

# Novel BUF2-magnetite nanobioconjugates with cell-penetrating abilities

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**Introduction:** One of the major challenges of modern pharmacology is the development of systems for the delivery of therapeutic molecules in a controlled and localized manner. One strategy is to use nanostructured supports, which are well suited to carry a large number of molecules on a per mass basis. A major challenge for these supports is, however, their limited ability to bypass the cell membrane. Recent studies propose that to overcome this issue, potent translocating cell-penetrating peptides (CPPs) can be conjugated to their surfaces.

**Methods:** Here, we conjugated the antimicrobial CPP buforin II (BUF2) to the surface of magnetite nanoparticles to enhance their cell penetration. Conjugates were characterized via Fourier transform infrared spectroscopy, dynamic light scattering, and thermogravimetric analysis, and their biocompatibility was corroborated. The conjugates were delivered in both bacterial and mammalian cells demonstrating the intracellular inclusion in THP-1 cells for the first time.

**Results:** Despite the promising outcome, our studies showed that the obtained conjugates failed to maintain the native antimicrobial activity of BUF2. We hypothesize that to overcome this issue, a flexible linker can be inserted prior to conjugation.

**Conclusion:** Our study highlights the potential of BUF2-magnetite conjugates as cell-penetrating vehicles for the targeted delivery of pharmacological agents. This provides support for the idea of a promising combined drug delivery and antimicrobial peptide therapy.

**Keywords:** BUF2-magnetite, antimicrobial activity, cell-penetrating peptides, nanomaterials, drug delivery

## Introduction

Drug delivery systems have attracted considerable attention in the medical arena due to possibilities for treating or eliminating cells of certain diseases in a targeted manner, ie, without affecting the surrounding tissues or cells. Despite their potential, implementation of most drug delivery systems at the clinical level has been largely limited due to challenges such as low membrane permeability and short circulation half-time.<sup>1</sup> Recent reports suggest that by incorporating cell-penetrating peptides (CPPs) into these systems, these issues can be successfully addressed. This approach takes advantage of CPPs' small size, ease of production, and particularly their ability to spontaneously translocate across the cell membrane or even complex structures such as the blood-brain barrier.<sup>1</sup>

Buforin II (BUF2) is a cationic (+6) and amphipathic 21-residue CPP capable of killing bacteria by efficiently crossing the cell membrane without damaging it and subsequently binding to DNA and RNA, thereby interrupting the replication cycle.<sup>2-4</sup> The C-terminal region of BUF2 is associated with the antimicrobial mechanism, while a proline hinge in the structure is responsible for aiding the cell membrane penetration without promoting significant disruption.<sup>3</sup> In spite of its potent antimicrobial activity,

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BUF2 is prone to proteolytic degradation by endogenous proteases, which significantly reduces its *in vitro* and *in vivo* half-life and consequently its therapeutic value.<sup>4-6</sup>

A strategy to maintain the CPPs biological activities as well as extending their stability, efficacy, and lifespan is the immobilization on nanomaterials.<sup>7</sup> Immobilization of CPPs on magnetite iron-oxide nanoparticle (MNP) surfaces has been exploited for the development of anti-bacterial coatings.<sup>8</sup> Additionally, MNPs have been widely used for a number of biological applications including bioseparations, agents for MRI contrasting, magnetic fluid hyperthermia for cancer treatment, and drug delivery for the treatment of various conditions.<sup>9-11</sup> MNPs generally exhibit low cytotoxicity depending on factors such as dose, chemical composition, size, structure, surface chemistry, and route of administration.<sup>12,13</sup> Moreover, they have been extensively used as immobilization supports to improve the stability of a variety of biomolecules including epidermal growth factor,<sup>10</sup> albumin,<sup>14</sup> and bacitracin.<sup>8</sup> Immobilization of biomolecules on MNPs is also advantageous mainly due to the increased number of active molecules per unit mass and the ease of transport and manipulation under magnetic fields.<sup>15</sup> Accordingly, this work was aimed at exploring the immobilization of BUF2 on magnetite as cell translocating vehicles to enable highly targeted drug delivery applications in mammalian cells. Here, we also worked on estimating possible changes in the antimicrobial activity of BUF2 upon immobilization.

## Materials and methods

### Materials

Iron (II) chloride tetrahydrate (98%) and sodium hydroxide (NaOH) (98%) were obtained from PanReac AppliChem. Iron (III) chloride hexahydrate (97%), dimethyl sulfoxide (99.5%), tetramethylammonium hydroxide (TMAH) (25%), (3-aminopropyl) triethoxysilane (APTES) (98%), carboxymethyl cellulose (CMC) (98%), N-[3-(dimethylamino)propyl]-N'-ethylcarbodiimide hydrochloride (EDC) (98%), and glutaraldehyde (25%) were purchased from Sigma-Aldrich Co. DMEM, RPMI 1640 medium, FBS, and trypsin EDTA were obtained from Biowest. Penicillin/streptomycin (P/S) was purchased from Lonza. BUF2 (BUF2-TRSSRAGLQFPVGRVHLLRK) and fluorescein isothiocyanate (FITC)-BUF2 were synthesized by the Peptide Synthesis Facility at Pompeu Fabra University and GL Biochem (Shanghai, China). Purification was performed by high-pressure liquid chromatography (>95%), and masses were confirmed via mass spectrometry as we did in a previous work.<sup>4</sup> Vero Cells (ATCC® CCL-81), THP-1 Cells (ATCC®

TIB-202) were used for cytotoxicity and delivery assays. Bacterial strain used was *Escherichia coli* (ATCC 25922).

### Synthesis and functionalization of magnetite nanoparticles

Nanoparticles were obtained by co-precipitating 500 mM ferric chloride and 250 mM ferrous chloride (molar ratio 1:2) in the presence of a 5 M NaOH aqueous solution at 90°C. The solution was left to cool for at least 30 minutes prior to use. Nanoparticles were subsequently washed with distilled water to remove excess reagents aided by a permanent magnet. The MNP suspension (100 mg) was adjusted to pH 11.3 and sonicated for 10 minutes. TMAH solution (2 mL, 25% (v/v)) was then added to the magnetite solution and sonicated for 10 minutes. APTES (200 µL) was added to the magnetite solution for silanization along with 100 µL of acetic acid. Silanized nanoparticles were washed five times with milli Q water to remove excess of APTES aided by a strong permanent magnet and subsequently resuspended in 30 mL of milli Q water. Silanization rendered free amine groups on the surface of the nanoparticles to subsequently conjugate the N-terminal and C-Terminal of BUF2 to the nanoparticles. Free amine groups were also used to conjugate Phalloidin Alexa Fluor 488 via glutaraldehyde as crosslinker (see the following section for details on the conjugation protocol).

### Nanoparticle characterization

Nanoparticle size was confirmed by dynamic light scattering (DLS, Zeta-Sizer Nano-ZS; Malvern Instruments, Malvern, UK), and the sample was sonicated for 6 hours in water at 37°C. Surface modifications of MNPs with APTES and BUF2 were confirmed via Fourier transform infrared spectroscopy (FTIR) using a Bruker Alpha II FTIR Eco-ATR (Bruker Optik GmbH, Ettlingen, Germany). Spectra were collected in the range of 4,000–400 cm<sup>-1</sup> with a spectral resolution of 2 cm<sup>-1</sup>. Thermogravimetric analysis (TGA) was carried out with 10 mg of samples in a simultaneous TGA/differential scanning calorimetry (TA Instruments, Newcastle, DE, USA). Analyses were performed by ramping up the temperature at a rate of 10°C/min from 25°C to 800°C in a nitrogen atmosphere with a gas flow rate of 100 mL/min. TGA data are shown as the percentage of mass loss as a function of temperature.

### In silico analysis of BUF2 in vitro and in vivo half-life

A prediction of BUF2 stability and half-life *in vivo* and *in vitro* was completed in silico via the Antimicrobial Peptide

Database (<http://aps.unmc.edu/AP/main.php>). ProtParam tool relies on the “N-end rule”, which relates the half-life of a protein to the identity of its N-terminal residue.<sup>16</sup>

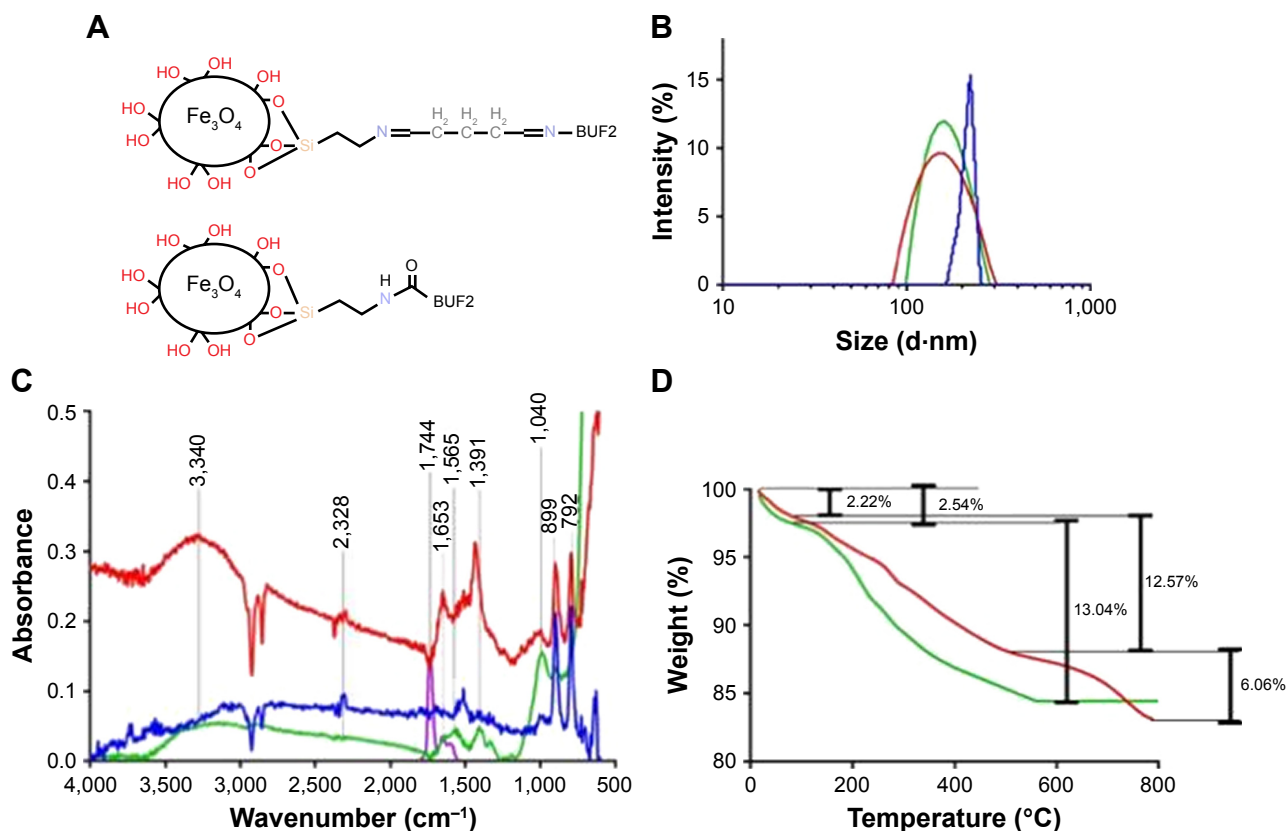
## Conjugation of BUF2 on magnetite nanoparticles

To obtain BUF2-magnetite conjugates (Figure 1A), 100 mg of magnetite-APTES nanoparticles were freeze dried for 24 hours and subsequently dispersed in 30 mL of distilled water and sonicated for 30 minutes. This was followed by addition of 1 mL of glutaraldehyde 2% (v/v). The reaction mixture was left at rest for 30 minutes. Excess of glutaraldehyde was removed by washing the nanoparticles five times with milli Q water with the aid of a strong permanent magnet. Finally, 500  $\mu$ L of BUF2 solution at 1 mg/mL in sterile PBS 1X was added to the nanoparticles solution and left to react overnight. This solution was then washed four times with distilled water and freeze dried for 24 hours. Alternatively, BUF2-FITC-magnetite conjugates were prepared by conjugation to the C-terminal with the aid of 2 equivalents of EDC and N-hydroxysuccinimide with respect to the free amine

groups on the surface of the nanoparticle. The reaction was conducted under continuous agitation at room temperature for 24 hours on 100 mg of well suspended magnetite-APTES nanoparticles in distilled water. After conjugation, samples were thoroughly washed with milli Q water with the aid of a strong permanent magnet.

## Cell culture and cytotoxicity assay

The African green monkey fibroblast-like kidney cells (Vero ATCC CCL-81), their maintenance, and subculture were described in a previous work.<sup>4</sup> Cell monolayers were trypsinized, washed with culture medium, and plated in a flat-bottomed 96-well microtiter plate with  $3 \times 10^4$  cells per well, and then cell toxicity was determined by lactate dehydrogenase (LDH) assay kit (Thermo Fisher Scientific, Waltham, MA, USA). Vero cells were exposed to magnetite and BUF2-magnetite conjugates in serially diluted samples starting at 100  $\mu$ g/mL down to 6.25  $\mu$ g/mL during 24 and 72 hours of incubation at 37°C and 5% CO<sub>2</sub>. Cells without treatment were used as a negative control, and for the positive control, cells were treated with LDH 10X lysis buffer.



**Figure 1** (A) Schematic of the BUF2-magnetite by conjugation with glutaraldehyde (top) and EDC (bottom) as crosslinkers. (B) DLS histograms for size distribution of magnetite nanoparticles at different functionalization steps. Bare magnetite (red), magnetite functionalized with APTES (green), and magnetite conjugated with BUF2 (blue). (C) FTIR spectra of magnetite (blue), magnetite with APTES (red), BUF2 (purple), and magnetite-APTES-BUF2 (green). (D) TGA of magnetite (green) and magnetite conjugated with BUF2 (red). The first weight loss steps (2.54% and 2.22%) represent the dehydration of the samples. Second weight loss steps (12.57% and 13.04%) correspond to physically absorbed organic solvents. The final weight loss step (6.06%) is attributed to the detachment of BUF2 from the nanoparticle's surface.

**Abbreviations:** BUF2, buforin II; EDC, N-[3-(dimethylamino)-propyl]-N'-ethylcarbodiimide hydrochloride; DLS, dynamic light scattering; APTES, (3-aminopropyl) triethoxysilane; FTIR, Fourier transform infrared spectroscopy; TGA, thermogravimetric analysis.

After incubation, plates were centrifuged at  $300\times g$  for 5 minutes, and the supernatant was transferred to a new plate followed by addition of 50  $\mu\text{L}$  of LDH reaction mixture during 30 minutes. The absorbance from the LDH released was quantified at 490 nm wavelength and 650 nm as reference wavelength, using a microplate reader. Cell viability was calculated by triplicate readings as the mean absorbance of LDH activity by the following formula: % Cell viability =  $(M_{\text{sample}} - M_{\text{ncontrol}})/(M_{\text{pcontrol}} - M_{\text{ncontrol}}) \times 100$ .

### Antimicrobial activity assay

Antimicrobial activity of conjugates and peptide was determined by the broth microdilution assay, as described by Park et al.<sup>3</sup> Two concentrations of each BUF2-magnetite (500 and 100  $\mu\text{g}/\text{mL}$ ) treatment were assayed against *E. coli* (ATCC 25922), as well as two concentrations of BUF2 (100 and 3.125  $\mu\text{M}$ ). Bacterial cells were also treated with a negative (no treatment) and a positive control (10  $\mu\text{M}$  gentamicin). Fifty microliters of each treatment was seeded in triplicate on a 96-well flat bottom microplate (TPP). Bacterial cells were cultured overnight in LB Broth at  $37^\circ\text{C}$ , and an aliquot of this culture was taken and incubated in fresh medium until mid-logarithmic phase was achieved. Cells were harvested by centrifugation, washed three times with a 10 mM sodium phosphate buffer (pH 7.4), and diluted 1:10,000 in the same buffer. Fifty microliters of the diluted cell suspension was added to 50  $\mu\text{L}$  of each of the treatments assayed and incubated for 3 hours at  $37^\circ\text{C}$ . After incubating the mixture, 100  $\mu\text{L}$  of fresh LB Broth was added to each well and incubated for 16 more hours at  $37^\circ\text{C}$ . Inhibitory growth effects were determined by absorbance at 620 nm. The presence of NaCl and CMC on the antimicrobial activity of the peptide and conjugates was also assayed.

### Hemolysis assay

Blood was obtained from a healthy donor in a vacutainer blood tube with EDTA. The hemolytic activities of magnetite and BUF2-magnetite conjugates were evaluated according to a method described previously.<sup>4</sup> Serially diluted concentrations of magnetite and BUF2-magnetite conjugates were prepared for the test in concentrations ranging from 1,500  $\mu\text{g}/\text{mL}$  to 2.93  $\mu\text{g}/\text{mL}$ . The hemolytic assay was performed in a 96-well microtiter plate; 100  $\mu\text{L}$  of either magnetite or BUF2-magnetite and 100  $\mu\text{L}$  of the diluted red blood cells were incubated for 1 hour at  $37^\circ\text{C}$ . The positive control was an erythrocyte suspension incubated with ultrapure water (100% hemolysis), and the negative control was an erythrocyte suspension incubated with PBS 1X (0% hemolysis).

## Delivery of BUF2-magnetite conjugates in mammalian cells and *E. coli*

Translocation of conjugates and peptide in mammalian cells and bacteria was imaged via confocal microscopy. Briefly, fluorescently labeled BUF2 (BUF2-FITC) was conjugated to magnetite as described earlier using both glutaraldehyde and EDC as crosslinkers. BUF2-FITC and BUF2-FITC-magnetite conjugates were then delivered to THP-1 cells at a dilution ratio of 1:100 in 1 mL of RPMI (10% FBS and 1% P/S). Samples were incubated for 1 hour at  $37^\circ\text{C}$  and 5%  $\text{CO}_2$ . Also, BUF2-FITC was delivered to *E. coli* (0.5 McFarland standard) at a dilution ratio of 1:100 in 1 mL of PBS 1X. Samples were incubated for 1 hour at room temperature. Imaging was conducted in a confocal laser scanning microscope Olympus FV1000 with a 60x/NA 1.35 oil immersion objective. Finally, image analysis was performed in ImageJ and Fiji.

## Results and discussion

### Nanoparticle characterization

Size distribution of MNPs after each functionalization step was determined by DLS of samples suspended in distilled water at 1 mg/mL. Figure 1B shows that the size distribution by intensity increases after each functionalization step. Bare MNPs exhibited a mean hydrodynamic diameter of 161 nm with a polydispersity index (PI) of 14.66%. After silanization, the mean hydrodynamic diameter increased to 170 nm with a PI of 23.08%. Finally, conjugation of BUF2 led to a mean hydrodynamic diameter of 213.5 nm and a PI of 12.48%. The sizes and PIs obtained here are comparable with those reported elsewhere for the same synthesis method.<sup>17,18</sup>

Surface modifications of MNPs were confirmed and shown in Figures 1C and D by FTIR and TGA. Figure 1C shows the FTIR spectra of bare, and APTES- and BUF2-conjugated MNPs. FTIR spectrum of bare nanoparticles exhibited absorption bands at around 632 and 585  $\text{cm}^{-1}$ , which can be attributed to the Fe–O bond of iron oxide.<sup>16,19</sup> Silanization with APTES was confirmed by the presence of the Si–O stretching vibration at about 1,040  $\text{cm}^{-1}$  as well as bands at 1,391 and 1,653  $\text{cm}^{-1}$ , which can be explained by bending vibrations of C–H and N–H, respectively.<sup>20</sup> Finally, conjugation of BUF2 was verified by the presence of the amide I band at 1,650  $\text{cm}^{-1}$  and NH and CN stretching vibrations of the peptide at 1,565  $\text{cm}^{-1}$ .<sup>20</sup>

The presence of BUF2 conjugates on the surface of the MNPs was also assessed via TGA (Figure 1D). Magnetite and BUF2-magnetite conjugates showed a first weight loss of 2.54% and 2.22% due to the dehydration of the samples.



**Table 1** APD in silico stability and half-life of the BUF2 peptide

Instability index	Half-life	In vivo		
		In vitro	In vivo	
		Mammalian reticulocytes	Yeast	<i>E. coli</i>
84.2	Unstable	7.2 hours	>20 hours	>10 hours

**Abbreviations:** APD, Antimicrobial Peptide Database; *E. coli*, *Escherichia coli*.

Second weight loss for bare magnetite approached 13.04%, while for the BUF2-magnetite conjugates was of 12.57%. These losses can be attributed to physically absorbed organic compounds left by the synthesis and functionalization processes. Finally, the detachment of BUF2 from the magnetite is estimated with the final weight loss step (6.06%). This roughly corresponds to  $3 \times 10^5$  BUF2 molecules/MNP.

## In silico analysis of BUF2

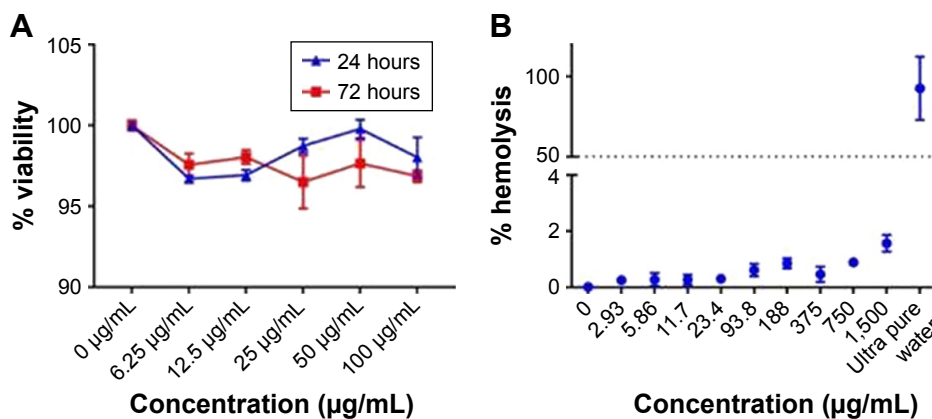
Table 1 summarizes the in silico stability and half-life in vitro and in vivo of BUF2 as predicted by the Antimicrobial Peptide Database. In this case, the instability index estimates the stability of peptide molecules in a test tube (reference value of 40 for stable molecule). For BUF2, the index was 84.20, which indicates low stability and thereby the necessity of finding a method to increase BUF2 stability without detrimentally impacting its antimicrobial capacities.<sup>21</sup> In contrast, half-life is a prediction of the time it takes for half of the amount of protein/peptide in a cell to disappear after its synthesis in the cell. This parameter is related with the N-end rule originated from the observations that the identity of the N-terminal residue of a protein plays an important role in determining its stability in vivo.<sup>22</sup> In this regard, the predicted half-life indicates a short life while it is tested in vitro, while its in vivo half-life is greater than 10 hours.

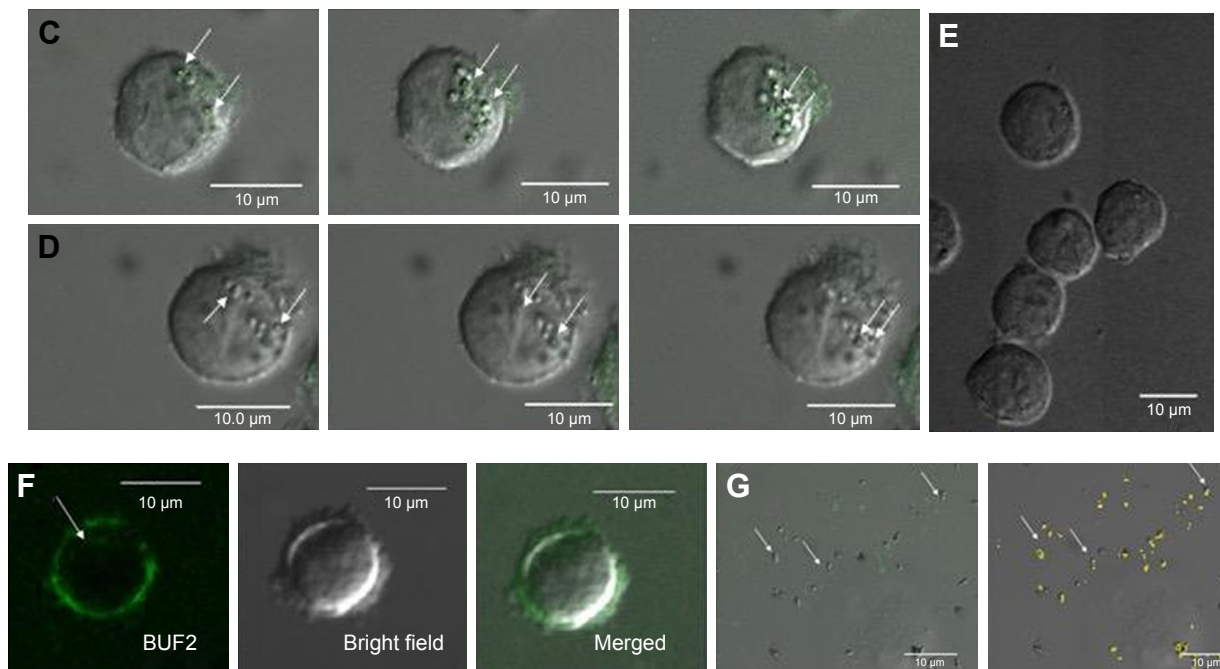
## Biocompatibility and cellular delivery of BUF2-magnetite conjugate

In accordance with the ISO 10993 standard about biocompatibility of nanomaterials, we assessed the cytotoxic (100–6.25  $\mu\text{g/mL}$ ) effect on Vero cells (ATCC® CCL-81) and hemolytic effect in red blood cells (1,500–2.93  $\mu\text{g/mL}$ ) of the BUF2-magnetite conjugate (Figure 2A and B). The results showed that the viability of Vero cells exceeded 95% in the presence of BUF2-magnetite conjugate for all evaluated concentrations. Similarly, hemolysis levels remained below 5%, which complied with the followed ISO standard. The well documented antimicrobial activity of BUF2 includes a translocation step across bacterial membranes without promoting disruption.<sup>23</sup> Here, we explored whether this translocation ability is preserved after immobilization by exposing THP-1 cells to the BUF2-magnetite conjugate. Our confocal images corroborated for the first time, to the best of our knowledge, that BUF2-FITC-magnetite conjugates are capable of translocating the membrane of THP-1 cells without significantly reducing their viability (Figure 2C and D). This is in contrast with the bare MNPs that failed to translocate (Figure 2E) and the peptide alone that remains in THP-1 cell membrane (Figure 2F). We also confirmed the entrance of our conjugate into *E. coli* (Figure 2G).

## Antimicrobial activity

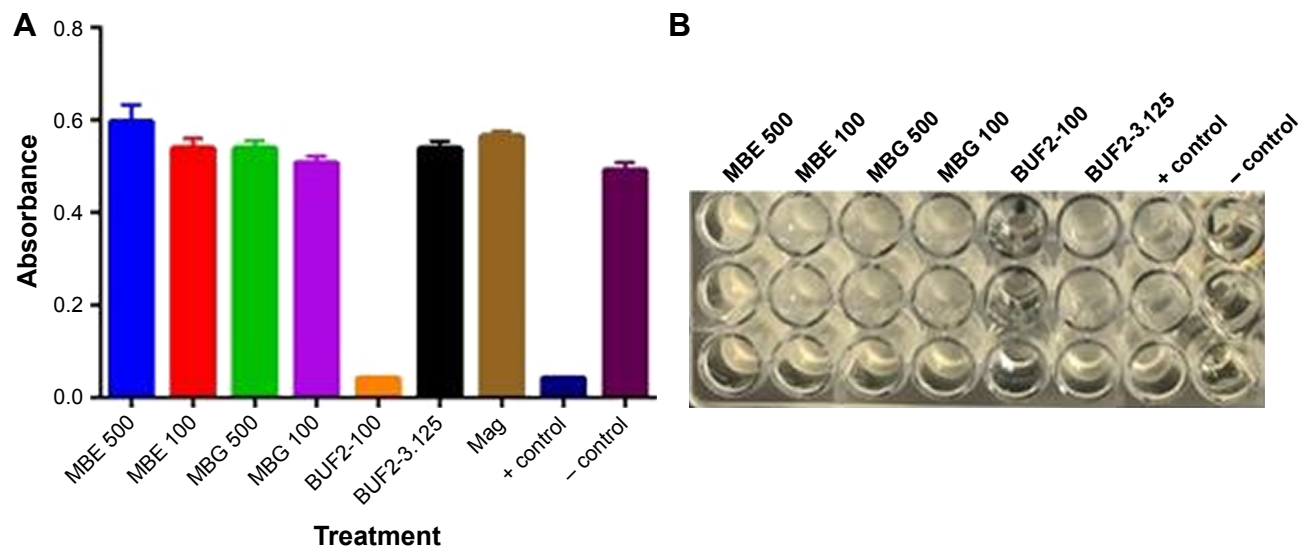
Preliminary results showed that as opposed to BUF2, BUF2-magnetite conjugates failed to produce any significant antimicrobial activity. This was the case for conjugates prepared with both EDC (MBE) and glutaraldehyde (MBA) as crosslinkers (Figure 3A and B). In an effort to understand the possible causes for this result, we conducted variations in the components of the growth medium. We first

**Figure 2** (Continued)



**Figure 2** (A) Cytotoxicity of BUF2-magnetite conjugates as tested by LDH assays after 24 and 72 hours. Cell viability remained above 95% for all studied concentrations. (B) Assessment of the hemolytic effect of BUF2-magnetite conjugates at different concentrations. Data points are mean data of each concentration. In all cases, hemolysis was below 5%. (C) Confocal microscopic images of effective cellular internalization (THP-1 cells) of BUF2-FITC-magnetite as prepared by conjugation via glutaraldehyde (arrow in inset). Scale bar corresponds to 10  $\mu$ m. (D) Confocal microscopic images of effective cellular internalization (THP-1 cells) of BUF2-FITC-magnetite as prepared by conjugation via EDC (arrow in inset). Scale bar corresponds to 10  $\mu$ m. (E) Confocal microscopic image showing failed internalization of fluorescently labeled bare magnetite (negative control). (F) Confocal microscopic images of effective cellular internalization of BUF2-FITC (arrow in inset). Scale bar corresponds to 10  $\mu$ m. (G) Confocal microscopic images of internalization of BUF2-FITC into *E. coli* (arrow in inset). The image on the right shows the threshold mask to BUF2-FITC to remove the fluorescence background. Scale bar corresponds to 10  $\mu$ m.

**Abbreviations:** BUF2, buforin II; LDH, lactate dehydrogenase; EDC, N-[3-(dimethylamino)-propyl]-N'-ethylcarbodiimide hydrochloride; FITC, fluorescein isothiocyanate; *E. coli*, *Escherichia coli*.



**Figure 3** (A) Antimicrobial activity against *Escherichia coli* of magnetite (brown) and BUF2-magnetite (500  $\mu$ g/mL, green; 100  $\mu$ g/mL, purple) as prepared by conjugation via glutaraldehyde. BUF2-magnetite as prepared by conjugation via EDC at the same concentrations (blue and red, respectively). Comparison with the antimicrobial activity of BUF2 at 100  $\mu$ M (orange) and 3.125  $\mu$ M (black). The positive control was *Escherichia coli* in gentamicin at 10  $\mu$ M, and suspension containing only cells was the negative control. (B) Actual image of the microplate where the antimicrobial assay was conducted. BUF2 at 100  $\mu$ M and the positive control showed insignificant turbidity when compared with the rest of the treatments.

**Abbreviations:** BUF2, buforin II; EDC, N-[3-(dimethylamino)-propyl]-N'-ethylcarbodiimide hydrochloride; Mag, magnetite; MBE, BUF2-magnetite conjugated with EDC; MBG, BUF2-magnetite conjugated with glutaraldehyde.

started by carrying out the tests in the presence of 0.1 M NaCl, which showed BUF2 inhibition and no impact for the conjugates, (data not shown). Next set of experiments responded to the notion that conjugates were not fully suspended in the medium during the test. Accordingly, CMC was added to the medium to increase its viscosity and maintain the conjugates suspended throughout the experiment. No inhibitory bacterial growth effect was observed in the presence of CMC for the conjugates, and a complete loss of activity was detected for BUF2 (data not shown). This was attributed to interactions of BUF2 with a charged moiety present in the CMC structure. Magnetite alone showed no antimicrobial or growth promoting activity. Taken all together, our findings suggest that upon immobilization, regions of BUF2 responsible for interaction with the replication machinery of *E. coli* were blocked, which led to a total loss of the antimicrobial functionality. A possible avenue to overcome this issue is to conjugate the peptide with the aid of a flexible spacer that helps to maintain the structural integrity.

## Conclusion

We presented a feasible approach for immobilization of BUF2 on MNPs. Upon immobilization, BUF2 loses antimicrobial activity most likely due to the blockage of C-terminal residues involved in abrogating the replication machinery of *E. coli*. BUF2-magnetite conjugates are still capable of bypassing mammalian and bacterial membranes very effectively without promoting disruption as evidenced by the high biocompatibility of LDH assays. Our study highlights the potential of BUF2-magnetite conjugates as cell-penetrating vehicles for the targeted delivery of pharmacological agents. This provides support for the idea of a promising combined drug delivery and antimicrobial peptide therapy.

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## Disclosure

The authors report no conflicts of interest in this work.

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