Caffeine vehiculation into alpha-zirconium phosphate: a multifunctional intercalation product and its application for modified topical release

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Caffeine possesses many important biological effects and is known to be a model drug for topical applications. Zirconium phosphate (ZrP) is able to intercalate basic molecules and it is an effective drug carrier. The vehiculation of drugs in lamellar solids can finely tune their properties. In this article the first intercalation of caffeine in ZrP is reported. The intercalation strategy was designed on the basis of the chemical properties of the host and guest, and the synthesized product was fully chemically–physically characterized and tested for topical release using the Franz cell diffusion system.

Introduction

Caffeine (1,3,7-trimethylxanthine, Fig. 1) has been a part of our global history for thousands of years; each country has its own story and source of caffeine: there are 63 different species of plants known to contain caffeine.1

Caffeine was first extracted from cocoa beans into its purest form, a white powder, in the 1820s by a German Scientist named Friedrich Ferdinand Runge.

It is a weak base belonging to the methylxanthine class, and it acts as a central nervous system stimulant. The biological effects of caffeine are mainly due to its ability to reversibly block the action of adenosine on its receptors.2

Caffeine (CAF) has many interesting activities, because it can treat and prevent bronchopulmonary dysplasia in premature infants;3 it may confer a modest protective effect against some diseases,4 including Parkinson’s disease,5 and it can improve the effect of anticancer drugs like 5-fluorouracil and cisplatinum.6 Recently Li et al.7 demonstrated that caffeine has an important activity in protecting skin from oxidative stress-induced senescence through the activation of autophagy. In topical application caffeine is used as an active ingredient in anti-cellulite products. CAF stimulates the degradation of fats during lipolysis through the inhibition of the phosphodiesterase activity preventing excessive accumulation of fat in cells. It also increases the microcirculation of blood in the skin.8 Moreover topical application of caffeine to mouse skin after UV irradiation promotes the deletion of DNA-damaged keratinocytes and may partially diminish photodamage as well as photocarcinogenesis.9

Caffeine is also recommended by the OECD as test substance for topical applications because it has been widely tested. As a hydrophilic drug, it possesses a log P of −0.07 that corresponds to a slow permeation in topical applications. Moreover it forms aggregates in aqueous solutions.10 Due to its high crystallization tendency, in formulation with concentration higher than 1% caffeine needs to be pre-solubilized in alcohol or an alcohol–water mixture.11

Zirconium phosphate of alpha type is an acidic, inorganic cation exchange material which has a layered structure with formula Zr(HPO4)2·H2O.12,13 Zirconium phosphate can be easily functionalized through intercalation reactions with bases.14 This solid has been widely studied in academic literature with the aim of vehiculating bioactive organic molecules, as a drug carrier for cancer therapy,15–18 anti-inflammatory drugs19 and also antibiotics.20 ZrP is composed of stacking layers made of planes of Zr atoms bonded, on both sides, to monohydrogen phosphate groups. Each phosphate group is bonded to three Zr atoms of the plane, while each zirconium is

Fig. 1 Chemical structure of caffeine.
octahedrally coordinated by six oxygens of six different mono-
hydrogen phosphate groups. The water molecules are located
in the zeolitic cavities of the interlayer region.\textsuperscript{21}

In the past, several authors have reported that caffeine
can be successfully intercalated in lamellar solids reaching
interesting goals.\textsuperscript{22–28} Herein we report the first intercalation of CAF on ZrP, obtaining an intercalation product that has been
successfully used to prepare a stable emulsion. The cream has
been tested for a topical application by the OEDC method.\textsuperscript{29}
The results demonstrate that the new intercalation compound
opens new frontiers in the development of high-tech products
for topical application.

Materials and methods

1. Chemicals and solvents

Propylamine, monobasic potassium phosphate (KH\textsubscript{2}PO\textsubscript{4}) and
dibasic sodium phosphate heptahydrate (Na\textsubscript{2}HPO\textsubscript{4}·7 H\textsubscript{2}O) were
purchased from Sigma-Aldrich. Zirconyl chloride octahydrate
was supplied by MEL Chemicals and 75% phosphoric acid was
purchased from C.M. Chimica. Anhydrous caffeine, xanthan
gum, cetearyl alcohol, paraffin oil, petrolatum, steareth-2,
ceteareth-20, glyceryl stearate, and sodium chloride were sup-
pplied by ACEF. Strat-M membranes were provided by Merck.
Solvents and materials were of reagent grade and used without
further purification.

2. ZrP synthesis

Crystalline zirconium phosphate Zr\textsubscript{2}(HPO\textsubscript{4})\textsubscript{3}·H\textsubscript{2}O of alpha type
was obtained by refluxing zirconyl chloride (ZrOCl\textsubscript{2}·8H\textsubscript{2}O) in a
10 M phosphoric acid solution for 48 h.\textsuperscript{13} The residual solid
was centrifuged, washed 3 times with water and dried in an
oven at 60 °C. The dried solid was properly dissolved and ICP-
OES elemental analysis, combined with thermogravimetric
analysis, allowed the determination of the following chemical
formula: Zr\textsubscript{2}(HPO\textsubscript{4})\textsubscript{3}·H\textsubscript{2}O.

3. Intercalation of CAF in ZrP

A colloidal dispersion of exfoliated zirconium phosphate was
prepared by titrating dropwise 5 g of solid ZrP suspended in
330 mL of water with 165 mL of 0.1 M propylamine solution
at room temperature and under vigorous stirring. Then, a
solution of HCl was added to the colloidal dispersion until
complete exchange of the propylammonium ions with protons,
regenerating the acid form of α-ZrP in a gel dispersion. In a
second step, a solution containing 6.4 g of CAF in 250 mL of
water was added to the gel and the mixture was stirred for 24 h
at room temperature. The reaction mixture was centrifuged,
washed twice with deionized water and dried in an oven at
60 °C. The white solid obtained (ZrP-CAF) was finely ground
with a mortar.

4. Extraction of CAF from ZrP-CAF

0.5 g of ZrP-CAF was suspended in 50 mL of 0.1 N aqueous
solution of NaCl. Then the mixture, kept under magnetic
stirring, was slowly titrated with 0.1 N NaOH until complete
formation of Na\textsubscript{2}Zr(PO\textsubscript{4})\textsubscript{2}.\textsuperscript{30} The obtained dispersion was
filtered and the solution was analyzed three times by UV-vis
in order to determine caffeine content in the solid. The
obtained value was 31.8 ± 0.6 wt%.

5. Cream preparation

A cream containing pure caffeine as a reference was prepared.
4% of caffeine was suspended in the following cream: xanthan
gum 0.2%, steareth-2 1%, ceteth-20 1.5%, ceteyrl alcohol 2%,
petrolatum 2%, glyceryl stearate 5%, paraffin oil 8%, and
water 76.3%. In the second cream an amount of ZrP-CAF
Corresponding to 4% of caffeine was dispersed in steareth-2
1%, ceteth-20 1.5%, ceteyrl alcohol 2%, petrolatum 2%,
glyceryl stearate 5%, paraffin oil 8%, and water 68.3%.

6. Analytical procedures and instrumentation

6.1 Inductively coupled plasma-optical emission spec-
trometry. The Zr and P content in the synthesized ZrP was
determined using an ICP-OES Varian 700-ES series, after
dissolving the sample using few drops of HF and deionized
water. The solution was analyzed after a proper dilution.

6.2 Thermal analysis. Thermogravimetric analysis (TGA)
and differential scanning calorimetry (DSC) of the samples
were carried out with an STD Q600 thermal analyzer TA
Instrument, in air flow with a heating rate of 10 °C min\textsuperscript{-1}.

6.3 UV-Vis spectrophotometry. UV-VIS analysis was carried out using a UV-visible double-beam spectrophotometer (Jasco
V-750). The CAF released in the permeation tests was detected
by UV-vis spectrophotometry at the wavelength of maximum
absorption of caffeine in the phosphate buffer (\(\lambda = 273\) nm).
A suitable calibration was carried out by dissolving known
amounts of caffeine in a suitable volume of phosphate buffer.
Eight standards and a blank sample were then prepared and
analyzed in order to obtain the calibration curve (Fig. 2). The
correlation coefficient of the calibration \(r\) was 0.99973. Tests
were carried out in triplicate and the results were reported as
average values. Cuvettes of 0.1 cm path length were filled with

![Fig. 2](image-url)
the 0.4 mL samples and the concentration was measured using the buffer solution as blank.

6.4 FT-IR analysis. FT-IR spectra were recorded at room temperature using a FT-IR-4600 Jasco. Typically, each spectrum was obtained in the spectral region from 600 to 4000 cm\(^{-1}\). For data collection an ATR crystal in ZnSe has been used.

6.5 X-ray powder diffraction. The X-ray powder diffraction (XRPD) patterns were recorded with a Bruker D2 Phaser diffractometer operating at 30 kV and 15 mA, a step size of 0.02 \(2\theta\) degrees and time per step of 1 s, using the Cu K\(\alpha\) radiation and a multistrip LYNXEYE SSD160 detector.

6.6 SEM analysis. The morphology of the samples was investigated with a FEG LEO 1525 scanning electron microscope (FE-SEM). FE-SEM micrographs were collected by depositing the samples on a stub holder and after a sputter coating with chromium for 20 s.

6.7 Permeation studies. Permeation studies from the creams containing pure caffeine and ZrP-CAF were performed using a Franz Diffusion Cell (PermeGear, Inc., Bethlehem, P.A., 20 mm diameter), following the OECD guideline for measuring skin absorption and dermal delivery in vivo.\(^{29}\) Buffer solution at pH 7.4 was chosen as acceptor solution, and prepared as follows: 0.6 potassium dihydrogen phosphate, 6.4 g disodium hydrogen phosphate, 5.85 g sodium chloride, and deionized water to 1 L.\(^{31}\) The system was equilibrated at 32 °C was chosen as acceptor solution, and prepared as follows: 0.6 potassium dihydrogen phosphate, 6.4 g disodium hydrogen phosphate, 5.85 g sodium chloride, and deionized water to 1 L.\(^{31}\) The system was equilibrated at 32 °C using an external circle bath. The solubility test of caffeine in the receptor fluid at 32 °C was carried out: the molecule was really soluble in the chosen fluid (46 g L\(^{-1}\)). The amount of caffeine to be placed in the donor chamber was fixed at 8 mg (equivalent to 200 mg of cream), so that the fluid would not act as a barrier to absorption. The amount of cream was also calculated in order to be in the range indicated in the method for mimicking human exposure. The amount of cream in the chosen lamellar solid, a ZrP gel in water was used: in the ZrP gel protons are solvated by water and more accessible than in the solid form.\(^{35}\) The gel was obtained regenerating the acid form by adding HCl to a colloidal dispersion of ZrP-PrNH\(_2\) in water, according to the work of Casciola et al.\(^{36}\) Then a water solution of CAF was added to the gel, stirred 24 h and recovered by centrifugation. The product was dried in an oven to obtain a fine micropowder and characterized by XRPD, FT-IR, TGA, and DSC. The chemical-physical characterization clearly demonstrated that CAF was successfully intercalated into ZrP. The results of the XRPD analysis are reported in Fig. 3. The ZrP-CAF diffraction pattern is compared with that of the pristine ZrP, which possesses an interlayer distance of 7.5 Å typical of the acidic form.\(^{37}\) In ZrP-CAF a reflection corresponding to 15.6 Å is detectable, compatible with the intercalation of caffeine, when compared with the interlayer distances obtained for the intercalation of aromatic and heterocyclic bases.\(^{38}\) The increment from the starting interlayer distance is consistent with the presence of a monolayer of molecules lying perpendicular, or nearly perpendicular, to the layers of the host structure,\(^{39}\) considering the total interlayer distance of 15.6 Å, the caffeine dimension of 7.8 Å \(\times\) 6.1 Å \(\times\) 2.1 Å\(^{40}\) and the layer thickness\(^{35}\) of about 7.6 Å (see Fig. 4). The intercalation reaction was

![Fig. 3 XRPD patterns of ZrP (black) and ZrP-CAF intercalation compounds (red).](image)

**Results and discussion**

The intercalation chemistry of ZrP is characterized by the presence of Brønsted acid groups in the interlayer region; the reaction is thus often driven by a topotactic acid–base reaction\(^{34}\) between the host and the guest. CAF is a weak base with a \(pK_a\) of 10.4, while protons in the phosphates of ZrP are a medium strong Brønsted acid.\(^{21}\) In order to intercalate CAF in

![Fig. 4 FT-IR spectra of ZrP (black), ZrP-CAF intercalation compound (red), and CAF (blue).](image)
effectively driven by an acid–base mechanism, as demonstrated by the FT-IR analysis.

In Fig. 4 the FT-IR spectra of ZrP, CAF and ZrP-CAF are reported. The typical signal detectable in a ZrP spectrum is the P–O stretching at ~1040 cm⁻¹, that is still evident in the ZrP-CAF spectrum. The spectrum of pure caffeine is characterized by a region between 1750 and 1400 cm⁻¹ that presents the signals generated by C–O, C–N, and C–C bond vibrations. In detail, the signals generated by the two carboxylates in the anhydrous caffeine as free base can be detected at 1698 and 1656 cm⁻¹ respectively, as observed in the ZrP-CAF spectrum. Also the shift from 1598 to 1646 cm⁻¹ demonstrates that the free base is protonated, confirming the intercalation of the molecule in the interlayer region of zirconium phosphate through an acid–base reaction.

The intercalation mechanism is better explained in the scheme reported in Fig. 5. Herein a simple representation of the reaction is reported: the basic sites of CAF react with the acidic protons of ZrP generating the intercalation compound. For the sake of simplicity, the layers composed of Zr atoms are represented by brown lines, and only one molecule of caffeine has been completely reported.

The TGA curves of ZrP (black) and ZrP-CAF intercalation compound (red) are shown in Fig. 6. ZrP shows two main weight losses: the loss of hydration water occurred in the range of 20–200 °C, while the step between 200 and 800 °C is attributed to the loss of water due to the condensation of the HPO₄ groups. The thermogravimetric curve of ZrP-CAF sample shows one main step in the range of 20–200 °C and two main steps between 200 and 800 °C: the first is due to the loss of water, while the other two can be attributed to the thermal decomposition of CAF and the condensation of the phosphate groups to produce zirconium pyrophosphate. The overall weight loss of the ZrP-CAF sample (red) is obviously higher when compared to the pristine ZrP material (black), because of the combustion of the organic part. From the difference in the weight loss it was possible to calculate the CAF content. The experimental caffeine content in the ZrP-CAF intercalation compound was about 31.8 wt % corresponding to the following chemical formula: Zr(HPO₄)₁.₂₇(PO₄)₀.₇₃(CAFH)₀.₇₃ + 2H₂O.

The extracted solution was analyzed by UV-vis spectroscopy and the result obtained confirmed the herein previously reported formula.

DSC analysis was carried out on pure caffeine and Zr-CAF intercalation compound samples. The DSC curve of pure caffeine (line blue, Fig. 7) exhibited an endothermic peak at 240 °C which corresponds to the crystalline CAF melting point. The DSC thermogram of the ZrP-CAF sample (line red) showed an endothermic peak at 145 °C due to the loss of crystallization water and two exothermic peaks due to the combustion of the organic part. In this DSC curve the lack of the CAF endothermic peak at 240 °C is a proof of the formation of a new compound.
as a consequence of the drug intercalation into ZrP. These results suggest that the caffeine is intercalated in the interlayer region of zirconium phosphate in a molecular level of incorporation, that is classically observed when drugs are vehiculated in clays.43 After the intercalation, CAF does not retain its crystalline structure, having a new chemical–physical behaviour as demonstrated later by the release studies.

The SEM micrograph of the intercalation compound is reported in Fig. 8. Zirconium phosphate shows the typical lamellar structure, with pseudo-hexagonal shaped crystals of micrometric dimensions, as expected from the synthesis method that has been chosen. As expected, no crystals of caffeine were detected in the sample.

As discussed in the introduction, among the multiple biological activities of caffeine, the aim of this study is to test a topical application. The release method chosen for this study followed OECD guidelines for skin absorption, that uses a Franz-Cell, a well-known system for rapid and accurate investigation of drug diffusion.29 The skin permeability was simulated using the Strat-M membrane, a synthetic, non-animal based model for transdermal diffusion testing that is predictive of diffusion in human skin without lot-to-lot variability, safety or storage limitations.33 The emulsions containing pure caffeine and ZrP-CAF were carefully prepared as described in the experimental part, and used for the tests. 200 mg of cream, corresponding to 8 mg of caffeine, were used for the test. The amount of CAF released at each time was detected by UV-vis spectrophotometry at the wavelength of maximum absorption of caffeine in the phosphate buffer (λ = 273 nm). All experiments showed a good reproducibility; the permeation profiles are reported in Fig. 9. The results are expressed in μg mL⁻¹ of permeated caffeine in the acceptor chamber. The caffeine permeation from the cream containing free caffeine is, as expected, a slow permeation that can be correlated to a hydrophilic drug, with a log P = -0.07.10 Similar results were obtained by Farner et al., for the permeation of the pure caffeine from a cream at 4%,44 using a Sartorius-cellulose acetate membrane.

The ZrP-CAF sample showed a slower permeation during the whole analyzed time range when compared to the pure caffeine.

The observed behavior can be attributed to the release mechanism expected from an intercalation compound: while pure CAF is ready for the permeation, in the intercalation compound the release is subsequent to a cationic exchange. Probably, the CAF release from the intercalation compound is due to Na⁺ and K⁺ cations that are able to permeate the membrane, reaching the cream, and activate the ion exchange in the ZrP-CAF. In fact, only after this exchange the caffeine can be ready to permeate the membrane. In order to validate this hypothesis XRD measurements were performed on the ZrP-CAF cream, and on the same cream after the permeation test. The analysis, reported in Fig. 10, shows that in the cream containing the ZrP-CAF the structure of the intercalation compound has been perfectly retained, while after the release test a new phase with an interlayer distance of 9.7 Å is present, probably due to the intercalation of Na⁺ and K⁺.

Thanks to the intercalation technology, the caffeine hosted in the interlayer region of ZrP can have a really slow release. ZrP-CAF possesses great potential in order to project slow release systems of caffeine, not only in cosmetics but also in

![Fig. 8 SEM micrograph of the ZrP-CAF intercalation compound.](image)

![Fig. 9 Permeation profile from pure CAF (blue) and from ZrP-CAF (red), μg mL⁻¹ vs. time.](image)

![Fig. 10 XRD patterns of the ZrP-CAF cream (red) and ZrP-CAF cream after the permeation test (light blue).](image)
other fields of application. Moreover, the intercalation strategy described in this study could be an helpful tool to avoid the well-known crystallization problems of caffeine. Zirconium phosphate, as a drug delivery system for topical application, could be really promising for developing slow release system, like transdermal patches that require long time of delivery (weeks or months).

**Conclusions**

Caffeine is a well-known drug and zirconium phosphate is a lamellar solid able to host basic molecules. In this article the first intercalation of caffeine into zirconium phosphate is described in this study could be an helpful tool to avoid the well-known crystallization problems of caffeine. Zirconium phosphate, as a drug delivery system for topical application, well-known crystallization problems of caffeine. Zirconium phosphate, as a drug delivery system for topical application, so intercalation compound has been tested for topical release as demonstrated by XRPD, FT-IR, TGA, and DSC analyses. The results reported. The designed intercalation strategy was successful, first intercalation of caffeine into zirconium phosphate is lamellar solid able to host basic molecules. In this article the intercalation confers to the drug a slower release if compared to the pure active.

**Conflicts of interest**

There are no conflicts to declare.

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