hassoun, 2017), while in
495 another study, the rosemary EO was found to be more efficient in preventing lipid oxidation
496 than oregano EO (Makri, 2013). In another investigation, three EOs, including clove, cumin,
497 and spearmint, have been evaluated in vapour phase for their efficacy in preventing quality
498 degradation and prolonging shelf life of red drum (*Sciaenops ocellatus*) fillets during 20 days
499 of refrigerated storage at 4 °C (Cai et al., 2015). Among other results, the authors
500 demonstrated that the addition of these EOs at 4 µl/L reduced biogenic amine contents and
501 microflora counts of various microorganisms, thereby prolonging the shelf life of the fish by
502 10 days as compared to the control sample; however, more effective activity was obtained for
503 spearmint EO compared to the two other ones. The difference effectiveness of the various
504 EOs could be attributed to the difference in their chemical composition, especially with regard

505 to the major components, which in turn are related to different conditions such as climatic,
506 genetic, etc.

507 **6. Synergy between EOs and other preservation methods**

508 Due to synergistic effects, some authors demonstrated that combined treatments of EOs
509 and other preservative method could have better antimicrobial and/or antioxidant activities
510 than either treatment alone (**Table 2**). According to the literature, it appears that EOs could be
511 applied in combination with various preservation methods, such as vacuum packaging,
512 modified atmosphere packaging, chitosan, nisin, and other factors.

513 EOs have been demonstrated to be synergistic with vacuum packaging and modified
514 atmosphere packaging, as verified by the following findings. The combined effect of oregano
515 EO at two concentrations; 0.2%, 0.4% and vacuum packaging was evaluated on
516 Mediterranean octopus (*Octopus vulgaris*) stored under refrigeration for a period of 23 days.
517 The results revealed significant antimicrobial and antioxidant stabilities of the vacuum packed
518 samples treated with 0.4% oregano EO as compared to the control. From the obtained results,
519 the authors concluded that the use of this EO in combination with vacuum packaging
520 achieved a shelf life extension of Mediterranean octopus of approximately 17 days compared
521 to untreated samples (Atrea, Papavergou, Amvrosiadis, & Savvaidis, 2009). These results
522 were in agreement with other studies conducted on refrigerated trout (*Oncorhynchus mykiss*)
523 fillets using the same EO at the similar concentration (Frangos, Pyrgotou, Giatrakou,
524 Ntzimani, & Savvaidis, 2010) as well as on common carp (*Cyprinus carpio*) fillets using 0.1%
525 cinnamon EO (Zhang et al., 2016).

526 In another investigation, the research group of one of us obtained similar results by combining
527 sage EO (2%, 4%) and vacuum packaging during refrigerated storage of rainbow trout
528 (*Oncorhynchus mykiss*) fillets stored at 4 °C (Emir Çoban et al. 2016). Based on some

529 microbiological (total aerobic mesophilic and psychrophilic bacteria) analyses, this combined
530 treatment showed a significant microbiological shelf life extension as shown in **Figure 2**. For
531 example, the total aerobic mesophilic bacteria (**Figure 2A**) exceeded the value of 7 log cfu/g,
532 which is considered as the upper acceptability limit for fish, on day 5 for air packed samples
533 (control) and on day 14 for vacuum packed ones, while the vacuum packaged samples with
534 added sage EO at the both concentrations did not reach this value throughout the whole
535 storage period. It can be concluded that although the use of vacuum packaging exhibited a
536 shelf life extension compared to air packed samples, and its combination with sage EO, in
537 particular at the higher concentration (i.e., 4%), achieved the optimal results, extending the
538 shelf life of fish up to 29 days as compared to only 3 days for the control samples (Emir
539 Çoban et al. 2016).

540 In another combined strategy, a research team from Greece provided evidence for synergistic
541 effects of thyme EO and MAP on the quality of sea bass (*Dicentrarchus labrax*) (Kostaki et
542 al., 2009) and swordfish (*Xiphias gladius*) fillets (Kykkidou, Giatrakou, Papavergou,
543 Kontominas, & Savvaidis, 2009). The same researchers also found that combination of
544 oregano EO with MAP in different gas mixtures was efficient in extending the shelf life of
545 fresh Swordfish (Giatrakou, Kykkidou, Papavergou, Kontominas, & Savvaidis, 2008) and
546 rainbow trout fillets (Pyrgotou, Giatrakou, Ntzimani, & Savvaidis, 2010).

547 Various research studies have proposed the use of EOs in combination with chitosan in
548 order to improve quality and extend the shelf life of fish and other seafoods (Alishahi &
549 Aïder, 2012; Yuan, Chen, et al., 2016). For example, the use of a coating chitosan enriched
550 with cinnamon EO delayed lipid oxidation in refrigerated rainbow trout and markedly reduced
551 the TBA and PV values compared with the control samples (Ojagh et al., 2010). In addition,
552 the authors reported that this combination strategy effectively decreased the TVC and
553 psychrotrophic bacteria in the fish during 16 days of cold storage. Similar results were also

554 found in other recent studies, in which chitosan films were enriched by EOs from rosemary
555 and thyme (Doğan & İzci, 2017), oregano (Vatavali et al., 2013), and garlic (Aşik &
556 Candoğan, 2014).

557 **7. Limitations and future trends**

558 Despite the promising antimicrobial and antioxidant activities observed for many EOs,
559 some limitations have been underlined in their application in fish and other seafood products.
560 For example, our review study showed that the efficiency of EOs as natural preservatives was
561 variable, changing from one study to another, possibly due to the differences in either the
562 composition of EOs or the nature and the type of seafood products treated with these EOs.
563 Indeed, on the one hand, many authors reported that the composition of EOs is dependent on
564 many factors, such as the harvesting season, the variety of herb spice, or plant, the part of
565 vegetables used for extraction of EOs, geographical origin, and the method used in the
566 extraction (Burt, 2004; Hyldgaard et al., 2012; Rodriguez-Garcia et al., 2016). On the other
567 hand, some authors (Tajkarimi et al., 2010) reported that the efficiency of EOs may be
568 affected by fat level of fish, since some EOs were found to be more effective on lean fish
569 (e.g., cod) than on fatty fish (e.g., salmon). Moreover, the presence of fats, carbohydrates,
570 proteins, and salts as well as the interaction between these compounds and EOs added to
571 seafood products could reduce the preservative activity of these oils when compared to *in*
572 *vitro* application. That is why higher concentrations of EOs are usually necessary to achieve
573 satisfactory antimicrobial and antioxidant activity in such products, which in turn may cause
574 negative organoleptic effects and even health problems (Burt, 2004; Calo et al., 2015;
575 Hyldgaard et al., 2012; Solórzano-Santos & Miranda-Navales, 2012).

576 Recently, some solutions based on the encapsulation of EOs in polymers of edible and
577 biodegradable coatings, or into micro- and nanoemulsions, or the use of EOs in active food

578 packaging, have been proposed to overcome drawbacks related to the possible negative
579 sensory effects of high concentrations of EOs (Acevedo-Fani et al., 2016; Atarés & Chiralt,
580 2016; Ribeiro-Santos et al., 2017). However, further work is still required in this research area
581 in order to optimize the effectiveness of EO for applications in preservation of fish and other
582 seafoods. This may include a better understanding of the exact mechanisms of action of EOs
583 as antimicrobial and antioxidant agents and the determination of the optimum dose needed to
584 get the desired impacts of this treatment without compromising the sensory property or the
585 safety of seafoods. Additionally, future research should also focus on synergism between EOs
586 and other compounds or preservative techniques, in order to provide the maximum beneficial
587 impact, thereby extending as much as possible the shelf life of fish and other seafood
588 products.

589 **8. Concluding remarks**

590 The information compiled in this review demonstrates that different EOs incorporated
591 directly into fish and other seafoods, or applied indirectly by other methods, can effectively
592 inhibit or reduce lipid oxidation and growth of various microorganisms. Many EOs could be
593 used alone or in combination with other preservative treatments to further prevent or retard
594 oxidation and microbial spoilage in food systems, especially in fish and fish products, thereby
595 extending the shelf life of these products. Indeed, while the importance of the use of EOs in
596 enhancing antioxidant and antimicrobial stability of seafoods is being widely recognised, their
597 combination with other preservation method is resulting in further superior results. Being the
598 principal constituents of EOs, many authors reported that phenolic compounds are mainly
599 responsible for their antimicrobial and antioxidant properties.

600 Our literature review revealed that EOs from plant materials, such as oregano, rosemary,
601 thyme, sage, clove, laurel, cumin, and basil could be used at different concentrations, and

602 often, the preservative effect was greater as the EO concentration was higher. Therefore,
603 natural additives such as EOs have the potential to replace or partly replace the synthetic
604 additives. However, it must be kept in mind that the application of EOs at high dose could
605 impart some undesirable organoleptic changes and may even induce serious health problems.
606 Hence, some considerations must be taken into account when using EOs in food preservation
607 in order to find a balance between the effective compound dose and the potential risk of
608 toxicity. Future research should thus focus on the safety and possible side effects of EOs
609 before a regularly approval for their use as natural additives of fish and other seafood
610 products.

611

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955 **Figure and Table Captions**

956 **Figure 1:** Effect of *Zataria multiflora* Boiss essential oil added at concentration of 0.2 % (0.2
957 ZMEO) and 0.4 % (0.4 ZMEO) on (A) peroxide value (PV) and (B) thiobarbituric acid (TBA)
958 of catfish burgers during storage at 4 °C.

959 **Figure 2:** Effect of sage essential oil in combination with vacuum packaging on (A) total
960 aerobic mesophilic bacteria and (B) Psychrophilic bacteria of rainbow trout stored in air
961 without sage essential oil (control), vacuum packaged (VP), vacuum packaged combined with
962 2% (VP-EO2) or 4% (VP-EO4) sage essential oil.

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964 **Table 1:** Summary of some relevant research results testing the antimicrobial and antioxidant
965 activities of common essential oils or their components in fish and other seafoods

966 **Table 2:** Relevant examples of antimicrobial and antioxidant properties of some common
967 essential oils combined with other preservation methods in fish and other seafoods

Figure 1: Hassoun and Emir Coban (2017)

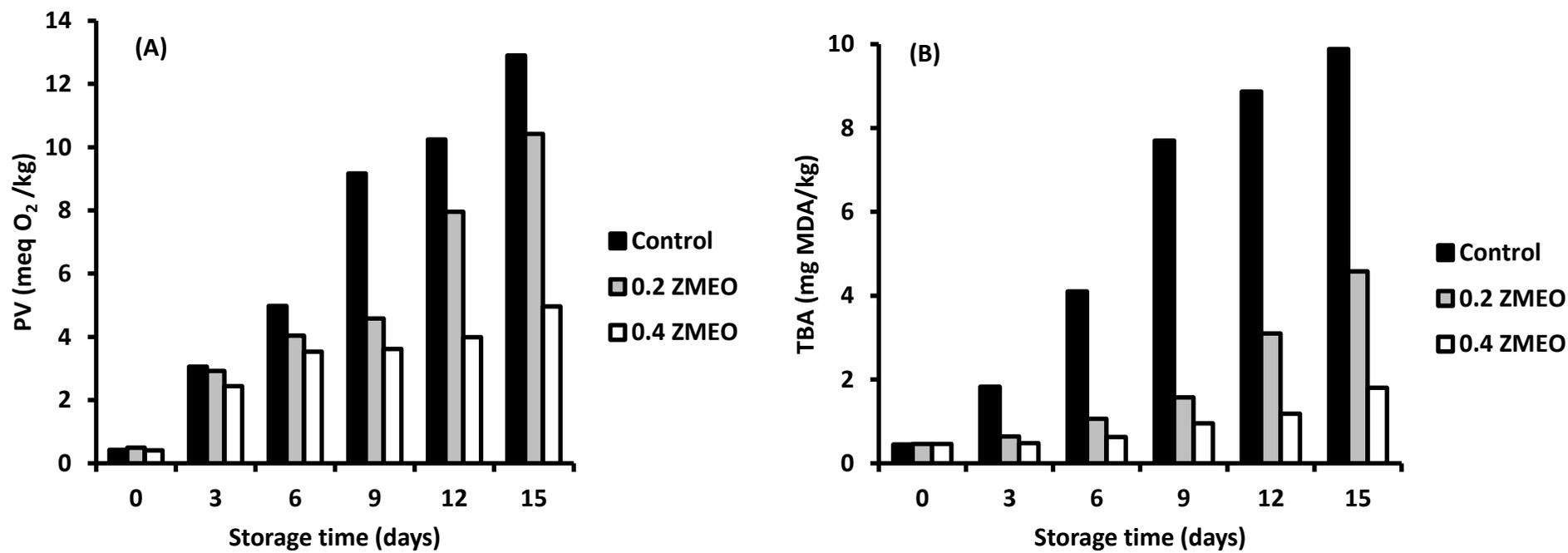


Figure 2: Hassoun and Emir Coban (2017)

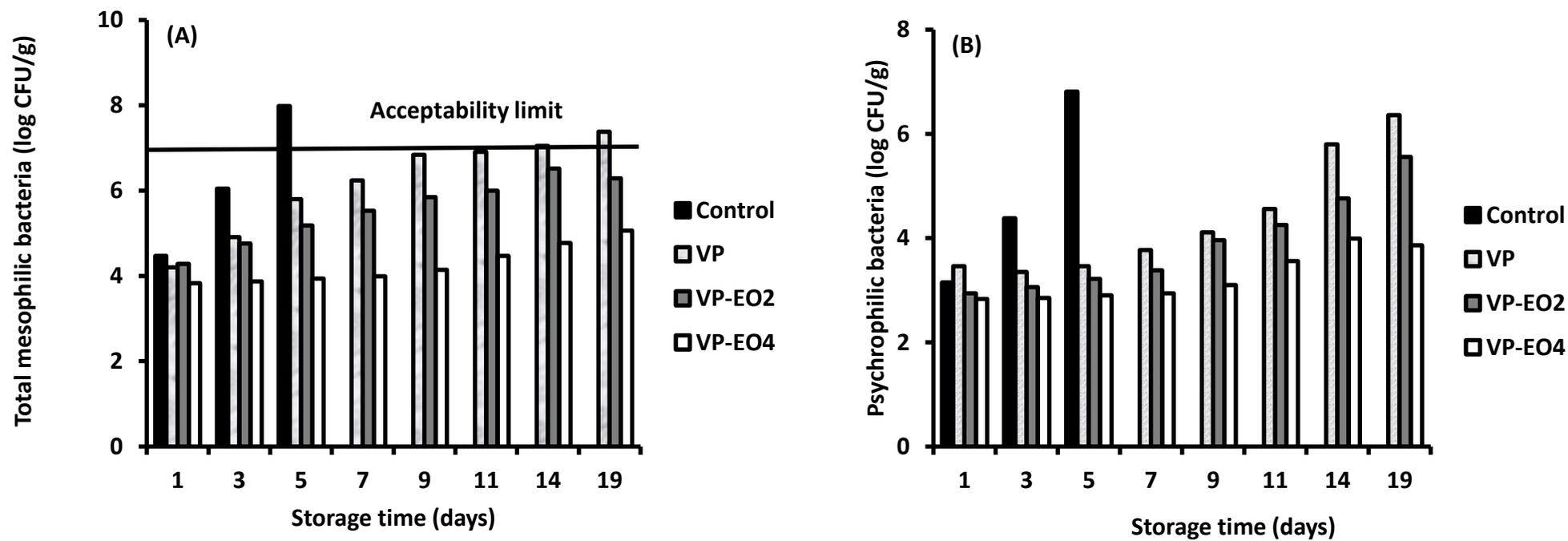


Table 1: Hassoun and Emir Coban (2017)

Seafood product	EOs or their compounds	Mode of application	Antimicrobial and antioxidant effects	Reference
Atlantic Mackerel (<i>Scomber scombrus</i>)	Rosemary and basil	Immersion (1%)	Treatment with these EOs resulted in lower contents of volatiles compounds and primary and secondary oxidation products, with a shelf life extension of 2-5 days compared to the control samples	(Karoui & Hassoun, 2017)
Carp (<i>Cyprinus carpio</i>)	Garlic EO and different constituents of EOs	Immersion (1% and 2%)	Dipping fish fillets into a solution containing both carvacrol and thymol led to a remarkable reduction in the microbial growth, consequently extending the shelf life of the fish	(Mahmoud et al., 2004)
Bluefish (<i>Pomatomus saltatrix</i>)	Thyme and laurel	Immersion (1%)	Treatment with both EOs resulted in a reduction of microbial growth and lower lipid oxidation rates, extending the shelf life from 9 days for the control to 13 days for treated samples	(Erkan, Tosun, Ulusoy, & Üretener, 2011)
<i>Sarda sarda</i>	Ginger	Direct addition (0.5% and 1%)	Treatment of fish fingers with 1% EO extended shelf life up to 17 days compared to only 5 days for untreated samples	(Emir Çoban, 2013)

Seafood product	EOs or their compounds	Mode of application	Antimicrobial and antioxidant effects	Reference
Rainbow trout (<i>Oncorhynchus mykiss</i>)	Clove	Immersion (0.1%, 0.5%, and 1%)	Treatment of smoked fillets with 0.5% and 1% clove EO decreased spoilage bacterial growth (TVC and psychrotrophic bacteria) and lipid oxidation (PV and TBA) and extended shelf life by 4–5 weeks compared to the control samples	(Emir Çoban & Patir, 2013)
Rainbow trout (<i>Oncorhynchus mykiss</i>)	Rosemary, laurel, thyme, and sage	Nanoemulsion	In addition to their antimicrobial and antioxidant properties, the encapsulation of these EOs, in particular rosemary and thyme ones, into nanoemulsions enhanced organoleptic quality of fish, giving a bitter taste	(Ozogul et al., 2017)
Shrimps (<i>Parapenaeus longirostris</i> <i>Lucas 1846</i>)	Orange	Coating (0.5%, 1%, and 2%)	Gelatin coating enriched with 2% orange leaf EO showed significant antioxidant and antimicrobial activities, achieving a shelf life extension in shrimps of about 10 days	(Alparslan et al., 2016)

EO: Essential Oil; MAP: Modified Atmosphere Packaging; PV: Peroxide Value; TBA: Thiobarbituric Acid; TVC: Total Viable Count

Table 2: Hassoun and Emir Coban (2017)

Seafood product and storage conditions	EOs or their compounds	Main results	Reference
Rainbow trout fillets packaged under vacuum in combination with salt	Oregano (0.2%, 0.4%)	The combination of oregano (0.2%) and vacuum packaging achieved a significant shelf life prolongation of fish fillets (11–12 days) compared to control samples packaged in air (5 days)	(Frangos et al., 2010)
Smoked and vacuum packed rainbow trout fillets	Rosemary, sage, thyme, and clove (0.06%)	In particular, the results demonstrated that the addition of clove EO had the highest preservation impact, resulting in a significant extension of shelf life of the product of about 6-7 weeks.	(Emir Çoban, Patir, & Yilmaz, 2012)
Common carp stored under vacuum packaging	Cinnamon (0.1%)	Based on sensory, microbial, and some physico-chemical parameters, it was reported that the combined treatment maintained good quality shelf life was extended by 2 days compared to untreated samples	(Zhang et al., 2016)
Sea bass fillets packaged under different MAP	Thyme (0.2%)	The use of this EO improved the quality of fish fillets when applied in combination with 60% CO ₂ , 30% N ₂ , and 10% O ₂ , extending the shelf life to 17 days compared to only 6 days for control samples	(Kostaki et al., 2009)
Mediterranean Swordfish fillets packed under MAP	Thyme (0.1%)	The combination of thyme EO and MAP reduced the TVC and H ₂ S-producing bacteria and inhibited lipid oxidation, extending the shelf life to about 20 days compared to 8 days for the control	(Kykkidou et al., 2009)

Seafood product and storage conditions	EOs or their compounds	Main results	Reference
Salted rainbow trout fillets stored under MAP	Oregano (0.2%, 0.4%)	The use of 0.2% oregano EO in combination with MAP reduced the main spoilage microorganisms and the content of some biochemical parameters (TVB-N, TMA-N) and extended the shelf life of the fish up to 21 days	(Pyrgotou et al., 2010)
Salted sea bream fillets stored under MAP	Oregano (0.4%, 0.8%)	The combination of oregano EO (0.8%) and MAP exhibited a strong antioxidant (measured as TBA value) and antimicrobial (estimated as volatile amines contents) activities, which extended the shelf life of fish fillets by more than 17 days	(Goulas & Kontominas, 2007)
Peeled shrimps packaged under MAP	Thymol (0.05%, 0.1%, 0.15%)	The authors obtained a shelf life of about 14 days for the active coating (0.1%) packaged under MAP compared to the samples stored in air (5 days)	(Mastromatteo, Danza, Conte, Muratore, & Del Nobile, 2010b)
Carp fillets pre-treated with electrolyzed NaCl solutions	Carvacrol and thymol (0.5%)	The combined treatment resulted in a significant reduction in the total microbial count and content of lipid oxidation products (determined by PV and TBA) which prolonged the shelf life of the treated fillets to 16 days compared to 4 days for the control	(Mahmoud, Yamazaki, Miyashita, Shin, & Suzuki, 2006)
Whole red Porgy coated with chitosan	Oregano (0.1 %)	The combination of oregano EO and chitosan improved antimicrobial and antioxidant properties of fish and achieved a shelf life extension of about 8 to 9 days	(Vatavali et al. 2013)

EO: Essential Oil; MAP: Modified Atmosphere Packaging; PV: Peroxide Value; TBA: Thiobarbituric Acid; TMA-N: Trimethylamine Nitrogen; TVB-N: Total Volatile Base Nitrogen