

## Volumetric Study Of Monosaccharides (D-Ribose And D-Mannose)-Saccharin Sodium Salt In Aqueous Solutions At $T = 298.15$ K.

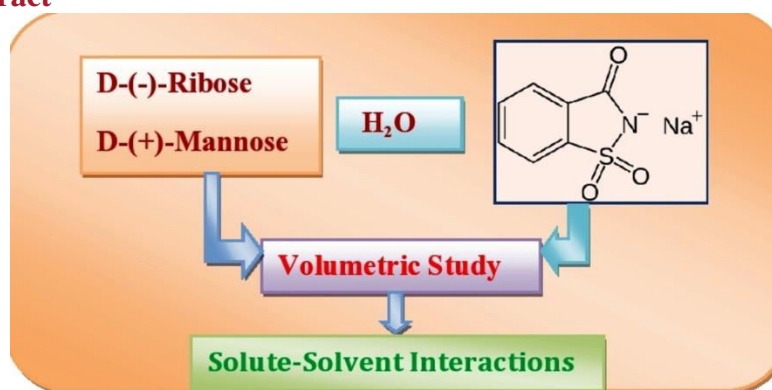
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### Graphical abstract



**ABSTRACT:** Using a volumetric approach, monosaccharides were investigated in terms of temperature and different concentrations. At  $T = 298.15$  K, the densities of D(-)-ribose, D(+)-mannose in H<sub>2</sub>O and in aqueous saccharin Na salt with molality,  $m = (0.05, 0.15, \text{ and } 0.3) \text{ mol} \cdot \text{kg}^{-1}$  were determined. The apparent molar volumes ( $V_\phi$ ), partial molar volumes ( $V_\phi^0$ ), and Masson's coefficient ( $S_v$ ) were evaluated by means of experimental density ( $\rho$ ) values. The data was further analysed in terms of transfer characteristics ( $\Delta_{trs}V_\phi^0$ ), apparent specific volumes (ASV), and volumetric interaction coefficients ( $V_{AB}, V_{ABB}$  using McMillan–Mayer theory). The significant interactions between the hydrophilic groups of solute (D-ribose and D-mannose) and the Na<sup>+</sup> ion of cosolute (saccharin sodium salt) at varied concentrations have been reported. Studied saccharide was found to have a sweet taste in mixed aqueous sodium saccharin.

**Keywords:** Apparent specific volume, Density, Monosaccharide, Partial molar volume, Sodium saccharin.

### INTRODUCTION

Recent work has examined the nature and degree of intermolecular interaction in aqueous solutions of combinations using thermophysical characteristics. Depending on the number of saccharide units present, sugars are divided into four different types. Simple sugars like glucose, fructose, galactose, mannose, ribose, xylose, arabinose, etc are called monosaccharides. Carbohydrates are the most prominent source of energy consumed by humans<sup>1</sup>. It also performs an important role in the regulation of living creatures throughout biologically based cycles<sup>2,3</sup>. The hydration abilities of saccharides are due to their hydrophilic hydroxyl groups (-OH)<sup>4,5</sup>. Sugar is important in confections and medicine because it not only adds sweetness but also improves the texture of foods, improves taste, exerts stability, and

thus protects vitamins during processing and storage. Saccharides in H<sub>2</sub>O electrolyte arrangements play an important role in a number of scientific fields<sup>6-10</sup>, including biology, pharmaceutical preparations, catalysis, climate, and food preparation. Recent data from the literature suggests that saccharide-metal ion blends have been extensively studied, further interpreted as interactions among solute-solute and solute-solvent<sup>11-14</sup>.

Today, a range of new sweeteners are available as a substitute for natural sugars. Many man-made sweeteners<sup>15-20</sup> are available, of which Aspartame, Sucralose, Saccharin Na salt, and Acesulfame K are approved by the "Food and Drug Administration" (FDA) for use in the food and pharmaceutical industry. Blending sweeteners<sup>21-23</sup> with sugars is to reduce the amount used, reduce the cost, and improve the taste. Molecular interactions between sugar and the receptor form via H<sub>2</sub>O molecules that surround them. The conduct of metal ions with saccharides in aqueous solutions is useful to comprehend solute-solvent (sugar-H<sub>2</sub>O) and solute-cosolute (sugar-saccharin) interactions along with taste.

The D-ribose and D-mannose (0.04-0.20) *m* densities were evaluated at *T* = 298.15 K in H<sub>2</sub>O and in aqueous saccharin Na salt with molalities of 0.05, 0.15, and 0.3 *m*. The interactions between saccharide (D-ribose and D-mannose)-saccharin sodium salt in an aqueous media were discovered through partial molar volumes, transfer characteristics, interaction coefficients, and apparent specific volumes.

## Materials and Methods

D(-)-ribose, D(+)-mannose, and saccharin Na salt were purchased with a purity of 99.0% from Sigma and used.

Aqueous solutions of solute and cosolute were prepared using triply distilled freshly prepared water and a stoppered airtight glass bottle. Similarly, measurements were done on a weight-by-weight (w/w) scale with an accuracy of ± 0.0001 g on a Dhona balance. H<sub>2</sub>O is used as a solvent and saccharide is used as a solute in both binary and ternary solutions with saccharin Na salt as a stock solution for subsequent usage.

The densities ( $\rho$ ) of D(-)-ribose and D(+)-mannose in H<sub>2</sub>O and in aqueous saccharin Na salt were measured experimentally using a bi-capillary Pycnometer<sup>24-29</sup> at *T* = 298.15 K. To submerge the pycnometer in a vertical position, a glass-walled water bath with a constant temperature of ± 0.01 K was utilized. At the studied temperature, the Pycnometer was calibrated by using organic solvents like ethyl acetate, carbon tetrachloride, and acetophenone. When the outcomes were compared to the reported results, it was revealed that they were in good agreement. Solvent density at the studied temperature was taken from published data<sup>30,31</sup>. The investigational variability in density determinations was 0.2 kg·m<sup>-3</sup>. The density of studied systems (saccharide with stock solution sodium saccharin) was investigated using the same method.

## Results and Discussion

### Volumetric study of D(-)-Ribose and D(+)-Mannose in aqueous solutions of saccharin sodium.

#### Apparent molar volume

At *T* = 298.15 K, the thermophysical characteristics and parameters of saccharides in aqueous sodium saccharin are studied. The apparent molar volumes ( $V_\phi$ ) of the studied saccharide in H<sub>2</sub>O and in saccharin Na were calculated with the equation<sup>5,32</sup>.

$$V_\phi = M/\rho - (\rho - \rho_o)/(m\rho\rho_o) \quad 1$$

In this equation, apparent molar volume is represented by  $V_\phi$ . Similarly, the density of a solution is denoted by  $\rho$  and its molality is denoted by  $m$ . The molar mass of a solute is also indicated by  $M$ .

The  $\rho$ , and  $V_\phi$  of D(-)-ribose, and D(+)-mannose in  $H_2O$  and in aqueous saccharin Na salt,  $m = (0.05, 0.15, \text{ and } 0.3) \text{ mol}\cdot\text{kg}^{-1}$  at  $T = 298.15 \text{ K}$  are shown in Table 1 and 2, respectively. The results show that, observed densities and apparent molar volumes depend on concentration and vary linearly with solute and cosolute concentrations.

By the least-square fit approach and Masson's equation<sup>33,34</sup>, the  $V_\phi$  values of binary and ternary mixtures are correlated with molality as shown below:

$$V_\phi = V_\phi^0 + S_v \cdot m \quad 2$$

The equation (2) is in the form  $y = mx + c$ , with an intercept,  $V_\phi^0$  (partial molar volume), and a slope,  $S_v$  (Masson's coefficient).

The results of  $V_\phi^0$  are summarized in Table 3, while variations of  $V_\phi$  of D-ribose, and D-mannose in  $H_2O$ , and in aqueous saccharin sodium with molalities,  $m = (0.05, 0.15, \text{ and } 0.3) \text{ mol}\cdot\text{kg}^{-1}$  are represented in Fig. 1, and 2. The reported values of  $V_\phi^0$  ( $\text{m}^3\cdot\text{mol}^{-1}$ ) of D-mannose in  $H_2O$  at  $T = 298.15 \text{ K}$  are  $(111.30^4, 111.7^{35,36}, \text{ and } 111.72^{37})\cdot 10^{-6}$ . Table 3 shows the  $V_\phi^0$  estimated at  $T = 298.15 \text{ K}$ , from the present study is  $(111.7)\cdot 10^{-6}/(\text{m}^3\cdot\text{mol}^{-1})$ . Similarly, for D-ribose the reported values at 298.15 K are  $(95.26^{38}, \text{ and } 95.3^{39,40})\cdot 10^{-6}$  while the calculated value at the same temperature is  $(95.3)\cdot 10^{-6}/(\text{m}^3\cdot\text{mol}^{-1})$ . The  $V_\phi^0$  of D-ribose and D-mannose in  $H_2O$  are in good accord with published figures in the literature at the investigated temperature.

The positive  $V_\phi^0$  values offer significant solute-solvent interactions for molality,  $m = (0.05, 0.15 \text{ and } 0.3)$  of the saccharide-saccharin sodium salt system<sup>41,42</sup>. Additionally, the Masson's coefficients ( $S_v$ ) are positive but less than the values of  $V_\phi^0$ . It shows that solute-solute interactions<sup>43</sup> are much lower than those between solute and solvent.

**Table 1: Densities ( $\rho/\text{kg}\cdot\text{m}^{-3}$ ), and apparent molar volumes ( $V_\phi/\text{m}^3\cdot\text{mol}^{-1}$ ) of monosaccharides in  $H_2O$  at  $T = 298.15 \text{ K}$ .**

D(-)-Ribose			D(+)-Mannose		
$m$ ( $\text{mol}\cdot\text{kg}^{-1}$ )	$\rho$ ( $\text{kg}\cdot\text{m}^{-3}$ )	$V_\phi$ $10^6/(\text{m}^3\cdot\text{mol}^{-1})$	$m$ ( $\text{mol}\cdot\text{kg}^{-1}$ )	$\rho$ ( $\text{kg}\cdot\text{m}^{-3}$ )	$V_\phi$ $10^6/(\text{m}^3\cdot\text{mol}^{-1})$
0.0000	997.1	95.3	0.0000	997.1	
0.0431	999.4	95.4	0.0372	999.6	111.8
0.0814	1002	95.4	0.0805	1002	111.9
0.1208	1004	95.5	0.1210	1005	112.0
0.1611	1006	95.5	0.1603	1008	112.1
0.2076	1008	95.3	0.2056	1011	112.3

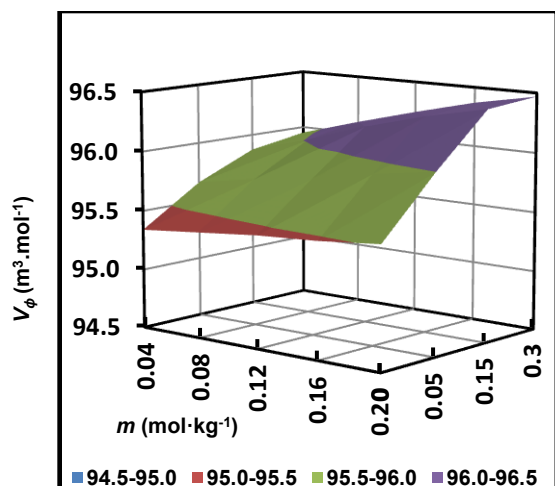
**Table 2: Densities ( $\rho/\text{kg}\cdot\text{m}^{-3}$ ), and apparent molar volumes ( $V_\phi/\text{m}^3\cdot\text{mol}^{-1}$ ) of monosaccharides in aqueous saccharin Na salt at  $T = 298.15 \text{ K}$ .**

$m$ (mol.kg <sup>-1</sup> )	$\rho$ (kg.m <sup>-3</sup> )	$V_\phi \cdot 10^6$ (m <sup>3</sup> .mol <sup>-1</sup> )	$m$ (mol.kg <sup>-1</sup> )	$\rho$ (kg.m <sup>-3</sup> )	$V_\phi \cdot 10^6$ (m <sup>3</sup> .mol <sup>-1</sup> )	$m$ (mol.kg <sup>-1</sup> )	$\rho$ (kg.m <sup>-3</sup> )	$V_\phi \cdot 10^6$ (m <sup>3</sup> .mol <sup>-1</sup> )
	0.05			0.15			0.3	
D(-)-Ribose + Sodium saccharin								
0.0000	1001		0.0000	1009		0.0000	1022	
0.0370	1004	95.7	0.0399	1011	95.8	0.0385	1024	95.9
0.0798	1006	95.7	0.0798	1013	96.0	0.0809	1026	96.1
0.1199	1008	95.8	0.1287	1016	96.2	0.1236	1028	96.2
0.1598	1010	95.9	0.1663	1018	96.3	0.1674	1031	96.4
0.1998	1012	96.0	0.2079	1020	96.4	0.2013	1032	96.5
D(+)-Mannose + Sodium saccharin								
0.0000	1002		0.0000	1008		0.0000	1022	
0.0395	1004	112.3	0.0421	1011	112.4	0.0409	1024	112.8
0.0790	1007	112.4	0.0810	1013	112.5	0.0812	1027	112.9
0.1190	1010	112.6	0.1237	1016	112.6	0.1199	1030	113.1
0.1597	1012	112.7	0.1662	1019	112.7	0.1632	1032	113.2
0.1901	1014	112.8	0.2133	1022	112.9	0.2010	1034	113.4

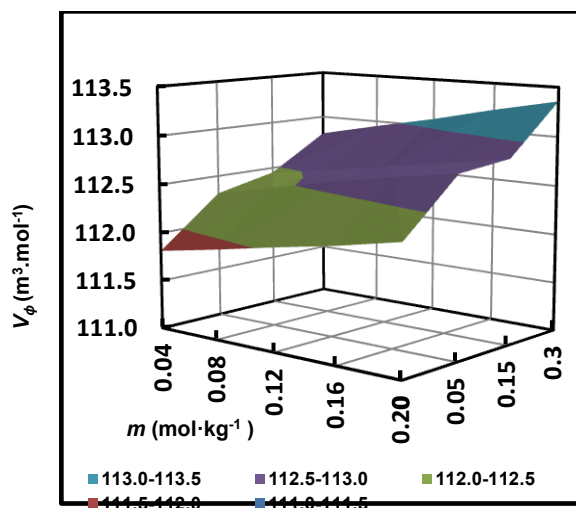
**Table 3:** ( $V_\phi^0$ ), ( $S_v$ ), ASV, and ( $\Delta_{trs}V_\phi^0$ ) of monosaccharides in H<sub>2</sub>O, and in aqueous saccharin Na salt at  $T = 298.15$  K.

System	Parameters			
	$V_\phi^0$ ·10 <sup>6</sup> /(m <sup>3</sup> .m ol <sup>-1</sup> )	$\cdot 10^6 S_v$ /(m <sup>3</sup> .kg.m ol <sup>-2</sup> )	ASV ·10 <sup>6</sup> /(m <sup>3</sup> .k g <sup>-1</sup> )	( $\Delta_{trs}V_\phi^0$ ) ·10 <sup>6</sup> /(m <sup>3</sup> .m ol <sup>-1</sup> )
D(-)-Ribose + H <sub>2</sub> O	95.3	1.1	0.63	
D(+)-Mannose + H <sub>2</sub> O	111.7	2.78	0.62	
D(-)-Ribose + 0.05 m Sodium saccharin	95.6	2.09	0.64	0.28
D(-)-Ribose + 0.15 m Sodium saccharin	95.7	3.40	0.64	0.44
D(-)-Ribose + 0.3 m Sodium saccharin	95.8	3.21	0.64	0.53

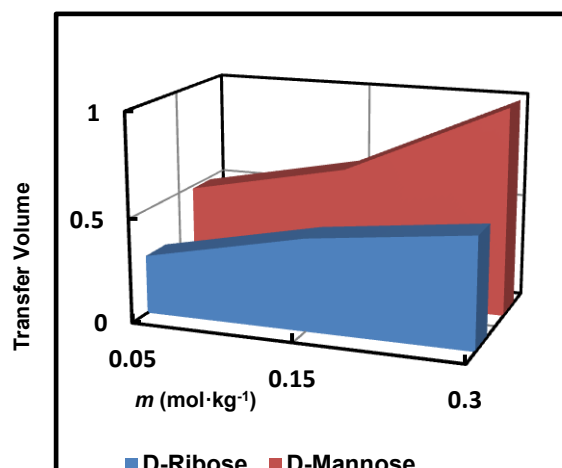
D(+)-Mannose + 0.05 <i>m</i> Sodium saccharin	112.2	3.23	0.62	0.49
D(+)-Mannose + 0.15 <i>m</i> Sodium saccharin	112.4	2.41	0.62	0.64
D(+)-Mannose + 0.3 <i>m</i> Sodium saccharin	112.7	3.41	0.62	0.97



**Figure 1:** Graph of  $V_{\phi}$  of D(-)-ribose in  $H_2O$  and in saccharin Na salt with molality,  $m = (0.05, 0.15, 0.3) \text{ mol}\cdot\text{kg}^{-1}$  at 298.15 K.



**Figure 2:** Graph of  $V_{\phi}$  of D(+)-mannose in  $H_2O$  and in saccharin Na salt with molality,  $m = (0.05, 0.15, 0.3) \text{ mol}\cdot\text{kg}^{-1}$  at 298.15 K.



**Figure 3:** Partial molar volume of transfer,  $(\Delta_{trs}V_{\phi}^0)$  of D(-)-ribose and D(+)-mannose in saccharin Na salt with molality,  $m = (0.05, 0.15, 0.3)$  mol·kg<sup>-1</sup> at 298.15 K.

**Transfer volume**

Using the following relation<sup>40</sup>, the transfer volume of  $(\Delta_{trs}V_{\phi}^0)$  monosaccharides from H<sub>2</sub>O to aqueous saccharin Na Salt was obtained for the investigated systems at infinite dilution.

$$\Delta_{trs}V_{\phi}^0 = V_{\phi}^0(\text{in sodium saccharin}) - V_{\phi}^0(\text{in H}_2\text{O}) \tag{6}$$

Table 3 shows the D-ribose and D-mannose  $V_{\phi}^0$  values that can be used to determine  $(\Delta_{trs}V_{\phi}^0)$  from H<sub>2</sub>O to aqueous saccharin sodium with molality  $m$ . The  $\Delta_{trs}V_{\phi}^0$  is positive at infinite dilution, often used to examine solute-cosolute interactions in aqueous medium<sup>43</sup> and increases with the concentration of cosolute<sup>36</sup>. Fig. 3 shows the variations in the transfer volume at infinite dilution  $(\Delta_{trs}V_{\phi}^0)$  with molalities,  $m$  of saccharin Na at experimental temperature.

In the present study, two possible types of interaction happening among monosaccharide (D-ribose and D-mannose) and aqueous saccharin sodium in a ternary system could be categorized as follows<sup>43</sup>:

- I) Hydrophilic-ionic interactions among monosaccharide hydrophilic groups (-C=O, -OH and -O-), and the Na<sup>+</sup> ion of saccharin Na salt.
- II) Hydrophobic-ionic interactions among monosaccharide and cosolute ion.

In accordance with the "co-sphere overlap model"<sup>44</sup>, type (I) interactions positively contribute to the  $\Delta_{trs}V_{\phi}^0$  whereas type (II) interactions negatively contribute to the  $\Delta_{trs}V_{\phi}^0$ . Kumar et al.<sup>45</sup> reported similar results.

**Volumetric Interaction Coefficients**

Kozak et al<sup>46,47</sup> established the "McMillan-Mayer theory" of solutions in order to study volumetric interaction coefficients. On the basis of this, Krishnan et al.<sup>48</sup> explored the interactions of solutes and cosolutes in solvation spheres. Numerous researchers have investigated interactions in aqueous solutions using the theory<sup>49-51</sup>.

The transfer volume is also related as:

$$\Delta_{trs}V_{\phi}^0 = 2V_{AB}m_B + 3V_{ABB}m_B^2 + \dots \tag{7}$$

D-ribose and D-mannose (solutes) are represented by A, while saccharin sodium (co-solute) is represented by B. These parameters were evaluated by analyzing investigational



figures and using the least-squares approach to equation 7. The calculated values of  $V_{AB}$  for D-ribose and D-mannose are  $(2.34, \text{ and } 3.38) \cdot 10^{-6}/(\text{m}^3 \cdot \text{mol}^{-2} \cdot \text{kg})$ , whereas those for  $V_{ABB}$  are  $(-3.25, \text{ and } -4.01) \cdot 10^{-6}/(\text{m}^3 \cdot \text{mol}^3 \cdot \text{kg}^2)$ , respectively, at  $T = 298.15 \text{ K}$ . Positive figures are contributed by the doublet interaction parameter ( $V_{AB}$ ), whereas negative figures are contributed by the triplet interaction parameter<sup>43</sup> ( $V_{ABB}$ ). Positive  $V_{AB}$  figures illustrate that, monosaccharide (D-ribose and D-mannose), and saccharin Na have strong synergy. Negative figures of  $V_{ABB}$  signify the nonexistence of D-ribose/mannose-saccharin-saccharin relations. Jiang and coworkers<sup>52</sup> noted a similar observation for ternary systems at 298.15 K.

The "Group Additivity Model" of Savage and Wood<sup>53</sup> examines the four forms of pair interactions involving saccharin Na (electrolyte), and D-ribose/mannose (non-electrolyte). The  $\text{Na}^+ - \text{R}$  (R is an alkyl group) makes a small negative input (type I interaction) to  $V_{AB}$  and Type II interaction of Anion - R contributes negatively to  $V_{AB}$  also Type III interaction between  $\text{Na}^+ - \text{O}$  is more dominant and contributes positively to  $V_{AB}$  (-O, hydrophilic groups in non-electrolyte) whereas type IV interaction of anion - O, contributes negatively to  $V_{AB}$  but smaller than type I (hydrophobic groups).

In an aqueous medium, the complete dissociation of ions of saccharin Na takes place. The pair interaction coefficient of sodium ion with hydrophilic groups of monosaccharides (-OH, C=O, and -O-) is positive. Both theories anticipate that interactions linking solutes (D-ribose and D-mannose) and cosolute (saccharin sodium) are taking place.

#### Apparent Specific Volume (ASV)

It is helpful for getting data with respect to the taste conduct of sugars in blended aqueous sodium saccharin solutions. The aqueous solutions are classified as salty, sweet, bitter, and sour with respect to taste by Shamil and Birch<sup>54</sup>. The ASV range suggested by Parke et al<sup>55</sup>, for sweet molecules is from  $(0.51 \text{ to } 0.71) \cdot 10^{-6}/(\text{m}^3 \cdot \text{kg}^{-1})$  with an ideal value of  $(0.618) \cdot 10^{-6}/(\text{m}^3 \cdot \text{kg}^{-1})$  at the centre. The equation (8) is used to calculate the ASV of solutes in solvent and cosolute.

$$\text{ASV} = V_{\phi}^0 / M \quad 8$$

The solute's molar mass is denoted by M and its partial molar volume is denoted by  $V_{\phi}^0$ . Table 3 shows the ASV values of monosaccharide in  $\text{H}_2\text{O}$ , and saccharin sodium at the examined concentrations and temperature. At experimental temperature, the ASV values for ribose range from  $(0.635 \text{ to } 0.638) \cdot 10^{-6}/(\text{m}^3 \cdot \text{kg}^{-1})$ , while those for mannose range from  $(0.620 \text{ to } 0.625) \cdot 10^{-6}/(\text{m}^3 \cdot \text{kg}^{-1})$ . The saccharides examined in this study when incorporated into saccharin sodium stock solutions kept their sweetness.

#### Conclusions

Volumetric data of monosaccharides (D(-)-ribose, and D(+)-mannose) in  $\text{H}_2\text{O}$  and aqueous saccharin sodium salt at  $T = 298.15 \text{ K}$  are included in this study. The investigational data were utilized to compute thermodynamic parameters such as  $V_{\phi}$ ,  $V_{\phi}^0$ ,  $S_v$ ,  $\Delta_{trs}V_{\phi}^0$ , ASV, and ( $V_{AB}$ , and  $V_{ABB}$ ). The corresponding  $\Delta_{trs}V_{\phi}^0$  of ribose and mannose shift from  $\text{H}_2\text{O}$  to saccharin Na salt is positive, and its degree rises as concentration increases. Positive values of  $V_{AB}$  indicate the strong interactions between the solute (D-ribose and D-mannose) and the saccharin sodium. The ASV  $(0.623\text{-}0.638) \cdot 10^{-6}/(\text{m}^3 \cdot \text{kg}^{-1})$  figures suggested a sweet taste for studied saccharides in blended saccharin sodium.

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