ABSTRACT

Concentrating milk by reverse osmosis (RO) has the potential to increase cheese yield but is known to impair cheese-making properties. The main compositional differences between ultrafiltration (UF) and RO concentrates are the high lactose and mineral contents of the latter. The objective of this work was to determine the distinct effects of high lactose and high minerals on the cheese-making properties of RO concentrate, by supplementing UF concentrate with lactose. The soluble colloidal equilibria of concentrates were studied as well as several other properties: rennet gelation behavior, cheese mass balance, composition, and microstructure. Rennet coagulation time was longer and gel firming rate was lower for RO concentrate than for UF concentrate. Lactose was mainly responsible for these differences. Lactose in RO concentrate was also responsible for the 7% increase of moisture-adjusted cheese yield, relative to UF concentrate. Compared with cheese made from UF concentrate, cheese made from RO concentrate showed higher moisture content, which could not be attributed to lactose but to the high mineral concentration. This study showed the potential of using RO instead of UF concentrate to maximize cheese yield. The approach is, however, limited to applications where post-acidification can be controlled, and will require appropriate strategies to reduce the negative effects of high mineral content in RO concentrate.

Key words: reverse osmosis, lactose, cheese making, mineral equilibrium, ultrafiltration

INTRODUCTION

The concentration of cheese milk by filtration is a common practice in the dairy sector. Milk concentration increases the production capacity of cheese plants and increases cheese yield (Mistry, 2003). Ultrafiltration is the main process used for milk concentration, and the increase of cheese yield is attributed to higher concentration of whey proteins in the aqueous phase of cheese. The use of reverse osmosis (RO) could be of interest, because it would increase not only the concentration of whey proteins but also the concentrations of lactose and soluble minerals in the aqueous phase of the cheese.

The effects of milk ultrafiltration on cheese yield and quality have been broadly studied. Conflicting results are reported regarding the effects of UF concentration on coagulation time. Depending on different parameters, such as pH or protein concentration, it has been shown that the rennet coagulation time of UF concentrate was nonaffected, negatively affected, or positively affected, compared with skim milk (SM; Dalgleish, 1980; Guinee et al., 1997; Waungana et al., 1998; Mistry, 2003; Karlsson et al., 2007a; Sandra et al., 2011). However, the scientific community agrees on the effect of milk ultrafiltration on rennet gel firming rate and final firmness: they are both increased with the use of concentrate (Waungana et al., 1998; Mistry and Maubois, 2004; Thomann et al., 2008; Sandra et al., 2011). Cheese properties are also modified by milk ultrafiltration (Lelievre and Lawrence, 1988; Mistry, 2013). Depending on the concentration factor, the main compositional differences between traditional cheese and UF cheese are the moisture and mineral contents (Soodam and Guinee, 2018). Cheeses made from UF milk usually have lower moisture content due to faster syneresis compared with traditional cheeses (Mistry, 2003), and higher mineral content due to the retention of micellar calcium phosphate during concentration. The latter is responsible for the high buffering capacity of UF milk, which affects the rate of acidification by starter bacteria and the mineral concentration in the cheese.

The use of RO concentrate, evaporated milk, or skim milk powder for cheese making has been reported in
the literature (Barbano and Bynum, 1984, 1985; Mayes, 1985; Kelly and Fox, 2016). These studies mainly describe practical approaches for use in cheese making. Increasing total milk solid concentration before cheese making was associated with cheese textural defects (Agbevai et al., 1983), but little information is available on the effects of specific RO concentrate constituents. In a previous study, we showed that RO concentrate impaired rennet gelation properties, increasing coagulation time and lowering gel firming rate, compared with UF concentrate (Lauzin et al., 2018). The main compositional differences between RO concentrate and UF concentrate are the lactose and mineral contents, but their specific effects on cheese-making properties have not been determined. Increasing ionic strength in milk has been shown to impair the rennet coagulation properties (Zoon et al., 1989; Awad, 2007; Karlsson et al., 2007b), but the effects of high lactose content have received little attention.

In the present study, UF concentrate was supplemented with lactose (UFL) to assess its specific effects on cheese-making properties. The milk minerals content was the main compositional difference between UFL concentrate and RO concentrate, and comparing these 2 treatments provided insight on the contribution of milk minerals to the cheese-making properties of RO concentrate.

**MATERIALS AND METHODS**

Production of Skim Milk and Concentrates

Bulk pasteurized skim milk was purchased from Natrel (Québec, Canada) and stored at 4°C until use. Milk concentration was performed using a filtration pilot system (model 1812 Lab Unit, Filtration Engineering Company, Champlin, MN) equipped with a 0.32 m² spiral-wound UF or RO membrane. The temperature was kept constant at 50°C during the process. The UF membrane used was made of polyethersulfone and had a molecular weight cutoff of 10 kDa (Synder Filtration, Vacaville, CA). The RO membrane used was made of polyamide and was characterized by a 99% average NaCl rejection (General Electric, Trevose, PA). Skim milk was concentrated until it reached a total protein concentration of 7.2% (as verified by infrared analysis; Milkoscan FT120, Foss Electric). After supplementation, the UFL concentrate was agitated at low speed for 3.5 h at 4°C. Samples were then warmed in a water bath at 32°C for 1 h before pH adjustment.

The pH of SM and UF, UFL, and RO concentrates was adjusted to 6.45 ± 0.02 at 32°C with either 1M NaOH or 10% lactic acid (Thermo Fisher Scientific). To account for the dilution factor from pH adjustment, the protein content of concentrates was then adjusted to 7.0% with water. After pH and concentration adjustments, samples were stirred at low speed at 4°C overnight. Samples were then warmed to 32°C for 1 h before readjusting the pH to 6.45 ± 0.01.

Composition of Skim Milk and Concentrates

After 2 h at 32°C, the sedimentable and nonsedimentable phases of SM and concentrates were separated by ultracentrifugation at 100,000 × g for 90 min (Optima XPN-90 ultracentrifuge, Beckman Coulter, Brea, CA). The supernatants were carefully collected, filtered through a 0.45-μm polyethersulfone filter and stored at 4°C until analysis. The concentration of diffusible minerals was determined on dialysates. Water of HPLC grade (10 mL) was loaded into dialysis tubing (6 to 8kDa, Thermo Fisher Scientific) and immersed in 500 mL of SM or concentrates for 48 h at 4°C under gentle agitation to reach equilibrium. The contents of the dialysis tubing were collected and stored at 4°C until mineral analysis.

Skim milk concentrates and their supernatants were analyzed for total nitrogen (NT), non-protein nitrogen (NPN), and non-casein nitrogen (NCN) fractions by the official micro-Kjeldahl method of analysis. A nitrogen-to-protein conversion factor of 6.38 was used. The mineral content (wt/wt; calcium, sodium, phosphorus, and magnesium) of SM, concentrates, and dialysates was determined by inductively coupled plasma (ICP) analyses on dry ashes (at 550°C overnight), as previously described by (Lauzin et al., 2018). Lactose contents of SM and concentrates were analyzed via HPLC using a Waters chromatograph (Waters Corp., Milford, MA) equipped with a Hitachi (Foster City, CA) differential refractometer detector L-7490, a 600E controller, a column oven, and a cooled 717Plus autosampler. An IC-Sep ICE-ION-300 column (Transgenomic, Omaha, NE) was used with 8.5 mM of H₂SO₄ (180 μL of H₂SO₄/L) as the mobile phase at a flow rate of 0.4 mL/min. The column temperature was kept at 40°C. Samples were prepared according to the reference method (ISO 22662, IDF...
Retention coefficient = 1 - \left(\frac{\% \text{ fat or protein in whey}}{\% \text{ fat or protein in milk}}\right).

\[1\]

Microstructure. The model cheeses were stored at 4°C for 11 d after production before analysis of their microstructure. A 2-cm-thick slice was removed from the top of the cheese, and samples (7 × 2 × 2 mm) were taken from the middle of the remaining piece of cheese. For each sample 5 pieces were taken. The pieces were put in a vial containing 3 mL of a 2% glutaraldehyde solution in 0.1 M cacodylate buffer at pH 7.3 and held at 4°C overnight. The next day, the fixation solution was removed and replaced with cacodylate buffer at pH 7.3. The samples were rinsed 3 times for 10 min in the 0.1 M cacodylate buffer (pH 7.3) before the postfixation step of a 90-min incubation in a solution containing 1% (wt/vol) osmium tetroxide in 0.1 M cacodylate buffer at pH 7.3. The rinsing step with buffer was repeated, and the samples were then dehydrated by successive steps in ethanol solution at 30, 50, 70, 95, and 100% (2 × 10 min for each concentration). The samples were then incubated for 40 min in 100% ethanol, followed by 2 × 20 min in hexamethyldisilazane, before being left to dry overnight. The dried samples were fractured and mounted on an observation frame with the fractured surfaces uppermost. They were coated with gold–palladium using a sputter coater (Nanotech Semprep II, Manchester, UK) and observed via scanning electron microscope (LSM-6360LV, JEOL Ltd., Tokyo, Japan).

Statistical Analyses

All experiments were repeated 3 times. Mean values and standard deviations are reported in the tables. Significant differences were evaluated by ANOVA with Tukey tests (α = 0.05).

RESULTS AND DISCUSSION

Composition of Milk and Concentrates

The volume of skim milk was reduced by a factor of approximately 2.1 using UF or RO processes. The overall composition of SM and concentrates is shown in Table 1. True protein and casein contents increased according to the concentration factor, but both were slightly lower (<0.015%) in RO concentrate than in UF concentrate. Concentrations being expressed on a weight basis, the difference was attributed to the higher density of RO concentrate. As expected, lactose concentration was 2.1 times higher in RO concentrate,
compared with SM. Lactose was added to UF concentrate to match the concentration in RO concentrate. Both lactose and total solids concentrations were similar in RO and UFL concentrates. Compared with SM, the ash content increased by a factor of 2.1 and 1.4, respectively, in RO and UF concentrates. All minerals are concentrated by reverse osmosis, whereas only colloidal minerals are concentrated by ultrafiltration. As expected, the concentration of sodium was not significantly affected by ultrafiltration, but the concentration of calcium, which is partly bound to casein micelles, significantly increased. The UF process did not modify the serum phase, and UF concentrate had similar ionic strength to SM.

The soluble and micellar phases of SM and concentrates were separated and the composition of the soluble phase analyzed (Table 2). Both UF and RO concentrates had similar soluble casein concentrations, which were higher than that of the original SM (P < 0.05). However, compared with the total casein content, SM had 6% soluble casein, as opposed to approximately 5% for RO and UF concentrates. Hence, the concentration process did not significantly affect the protein equilibrium between the soluble and micellar phase. The elevated temperature during filtration (50°C) certainly promoted a shift of calcium from the serum phase to the micellar phase (Gaucheron, 2005). The concentration of micellar calcium was further increased in RO concentrate. The aqueous phase of milk is saturated with calcium phosphate, and when concentrated by reverse osmosis, precipitation is expected (Lauzin et al., 2019). Supplementation of UF concentrate with lactose (UFL) did not affect the soluble casein content or the mineral equilibrium, and the ratio of soluble to total milk minerals was similar to those of UF and RO concentrates.

### Table 1. Overall composition (mean ± SD) of skim milk and concentrates

<table>
<thead>
<tr>
<th>Item</th>
<th>SM</th>
<th>UF</th>
<th>UFL</th>
<th>RO</th>
</tr>
</thead>
<tbody>
<tr>
<td>True protein (%)</td>
<td>3.25 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.97 ± 0.03&lt;sup&gt;d&lt;/sup&gt;</td>
<td>6.57 ± 0.03&lt;sup&gt;b&lt;/sup&gt;</td>
<td>6.84 ± 0.04&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Casein (%)</td>
<td>2.74 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.85 ± 0.04&lt;sup&gt;d&lt;/sup&gt;</td>
<td>5.52 ± 0.04&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.79 ± 0.03&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Lactose (%)</td>
<td>4.88 ± 0.05&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.61 ± 0.05&lt;sup&gt;b&lt;/sup&gt;</td>
<td>10.67 ± 0.53&lt;sup&gt;c&lt;/sup&gt;</td>
<td>10.15 ± 0.10&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>TS (%)</td>
<td>8.5 ± 0.2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>12.2 ± 0.1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>17.7 ± 0.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>17.8 ± 0.2&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ash (%)</td>
<td>0.71 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.03 ± 0.01&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.96 ± 0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1.53 ± 0.01&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Calcium (mg/kg)</td>
<td>1,067 ± 33&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2,102 ± 25&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1,935 ± 69&lt;sup&gt;c&lt;/sup&gt;</td>
<td>2,561 ± 61&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Sodium (mg/kg)</td>
<td>523 ± 88&lt;sup&gt;a&lt;/sup&gt;</td>
<td>610 ± 54&lt;sup&gt;b&lt;/sup&gt;</td>
<td>542 ± 9&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1,029 ± 60&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ionic strength (mM)</td>
<td>70 ± 2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>70 ± 1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>66 ± 1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>106 ± 9&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a–d</sup>Values with different superscript letters within a row are significantly different (P < 0.05).

<sup>1</sup>SM = skim milk; UF = ultrafiltration concentrate; UFL = ultrafiltration concentrate supplemented with lactose; RO = reverse osmosis concentrate.

### Table 2. Composition (mean ± SD) of the soluble phase of skim milk and concentrates

<table>
<thead>
<tr>
<th>Item</th>
<th>SM</th>
<th>UF</th>
<th>UFL</th>
<th>RO</th>
</tr>
</thead>
<tbody>
<tr>
<td>Casein (%)</td>
<td>0.15 ± 0.04&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.26 ± 0.05&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.30 ± 0.10&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.29 ± 0.06&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ash (%)</td>
<td>0.46 ± 0.01&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.45 ± 0.02&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.40 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.89 ± 0.05&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ca (% of total Ca)</td>
<td>37.7 ± 2.5&lt;sup&gt;b&lt;/sup&gt;</td>
<td>20.8 ± 0.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>21.0 ± 1.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>22.4 ± 2.8&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Mg (% of total Mg)</td>
<td>69.4 ± 4.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>53.0 ± 0.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>53.3 ± 2.2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>54.5 ± 7.2&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Na (% of total Na)</td>
<td>93.3 ± 6.1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>82.8 ± 15.4&lt;sup&gt;d&lt;/sup&gt;</td>
<td>75.6 ± 9.2&lt;sup&gt;b&lt;/sup&gt;</td>
<td>78.8 ± 6.5&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>P (% of total P)</td>
<td>48.8 ± 1.1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>28.9 ± 1.4&lt;sup&gt;b&lt;/sup&gt;</td>
<td>26.2 ± 0.8&lt;sup&gt;a&lt;/sup&gt;</td>
<td>32.0 ± 4.2&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Micellar calcium (mg of Ca/g of casein)</td>
<td>25.6 ± 1.0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>29.8 ± 0.1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>27.5 ± 1.6&lt;sup&gt;b&lt;/sup&gt;</td>
<td>36.2 ± 1.3&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a–c</sup>Values with different superscript letters within a row are significantly different (P < 0.05).

<sup>1</sup>SM = skim milk; UF = ultrafiltration concentrate; UFL = ultrafiltration concentrate supplemented with lactose; RO = reverse osmosis concentrate.

### Rennet Coagulation Properties

The coagulation profiles of SM and concentrates are presented in Figure 1, and the coagulation parameters are reported in Table 3. The formation of rennet gel was faster in UF concentrate, with a T<sub>lag</sub> of 4.7 min, compared with 8.6 min for SM (P < 0.05). The influence of UF concentration of milk on rennet coagulation time has proportion of soluble multivalent ions than for SM. On average, the proportions of calcium, magnesium, and phosphorus in the soluble phase of concentrates were respectively 45, 22, and 38% lower than in SM. The micellar calcium concentration (milligrams per gram of casein) was higher in UF concentrate than in SM (P < 0.05).
been widely studied, and some authors have found that coagulation time was reduced (Guinee and Mulholland, 1994; Upreti et al., 2011; Soodam and Guinee, 2018), whereas others found no effect (Guinee et al., 1997; Waungana et al., 1998; Sandra et al., 2011). However, concentrating milk by RO increased Tlag (12.07 min) compared with SM and UF concentrate (\(P < 0.05\)). This is in agreement with our previous study (Lauzin et al., 2018), where increased Tlag was attributed to high ionic strength and viscosity of RO concentrate.

The maximal curd firming rate (Vmax) was also affected by milk concentration. Concentrate obtained via UF is known to have a higher Vmax than SM due to a crowding effect, resulting in higher collision frequency and an increased number of bonds between casein micelles (Sandra et al., 2011). We found that RO concentrate showed higher Vmax than SM (\(P < 0.05\)) but lower Vmax than UF concentrate (\(P < 0.05\)). The casein content in UF and RO concentrates were similar, but the higher viscosity of RO could be responsible for slower micelle diffusion, resulting in a lower Vmax (Karlsson et al., 2007a). As previously shown (Table 2), the micellar calcium concentration in RO concentrate was significantly higher than in SM or in UF concentrate, which could also contribute to reduced curd firming rate. According to Malacarne et al. (2014), excessive content of micellar calcium could reduce the number of phosphate groups available for curd formation in the secondary phase of rennet coagulation. The lower curd firming rate for RO concentrate could represent an advantage over UF concentrate, because it increases the cutting window. The cutting window corresponds to the period during which the curd firmness is appropriate for cutting and was defined as the time between storage modulus values of 35 and 70 Pa (Panthi et al., 2019). As shown in Figure 1, the cutting window was 25% longer for RO concentrate than for UF concentrate. The gel firmness 30 min after coagulation (\(G'_{Tlag+30}\)) was higher for the concentrates than for SM (\(P < 0.05\)), due to the increased number of bonds between casein micelles and the higher casein volume fraction (Sandra et al., 2011). However, curd from RO concentrate showed lower \(G'_{Tlag+30}\) (341 Pa) than did UF concentrate (431 Pa).

The addition of lactose increased Tlag of UF concentrate from 4.7 to 9.9 min and reduced the Vmax from 26 to 19 Pa/min (\(P < 0.05\)). The lactose concentration in UFL was twice that in UF concentrate (Table 1), and according to Schorsch et al. (2002) and Famelart (1994), increasing the concentration of sugar retards both the enzymic and the aggregation steps during rennet-induced coagulation of casein micelles. It has been suggested that sugar reduces solvent quality and promotes the collapse of κ-casein molecules onto the casein micelle surface, reducing both the accessibility to chymosin and the rate of paracasein aggregation (Schorsch et al., 2002). These authors also observed lower curd firmness with increasing lactose concentration. Interestingly, the coagulation profile (Figure 1) and coagulation parameters (Table 3) of UFL concentrate fell between those of UF and RO concentrates. Lactose was, then, only partly responsible for the difference in coagulation kinetics between UF and RO concentrates. As we see in Table 3, the contribution of lactose to the differences for Tlag, Vmax, and \(G'_{Tlag+30}\) between UF and RO concentrates.

![Figure 1. Elastic moduli (G') during renneting of skim milk (SM), ultrafiltration concentrate (UF), ultrafiltration concentrate supplemented with lactose (UFL), and reverse osmosis concentrate (RO).](image)

Table 3. Rennet coagulation parameters\(^1\) (mean ± SD) of skim milk and concentrates\(^2\)

<table>
<thead>
<tr>
<th>Item</th>
<th>SM</th>
<th>UF</th>
<th>UFL</th>
<th>RO</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tlag (min)</td>
<td>8.59 ± 0.95(^a)</td>
<td>4.74 ± 0.81(^a)</td>
<td>9.93 ± 0.92(^ac)</td>
<td>12.07 ± 0.21(^c)</td>
</tr>
<tr>
<td>Vmax (Pa/min)</td>
<td>5.4 ± 0.8(^a)</td>
<td>26.4 ± 0.5(^a)</td>
<td>19.0 ± 1.4(^c)</td>
<td>16.4 ± 0.8(^b)</td>
</tr>
<tr>
<td>(G'_{Tlag+30}) (Pa)</td>
<td>95 ± 3(^a)</td>
<td>431 ± 33(^c)</td>
<td>381 ± 34(^b)</td>
<td>341 ± 19(^b)</td>
</tr>
</tbody>
</table>

\(^a\)Values with different superscript letters within a row are significantly different (\(P < 0.05\)).

\(^1\)Tlag = coagulation time; Vmax = maximum firming rate; \(G'_{Tlag+30}\) = elastic modulus (G’) 30 min after the start of coagulation.

\(^2\)SM = skim milk; UF = ultrafiltration concentrate; UFL = ultrafiltration concentrate supplemented with lactose; RO = reverse osmosis concentrate.
RO concentrates was estimated at 71, 74, and 56%, respectively. This suggests that the high mineral content in RO concentrate also affects the rennet-induced coagulation kinetics of casein micelles, but to a lower extent than lactose.

Model Cheeses

Small cheeses were produced from SM and concentrates at a laboratory scale. Cheese yield, protein and fat retention coefficients, and cheese composition are reported in Table 4. The moisture content of cheese produced from skim milk was 46.1% and cheese yield, adjusted at 50% moisture, was 11.2%. Milk concentration by UF increased cheese yield by a factor of 2.11, which roughly corresponds to the UF concentration factor. In comparison, milk concentration by RO increased cheese yield by a factor of 2.26, corresponding to a 7% relative increase relative to UF concentrate ($P < 0.05$). Similar protein and fat retention coefficients were observed in cheeses made from UF and RO concentrates, and the higher yield for RO cheese was attributed to the high concentration of solids in the aqueous phase of cheese. As shown in Table 4, RO cheese contained twice the amount of other solids compared with UF cheese. Adding lactose to UF concentrate increased cheese yield to a value similar to that of RO concentrate, confirming the significant contribution of lactose to cheese yield. Milk concentration by UF or RO had no significant effect on fat retention ($P > 0.05$) but slightly increased protein retention ($P < 0.05$). Interestingly, a higher protein retention coefficient was observed in UFL cheese than in UF or RO cheeses. Schorsch et al. (2002) suggested that sugar reduces solvent quality, which could prevent casein dissociation or caseinomacropeptide release in cheese whey. However, despite high lactose concentration in RO cheese, its protein retention coefficient was lower than that of UFL cheese. This result suggests that high mineral concentration in RO concentrate could counteract the positive effect of lactose on protein retention in cheese.

Milk concentration by ultrafiltration had no significant effect on cheese moisture, but the moisture content of RO cheese was significantly higher (50.6% vs. 47.2%; $P < 0.05$). The increase of cheese moisture cannot be attributed to the higher lactose concentration in RO concentrate, because the moisture content of UFL cheese was similar to that of SM or UF cheeses ($P > 0.05$). As suggested by Fagan et al. (2017), the high mineral concentration is likely responsible for higher moisture content in cheese. According to Malacarne et al. (2014), excessive mineral concentration can reduce the number of phosphate groups available for curd formation and may reduce curd contraction and syneresis during cooking. The protein-to-fat ratio was similar in UF, UFL, and RO cheeses ($P > 0.05$) and averaged 1.20 ± 0.02. However, it was lower in SM cheese, as expected from the lower protein retention coefficient (Table 4). The main difference among cheese compositions was the proportion of other solids, essentially composed of lactose and ash. Ash content was slightly higher in UF cheese than in SM cheese ($P < 0.05$). Because for all cheeses, whey was drained at the same pH (6.2), a lower proportion colloidal minerals could be solubilized when cheese was made from UF concentrate. As expected, the highest ash content was observed in RO cheese. The total milk minerals were concentrated by RO, and only a small proportion was released in whey at pH 6.2. The estimated lactose concentration (corresponding to the difference between other solids and ash) increased from approximately 3.0% in SM and UF cheeses to 8.9% in UFL and RO cheeses (Table 4). As previously men-

| Table 4. Moisture-adjusted yield,1 retention coefficients, and composition (mean ± SD) of model cheeses made from skim milk and concentrates2 |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| Item            | SM              | UF              | UFL             | RO              |
| Yield and recovery                                  |
| Moisture-adjusted yield (%)                        | 11.18 ± 0.21a   | 23.62 ± 0.12b   | 25.73 ± 0.13c   | 25.32 ± 0.34c   |
| Protein retention coefficient                      | 0.736 ± 0.005a  | 0.771 ± 0.004b  | 0.788 ± 0.002c  | 0.763 ± 0.007c  |
| Fat retention coefficient                           | 0.923 ± 0.017   | 0.935 ± 0.012a  | 0.934 ± 0.005b  | 0.922 ± 0.008a  |
| Cheese composition                                 |
| Moisture (%; 19.6/wt)                              | 46.1 ± 1.0a     | 47.2 ± 1.5a     | 47.7 ± 2.0a     | 50.6 ± 1.4b     |
| Protein (%; dry basis)                             | 47.28 ± 0.28b   | 49.98 ± 0.49c   | 47.20 ± 0.56b   | 45.82 ± 0.48a   |
| Fat (%; dry basis)                                 | 45.00 ± 0.33c   | 42.24 ± 0.30b   | 39.00 ± 0.51a   | 38.23 ± 0.98a   |
| Protein-to-fat ratio                                | 1.05 ± 0.01a    | 1.18 ± 0.02a    | 1.21 ± 0.01b    | 1.20 ± 0.02b    |
| Other solids (%; dry basis)                         | 7.72 ± 0.17a    | 7.78 ± 0.53a    | 13.8 ± 1.3b     | 15.9 ± 1.4c     |
| Ash (%; dry basis)                                 | 4.15 ± 0.50a    | 5.30 ± 0.06b    | 4.90 ± 0.07ab   | 7.02 ± 0.21c    |

1Yield adjusted at 50% moisture.
2SM = skim milk; UF = ultrafiltration concentrate; UFL = ultrafiltration concentrate supplemented with lactose; RO = reverse osmosis concentrate.

a–cValues with a different superscript letter within a row are significantly different ($P < 0.05$).
tioned, lactose retention in RO cheese is responsible for the significant increase of cheese yield. Indeed, cheese with high lactose content is prone to post-acidification, which could result in texture and flavor defects (Moynihan et al., 2016). However, for specific applications, such as thermally treated or direct-acid cheeses, the use of RO concentrate could be a viable approach to increase cheese yield.

**Cheese Microstructure**

The protein networks of cheeses made from SM and concentrates were observed via scanning electron microscope, and representative micrographs are presented in Figure 2. We found that SM cheese was characterized by a continuous protein network, with homogeneously distributed void spaces corresponding to fat droplets and serum pockets removed during sample preparation (Figure 2a). The protein network was quite uniform, but individual casein micelles, forming aggregates, could be observed. In cheese made from UF concentrate (Figure 2b), larger void spaces and increased coarseness were observed. According to Green et al. (1981), increasing protein concentration via UF alters the initial curd-forming process, which determines cheese microstructure. The size of void spaces and coarseness were further increased in RO cheese (Figure 2d). The presence of aggregated casein also increased, and the individual micelles were smaller. Water in the cheese exists in different forms, and during the first weeks of storage, the bulk water decreases and the water bound to casein increases, due to casein swelling and hydration (McMahon et al., 1999). Despite higher moisture content in RO cheese (Table 4), the protein network appeared dehydrated, suggesting a higher proportion of bulk water, held within larger serum pockets. The swelling of casein micelles seemed to be very limited in cheeses made from RO concentrate.

**Figure 2.** Scanning electron microscope images of cheese made from (a) skim milk, (b) UF concentrate, (c) UF concentrate supplemented with lactose, and (d) reverse osmosis concentrate. Bar = 5 μm.
The addition of lactose to UF concentrate had little effect on cheese microstructure (Figure 2c), and the size of void spaces and coarseness were similar to those observed for UF cheese. The dense protein network and larger void spaces observed in RO cheese are therefore attributable to its higher mineral concentration, which promoted casein–casein interactions and prevented swelling and hydration (Joshi et al., 2004). Excessive mineralization was shown to reduce casein micelle voluminosity (De Kort et al., 2011) and is likely responsible for the lower solvation of the protein network in RO cheese. The microstructural characteristics of cheese made from RO concentrate may negatively affect the texture; to improve the quality of RO cheese, the cheese-making process would require adaptations to promote demineralization.

CONCLUSIONS

Making cheese from RO concentrate increased moisture-adjusted cheese yield by 7%, relative to UF concentrate, due to the high lactose concentration in cheese. The kinetics of curd formation were slower in RO than in UF concentrate, and, again, lactose was mainly responsible for the difference. As a consequence, the curd cutting window was increased by 20% in RO concentrate. Cheese moisture was about 3% higher in RO cheese than in UF cheese, but the difference could not be attributed to lactose. Higher mineral concentration increased the size of serum pockets in RO cheese microstructure, which could explain the higher moisture content. Similar protein and fat retention coefficients were observed in cheeses made from RO and UF concentrates, leading to cheese compositions with the same protein-to-fat ratio. The use of RO concentrate is a viable approach to increase cheese yield. However, this approach is limited to specific applications, where post-acidification can be controlled. Our results suggest that the quality of RO cheeses could be improved by promoting demineralization during the cheese-making process.

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