

**This item is the archived peer-reviewed author-version of:**

The effect of the buffer solution on the adsorption and stability of horse heart myoglobin on commercial mesoporous titanium dioxide : a matter of the right choice

**Reference:**

Loreto Stefano, Cuypers Bert, Brokken Jacotte, Van Doorslaer Sabine, De Wael Karolien, Meynen Vera.- The effect of the buffer solution on the adsorption and stability of horse heart myoglobin on commercial mesoporous titanium dioxide : a matter of the right choice

Physical chemistry, chemical physics / Royal Society of Chemistry [London] - ISSN 1463-9076 - 19:21(2017), 13503

Full text (Publisher's DOI): <https://doi.org/10.1039/C6CP08585G>

To cite this reference: <http://hdl.handle.net/10067/1435140151162165141>



## Journal Name

## ARTICLE

## The effect of the buffer solution on the adsorption and stability of horse heart myoglobin on commercial mesoporous titanium dioxide: a matter of the right choice.

Received 00th January 20xx,  
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

www.rsc.org/

Stefano Loreto,<sup>a,c</sup> Bert Cuypers<sup>b</sup>, Jacotte Brokken<sup>a</sup>, Sabine Van Doorslaer<sup>b</sup>, Karolien De Wael<sup>c</sup> and Vera Meynen<sup>a\*</sup>

Despite the numerous studies on the adsorption of different proteins into mesoporous titanium dioxide and indications on the important role of buffer solutions in bioactivity, a systematic study on the impact of the buffer on the protein incorporation in porous substrates is still lacking. We here studied the interaction between a commercial mesoporous TiO<sub>2</sub> and three of the most used buffers for proteins incorporation, *i.e.* HEPES, Tris and phosphate buffer. In addition, this paper analyzes the adsorption of horse heart myoglobin (hhMb) onto commercial mesoporous TiO<sub>2</sub> as a model system to test the influence of buffers on the protein incorporation behavior in mesoporous TiO<sub>2</sub>. N<sub>2</sub> sorption analysis, FT-IR and TGA/DTG measurements were used to evaluate the interaction between the buffers and the TiO<sub>2</sub> surface, and the effect of such interaction on hhMb adsorption. Cyclic voltammetry (CV) and electron paramagnetic resonance (EPR) were used to detect changes in the microenvironment surrounding the heme. The three buffers show a completely different interaction with the TiO<sub>2</sub> surface, which drastically affects the adsorption of myoglobin as well as its structure and electrochemical activity. Therefore, special attention is required while choosing the buffer medium to avoid misguided evaluation of protein adsorption on mesoporous TiO<sub>2</sub>.

### 1. Introduction

In the last decades the interest towards the adsorption of proteins on a solid surface has tremendously increased<sup>1–6</sup>. The use of porous materials to incorporate biomolecules has attracted great attention because of their large surface area and tunable textural properties such as their pore size and volume. This permits to vary the amount of immobilized proteins and enzymes in a wide range of molecular sizes, making these materials suitable in the field of biosensors<sup>7,8</sup>, bio(electro)catalysis<sup>9–11</sup>, bioelectrochemistry<sup>12</sup> and drug delivery<sup>13–17</sup>. Among all these materials, mesoporous silica is frequently used for the incorporation of proteins in a wide range of applications because of its ease of synthesis resulting in materials with a highly

ordered and tunable mesoporous structure, narrow pore-size distribution, high pore volume and surface area<sup>18–25</sup>. Nevertheless, the electronic properties of silica, considerably limit its applicability in (bio)electrochemistry<sup>26</sup>. The development of well-structured non-siliceous metal oxides<sup>27,28</sup> such as Nb<sub>2</sub>O<sub>5</sub><sup>7</sup>, ZrO<sub>2</sub><sup>29,30</sup>, ZnO<sup>31</sup>, and TiO<sub>2</sub><sup>32,33</sup> has led to a huge increase in the use of these materials. Mesoporous TiO<sub>2</sub> is a promising substrate for the adsorption of biomolecules because of its bio and eco compatibility<sup>34–36</sup>. Its applications in electrochemistry<sup>37</sup> and biochemistry<sup>38</sup> are therefore continuously increasing. Despite the many investigations performed on the encapsulation of proteins and biomolecules onto mesoporous metal oxide, there are only few studies describing the effect of the buffer solution on the adsorption. Parkes et al.<sup>39</sup> studied the effect of different buffers on the adsorption of bovine serum albumin (BSA), showing how the pH greatly affects the tribofilm formation. Wei et al.<sup>40</sup> and Moulton et al.<sup>41</sup> characterized the effect of the phosphate buffer (phosphate buffer saline, PBS) on the adsorption of proteins (immunoglobulin, BSA, fibrinogen and

<sup>a</sup> Department of Chemistry, University of Antwerp, 2610 Wilrijk, Belgium

<sup>b</sup> Department of Physics, University of Antwerp, 2610 Wilrijk, Belgium

<sup>c</sup> Department of Chemistry, University of Antwerp, 2010 Antwerpen, Belgium

Email: vera.meynen@uantwerpen.be

Electronic Supplementary Information (ESI) available. See DOI: 10.1039/x0xx00000x

## ARTICLE

## Journal Name

lysosome) on a germanium crystal surface and TiO<sub>2</sub> films respectively. They both concluded that the use of PBS decreases the adsorption of proteins. B. Fubini and co-workers<sup>42</sup> compared the use of 4-(2-hydroxyethyl)piperazine-1-ethanesulfonic acid (HEPES) and PBS for the adsorption of BSA on commercial non-porous TiO<sub>2</sub>. The results clearly indicate a great influence of the buffer choice on the surface properties of the titanium dioxide. To the best of our knowledge, there are no previous systematic studies on the impact of buffer choice on the adsorption of proteins on mesoporous TiO<sub>2</sub> (adsorption capacity and kinetics). This is, however, important as many applications involving the immobilization of biomolecules imply their encapsulation in porous systems. On the one hand, the stability of the biomolecule itself is largely dependent on the buffer solution. On the other hand, the control of the pH of the protein solution plays a key role in the adsorption process<sup>39</sup> and it is a crucial step in their incorporation. In fact, a higher amount of proteins can be adsorbed if the solution pH is close to the isoelectric point (IP) of the proteins and when the proteins and the mesoporous surface are electrostatically attractive<sup>43</sup>. Furthermore, one can also imagine that the interaction of the buffer solution with the surface will alter the surface properties of the mesoporous materials, influencing the protein adsorption and its correlated performance in application<sup>44</sup>.

Buffers play a dual role in protein incorporation: they should ensure protein stability and allow for an optimal sorption capacity and kinetics. In order to gain understanding in this role, we have investigated the impact of the interaction between a commercial mesoporous titanium dioxide (Millennium PC 500) and three of the most applied buffers in protein incorporation: HEPES, 2-amino-2-hydroxymethyl-propane-1,3-diol (Tris) and PBS.

Although those three buffers are being applied for the same intent, they have large differences in chemical structure. Therefore, they may interact in a different way with the TiO<sub>2</sub> surface, forming bonds with different stability and altering the surface properties to some extent. As a consequence, they may have different degrees of impact on the rate of protein adsorption in the mesoporous material as well as on the total capacity of protein loading and on the protein structure upon adsorption. To evaluate the impact of the buffer on the mesoporous TiO<sub>2</sub> surface, infrared (IR) and X-ray fluorescence

(XRF) spectroscopy, thermogravimetric analysis (TGA) and N<sub>2</sub> sorption have been employed. Horse heart myoglobin (hhMb) has been used as model protein to examine the influence of the three buffer solutions on protein adsorption. The protein incorporation has been monitored by UV-visible absorption spectroscopy of the supernatant and the successful incorporation into the pores has been confirmed by nitrogen sorption. The fate of the proteins after incorporation has been analyzed by IR spectroscopy and TGA. Electron paramagnetic resonance (EPR) spectroscopy was performed to monitor changes in the heme pocket of hhMb upon the adsorption of the protein. The electrochemical activity of hhMb adsorbed in the three buffers has been tested by cyclic voltammetry (CV). A better understanding of the interaction between biomolecules-buffer-titania, and thus a better control of the immobilization of proteins onto surfaces, is envisioned. In addition, knowledge of structural modification and activity changes due to such interactions, will avoid misinterpretation of biomolecule-adsorption results.

## 2. Experimental methods

### 2.1 Materials

Myoglobin from equine heart (≥90%, essentially salt-free, lyophilized powder), 4-(2-hydroxyethyl)piperazine-1-ethanesulfonic acid (HEPES buffer, ≥99%), 2-amino-2-hydroxymethyl-propane-1,3-diol (Tris buffer, ≥99.9%), phosphate buffer saline (PBS) and sodium hydroxide (ACS reagent, ≥97%) were all purchased from Sigma Aldrich and used without further purification. Titanium dioxide Millennium PC500 was obtained from Cristal Global. Non-porous titanium dioxide P25 was purchased from Degussa-Hüls GmbH.

Prior to use, the commercial titanium-dioxide material was pretreated to enhance its pore size, required for protein adsorption. Millennium PC500 was calcined as received at 350°C for 6h (1°C/min) in a Lenton chamber furnace in order to obtain an enlarged pore size<sup>45</sup>. The surface area and the pore size of the calcined and of the myoglobin-incorporated material were measured via N<sub>2</sub>-sorption. All buffers used in this work are aqueous solutions of the three buffers discussed above. Considering the PZC (point of zero charge) of titanium dioxide (~6.2)<sup>46</sup> and the isoelectric points of hhMb (7.2)<sup>43</sup>, the pH of all the solution was adjusted to 7 by adding NaOH in order to maximize the adsorption<sup>43</sup>. Buffer

solutions without proteins are denoted in the text as "(buffer) solutions", while the buffer solution with proteins added is denoted as "protein solution".

## 2.2 Study of the interaction between the buffers and the TiO<sub>2</sub>

In a typical experiment 10 mg of mesoporous TiO<sub>2</sub> were dissolved in 4 mL of the buffer solution. Then, the mixture was left shaking for 2h at 300 rpm on a VWR ADV 3500 shaker. Unless stated otherwise, the buffer concentration is 10 mM. After shaking, the solution was centrifuged at 4000 rpm for 5 min and the precipitate washed and filtrated under vacuum with distilled water in order to remove the physisorbed molecules. Finally, the washed samples were dried at ambient conditions for 1 day. The dried powders were analyzed using TGA/DTG analysis, FT-IR and XRF spectroscopy.

## 2.3 Measurement of proteins adsorption

The protein adsorption was achieved dissolving 10 mg of calcined Millennium in 4 mL of buffered myoglobin solution (0.25 mg protein/mL), then the mixture was left shaking for at least 24 h. Higher concentrations of hhMb (0.5 mg/mL) have also been tested to exclude the effect of protein depletion. The amount of adsorbed proteins was measured by transferring 1 mL of the solution in an Eppendorf tube and centrifuging it at 4000 rpm for 5 min. The concentration of the proteins was calculated analyzing this supernatant with a Thermo Electron Evolution 500 UV-vis spectrophotometer at the wavelength maximum of the Soret band of hhMb ( $\lambda = 408$  nm). As different values are reported in literature<sup>47,48</sup>, the absorption extinction coefficient  $\epsilon$  for hhMb at 408 nm was calculated by the calibration line at different concentrations in the three buffers. The average value of  $129.000\text{M}^{-1}\text{cm}^{-1}$  was used for all the UV-vis measurements (the error occurring from the use of this value for all the three buffers is within the experimental error included in the error bars). The amount of proteins loaded on the mesoporous TiO<sub>2</sub> was calculated subtracting the concentration of proteins still in solution from the initial concentration of the solution. Myoglobin solutions without the mesoporous material were also analyzed to exclude concentration decrease due to precipitation or degradation of proteins out of the solution. The solid samples were also dried and washed with the same procedure as described above, using fresh buffer solution to remove the physisorbed proteins. Subsequently, the myoglobin

incorporated powders were analyzed by IR spectroscopy, TGA/DTG analysis and EPR. Each measurement was repeated at least three times.

## 2.4 Characterization methods

The N<sub>2</sub>-sorption analysis was carried out on a Quantachrome Quadrasorb SI automated gas adsorption system. Before starting the measurement the samples were degassed with an AS-6 degasser for 16 h at 150°C, then the analysis was performed at -196°C. When adsorbed molecules were present, the samples were degassed at 25°C to prevent changes in the loading of the materials. It has to be mentioned that this will leave some residual solvent at the surface, lowering the available surface to some extent. The surface of the sample ( $S_{\text{BET}}$ ) was calculated via a multipoint BET. The pore diameter ( $D_p$ ) was determined via the BJH method on the desorption branch of the isotherm, while the total pore volume ( $V_p$ ) was determined from the adsorption at  $P/P_0$  0.95. A Nicolet 6700 FT-IR spectrophotometer was used for the FT-IR measurements; taking 200 scans per samples with a resolution of  $4\text{cm}^{-1}$ . The samples were diluted with dry KBr to 2 wt% and pure KBr was used as reference. The TGA/DTG analysis was performed using a Mettler Toledo TGA/SDTA 851. All the curves were acquired in O<sub>2</sub> flow in the temperature range 25-600°C with a heating rate of 5°C/min. X-band continuous-wave (CW) EPR measurements were performed on a Bruker ESP300E spectrometer (microwave frequency  $\sim 9.45\text{GHz}$ ) equipped with a gas-flow cryogenic system (Oxford Inc.), allowing for operation from room temperature down to 2.5K. The magnetic field was measured with a Bruker ER035M NMR Gauss meter. During the experiments, a vacuum pump was attached to the EPR tube in order to remove paramagnetic oxygen from the sample. The spectra of heme proteins are typically measured with a microwave power of 0.5mW, a modulation amplitude of 0.5mT, and a modulation frequency of 100kHz at a temperature of 10K. Simulation of the spectra was performed using Easyspin<sup>49</sup>, a toolbox for MATLAB (Mathworks, Natick, Massachusetts, USA). The X-ray fluorescence measurements were carried out on an Epsilon 5 XRF from Panalytical. Titanium was used as secondary target with a tube anode voltage and current of 25 kV and 24 mA, respectively. The lifetime used was 1500 s. For the CV measurements, a 4 mL three-electrode electrochemical cell was used with a calomel reference electrode (SCE) connected to

## ARTICLE

## Journal Name

an Autolab potentiostat. A graphite disk electrode prepared from spectroscopic pure graphite rods with the side wall isolated by epoxy resin was used as working electrode and a glassy carbon rod counter electrode was used as auxiliary electrode. All experiments were carried out in phosphate buffer 0.1 M at pH 7 purged by pure N<sub>2</sub> gas for 30 min before the measurement and the solutions were maintained under nitrogen atmosphere during the whole electrochemical experiment. All samples were analyzed at room temperature.

### 3. Results and discussion

#### 3.1 Characterization of calcined Millennium

Millennium PC500 is a porous, industrially produced titanium-dioxide powder. It has a high surface area (Table 1) and consists of both micro and mesopores. As hhMb is a globular protein with a mean diameter of 4 nm<sup>22</sup>, the pores of the non-calcined Millennium are not large enough to host this protein. Nevertheless, the pores can be enlarged by calcinations. As shown in Table 1, the calcination at 350°C for 6h leads to a dramatic reduction of the surface area, which is nonetheless still significant comparable to non-porous materials. The obtained pore size (5.6 nm) is suitable for hhMb incorporation. Furthermore, as shown elsewhere<sup>45</sup>, the thermal process increases the crystallinity of the Millennium PC500. From now on we will refer only to the mesoporous Millennium PC500 as the material calcined at 350°C with a pore size of 5.6 nm (Mil).

Table 1. Results of nitrogen-sorption analysis of Millennium PC500 as received (non-calcined) and after calcination at 350°C.

	Calc. Temp. (°C)	S <sub>BET</sub> (m <sup>2</sup> /g)	Tot. V <sub>p</sub> (cc/g)	D <sub>p, meso</sub> (nm)
Non-calcined Millennium PC500		350	0.38	3.4
Calcined Millennium PC500	350	122	0.31	5.6

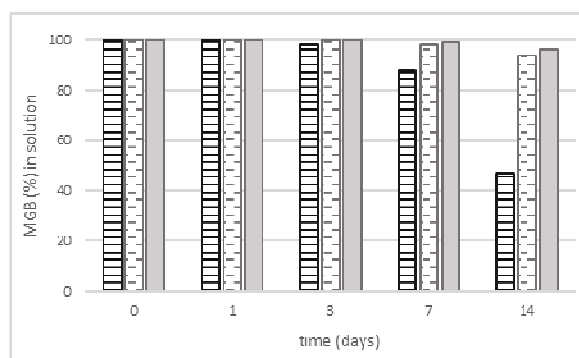
#### 3.2 Buffer effect on protein stability in solution

As stability is one of the key issues in the use of biomolecules, the protein stability has been evaluated for the three different buffers. HhMb (1 mg) has been dissolved in 4 mL of each of the three buffers, and the solutions were left shaking at room temperature for two weeks. Every 24 hours, an UV-vis analysis of the

supernatant, obtained after centrifugation (see Experimental section) was performed to determine the protein stability over time. Figure 1 represents the amount of hhMb still in solution as a function of time. Only 6 % of the initial concentration is lost after two weeks of shaking in Tris solution, this value is even smaller in the case of PBS (4%). In contrast, the concentration of hhMb decreases dramatically in the HEPES solution; more than half of the initial amount of proteins is lost after two weeks through denaturation and/or precipitation.

It has been shown that the presence of specific ligands<sup>50</sup> or ions<sup>51</sup> increases protein stability in solution. As a consequence of their different chemical structure, the three applied buffers may interact differently with hhMb leading to a higher or lower stability than the native protein. Both PBS<sup>52</sup> and Tris buffer<sup>53</sup> are known to stabilize and protect the native structure of the proteins against thermal denaturation, as also confirmed by our experiments in solution (Figure 1).

Fig. 1. Amount (%) of hhMb in HEPES (solid line), Tris (dashed line) and PBS (solid fill) solution versus time of shaking based on UV-vis analysis of the supernatant.



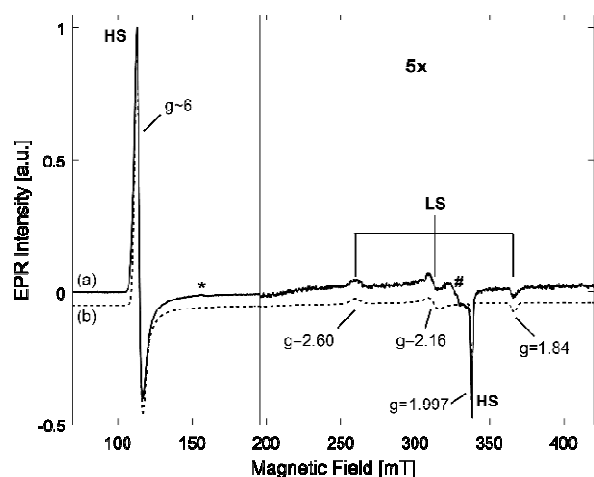
Non-native forms, which may arise in HEPES, have a tendency to agglomerate<sup>54</sup>, forming proteins agglomerates with larger size and higher molecular weight than the folded proteins. Therefore the centrifugation process will remove those aggregates (a precipitate has been observed after this process) leading to a decrease in the amount of proteins in solution detectable by UV-vis spectroscopy. The position of the Soret band (410 nm), typical of the ferric aquomet form of hhMb, was identical for all buffer systems and remained unchanged throughout the whole stability test.

As mentioned, the heme iron of the lyophilized hhMb used here is in its ferric (Fe(III)) state, with the heme iron binding a His residue at

position 8 of the F helix from the proximal side and water from the distal side of the heme (so-called aquomet form). This results in a paramagnetic high spin (HS) state of the heme iron with a very typical EPR spectrum. Changes in the heme pocket and heme iron ligation will have an immediate impact on the EPR spectra.

The EPR spectra of frozen solutions of hhMb in the three different buffers (Figure S.I. 1) show the expected feature of the HS heme iron with effective  $g$  values ( $g_{\text{eff},x} = 5.98 \pm 0.02$ ,  $g_{\text{eff},y} = 5.87 \pm 0.02$  and  $g_{\text{eff},z} = 1.997 \pm 0.002$ ). Furthermore, a (rather small) signal at  $g = 4.28$ , which is ascribed to non-heme iron and observed in most heme-protein solutions, and a background signal due to a Cu(II) cavity impurity are observed. Additionally, the EPR spectrum of hhMb in Tris buffer (Figure 2) shows a contribution of a low-spin (LS) Fe(III) heme center ( $g_x = 1.84 \pm 0.01$ ,  $g_y = 2.16 \pm 0.01$  and  $g_z = 2.60 \pm 0.001$ ). This change in the heme iron from a high to a low spin state indicates a change in distal ligand.

Fig. 2. CW-EPR spectra of a frozen solution of 0.5 mM hhMb in 10 mM Tris pH7 (solid line (a)). The simulated spectrum (dotted line (b)) shows the contributions of the HS and LS forms respectively. In (a), the contribution of non-heme iron is indicated with an asterisk and the Cu(II) background signal is indicated with a hash.



More specific, the here observed LS signal can be ascribed to hydroxide-coordinated hhMb, i.e. a low-spin form produced by the ionization of the distal water<sup>55</sup>. This form tends to become more prominent as pH increases. The EPR results thus show that, for none of the investigated buffers, major changes occur in the heme-pocket region, apart from the known deprotonation of the heme-ligating water occurring at pH>7.

Nevertheless, the UV-vis results clearly show that HEPES is not a very good buffer to preserve hhMb in solution over time.

### 3.3 Interaction between the different buffers and mesoporous TiO<sub>2</sub>

The interaction between the TiO<sub>2</sub> surface and the buffer solution can be monitored by FT IR. Figure S.I. 2 shows the FT-IR spectra of dried TiO<sub>2</sub> samples after dissolution in distilled water (Mil-H<sub>2</sub>O), HEPES (Mil-HEPES), Tris (Mil-Tris) and PBS (Mil-PBS) solution, followed by rinsing with distilled water. Weakly interacting molecules will be removed from the surface by washing. All four spectra show two broad intense bands, in the region between 3800 and 2600 cm<sup>-1</sup> and at 1630 cm<sup>-1</sup>, assigned to adsorbed water<sup>56</sup>, and a strong absorption band below 1000 cm<sup>-1</sup> due to the bulk TiO<sub>2</sub><sup>57</sup>. Furthermore, all spectra show a sharp peak at 3690 cm<sup>-1</sup> assigned to the OH group of the upmost adsorbed water layer<sup>56</sup>. The change in the shape of the broad band in the region between 3800 and 2600 cm<sup>-1</sup> in Mil-PBS can be ascribed to differences in the amount and structural features of the adsorbed water<sup>58</sup>. This might be induced by the presence of phosphonic acid or phosphoryl groups present in PBS.

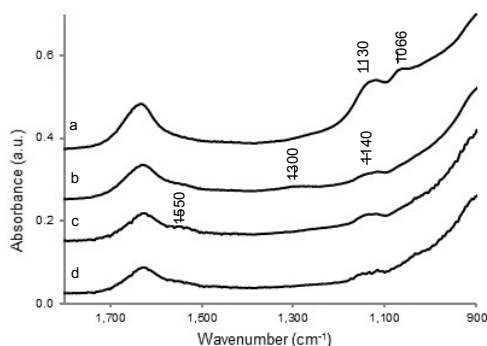
A magnification in the region 900-1800 cm<sup>-1</sup> of the IR spectra is shown in Figure 3. The weak bands at 1550 and 1140 cm<sup>-1</sup> are ascribed to impurities of the bulk material (presumably carbonate and sulfate according to the producer). A small broad peak centered at 1300 cm<sup>-1</sup>, assigned to the C-N stretching<sup>59</sup>, is visible in the case of the sample dissolved in Tris solution. Two bands, associated with the symmetric and asymmetric vibrations of the PO<sub>4</sub> groups<sup>60,61</sup>, appear at 1130 and 1066 cm<sup>-1</sup> in the spectrum of calcined Millennium treated with the PBS solution. The IR results thus show that washing with distilled water is not able to remove phosphate anions of the PBS solution or Tris molecules from the TiO<sub>2</sub> surface. In contrast, there is no evidence for the presence of HEPES molecules on the sample, at least not at concentrations detectable in IR. Indeed, the typical vibrations between 1050 and 1130 cm<sup>-1</sup> of the sulfate group<sup>62</sup> are not detected, confirming that if anything is present, the amount is small.



## ARTICLE

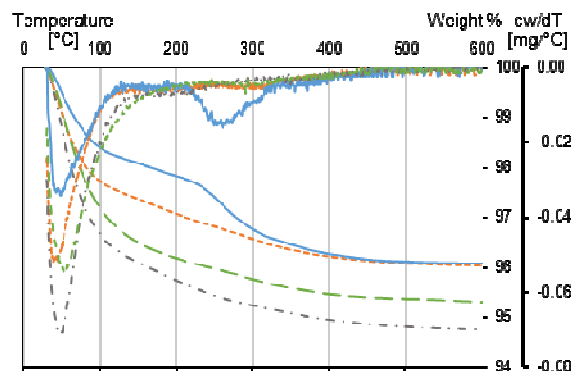
Journal Name

Fig. 3. Magnification of the IR spectra (offset 0.1) in the region 900–1800  $\text{cm}^{-1}$  of Mil-PBS (a), Mil-Tris (b), Mil-H<sub>2</sub>O (c) and Mil-HEPES (d) after washing with distilled water and drying at room temperature.



To highlight the presence of buffer molecules on the Millennium surface, a TGA/DTG analysis of Millennium treated with different buffer solutions has been performed (Figure 4). In all three TGA curves the weight loss occurs in two main steps. A first step, below 200 °C, due to the adsorbed water<sup>63</sup>, and a second one, between 200 °C and 500 °C, related to the burning or desorption of organic molecules<sup>64</sup>. Samples dissolved in PBS or HEPES show high weight losses below 200 °C (3.8% and 3%, respectively) although less than Mil-H<sub>2</sub>O (4.5%). The weight loss in Mil-Tris is even less (2.2%). Furthermore, the DTG profile of Mil-PBS indicates a shift toward higher temperature, when compared with the Millennium dissolved in distilled water. The shift in the DTG peak can be ascribed to water molecules linked to the phosphate group<sup>65</sup>. On the other hand, the weight loss of Mil-HEPES and Mil-Tris below 200 °C occurs at lower temperature compared to Mil-H<sub>2</sub>O or PBS. Mil-Tris has the most prominent (1.8%) weight loss above 200 °C, with a clear, broad DTG peak centered at 260 °C and a small shoulder up to 400 °C. Both samples treated with HEPES and PBS have weight losses above 200 °C comparable to Mil-H<sub>2</sub>O, indicating the absence of a large amount of surface adsorbed species that can burn and/or desorb. This suggests, like the IR spectrum, that no HEPES molecules are bound to the surface, or at least not in significant concentration. In contrast, the difference in weight loss at high temperatures observed for Mil-Tris (1.8%) versus Mil-H<sub>2</sub>O (0.7%) indicates the presence of buffer molecules bound to the surface in the former case.

Fig. 4. TGA and DTG profiles of Mil-H<sub>2</sub>O (dashed dotted gray line), Mil-HEPES (short-dashed orange line), Mil-Tris (solid blue line), Mil-PBS (dashed green line) after washing with distilled water and drying at room temperature.



The XRF analysis (Table 2) performed on Mil-H<sub>2</sub>O and Mil-PBS confirms the hypothesis that phosphate groups are bound on the surface. Additionally, other ions (mainly potassium and chloride) from the PBS buffer solution are detected in elevated amounts. A strong presence of chloride anions, one of the component of the Tris buffer solution, is also detected in Mil-Tris. This suggests, once again, interactions between the Tris buffer and the TiO<sub>2</sub> surface. The results for Mil-HEPES (element dosed: S) confirm that no or very few HEPES molecules are bound to the surface.

Table 2. Intensities ratio obtained from the XRF analysis of Mil-H<sub>2</sub>O, Mil-PBS, Mil-Tris and Mil-HEPES.

	X-Ray Intensities Ratio					
	P	S	K	Cl	Al	Ca
Mil/Mil-H <sub>2</sub> O	0.9	1	1	1	1	1
Mil-HEPES/Mil-H <sub>2</sub> O	0,9	0,5	1	1,6	1	1,5
Mil-PBS/Mil-H <sub>2</sub> O	5,5	0,3	51	631	0,8	1,9
Mil-Tris/Mil-H <sub>2</sub> O	0.9	0.5	1	16.5	1	1

All the observations discussed above are confirmed by N<sub>2</sub> sorption analysis (Figure S.I. 3). Samples dissolved in Tris and phosphate buffer show a remarkable decrease in free pore volume when compared with Mil-H<sub>2</sub>O dissolved in water and degassed at the same temperature (25 °C). The volume reduction is much less prominent in samples treated with HEPES buffer. This observation confirms the presence of Tris molecules and phosphate anions, together with possible residual adsorbed water, in high concentration on the TiO<sub>2</sub> surface. The presence of water cannot be excluded as degassing was only done at 25 °C. Nevertheless, TGA

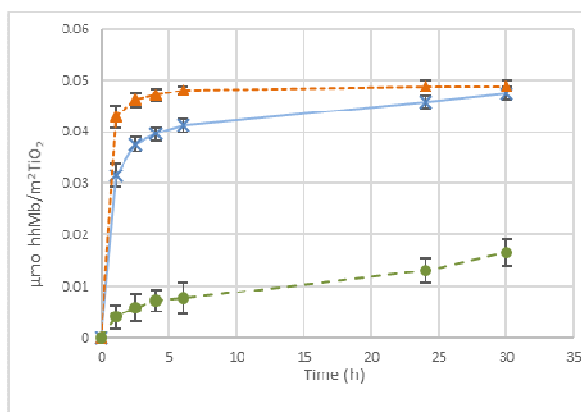
below 200°C shows that water adsorption alone is not responsible for these differences (Figure 4). In all the samples the effect is more prominent in pores with larger pore size (higher  $P/P_0$ ).

The different behavior observed can be explained considering the distinct differences in chemical structure of the three buffers. It is well known that phosphate anions can strongly interact with titanium dioxide<sup>66</sup>. With respect to the presence of anions on the surface, our current results are consistent with the XRF analysis reported by A. Marucco et al.<sup>42</sup> for TiO<sub>2</sub> nanoparticles dissolved in PBS buffer. The Tris molecule has three OH groups that can bind to the TiO<sub>2</sub> on top of a possible interaction via its amino function. It is thus not surprising that Tris has a strong and clear interaction with the surface. HEPES is only able to interact via the single OH group or via the sulfate group, which has a chelating effect as well although much weaker than the phosphate group<sup>67</sup>. The results of the thermal analysis and IR spectroscopy clearly show different interactions between the three buffers and the surface. Therefore, it is expected that the use of different buffers will also induce differences in the interaction of hhMb with Millennium, possibly influencing the protein adsorption rate and loading capacity.

### 3.4 Buffer effect on the adsorption of horse heart myoglobin

Figure 5 presents the time evolution of the adsorption of 1 mg ( $5.7 \times 10^{-8}$  mol) of hhMb on 10 mg of Millennium PC 500 in the three different buffers (hhMb-Mil-HEPES, Tris or PBS). The plot of  $\ln C/C_0$  versus time (Figure S.I. 4) allows the estimation of the kinetic constant  $k$  (assuming a first order kinetic for the adsorption) and thus an assessment of the initial adsorption rate (within 1h of shaking) of hhMb on the mesoporous Millennium.

Fig. 5. Adsorption of hhMb on the mesoporous Millennium in HEPES (short-dashed orange line), Tris (solid blue line) and PBS (dashed green line) solution. The concentration of the buffer is 10 mM. The results are expressed as amount of proteins (in  $\mu\text{mol}$ ) per  $\text{m}^2$  of TiO<sub>2</sub> versus shaking time. Error bars are calculated on set of three measurements.



The graph clearly illustrates a different trend in adsorption capacity and adsorption rate of hhMb on TiO<sub>2</sub> due to the nature of the buffer solution. Proteins in HEPES solution are most efficiently adsorbed on titania ( $k = 1.3$ ), reaching a maximal capacity after 6 hours. The encapsulation of proteins in Tris buffer differs basically in the initial rate of incorporation ( $k = 0.6$ ), but reaches a similar protein loading, albeit after a much longer adsorption time (factor of 5). The adsorption from PBS ( $k = 0.03$ ) shows a completely different behavior; only a small amount of proteins are able to bind to the surface, within the first 24 h of incorporation, with a very slow initial adsorption rate. In both hhMb-Mil-HEPES and hhMb-Mil-Tris it is possible to incorporate the same amount of proteins within a time frame of 24 h, therefore the use of these two buffers only seems to have an impact on the kinetics of the process. However, phosphate anions dramatically affect the loading capacity. In fact, the total amount of incorporated proteins in hhMb-Mil-PBS is about four times smaller than the one adsorbed for hhMb-Mil-HEPES or hhMb-Mil-Tris. Therefore, it is reasonable to suggest an inhibiting effect.

We have already shown that both Tris molecules and phosphate anions have a clear interaction with the TiO<sub>2</sub> surface, suggesting that the buffer-dependent protein-incorporation arises from these differences in interaction of the buffer molecules/ions with the mesoporous material. In order to evaluate the effect of the different buffer-TiO<sub>2</sub> interactions on hhMb adsorption, leaching tests with the three different buffers have been performed. After



## ARTICLE

Journal Name

the incorporation of 1 mg of hhMb in HEPES and Tris solution, samples have been collected and dissolved again in a fresh buffer solution to study the displacement of hhMb by the added buffer.

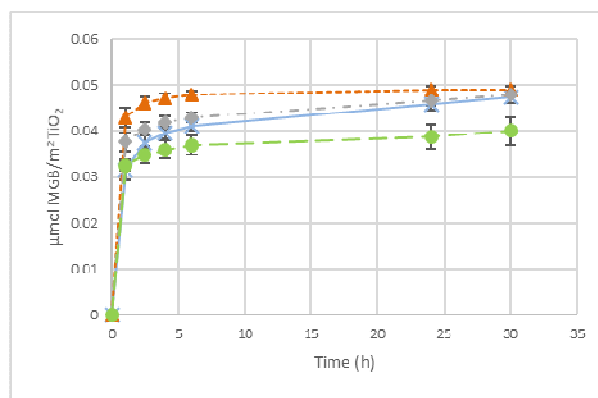
The results (Figure S.I. 5) clearly indicate a different effect of the PBS in relations to the other buffers. In fact, the phosphate anions and Tris molecules interact with the surface as also derived using IR and/or XRF, but in contrast to Tris, the phosphate interaction is stronger as the phosphate molecules have the ability to displace part of the adsorbed proteins. This indicates a higher affinity of the PBS for the mesoporous Millennium than the globin and a competitive adsorption between anions and proteins inducing a partial desorption of hhMb. The ability of the phosphate anions to displace proteins from the surface was observed also in previous studies on  $\text{TiO}_2$  thin films<sup>41</sup>. Even though leaching was observed in the presence of the phosphate anions, not all the proteins (37%) have been removed from the surface and a stable equilibrium is obtained after 5 hours.

As revealed by IR and TGA, the interaction of Tris molecules with the surface seems to be less competitive as no displacement is observed for hhMb-Mil-HEPES when dissolved in Tris buffer. However, hhMb-Mil-Tris appears to be more strongly bonded since the leaching by phosphate anions is weaker than in hhMb-Mil-HEPES (factor of 3 (Figure S.I. 5)). It may be possible that part of adsorbed hhMb interacts with the Tris molecules bonded to the Millennium (see Figure 3 and 4) via the amino function of the buffer molecules. As a consequence of such interaction, the displacement of the adsorbed hhMb by phosphate anions is more difficult and less proteins are leached out of the surface. Besides, this may explain the slower adsorption observed for hhMb-Mil-Tris (Figure 5) as the diffusion of hhMb inside the pores would be more challenging due to the stronger interaction to the surface (*via* the amino group of the Tris molecule), diminishing desorption from the surface and thus slowing down surface diffusion.

This is confirmed by hhMb incorporation (Figure 6) performed using HEPES and Tris buffers at higher concentration (0.1M). HhMb-Mil-HEPES(0.1M) and hhMb-Mil-Tris(0.1M) follow a different trend upon increase of the buffer concentration. On the one side, increase of HEPES concentration leads to a less efficient hhMb uptake, although the same amount of adsorbed proteins is

observed after 24h. On the other side, with an increase of Tris concentration a reduced hhMb adsorption is accomplished.

Fig. 6. Adsorption isotherms of hhMb-Mil-HEPES(0.01M) (short-dashed orange line,  $\Delta$ ), hhMb-Mil-HEPES(0.1M) (dotted dashed gray line,  $\diamond$ ), hhMb-Mil-Tris(0.01M) (solid blue line,  $\times$ ) and hhMb-Mil-Tris(0.1M) (dashed green line,  $\bullet$ ). The results are expressed as amount of proteins (in  $\mu\text{mol}$ ) per per  $\text{m}^2$  of  $\text{TiO}_2$ . Error bars are calculated on a set of three measures.



Again we explain this difference by differences in buffer-surface interactions influencing adsorption and diffusion. The slower uptake for hhMb-Mil-HEPES(0.1M) points to interactions also between HEPES and the mesoporous  $\text{TiO}_2$ , although this interaction seems to be the weakest as it only influences the adsorption rate (and not the adsorption capacity) at high buffer concentration. The finding for hhMb-Mil-Tris(0.1M) confirms the previous hypothesis about hhMb strongly interacting with Tris molecules adsorbed on the surface. As we already discussed, the diffusion inside the pores is more challenging in the presence of strong interactions between hhMb and the surface. It is straightforward to assume that multiple interaction between hhMb and different adsorbed Tris molecules are possible when the buffer concentration is increased by a factor of 10. As a consequence, hhMb diffusion in hhMb-Mil-Tris(0.1M) is more difficult (if not even prohibited), resulting in a lower adsorption capacity.

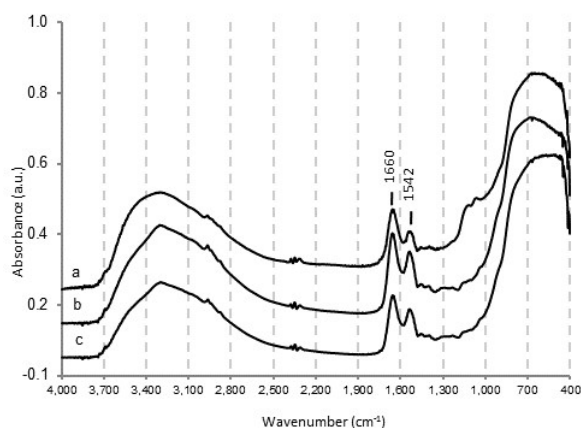
### 3.5 Structural stability of horse heart myoglobin upon adsorption in different buffers

In order to investigate the structural stability of the proteins after adsorption, the IR spectra of the samples have been recorded after protein incorporation (Figure 7). The peaks at  $1660$  and  $1542\text{ cm}^{-1}$  have been assigned to the amide band I (C=O stretching mode) and to the amide band II (N-H deformation) of hhMb, respectively<sup>68</sup>. The intensity ratio between these two band is  $1.2 \pm 0.1$  in the case of

proteins adsorbed from HEPES or Tris solution, and  $1.1 \pm 0.1$  in the case of pure hhMb, indicating that no important denaturation affects the secondary structure after the adsorption<sup>69</sup>. This ratio is quite different in hhMb-Mil-PBS (1.6), suggesting a structural change of hhMb as a result of the interaction with the surface and/or phosphate group. This result is consistent with the changes in the amino-acid chain observed in bovine and human serum albumin upon adsorption on TiO<sub>2</sub> surface in PBS<sup>70</sup>. Notwithstanding this conclusion, an influence of the adsorbed water (peak at 1630 cm<sup>-1</sup> see Figure S.1.2 or 3) on the ratio between the two amino bands in hhMb-Mil-PBS cannot be excluded.

Again, in the IR spectra depicted in Figure 7, the clear interaction of the phosphate groups with the surface is indicated by the peaks at 1130 and 1066 cm<sup>-1</sup> (compare with Figure 3).

Fig. 7. IR spectra of hhMb-Mil-PBS (a), hhMb-Mil-HEPES (b) and hhMb-Mil-Tris (c) after washing with distilled water and drying at room temperature.



Furthermore, the TGA/DTG curve (Figure 8) evidences differences in the deposition of the proteins on the surface. As already discussed earlier (Figure 4), the first weight loss below 200°C is due to adsorbed water and it occurs at higher temperature for hhMb-Mil-PBS. The subsequent weight losses above 200°C are due to the degradation of the hhMb and in the case of Tris also due to the loss of buffer. Samples hhMb-Mil-Tris and hhMb-Mil-HEPES show the same weight loss (5.8%) in the region 200-600°C. This is consistent with the UV-vis results, showing a similar amount of adsorbed proteins in the two buffers after 24 hours. The TGA curve of hhMb-Mil-PBS shows a smaller weight loss (4.6%), again pointing to its lower efficiency in adsorption. The thermal degradation of proteins is a complex process taking place in different steps, leading to broad

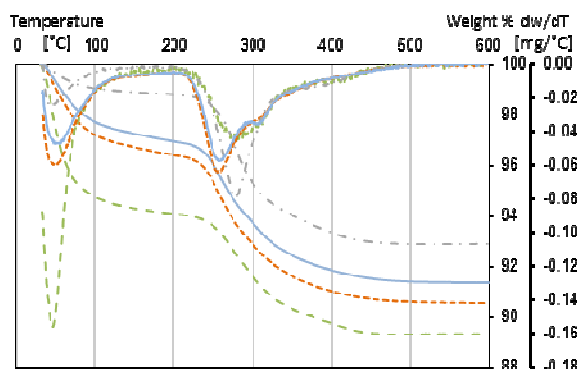
DTG peaks, in both *in vitro*<sup>71</sup> and *in vivo*<sup>72</sup> experiments. The DTG profiles of samples with hhMb adsorbed to Millennium in the three buffers differ from each other, pointing to changes in their local structure and/or environment. The degradation of the proteins in hhMb-Mil-Tris and HEPES starts at lower temperature and clearly shows two different steps at 260°C and a shoulder at 310°C. hhMb-Mil-Tris presents a slightly more intense shoulder at 310°C; a possible consequence of the interaction between hhMb and Tris molecules adsorbed on the surface as discussed above. However, hhMb-Mil-PBS presents a different degradation mechanism as can be deduced from the shift of the DTG peak and TGA curve to higher temperature with a maximum at 280°C and a small weight loss above 400°C. The shifts in the DTG peaks usually reflect the existence of competing steps in polymer degradation<sup>73</sup>.

The thermal analysis of non-porous titanium dioxide (P25) with hhMb adsorbed in HEPES buffer (hhMb-P25-HEPES) has also been performed to further elucidate the observed differences for the Millennium case (Figure 8) and to look in more detail to the role of the pores. Of course, differences induced by divergent surface topology and chemistry cannot be excluded. Nevertheless, the use of non-porous TiO<sub>2</sub> may reveal possible correlations between the different degradation steps observed and the position of the proteins on the surface, i.e. on the external surface or confined in the pores. Again the two weight losses due to 1) the adsorbed water and 2) the proteins are observed. The DTG profile, with the maximum degradation rate centered at 280°C, resembles the DTG of hhMb-Mil-PBS. As P25 is a non-porous titanium dioxide, the proteins are adsorbed only on the external surface. Therefore, we assume that the use of PBS mainly leads to the adsorption of proteins on the external surface of the samples rather than into the pores. This would explain the lower hhMb adsorption capacity (Figure 5). The different hhMb degradation observed for hhMb-Mil-Tris and HEPES may arise from differences in protein structure and conformation due to the pore confinement.

## ARTICLE

## Journal Name

Fig. 8. TGA and DTG profiles of hhMb-Mil-HEPES (short-dashed orange line), hhMb-Mil-Tris (solid blue line), hhMb-Mil-PBS (dashed green line) and hhMb-P25-HEPES (dashed dotted gray line) after washing with distilled water and drying in air at room temperature. The concentration of the three buffers was 10mM.



The  $N_2$  sorption analysis (Figure S.I. 6) of Millennium with incorporated hhMb shows similar values for the free pore volume irrespective of the buffer. The pore-volume reduction of the incorporated samples (0.08 cc/g) is comparable with the volume of 1 mg of hhMb (0.07 cc/g<sup>23</sup>). Therefore, on one side, this result strengthens the hypothesis that hhMb has diffused into the pores, when the incorporation takes place in Tris or HEPES solution. On the other side, the adsorption studies (Figure 5) show that the amount of adsorbed proteins in hhMb-Mil-PBS is too small to be causing such a large volume decrease. A possible explanation might be the blocking effect of the pores by hhMb proteins, or there may be a stronger water adsorption in case of the phosphate buffer due to the presence of phosphate groups on the  $TiO_2$  surface (compare with Figure S.I. 3).

As discussed above PBS stabilizes the native structure of the protein preventing hhMb from unfolding and subsequent aggregation. Thus, the diverging behavior observed for this buffer cannot be ascribed to full denaturation of the proteins inside the solution, as proved by the stability test shown above. As discussed earlier, the phosphate anions and hhMb seem to have a competitive adsorption, which causes a strong decrease of the surface coverage. In addition, some modifications on the secondary structure of the protein occur when the adsorption is performed in phosphate buffer, as shown by the IR spectrum (Figure 7). This is in agreement with structural rearrangements of hhMb observed upon adsorption on zirconia nanoparticles due to the phosphate anions<sup>74</sup>.

We propose that the phosphate anions ( $HPO_4^{2-}$  and  $H_2PO_4^-$  at pH 7) are responsible for the divergent incorporation behavior observed when PBS is used, generating on the  $TiO_2$  surface (which tends to be slightly positively charged at pH 7<sup>46</sup>) a net negative charge. This may have two main effects, in addition to the competition for the binding sites on the mesoporous  $TiO_2$  already discussed. On the one hand the negative amino acids residues can be repulsed by the negative charge on the surface, as a consequence, only a small amount of proteins is able to approach the surface. This is confirmed by the adsorption of hhMb on the non-porous P25 (Figure S.I. 7). The electrostatic repulsion between negative charges may account for the difference observed in the amount of hhMb adsorbed in HEPES and Tris and the one in PBS. The adsorption behavior is in agreement with the results reported for BSA on P25 in PBS and HEPES<sup>42</sup>.

On the other hand, similar to what is observed for the Tris case, the hhMb may have stronger interaction with the phosphate groups (possibly via the positively charged residues). Therefore the diffusion of the proteins on the surface and inside the pores might be strongly limited. In particular, proteins adsorbed strongly on the edge of the pore channel that are unable to diffuse further inside the pore may induce pore blocking that would explain the observed reduction in the pore volume. This hypothesis is strengthened by the TGA/DTG curves, according to which hhMb-Mil-PBS shows the same DTG profile of hhMb-P25-HEPES (Figure 8), suggesting adsorption of hhMb mainly on the external surface.

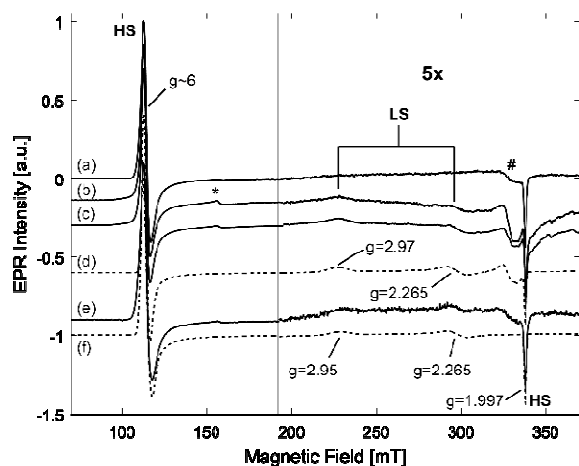
### 3.6 Heme pocket structure upon adsorption in different buffers

As discussed above, the EPR analysis gives information on the stability and the conformation of the heme center of proteins. Figure 9 shows the EPR spectra of a frozen solution of hhMb in HEPES buffer and of hhMb-Mil-HEPES, before and after drying. All spectra show the EPR feature typical for the HS aquomet form of hhMb (see also Figure 2). After incorporation of the hhMb in Millennium, this EPR spectrum (both for dried and wet materials) changes slightly when compared with the spectrum of hhMb in a frozen HEPES buffer solution (Figure S.I. 8), indicating a local change/pressure on the heme-pocket structure (inducing a change in zero-field splitting parameters). Additionally, in all spectra the non-heme iron (\*) and the Cu (II) background (#) signal are also

present (see also Figure 2). There are no clear differences in the HS feature of dry and wet incorporated  $\text{TiO}_2$ , which indicates that myoglobin is remaining strongly inside the pores, and not redissolving after addition of buffer (as already showed by the leaching tests) and that the here applied drying procedure does not cause a major loss of the water molecule axially ligating to the heme iron.

The EPR spectrum of hhMb incorporated in phosphate buffer (Figure S.I. 9) shows a large non-heme iron signal and a worse signal to noise ratio when compared to the other buffers. This indicates that a smaller amount of myoglobin is present in the pores, confirming the previous observations (see Figure 5).

Fig. 9. CW-EPR spectra of a frozen solutions of hhMb (a) and hhMb-Mil-HEPES (before (b) and after (c) drying of the final powder). (d) Simulation of the dried Millennium in 10 mM HEPES. (e) CW-EPR spectrum of frozen solution of 0.5 mM hhMb in HEPES after addition of SDS to a final concentration of  $9.5 \times 10^{-4}$  M. (f) shows the corresponding simulation. The contributions of high-spin features (HS), non-heme iron (\*) and a Cu(II) background signal (#) are shown. All spectra are rescaled to the same microwave frequency and normalized to equal intensity for facile comparison.



When using HEPES (Figure 9) or Tris buffer (Figure S.I. 10) for the protein incorporation, the EPR spectra of the hhMb incorporated in mesoporous  $\text{TiO}_2$  show a low-spin contribution with  $g_z = 2.97 \pm 0.01$  and  $g_y = 2.265 \pm 0.005$  (Figure 9). This is in contrast with hhMb-Mil-PBS, where no such contribution arises (Figure S.I.9). This again suggests that only a small amount of hhMb is able to enter the pores of mesoporous  $\text{TiO}_2$  in the presence of PBS; agreeing with earlier results. The extra low-spin component spectrum has similar  $g$ -values as found for the imidazole complex of Mb and other heme systems with bis-imidazole ligation of the heme iron<sup>75</sup>.

Tofani et al.<sup>76</sup> showed using UV-vis absorption spectroscopy that, upon addition of a low concentration of sodium dodecyl sulfate

(SDS), myoglobin forms hexacoordinated LS species through binding of the distal histidine (His-64) to the central iron atom. It is suggested that this occurs through breaking of important salt bridges, which renders the  $\alpha$  helix structure more flexible. Figure 9e shows the EPR spectrum of a frozen solution of hhMb in HEPES buffer where SDS was added to a final concentration of  $9.5 \times 10^{-4}$  M. This spectrum also shows an LS contribution with  $g_z = 2.95 \pm 0.02$  and  $g_y = 2.265 \pm 0.005$  which are similar values to those found for the LS contribution in incorporated mesoporous  $\text{TiO}_2$  incorporated with hhMb. This indicates that, upon incorporation into the pores, there is a strain on hhMb resulting in a shift of the E-helix of the protein and subsequent coordination of the distal histidine to the central heme iron, possibly related to similar breaking of the salt bridges as when SDS is added.

The apparent strong difference in the intensity of the HS and LS EPR contributions is mainly due to the larger magnetization of the HS iron centre. In fact, simulations reveal that about 30-35% of the spectrum can be ascribed to the LS form. This indicates a changed strained conformational for a significant fraction of the adsorbed hhMb.

### 3.7 Electrochemical activity of proteins incorporated in different buffers

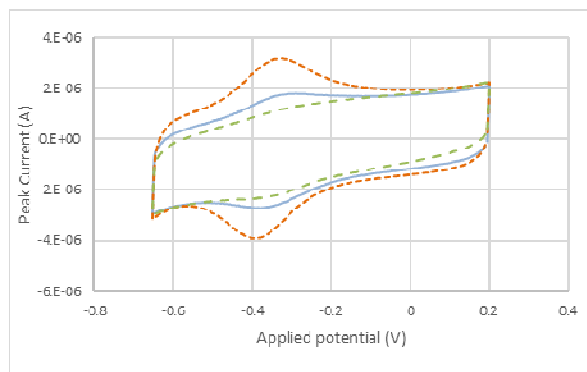
Cyclic voltammetry is a useful tool to investigate the microenvironment surrounding the heme center through evaluation of the redox activity. Differences in the electrochemical activity may arise from the pores confinement and/or different interaction with the  $\text{TiO}_2$  surface.

Figure 10 shows the cyclic voltammograms at scan rate of 0.15 V/s of mesoporous  $\text{TiO}_2$  with incorporated hhMb prepared in the three different buffers.

## ARTICLE

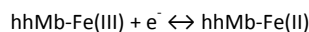
## Journal Name

Fig. 10. Cyclic voltammograms in PBS 0.1M at pH 7 of hhMb-Mil-HEPES (short-dashed orange line), Tris (solid blue line) and PBS (dashed green line) buffer at scan rate of 0.15 V/s.



The voltammogram of hhMb-Mil-HEPES shows a couple of stable and well-defined redox peaks at -0.39 and -0.33 V vs SCE, with an average middle point potential ( $\frac{E_c + E_a}{2}$ ) of -0.36 V. The value is in good agreement with the results obtained for myoglobin deposited in gold nanoparticles<sup>77</sup> and polymer matrix<sup>78</sup>, suggesting that mesoporous titanium dioxide provides a friendly environment for the adsorbed proteins. The midpoint potential in proteins adsorbed in Tris and PBS shifts toward more positive potential (-0.33 V). The cathodic and the anodic peak are nearly symmetric in all the three samples, indicating a thin layer electrochemical behavior<sup>79</sup> with equal reduction and oxidation peak height. This confirms that the electroactive hhMb-Fe(III) is reduced to hhMb-Fe(II) during the forward scan and then fully reoxidized to Fe(III) during the back scan. The peak current is proportional to the scan rate (Figure S.I. 11A) in the range 50-200 mV/s ( $R=0.9999$ ) indicating a quasi-reversible surface controlled thin layer process. This observation is confirmed by plotting the current logarithm versus the logarithm of the scan speed, the linear relationship gives a slope of 1.02, close to theoretical value of 1 for surface controlled electrochemical process<sup>79</sup>. At higher scan rate (0.2-1 V/s) the peak current becomes proportional to the square root of the scan rate  $v^{1/2}$  (Figure S.I. 11B), indicating a diffusion controlled process. It has been observed<sup>77</sup> that the redox reaction of myoglobin incorporated in matrices involves the participation of protons from the solution in order to neutralize the excess charge in the electrochemical process. The involvement of a proton gradient, generated at high scan speed, results in a diffusion-controlled behavior with scan speed in the range 0.2-1 V.

At low scan rate the peak-to-peak separation ( $\Delta E$ ) is  $55 \pm 4$  mV for all the samples, close to the theoretical value of  $\frac{59}{n}$  mV for processes involving one electron ( $n = 1$ ). This confirms that the electrochemical reaction of hhMb adsorbed into mesoporous TiO<sub>2</sub> follows the scheme



With the increase of the scan rate, an increase of  $\Delta E$  is observed in all the three samples. Thus the electrochemical parameters of the redox process of the encapsulated hhMb were calculated using the Laviron's equations<sup>80</sup>. The plot of the cathodic peak potential versus the logarithm of the scan rate gives a charge transfer coefficient  $\alpha$  of  $0.58 \pm 0.08$  for hhMb-Mil-HEPES (Figure S.I. 12) and Tris. This value dramatically decreases hhMb-Mil-PBS ( $0.15 \pm 0.02$ ). Since the peak-to-peak separation is less than 200 mV, the electron transfer rate constant  $k_s$  can be estimated according to the formula<sup>80</sup>

$$\log k_s = \alpha \log(1 + \alpha) + (1 - \alpha) \log \alpha - \log \frac{RT}{nFv} - \frac{(1 - \alpha) \alpha F n \Delta E}{2.3RT}$$

where  $F$  is the Faraday constant (96846 C/mol),  $v$  is the scan rate,  $R$  is the gas constant,  $T$  the absolute temperature,  $n$  the number of electrons involved (here  $n=1$ ) and  $\Delta E$  is the peak to peak separation.

For hhMb-Mil-HEPES and Tris  $k_s$  was calculated to be  $(2 \pm 0.1) \text{s}^{-1}$ , a value larger than the one observed for hhMb adsorbed on carbon coated Fe<sub>2</sub>O<sub>3</sub><sup>81</sup>, on graphene-SnO<sub>2</sub><sup>82</sup>, on ZnO<sup>83</sup> and on mesoporous silica<sup>84</sup>. This indicates a faster electron transfer rate in case of hhMb encapsulated in mesoporous TiO<sub>2</sub> in HEPES and Tris buffer. In case of hhMb-Mil-PBS the value of  $k_s$  is  $(0.7 \pm 0.1) \text{s}^{-1}$ , confirming the different situation for proteins adsorbed in phosphate buffer. In addition, for the observed redox process the concentration of active molecules on the electrode ( $\Gamma$ ) can be estimated using the Laviron's equation<sup>80</sup>:

$$I_p = \frac{n^2 F^2 v \Gamma A}{4RT}$$

where  $I_p$  is the peak current,  $A$  is the area of the electrode (0.072 cm<sup>2</sup>) and all the other terms have the meaning explained above. According to the slope of  $I_p$  vs  $v$ , the surface concentration of active molecules is calculated to be  $1.7 \times 10^{-10}$  mol/cm<sup>2</sup> for hhMb-Mil-HEPES,  $7 \times 10^{-11}$  mol/cm<sup>2</sup> for hhMb-Mil-Tris and  $2.3 \times 10^{-11}$  mol/cm<sup>2</sup> for hhMb-Mil-PBS.

It has been reported that the electrochemical activity of hhMb is highly influenced by the presence of inorganic anions<sup>85</sup>. Thus, it is reasonable that the reduced electrochemical activity for hhMb-Mil-PBS is due to the presence of phosphate anions adsorbed on the TiO<sub>2</sub> surface.

The CV experiments confirm the strong influence of the buffer solution on the structure of the adsorbed proteins, influencing their electrochemical activity. Although the electrochemical process follows the same scheme, regardless the buffer used for the incorporation, the shifting in the potential indicates that the electron transfer is favorable for hhMb-Mil-HEPES. Since the amount of proteins in hhMb-Mil-HEPES and Tris is the same (Figure 5), the lower peak current and concentration of active molecules for hhMb-Mil-Tris probably arises from the strong interaction between the adsorbed buffer molecules and the proteins.

In addition, the values of  $k_s$  show a much faster electron transfer for samples with hhMb incorporated in HEPES and Tris solution than in PBS buffer. Those results suggest, once again, that very different outcomes can be achieved depending on the choice of the buffer solution used for the incorporation.

#### 4. Conclusions

The impact of the choice of the buffer medium on the stability and the adsorption of hhMb on a commercial mesoporous titanium dioxide was evaluated. Tris and PBS were revealed to be a good choice to preserve the proteins in solution while a strong decrease of the concentration was observed after two weeks in HEPES solution. Nevertheless, the benefit or drawback of the buffer is quite different when incorporating hhMb in mesoporous TiO<sub>2</sub>. Very different interactions between the three buffers and the titania surface were observed. All the results indicate a much stronger interaction between mesoporous titanium, phosphate anions and Tris molecules than for the HEPES molecules (in sequence of interaction strength).

This results in a large impact of the medium choice on protein incorporation. The same hhMb adsorption capacity for Millennium TiO<sub>2</sub> has been observed using Tris and HEPES buffer. Nevertheless the use of Tris leads to a slower adsorption, probably due to the strong interaction between hhMb and the adsorbed buffer

molecules resulting in a slow proteins diffusion inside the pores. As it is reasonable to expect, the effect is amplified by an increase in the buffer concentration.

The adsorption from PBS solution exhibits a slower kinetic with a much lower adsorption capacity, which inhibits the incorporation of a high amount of proteins. Additionally, it has been shown how the different interaction (hhMb-Mil-Tris) and the adsorption mainly on the external surface (hhMb-Mil-PBS) create correlated changes in electrochemical activity of the adsorbed proteins. It is important to note that, irrespective of the buffer, the adsorption of the hhMb induces changes in its heme-pocket structure.

As our results clearly indicate, the choice of the buffer medium is a crucial step in protein incorporation. Protein stability in solution, adsorption kinetic, structural modification and activity of the adsorbed proteins are all parameters strictly connected to the selected buffer. Moreover, a clear impact specific for mesoporous materials was observed. Further analysis are required in order to clarify the type and quantify the strength of the interaction between buffer-mesoporous material-biomolecule, as well as possible changes in the conformation of the adsorbed proteins. Nevertheless, we have showed that special attention in the choice of the buffer medium is indispensable in order to avoid misunderstanding of the results about adsorption of biomolecules on mesoporous TiO<sub>2</sub>.

#### Acknowledgements

We are grateful to Gert Nuyts, for performing the XRF measurements, and Dr. Stanislav Trashin for his assistance during the electrochemical experiments. This work is supported by the Research Foundation – Flanders (FWO) (grant G.0687.13) and the University of Antwerp (BOF project).

#### References

- 1 D. I. Fried, F. J. Brieler and M. Fröba, *ChemCatChem*, 2013, **5**, 862–884.
- 2 M. Rabe, D. Verdes and S. Seeger, *Adv. Colloid Interface Sci.*, 2011, **162**, 87–106.
- 3 I. Fenoglio, B. Fubini, E. M. Ghibaudi and F. Turci, *Adv. Drug Deliv. Rev.*, 2011, **63**, 1186–1209.
- 4 J. J. Gray, *Curr. Opin. Struct. Biol.*, 2004, **14**, 110–5.
- 5 K. Nakanishi, T. Sakiyama and K. Imamura, *J. Biosci.*

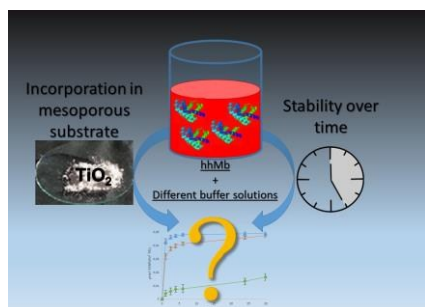


- Bioeng.*, 2001, **91**, 233–44.
- 6 R. A. Latour, 2005, 1–15.
- 7 X. Xu, B. Tian, S. Zhang, J. Kong, D. Zhao and B. Liu, *Anal. Chim. Acta*, 2004, **519**, 31–38.
- 8 S. Zong, Y. Cao, Y. Zhou and H. Ju, *Anal. Chim. Acta*, 2007, **582**, 361–6.
- 9 S. Hudson, J. Cooney and E. Magner, *Angew. Chem. Int. Ed. Engl.*, 2008, **47**, 8582–94.
- 10 Y. Chen, V. Lykourinou, T. Hoang, L.-J. Ming and S. Ma, *Inorg. Chem.*, 2012, **51**, 9156–8.
- 11 J.-J. Feng, J.-J. Xu and H.-Y. Chen, *Electrochem. commun.*, 2006, **8**, 77–82.
- 12 X. Xu, B. Z. Tian, J. L. Kong, S. Zhang, B. H. Liu and D. Y. Zhao, *Adv. Mater.*, 2003, **15**, 1932–1936.
- 13 M. Moritz and M. Geszke-Moritz, *Mater. Sci. Eng. C*, 2015, **49**, 114–151.
- 14 M. Vallet-Regí, a. Rámila, R. P. Del Real and J. Pérez-Pariente, *Chem. Mater.*, 2001, **13**, 308–311.
- 15 M. Vallet-Regí, *Chem. - A Eur. J.*, 2006, **12**, 5934–5943.
- 16 M. Vallet-Regí, F. Balas and D. Arcos, *Angew. Chemie Int. Ed.*, 2007, **46**, 7548–7558.
- 17 S. Wang, *Microporous Mesoporous Mater.*, 2009, **117**, 1–9.
- 18 D. Moelans, P. Cool, J. Baeyens and E. F. Vansant, *Catal. Commun.*, 2005, **6**, 307–311.
- 19 Y. J. Han, J. T. Watson, G. D. Stucky and A. Butler, *J. Mol. Catal. - B Enzym.*, 2002, **17**, 1–8.
- 20 Y. Xian, Y. Xian, L. Zhou, F. Wu, Y. Ling and L. Jin, *Electrochem. commun.*, 2007, **9**, 142–148.
- 21 J. Deere, E. Magner, J. G. Wall and B. K. Hodnett, 2003, **85**, 19–23.
- 22 Y. S. Chaudhary, S. K. Manna, S. Mazumdar and D. Khushalani, *Microporous Mesoporous Mater.*, 2008, **109**, 535–541.
- 23 M. Miyahara, A. Vinu and K. Ariga, *Mater. Sci. Eng. C*, 2007, **27**, 232–236.
- 24 A. Katiyar, S. Yadav, P. G. Smirniotis and N. G. Pinto, *J. Chromatogr. A*, 2006, **1122**, 13–20.
- 25 J. Deere, E. Magner, J. G. Wall and B. K. Hodnett, *J. Phys. Chem. B*, 2002, **106**, 7340–7347.
- 26 Z. Zhou and M. Hartmann, *Chem. Soc. Rev.*, 2013, **42**, 3894–912.
- 27 P. Yang, D. Zhao, D. I. Margolese, B. F. Chmelka and G. D. Stucky, *Chem. Mater.*, 1999, **11**, 2813–2826.
- 28 D. Gu and F. Schüth, *Chem. Soc. Rev.*, 2014, **43**, 313–44.
- 29 M. Guncheva, K. Paunova, M. Dimitrov and D. Yancheva, *J. Mol. Catal. B Enzym.*, 2014, **108**, 43–50.
- 30 L. T. Izrael Živković, L. S. Živković, B. M. Babić, M. J. Kokunešoski, B. M. Jokić and I. M. Karadžić, *Biochem. Eng. J.*, 2015, **93**, 73–83.
- 31 M. Zhao, Y. Zhou, B. Cai, Y. Ma, H. Cai, Z. Ye and J. Huang, *Ceram. Int.*, 2013, **39**, 9319–9323.
- 32 R. An, W. Zhuang, Z. Yang, X. Lu, J. Zhu, Y. Wang, Y. Dong and N. Wu, *Chem. Eng. Sci.*, 2014, **117**, 146–155.
- 33 H. Kim, S. S. Park, J. Seo, C. S. Ha, C. Moon and Y. Kim, *ACS Appl. Mater. Interfaces*, 2013, **5**, 6873–6878.
- 34 M. Heinlaan, A. Ivask, I. Blinova, H. C. Dubourguier and A. Kahru, *Chemosphere*, 2008, **71**, 1308–1316.
- 35 M. Geetha, A. K. Singh, R. Asokamani and A. K. Gogia, *Prog. Mater. Sci.*, 2009, **54**, 397–425.
- 36 M. C. Lomer, R. P. Thompson, J. Commisso, C. L. Keen and J. J. Powell, *Analyst*, 2000, **125**, 2339–2343.
- 37 N. Jia, Y. Wen, G. Yang, Q. Lian, C. Xu and H. Shen, *Electrochem. commun.*, 2008, **10**, 774–777.
- 38 X. Liu, P. K. Chu and C. Ding, *Mater. Sci. Eng. R Reports*, 2004, **47**, 49–121.
- 39 M. Parkes, C. Myant, P. M. Cann and J. S. S. Wong, *Tribol. Int.*, 2014, **72**, 108–117.
- 40 T. Wei, S. Kaewtathip and K. Shing, *J. Phys. Chem. C*, 2009, **113**, 2053–2062.
- 41 S. E. Moulton, J. N. Barisci, a. J. McQuillan and G. G. Wallace, *Colloids Surfaces A Physicochem. Eng. Asp.*, 2003, **220**, 159–167.
- 42 A. Marucco, F. Catalano, I. Fenoglio, F. Turci, G. Martra and B. Fubini, *Chem. Res. Toxicol.*, 2015, **28**, 87–91.
- 43 L. C. Sang, A. Vinu and M.-O. Coppens, *Langmuir*, 2011, **27**, 13828–37.
- 44 Y. Satyawali, S. V. Roy, A. Roevens, V. Meynen, S. Mullens, P. Jochems, W. Doyen, L. Cauwenberghs and W. Dejonghe, *RSC Adv.*, 2013, **3**, 24054–24062.
- 45 S. Ribbens, E. Beyers, K. Schellens, M. Mertens, X. Ke, S. Bals, G. Van Tendeloo, V. Meynen and P. Cool, *Microporous Mesoporous Mater.*, 2012, **156**, 62–72.
- 46 G. A. Parks, *Chem. Rev.*, 1965, **65**, 177–198.
- 47 L. Antonini and M. Brunori, *Hemoglobin and Myoglobin in Their interactions with Ligands.*, North Holland Pub., Amsterdam, 1971.
- 48 W. J. Bowen, *J. Biol. Chem.*, 1949, **179**, 235–246.
- 49 S. Stoll and A. Schweiger, *J. Magn. Reson.*, 2006, **178**, 42–55.
- 50 D. Matulis, J. K. Kranz, F. R. Salemme and M. J. Todd, *Biochemistry*, 2005, **44**, 5258–5266.
- 51 K. Arai, S. Yasuda and A. Kornberg, *J. Biol. Chem.*, 1981, **256**, 5247–5252.
- 52 P. H. Von Hippel and K. Y. Wong, *J Biol Chem*, 1965, **240**, 3909–3923.
- 53 M. Taha and M. J. Lee, *Phys. Chem. Chem. Phys.*, 2010, **12**, 12840–50.
- 54 T. R. Jahn and S. E. Radford, *Arch. Biochem. Biophys.*, 2008, **469**, 100–117.
- 55 D. A. Svistunenko, M. A. Sharpe, P. Nicholls, C. Blenkinsop, N. A. Davies, J. Dunne, M. T. Wilson and C. E. Cooper, *Biochem. J.*, 2000, **351 Pt 3**, 595–605.
- 56 M. Takeuchi, L. Bertinetti, G. Martra, S. Coluccia, M. Anpo, *Appl. Catal. A Gen.* 2006, **307**, 13–20.
- 57 J. Navío, G. Colón, M. I. Litter and G. N. Bianco, *J. Mol. Catal. A Chem.*, 1996, **106**, 267–276.
- 58 C. Morterra, *J. Chem. Soc. Faraday Trans. I* 1988, **84**, 1617–1637.
- 59 G. Socrates, *Infrared and Raman Characteristic Group Frequencies*, Wiley&Sons, New York, 3rd edn., 2001.
- 60 L. Körösi, S. Papp, I. Bertóti and I. Dékány, *Chem. Mater.*, 2007, **19**, 4811–4819.
- 61 W. Gong, *Int. J. Miner. Process.*, 2001, **63**, 147–165.
- 62 G. Parfitt and A. J. Sarkany, *J. Catal.*, 1979, **57**, 476–493.
- 63 K. Madhusudan Reddy, C. V. Gopal Reddy and S. V. Manorama, *J. Solid State Chem.*, 2001, **158**, 180–186.
- 64 H. Pan, X. Wang, S. Xiao, L. Yu and Z. Zhang, 2013, **20**, 561–567.
- 65 H. J. De Jager and L. C. Prinsloo, *Thermochim. Acta*, 2001, **376**, 187–196.
- 66 R. Hofer, M. Textor and N. D. Spencer, *Langmuir*, 2001, **17**, 4014–4020.
- 67 Y. Zhu, T. Ren and Z. Yuan, *New J. Chem.*, 2014, **38**, 1905.
- 68 F. Meersman, L. Smeller and K. Heremans, *Biophys. J.*,

- 2002, **82**, 2635–44.
- 69 M. Miyahara, A. Vinu, K. Z. Hossain, T. Nakanishi and K. Ariga, *Thin Solid Films*, 2006, **499**, 13–18.
- 70 R. Kurrat, J. E. Prenosil and J. J. Ramsden, *J. Colloid Interface Sci.*, 1997, **185**, 1–8.
- 71 T. Itoh, Y. Wada and T. Nakanishi, *Agr. Biol. Chem.*, 1976, **6**, 1083–1086.
- 72 A. P. Snyder, A. Tripathi, J. P. Dworzanski, W. M. Maswadeh and C. H. Wick, *Anal. Chim. Acta*, 2005, **536**, 283–293.
- 73 V. Schmidt, C. Giacomelli and V. Soldi, *Polym. Degrad. Stab.*, 2005, **87**, 25–31.
- 74 F. Bellezza, A. Cipiciani, M. A. Quotadamo, S. Cinelli, G. Onori and S. Tacchi, *Langmuir*, 2007, **23**, 13007–13012.
- 75 S. Van Doorslaer, M. van den Bosch, L. Tilleman and S. Dewilde, *Appl. Magn. Reson.*, 2015, **46**, 421–433.
- 76 L. Tofani, A. Feis, R. E. Snoke, D. Berti, P. Baglioni and G. Smulevich, *Biophys. J.*, 2004, **87**, 1186–95.
- 77 S. Liu and H. Ju, *Electroanalysis*, 2003, **15**, 1488–1493.
- 78 N. F. Hu and J. F. Rusling, *Langmuir*, 1997, **13**, 4119–4125.
- 79 A. J. Bard and L. R. Faulkner, *Electrochemical Methods Fundamentals and Applications*, John Wiley & Sons, Inc., New York, 2nd edn., 1984.
- 80 E. Laviron, *J. Electroanal. Chem. Interfacial Electrochem.*, 1979, **101**, 19–28.
- 81 Y. Ke, Y. Zeng, X. Pu, X. Wu, L. Li, Z. Zhu and Y. Yu, *RSC Adv.*, 2012, **2**, 5676.
- 82 W. Wang, L. Dong, S. Gong, Y. Deng, J. Yu, H. Dong, T. Wang and W. Sun, *Arab. J. Chem.*, 2015.
- 83 G. Zhao, J. J. Xu and H. Y. Chen, *Anal. Biochem.*, 2006, **350**, 145–150.
- 84 Z. Dai, S. Liu, H. Ju and H. Chen, *Biosens. Bioelectron.*, 2004, **19**, 861–867.
- 85 C. H. Taboy, K. M. Faulkner, D. Kraiter, C. Bonaventura and A. L. Crumbliss, *J. Biol. Chem.*, 2000, **275**, 39048–39054.

## Table of Contents

View Article Online  
DOI: 10.1039/C6CP08585G



An insight in the buffer role in proteins stability and adsorption on mesoporous materials.