Research Article

Kinetics of Quality Changes of Pangasius Fillets at Stable and Dynamic Temperatures, Simulating Downstream Cold Chain Conditions

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This study was about the quality changes of Pangasius fillets during storage under simulated temperature conditions of downstream cold chain. Sensory, chemical, and microbiological analyses were conducted over storage time and bacterial growth was modelled. Sensory quality index (QI), at five stable (1, 4, 9, 15, and 19 ± 1°C) and three dynamic temperatures, progressed faster at higher temperatures, especially with sooner temperature abuses. Total volatile basic nitrogen remained under the acceptable limit throughout all the storage conditions. Total viable psychrotrophic counts (TVC) were around 5.68 ± 0.24 log CFU g⁻¹ at the beginning and exceeded the limit of 6 log CFU g⁻¹ after 216, 96, 36, 16, and 7 h at 1, 4, 9, 15, and 19 ± 1°C, respectively. Meanwhile, Pseudomonas counts started at 3.81 ± 0.53 log CFU g⁻¹ and reached 4.60–6.36 log CFU g⁻¹ by the time of TVC rejection. Since lower shelf lives were given by TVC rather than QI, it should be appropriate to base the product shelf life on the TVC acceptable limit. Kinetics models based on the Baranyi and Roberts and square root models, developed for TVC and Pseudomonas spp., gave acceptable bacterial estimations at dynamic temperatures, with over 80% of observed counts within the acceptable simulation zone, revealing promising model applicability as a supporting tool for cold chain management. However, further improvement and validation of the models are needed.

1. Introduction

Temperature is among the most important factors affecting the quality of aquatic products [1, 2]. At the end of a cold supply chain, aquatic products are normally subjected to chill-stored conditions with temperature fluctuations and/or abuses [3–5], which affect their quality and shorten their shelf life [2]. Continuous monitoring of the temperatures and assessing the sensory, chemical, and microbiological changes of aquatic products over time are useful tools for determining their quality, estimating shelf life, and supporting quality management [2, 6–8]. Quality index (QI), a total sensory score obtained from the quality index method (QIM), and total volatile basic nitrogen (TVB-N), produced by enzymatic and microbiological activities, are commonly used as freshness indicators of fish [6, 7]. Furthermore, TVB-N contents along with microbial counts are frequently used to endorse the rejection time estimated by QIM [9]. Total viable psychrotrophic counts (TVC) and specific spoilage organisms (SSOs) such as Pseudomonas spp. during cold storage of fish also serve as good indicators for fish freshness [10–12], since Pseudomonas spp. are a common SSO of iced fresh water fish [13]. Five species of Pseudomonas (P. otitidis, P. hibiscicola, P. geniculate, P. beteli, and P. aeruginosa) were found from Pangasius at filleting and trimming steps of a large-scale factory in Mekong Delta of Vietnam [14]. SSOs often cause sensory spoilage, with off-odors and off-flavors, when they reach concentrations above 7 log CFU g⁻¹ in fresh fish [15].

Kinetic modelling of microbial growth under simulated supply chain conditions is becoming more and more important for monitoring food contamination, shelf life, and risk assessment [4, 8, 16]. Several models considered appropriate for food systems, for example, Baranyi and Roberts model [17], Gompertz model [18], Ratkowsky equation [19], and the Arrhenius equation [20], have been parameterized for SSOs...
and/or pathogens in certain food/seafood products under isothermal and nonisothermal conditions. For example, they have been fitted for the growth of *Pseudomonas psychrophila* and *Carnobacterium maltaromaticum* in tropical shrimp (*Penaeus notialis*) [8], TVC and lactic acid bacteria (LAB) in vacuum packed chilled tuna [5], and pseudomonads in aerobically stored gilt-head sea bream [12].

Tra catfish (*Pangasius hypophthalmus*) is a popular commercial farmed aquatic species traded worldwide with large markets like the US, EU, Asia, and Latin America [21]. Vietnam remains the largest exporter of *Pangasius* with a value of 1.715 billion USD in 2016 [22]. Skinless and boneless frozen/chilled *Pangasius* fillets are one of the most common product types [23, 24]. Therefore, it is of outmost importance to clearly understand the kinetics of quality changes of this product under simulated supply chain condition to support decision-making in cold chain management. Several researches with this product type have been focused on its pH and sensory changes during storage at 0, 3, 5, and 10°C [25], sensory QI changes at 0–2°C [26], microflora during processing [14], microbiological spoilage with vacuum and modified atmosphere packaging [27], import risk analysis of this product to New Zealand from Vietnam [28], composition and quality attributes of conventionally and organically farmed *Pangasius* fillets on the German market [23], and so on. However, sensory, chemical, and SSO quality changes of *Pangasius* fillets under stable and dynamic temperatures at downstream steps of the supply chain have not been systematically studied. To the best of our knowledge, no microbiological models have been developed for the product so far.

This work was to find out the quality changes of *Pangasius* fillets under simulated temperature conditions of cold chain downstream steps by assessing sensory, chemical, and microbiological changes and kinetic modelling of bacterial growth. This would enable the estimation of the product quality and shelf life based on temperature history to support cold chain management.

2. Materials and Methods

2.1. Preparation of Fish Samples. Farmed Tra catfish (*Pangasius hypophthalmus*) fillets of size 170/220 were used. Each stable temperature regime worked with 5 batches (batches 1–5), while every dynamic temperature regime did with 3 batches (batches 1–3); each consisted of 105–112 fillets.

Frozen fillets in semiblocks in plastic bags, fully covered with 500 g cooling mats in 30 kg insulated boxes, were transported by car from a factory, located in An Giang province of Vietnam, on the day of processing to the laboratories in Nha Trang city within 1 day. A 3M TL30 temperature logger (3M, Saint Paul, MN, USA) was put on top of the fish blocks to record the product temperature at 10 min intervals with a precision of ±0.5°C. At the laboratories, fish was stored in a freezer at 1 ± 1°C, packed 1 fillet per polystyrene tray, and covered with thin polyethylene film. This was to simulate a real supply chain situation in Vietnam and elsewhere that frozen and/or fresh fish is commonly repacked into retail package at retailers [2, 24].

2.2. Storage Conditions. Fish trays were stored in a refrigeration with five stable temperature conditions of 1, 4, 9, 15, and 19 ± 1°C and three dynamic temperature regimes, namely, Dynamic 1, Dynamic 2, and Dynamic 3 (Table 1), simulating downstream conditions of the supply chain. Starting point of the storage right after repack was considered as time 0 (0 h).

At Dynamic 1, fish fillets were stored at ± 1°C, assuming that they were kept in a retail refrigerator/cold store, for 48 h (2 days). Then they were placed outside at room temperature (of 28–30°C) for 2 h to simulate that they were bought and carried home by an end customer/consumer. Finally, the product was stored at 9 ± 1°C, simulating a household refrigerator condition, until unfit for human consumption.

At Dynamic 2, fish fillets were firstly stored at ± 1°C at 7:00 AM of day 0; from 9 AM to 12 AM and from 4 PM to 7 PM of day 0, refrigerator door was opened for 1 min every 30 min, simulating the product kept in a retail refrigerator at ± 1°C for 27 h with 1 min door opening every 30 min at busy shopping hours. Then the fish was placed outside at ambient temperature (of 28–30°C) for 2 h, assuming that the product was purchased at 10 AM of day 1 (after 27 h of storage in retail refrigerator) and carried home. Finally, the product was stored at 9 ± 1°C, resembling a household refrigerator condition, until it is unfit for human consumption.

Dynamic 3 assumed that fish fillets were kept in a chilled store of a retailer at ± 1°C, started from 8 AM of day 0 until being unfit for human consumption. Every day, from 8 AM to 8 PM, store door was opened for 5 min every 2 h, simulating conditions of loading and unloading the product at the retailer store.

Samples were taken over storage time from each batch for quality assessment. Each sample included 2 fillets for QIM, 2 fillets for TVB-N, and another one for microbiological analyses.

<table>
<thead>
<tr>
<th>Regime</th>
<th>Temperature</th>
<th>Total storage time (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(1) Dynamic 1</td>
<td>48 h 2°C 20 h</td>
<td>70</td>
</tr>
<tr>
<td>(2) Dynamic 2</td>
<td>27 h 1 min/30 min, 2 h*</td>
<td>47</td>
</tr>
<tr>
<td>(3) Dynamic 3</td>
<td>50 h 5 min/2 h**</td>
<td>50</td>
</tr>
</tbody>
</table>

* Abuses from 1°C to 28–30°C for 1 min every 30 min during storage hours 2–5 and 9–12 and for 2 h during storage hours 48–50. ** Abuses from 1°C to 28–30°C for 5 min every 2 h during storage hours 0–12 and 24–36.
For continuous monitoring of the temperatures, DSI922L-F5 iButton* loggers (Maxim Integrated Products, Inc., CA) were put on some tray surface and tray bottom and inside the trays in direct contact with the fish recording temperature at 10 min intervals, while EC850A loggers (MicroLogPRO II, Israel) were used for refrigerator temperature monitoring.

2.3. Sensory Evaluation. Sensory evaluation of raw fillets was conducted by 3 panelists familiar with QIM, selected and trained in accordance with ISO 8586: 2012 [29]. The QIM scheme for chill-stored Pangasius fillets consisted of 6 attributes (skin side colour, backbone side mucus, backbone side colour, backbone side texture, backbone side odour, and backbone side stickiness) of 1–3 demerit scores with the total QI of 13 [26]. The QIM evaluation was carried out with 2 fillets from each batch per sampling point; each session worked with fish from 2 different batches. The fillets were coded with random 3-digit numbers, placed in a random order, and evaluated individually.

2.4. Chemical Analysis. Analysis of TVB-N was performed as described by Malle and Tao [30] in duplicate. Fish fillets were minced and extracted with a 7.5% aqueous trichloroacetic acid solution and titrated with sulphuric acid solution. TVB-N was steam distilled into boric acid solution. TVC and Pseudomonas −1 (log CFU g ) counts were done on plate count agar (Merck, Germany). Enumeration of presumptive pseudomonads was performed using Pseudomonas Agar Base (Merck, Germany) with Cephaloridine Fucidin Agar Base (Merck, Germany) with Cetrimide-selective Agar Supplement (Merck, Germany) [31]. Spread-plating was used for all media. Plates were incubated at 19 ± 1°C for 5 days. Counts are reported as decimal logarithmic average values of colony-forming units per gram (log CFU g ).

2.5. Microbiological Analysis. Fillet was aseptically minced. Two to five replicate samples were evaluated for each storage regime. Minced flesh (25 g) was mixed with 225 mL of chilled saline peptone diluent in a stomacher for 1 minute. Successive 10-fold dilutions were done as required. TVC was done on plate count agar (Merck, Germany) with random 3-digit numbers, placed in a random order, and evaluated individually.

2.6. Microbiological Growth Modelling. Measured data of TVC and Pseudomonas spp. at five stable temperature conditions was fitted, using DMFIT available software on ComBase website (https://browser.combase.cc/DMFit.aspx), with the primary Baranyi and Roberts model [17, 32] as follows (see (1)):

\[
y = y_{\text{max}} + \ln \left( \frac{1 + \exp(\mu_{\text{max}} t - h_0) - \exp(-h_0)}{\exp(y_{\text{max}} - y_0) + \exp(\mu_{\text{max}} t - h_0) - \exp(-h_0)} \right)
\]

\[h_0 = \ln \left( \frac{1 + q_0}{q_0} \right)
\]

\[\lambda \cdot \mu_{\text{max}} = \ln \left( \frac{1 + q_0}{q_0} \right) = h_0,
\]

where \( y \) is the decimal logarithm of bacterial counts at time \( t \) (log CFU g ); \( y_0 \) is the decimal logarithm of initial bacterial counts (log CFU g ); \( y_{\text{max}} \) is the decimal logarithm of maximum bacterial counts (log CFU g ); \( \mu_{\text{max}} \) is the maximum specific growth rate (h ); \( \lambda \) is the lag time (h); \( q_0 \) represents physiological state of the bacterial cells introduced; \( h_0 \) is considered as a statically stable transformation of \( q_0 \).

The dependency of bacterial growth on temperatures was modelled with the secondary square root model of Ratkowsky [19] as follows (see (2)):

\[\sqrt{\mu_{\text{max}}} = b (T - T_{\text{min}}),\]

where \( T_{\text{min}} \) is the conceptual minimum temperature for the bacterial growth (°C); \( T \) is the temperature (°C); \( b \) is the regression coefficients (°C h ).

2.7. Validation of the Kinetics Models. The accuracy of the fittings was evaluated based on the coefficient of determination \( R^2 \) and standard error of fit (SE of fit) [20, 32].

For further validation of the developed kinetics models, data from three dynamic temperature regimes were used. The primary and secondary models were combined to predict the bacterial growth at dynamic temperature regimes within time-temperature history intervals of constant temperatures [20]. The acceptable simulation zone (ASZ), defined as ±0.5 log CFU g from the predicted counts, was used for comparison with bacterial observed growth; and the models would be considered acceptable if at least 70% of the observed counts was within the ASZ of prediction [33–36].

2.8. Data Analysis. Microsoft Excel 2010 (Microsoft, Redmond, WA, USA) was used to calculate means and standard deviations to build graphs. One-way ANOVA (analysis of variance) and post hoc Tukey’s test were conducted with the software SPSS version 17.0 (SPSS, Chicago, IL, USA) to compare the mean values for a statistical significant level of 0.05.

3. Results and Discussion

3.1. Sensory Changes of Pangasius Fillets at Different Storage Conditions. At stable temperature conditions, QI of Pangasius fillets has increased over time at each storage temperature condition and with faster changes at higher temperatures, well observed from trendlines (Figure 1). The fact that QI is close to zero for very fresh products and went up as fish deteriorated is in good agreement with the QIM results of other fishery products stored at low temperatures [6, 10, 37, 38]. At the end of the study period, QI reached mean scores of 10.4, 6.9, 7.2, 6.2, and 6.4, accounting for 80%, 53%, 55%, 47%, and 49% of the total QI of the scheme after 360 h at 1 ± 1°C, 168 h at 4 ± 1°C, 72 h at 9 ± 1°C, 26 h at 15 ± 1°C, and 12 h at 19 ± 1°C, respectively. According to Sykes et al. [39], the storage of aquatic products should be ended when QI value is around 75% of the total QIM scale (i.e., when QI = 75% * 13 ≈ 9.8 in this study), which indicates that the shelf life of Pangasius fillets was around 336 h (14 days) at 1 ± 1°C, >168 h at 4 ± 1°C, >72 h at 9 ± 1°C, >26 h at 15 ± 1°C,
and >12 h at 19 ± 1°C. The QIM results at 1 ± 1°C cofounded well with the findings from a previous study demonstrating that the maximal shelf life of *Pangasius* fillets stored at 0–2°C was 15 days as evaluated by QDA [26]. For skinless *Pangasius* fillets from Malaysia, Abbas et al. [25] predicted shelf lives of 16–18 days in ice storage at 0–2°C. For skin-on *Pangasius* fillets stored in ice, Bao [40] found a shelf life of about 12 days. For whole *Pangasius* from Bangladesh stored in ice, Azam et al. [41] found shelf lives of medium size (550–650 g) fish around 12 days in summer season and 10 days in winter season based on sensory evaluation. The difference in shelf life of *Pangasius* products might be due to the variation in product types (skinless versus skin-on fillets, as well as whole fish), fish origin (Vietnam versus Malaysia, as well as Bangladesh), storage conditions (chilled air versus in ice), and so on.

At all five stable temperatures, QI was well linearly correlated with storage time, which agreed well with other studies about the linear correlation of QI and storage time [6, 38–40]. The presented regression equations (Figures 1(a) and 1(b)) are good tools for freshness evaluation and shelf life estimation of the product.

At dynamic temperature conditions, an increasing trend of QI over time was also observed (Figure 1(c)). When comparing the changes of *Pangasius* fillet QI at stable 1 ± 1°C (Figure 1(a)) and those at dynamic 1°C (Dynamic 3, Figure 1(c)), it could be seen that QI in the latter case evolved faster than the first one. This difference could be explained by the fact that temperature abuse in the latter case caused faster rate of spoilage [2, 20]. Comparing QI from the dynamic regimes 1 and 2 (Figure 1(c)), it was found that the sooner temperature abuse occurred (after 48 h of Dynamic 1 and after 27 h of Dynamic 2), the faster QI changed, clearly observed after the temperature shifting. The results from dynamic conditions of storage show that QIM has potential for detecting the sensory changes in temperature fluctuation/abuse situations.

3.2. TVB-N Changes of Pangasius Fillets at Different Storage Conditions. At all studied temperature storage regimes, including five stable and three dynamic temperature conditions, TVB-N contents in fish fillets were initially as low as 4.41 mg N/100 g and remained well below the acceptable

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**Figure 1**: Changes in quality index (QI) of *Pangasius* fillets during storage at (a) 1, 4, and 9 ± 1°C, (b) 15 and 19 ± 1°C, and (c) dynamic temperature conditions. Different letters (A, B, C, D, and E) with numbers (1, 2, 3, 4, and 5) indicate significantly different QI mean values \((P < 0.05)\) between samples over storage time of each temperature regime. In the regression equations, \(y\) is QI and \(x\) is storage time (h).
limit for Pangasius fillet of 25 mg N/100 g [42] throughout storage time (Figure 2). No significant changes (P > 0.05) were observed in TVB-N level during storage periods at 4, 9, 15, and 19 ± 1°C and at three dynamic temperature regimes. At 1 ± 1°C, TVB-N remained around 4.31–8.09 mg N/100 g until the end of the storage (360 h), where its contents significantly rose (P < 0.05) up to 11.97 ± 3.30 mg N/100 g. This agreed well with previous studies demonstrating that TVB-N level only increases at the later time of storage when spoilage becomes evident [2, 43]. Ammonia contents in skin-on Pangasius fillets stored in ice were 4.5 mg N/100 g at initial time and 49 mg N/100 g at day 14 [40]. Meantime, for whole Pangasius of medium size, the initial content of TVB-N was 4.05 ± 0.25 mg N/100 g, and it exceeded the level of 30 mg N/100 g after 14 days in ice [41]. In a study on cod loin air stored at 0.5 ± 0.5°C, TVB-N remained low during the first 10 days of storage, after which its level increased rapidly, exceeding the human consumption allowed limit for gadoids of 35 mg N/100 g [44] after 13 days [2]. To summarize, TVB-N contents in Pangasius fillets in the current study were still within the limit for human consumption at the end of the storage periods of different temperature conditions, which reveals that TVB-N could not be a useful indicator for shelf life determination of this product.

3.3. Kinetics of TVC and Pseudomonas Growth in Pangasius Fillets at Stable Temperature Conditions. At stable temperatures, the TVC was around 5.68 ± 0.24 log CFU g⁻¹ at the beginning and increased over time more rapidly at higher storage temperatures, exceeding the acceptable limit of 6 log CFU g⁻¹ [42] after 216, 96, 36, 16, and 7 h at 1, 4, 9, 15, and 19 ± 1°C, respectively (Figure 3(a)). Meanwhile, Pseudomonas counts started at 3.81 ± 0.53 log CFU g⁻¹ and grew with the similar trend, reaching 4.60–6.36 log CFU g⁻¹ by the time of TVC rejection (Figure 3(b)). These levels of Pseudomonas spp. were much lower than the level of 7 log CFU g⁻¹, which caused spoilage for fresh gilthead sea bream Mediterranean fish (Sparus aurata) [15], or the set level of 7.5 log CFU g⁻¹ for the shelf life of fresh pork or poultry [20].

When comparing the shelf life limits of the studied product based on QI and TVC, it can be seen that TVC gave much lower shelf lives than sensory QI did. In contrast, some other studies found that sensory rejection time was well associated with a TVC level of 6 log CFU g⁻¹, for example, for
wild turbot [45] or for farmed turbot stored in flake ice [46]. Meanwhile, it was reviewed that there have been no common rules on the bacterial counts and the sensory rejection time [4]; for example, the bacterial level at the sensory shelf life limit was ca. 7 log CFU g⁻¹ [12] or ca. 8 log CFU g⁻¹ [47, 48]. Findings of this study indicate that the shelf life of Pangasius fillets at low temperatures would rather be based on the limit level of TVC (6 log CFU g⁻¹) than on the sensory QI (75% of the total scale).

To observe propagation trends of TVC and Pseudomonas spp. over time at stable temperatures, Baranyi and Roberts model was used to fit the microbial data, and the results were shown with fitted curves in Figure 3. The maximum growth rate, generally, increased with storage temperatures, clearly observed for TVC and Pseudomonas counts at temperatures of 1, 9, 15 and 19°C, which agreed well with the growth rate increasing trend with temperatures from other studies, for example, for total bacteria and Pseudomonas in turbot (Psetta maxima) at 0, 5, 10, and 15°C [4] or for Pseudomonas counts in poultry at temperatures of 2, 4, 10, 15, and 20°C [32]. Furthermore, the growth of psychrotrophic bacteria was shown to have very short lag phase (estimated about 6.31±4.56 h for TVC) or no lag (for Pseudomonas spp.) at 19±1°C (Figure 3), which can be explained by the fact that the indicated temperature is close to the optimum range for their growth [49]. The maximum bacterial counts at stationary phase were found to converge to about 9.20 log CFU g⁻¹ for TVC and 8.24 log CFU g⁻¹ for Pseudomonas spp. (Figure 3), which agreed well with the findings for other fish products; for example, maximum bacterial concentrations of total bacteria and Pseudomonas in turbot (Psetta maxima) at isothermal range of 0–15°C were 7.55–10.20 and 7.05–9.07 log CFU g⁻¹, respectively [4].

### 3.4. Modelling the Influence of Temperature on the Growth of TVC and Pseudomonas spp.

To exploit the influence of temperatures on the growth of the bacteria, the square root model [19] was used as secondary model and fit parameters were shown in Table 2. The parameters were $T_{\text{min}} = -10.87829 ± 1.38524°C$ and $b = 0.011059 ± 0.00131 (\text{C} \cdot \text{h}^{-\frac{1}{2}})$ for TVC and $T_{\text{min}} = -13.36799 ± 1.19593°C$ and $b = 0.144475 ± 0.01293 (\text{C} \cdot \text{h}^{-\frac{1}{2}})$ for Pseudomonas spp. The results were very similar to those of other seafood; for example, for the growth of Pseudomonas spp. in tropical shrimp (Penaeus notialis) stored at 0 to 28°C, the model parameters were $T_{\text{min}} = -12.1°C$ with 95% confidence interval (CI) (−13.1, −10.9°C) and $b = 0.019$ with 95% CI (0.017, 0.021) [8]. Meanwhile, estimated parameters of the square root approach in the Baranyi and Roberts model for Pseudomonas in poultry within a temperature range of 2–20°C were $T_{\text{min}} = -5.8812 ± 2.466°C$ and $b = 0.02045 ± 0.0029 (\text{C} \cdot \text{h}^{-\frac{1}{2}})$ [32], which are somewhat different from those of aquatic products.

The coefficient of determination ($R^2$) was relatively high (0.85 for TVC and 0.88 for Pseudomonas spp.) (Table 2), which indicates the possibility to further apply the models in estimation of the bacterial counts at dynamic temperature conditions.

### 3.5. Validation of the Kinetics Models in Dynamic Temperature Conditions

Data from the three dynamic regimes were used for validation of the model applicability, and results were shown in Figure 4. For dynamic temperature regime 1, the total bacterial count ($n_{\text{TVC}}(T, t)$) was studied during storage at stable temperatures, with observed values (markers) and fitting curves (lines) using primary Baranyi and Roberts model.

**Table 2**: Estimated parameters with standard error (SD) of the square root model for TVC and Pseudomonas growth in Pangasius fillets stored at low temperatures.

<table>
<thead>
<tr>
<th>Microorganisms</th>
<th>$b$ (°C⁻¹ h⁻¹/²)</th>
<th>SD of $b$ (°C⁻¹ h⁻¹/²)</th>
<th>$T_{\text{min}}$ (°C)</th>
<th>SD of $T_{\text{min}}$ (°C)</th>
<th>$R^2$</th>
<th>SE of fit</th>
</tr>
</thead>
<tbody>
<tr>
<td>TVC</td>
<td>0.011059</td>
<td>0.00131</td>
<td>-10.87829</td>
<td>1.38524</td>
<td>0.84577</td>
<td>0.033892</td>
</tr>
<tr>
<td>Pseudomonas spp.</td>
<td>0.144475</td>
<td>0.01293</td>
<td>-13.36799</td>
<td>1.19593</td>
<td>0.88034</td>
<td>0.028596</td>
</tr>
</tbody>
</table>

**Figure 3**: Changes in (a) TVC and (b) Pseudomonas counts of Pangasius fillets during storage at stable temperatures, with observed values (markers) and fitting curves (lines) using primary Baranyi and Roberts model.
Figure 4: Observed and predicted growth of TVC and *Pseudomonas* spp. in *Pangasius* fillets during storage at dynamic temperature regimes: (a) Dynamic 1, (b) Dynamic 2, and (c) Dynamic 3.

(Dynamic 1, Figure 4(a)), most of the measured TVC (20 out of 21 observations, or 95.24%) fell within the ASZ, and the left one (1/21, or 4.76%) was overestimated, which could be considered as a “fail-safe” prediction. For *Pseudomonas* counts, 18/21 (85.71%) measured counts lied within the ASZ, 2/21 were underestimated (9.52% “fail-safe” prediction), and 1/21 (4.76%) was underestimated, which could be considered as a “fail-dangerous” prediction. Similarly, for dynamic temperature regime 2 (Dynamic 2, Figure 4(b)), 100% of observed TVC and 80.95% of observed *Pseudomonas* counts fell within the ASZ; and 19.05% *Pseudomonas* counts were overestimated. Comparably, at dynamic temperature regime 3 (Dynamic 3, Figure 4(c)), 87.50% of observed TVC and *Pseudomonas* counts were within the ASZ of prediction, and the other 12.5% of TVC was underestimated and 12.5% of *Pseudomonas* spp. were overestimated.

The findings are comparable with those of other studies; for example, on average, about 83% of *Listeria monocytogenes* and 75% of LAB observed in cottage cheese were within the ASZ [35]; and 78% of measured psychrotolerant pseudomonads in milk and cottage cheese lied in the ASZ [33].

The above results indicate that the developed kinetics models were acceptable for further application in prediction of TVC and *Pseudomonas* growth in *Pangasius* fillets at low temperatures, since more than 70% of the observed counts were within the ASZ [33–36].

In this study, due to short duration of storage, at the end, the TVC and *Pseudomonas* spp. have not reached the stationary phase yet. Therefore, further investigation with longer time of storage is needed in order to better validate the applicability of the models. In addition, to improve the models, more studies at other stable storage temperatures and
more replications would be desired. Moreover, it is important to bear in mind that continuous improvement and validation of kinetics models used as a quality/shelf life prediction tool are essential in quality management [20].

4. Conclusions

*Pangasius* fillet QI showed a linear correlation with storage time at 1, 4, 9, 15, and 19 ± 1°C with speedier QI rise at higher temperatures. Furthermore, QI changed faster if there were temperature abuses and if abuses occurred sooner. The findings revealed that QIM is capable of detecting sensory changes in temperature fluctuation/abuse situations. Meanwhile, TVB-N contents in fish fillets remained under the acceptable limit for human consumption till the end of all the storage conditions, which indicates that TVB-N could not be a useful indicator for shelf life determination of this product.

The TVC of *Pangasius* fillets exceeded the acceptable level of TVC (6 log CFU g⁻¹) in less than 216 h at 1 ± 1°C, 96 h at 4 ± 1°C, 36 h at 9 ± 1°C, 16 h at 15 ± 1°C, and 7 h at 19 ± 1°C, showing much lower shelf lives compared to those given by sensory QI. This reveals that the shelf life of *Pangasius* fillets stored at low temperatures should be based on the acceptable level of TVC.

Most observed bacterial counts (≥87.50% of TVC and 80.95% of *Pseudomonas* spp.) at dynamic temperature regimes fell within the ASZ of the developed kinetics models, revealing the applicability of the model as a decision supporting tool in quality management. However, further improvement and validation of the models are needed.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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