Iron oxide encapsulated by copper-apatite: an efficient magnetic nanocatalyst for N-arylation of imidazole with boronic acid†

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N-Arylation of imidazole was carried out with various arylboronic acids on iron oxide encapsulated by copper-apatite (Fe3O4@Cu-apatite), producing excellent yields. Firstly, the iron nanoparticles were prepared using a solvothermal method, and then they were encapsulated by copper-apatite to obtain magnetic Fe3O4@Cu-apatite nanocatalysts. Several physico-chemical analysis techniques were used to characterize the prepared nanostructured Fe3O4@Cu-apatite catalyst. The prepared Fe3O4@Cu-apatite was used as a nanocatalyst for N-arylation of imidazole with a series of arylboronic acids with different substituents to reveal the effectiveness of this magnetic nanocatalyst. The Fe3O4@Cu-apatite nanocatalyst can also be easily separated from the reaction mixture using an external magnet. More importantly, the as-prepared Fe3O4@Cu-apatite exhibited good reusability and stability properties in successive cycles. However, there was a notable loss of its catalytic activity after multiple cycles.

1. Introduction

Over the years, N-arlyheterocycles have gained prominence due to their presence in a wide variety of natural products and bioactive compounds. They have also played an important role as building blocks in organic synthesis. There is a special emphasis on N-aryl imidazole because it exhibits antipsychotic, antiallergic, and herbicidal properties, as well as others. The development of a mild and highly efficient method for the synthesis of N-arlyheterocycles over classical Ullman conditions (room temperature, weak base and ambient atmosphere) has also received a lot of attention as a solid catalyst separation technology, since it can be very efficient and fast. Numerous studies have focused on the immobilization of copper catalytic systems on a magnetic medium based on iron in order to separate them by the simple application of a magnet. In the same way, Alper and co-workers have immobilized Cu(i) catalyzed on the surface of Fe3O4 magnetic nanoparticle-supported l-proline as a recyclable and recoverable catalyst for N-arylation of heterocycles. Calcium phosphate, such as hydroxyapatite and its derivates appear very attractive due to their ion-exchange capability, adsorption capacity, and acid–base properties. Hydroxyapatite based materials have been used in the biomedical field lately and in some important chemical transformations. The latter applications are related to its well-known ability to immobilize divalent and trivalent metal ions, by partial cationic exchange with calcium. Recently, the possibility to combine the properties of apatite with other inorganic phases within core-shell nanostructures has attracted great attention. For instance, hydroxyapatite/TiO2, hydroxyapatite/carbon nanotubes, and...
hydroxyapatite/SiO₂ (ref. 36) core–shell nanoparticles have been studied for different applications, such as photocatalytic processes, fluorescence imaging and drug delivery. Indeed, the combination of hydroxyapatite with iron oxide particles could be useful to design sorbents that combine large activity and easy recovery via magnetic separation.37-38 Hence, several iron oxide–hydroxyapatite nanocomposites have been described in the literature.39-45 However, the Fe₃O₄@Cu-apatite has only rarely been explored, and to the best of our knowledge, the use of Fe₃O₄@Cu-apatite as a heterogeneous catalyst for the N-arylation of imidazole has not been reported in literature. Fe₃O₄ is one of the cheaper magnetic oxide, which act in this study as a high surface area framework that is coated by the growth of Cu-apatite shell. The Fe₃O₄ core has to facilitate the recycle of nanocatalysts through magnetic separation. This present work was achieved within our progressive program to develop an eco-friendly and efficient approach for the synthesis of various products.46-50 This is the background upon which this work is situated upon, and the objective is based on the synthesis and characterization of Fe₃O₄@Cu-apatite nanoparticles, and then their use as magnetic catalysts for the N-arylation of imidazole with arylboronic acids (Scheme 1).

2. Experimental

2.1. Materials and apparatus

Copper(ii) nitrate ([Cu(NO₃)₂·3H₂O, ≈98.0%]), ferric chloride (FeCl₃·6H₂O), calcium nitrate (Ca(NO₃)₂·4H₂O, ≈99.0%), ammonium hydrogen phosphate ((NH₄)₂HPO₄), sodium acetate (CH₃COONa), ethylene glycol (EG), sodium hydroxide (NaOH), ammonia solution (NH₃·H₂O, 30%), poly (ethylene glycol) (PEG, 1000) were purchased from Aldrich Chemical Company and were used as received without any further purification. Deionized water was used in all experiments. X-ray diffraction (XRD) patterns of the catalysts were obtained at room temperature on a Bruker AXS D-8 diffractometer using Cu-Kz radiation in Bragg-Brentano geometry (θ–2θ). SEM and STEM micrographs were obtained on a Tecnai G2 microscope at 120 kV. The average diameter of sample was quantified from the SEM image using ImageJ software. The gas adsorption data was collected using a Micromeritics 3Flex Surface characterization analyzer, using nitrogen. Prior to nitrogen sorption, all samples were degassed at 150 °C overnight. The specific surface areas were determined from the nitrogen adsorption/desorption isotherms (at –196 °C), using the BET (Brunauer–Emmett–Teller) method. Pore size distributions were calculated from the N₂ adsorption isotherms with the “classic theory model” of Barret, Joyney and Halenda (BJH). Fourier transformation infrared (FT-IR) spectra of samples in KBr pellets were measured on a Bruker Vector 22 spectrometer. Magnetic properties of Fe₃O₄ and Fe₃O₄@Cu-apatite were investigated in a MPMS-XL-7AC superconducting quantum interference device (SQUID) magnetometer. The magnetic measurements were performed from –15 000 to 15 000 Oe at room temperature. The element content of Fe₃O₄@Cu-apatite materials was determined by an Elemental analysis was realized using inductively coupled plasma atomic emission spectrometry (ICP AES; Ultima 2 – Jobin Yvon). The ¹H NMR spectra were recorded in CDCl₃ or DMSO d₆, using a Bruker Avance 600 spectrometer.

2.2. Catalyst preparation

2.2.1. Preparation of Fe₃O₄ nanoparticles. The magnetic iron oxide nanoparticles used in this study were synthesized using a solvothermal method, adapting methodologies described in the literature.51 In a typical procedure, 5.0 mmol of the FeCl₃·6H₂O was dissolved in 20 mL of ethylene glycol, and then 20 mmol of NaOH was added to the resultant mixture with vigorous stirring. Then, the obtained EG solution was transferred to a Teflon-lined stainless-steel autoclave and sealed. After reacting at 180 °C for 6 h, the autoclave was cooled down to ambient temperature and the mixture was withdrawn from the reactor with a magnet bar to get the Fe₃O₄ magnetic NPs, which were then washed with water and ethanol and dried at 40 °C for 24 h.

2.2.2. Fe₃O₄@Cu-apatite nanocomposite. The Fe₃O₄@Cu-apatite nanocomposite was prepared along a hydrothermal route.52 Firstly, 0.30 g of Fe₃O₄ NPs was dispersed in 100 mL distilled water, and then 12 g of urea was added. Then, 10 mL of a solution produced from a mixture of Ca(NO₃)₂·4H₂O (0.5904 g) and Cu(NO₃)₂·4H₂O (0.604 g) was added into the above solution in drops (molar ratio of Ca : Cu 1 : 1). After heating at 90 °C for 2 h, 10 mL of (NH₄)₂HPO₄ (0.1981 g) solution was also added in drops and the pH value of the solution was tuned to 10 with ammonia water (molar ratio of Ca : P 10 : 6). After 0.5 h, the solution was transferred into a Teflon-lined stainless-steel autoclave, where it was sealed and heated at 165 °C for 12 h. Finally, the Fe₃O₄@Cu-apatite NPs were collected using a magnet bar.

2.3. General procedure of imidazole’s N-arylation with arylboronic acid

In a 50 mL round-bottomed flask, imidazole (1 mmol), phenylboronic acid (1.2 mmol), K₂CO₃ (1.5 mmol) and Fe₃O₄@Cu-apatite (15 mol%) were added and stirred in MeOH under air at 60 °C for the required time, monitoring by TLC. After completion, the mixture was diluted with H₂O and the product was extracted with EtOAc (3 times). The combined extracts were washed with brine (3 times) and dried over Na₂SO₄. The product was purified using column chromatography (60–120 mesh silica gel, eluting with EtOAc–hexane). The structures of the prepared products were confirmed by ¹H NMR and assigned on the basis of their spectral data in comparison with those reported in the literature.

Scheme 1 N-Arylation of imidazole with boronic acid by a nanocatalyst (Fe₃O₄@Cu-apatite).
3. Results and discussion

3.1. Catalyst characterization

XRD was used to analyze the crystalline phases of the as-prepared Fe₃O₄ and Fe₃O₄@Cu-apatite samples (Fig. 1). The XRD pattern of Fe₃O₄ nanoparticles observed clearly at 2θ = 30.2, 35.4, 43.3, 53.7, 57.3, and 62.98, which are attributed to the (220), (311), (400), (422), (511), and (440) Bragg reflections of the face-centered cubic lattice of Fe₃O₄ nanoparticles, respectively (JCPDS no. 19-0629). In the XRD pattern of Fe₃O₄@Cu-apatite (Fig. 1b), we can observe the presence of a secondary phase in which the diffraction peaks correspond to calcium copper phosphate Ca₂₀Cu(PO₄)₁₄ (JCPDS 49-1080). The average crystallite size of Fe₃O₄ and Fe₃O₄@Cu-apatite was estimated using the Debye–Scherer formula, which gave 21 and 33 nm for Fe₃O₄ and Fe₃O₄@Cu-apatite, respectively.

The FTIR spectra of Fe₃O₄ and Fe₃O₄@Cu-apatite are presented in Fig. 2. This figure shows the FTIR spectrum of Fe₃O₄@Cu-apatite nanoparticles, where there are observable peaks around 1020, and 1090, 964, 600, 561 and 470 cm⁻¹, which are attributed to the asymmetric and symmetric stretching mode of P–O group and bending mode of O–P–O group, respectively. The characteristic bands at about 3571 and 3641 cm⁻¹ are related to the stretching vibrations of the OH group. The weak and broad bands at around 1470 cm⁻¹ and 873 cm⁻¹ are most likely assigned to carbonate groups adsorbed during the synthesis process. In addition, the fingerprint of iron

Table 1 | Lattice parameter, crystallite size, BET surface area and pore volume for all as-synthesized samples catalysts

| Samples            | Crystallite size (nm) | S_BET (m² g⁻¹) | V_meso (mL g⁻¹) | D_meso (nm) |
|--------------------|-----------------------|----------------|-----------------|-------------|
| Fe₃O₄             | 21                    | 22             | 0.0305          | 6.5391      |
| Fe₃O₄@Cu-apatite   | 33                    | 84             | 0.1350          | 7.1465      |

*Calculated by Debye–Scherer equation. †Calculated from adsorption of the N₂ isotherm. ‡Calculated from desorption of the N₂ isotherm.
oxide is assigned in the FTIR spectroscopy to the Fe–O stretching vibration, and detected towards 587 cm$^{-1}$.

The calculated texture parameters of Fe$_3$O$_4$ and Fe$_3$O$_4$@Cu-apatite are summarized in Table 1. The calculation of the BET surface area for Fe$_3$O$_4$ is 22 m$^2$ g$^{-1}$ and the pore volume is 0.0305 cm$^3$ g$^{-1}$. The surface area and pore volume of the Fe$_3$O$_4$@Cu-apatite are 84 m$^2$ g$^{-1}$ and 0.1350 cm$^3$ g$^{-1}$, respectively. This change in surface properties was assumed to be due to the coating of Cu-apatite on the surface of Fe$_3$O$_4$ nanoparticles. The rough surface provides more pores for nitrogen adsorption. The nitrogen sorption isotherm of Fe$_3$O$_4$ is type IV, displaying a hysteresis loops type H3, indicating the mesoporous nature of the material (Fig. 3a). For the Fe$_3$O$_4$@Cu-apatite, the nitrogen sorption isotherm is type (IV) with a hysteresis loop type H3, which proves the existence of mesopores according to the IUPAC manifests (Fig. 3b).

The corresponding pore size distribution curve indicates that Fe$_3$O$_4$ and Fe$_3$O$_4$@Cu-apatite have a centralized pore size distribution around 4 and 11 nm, respectively, which corresponding to mesoporous materials (inset Fig. 3a and b).

SEM was used to study the surface morphology of Fe$_3$O$_4$ and Fe$_3$O$_4$@Cu-apatite (Fig. 4). The micrographs show that Fe$_3$O$_4$ is composed of spherical nanoparticles. Furthermore, in the case of Fe$_3$O$_4$@Cu-apatite, some particles were observed on the surface of Fe$_3$O$_4$, which may be due to the coating of the magnetic nanoparticles with Cu-apatite nanoparticles. From the SEM images of Fe$_3$O$_4$, we determined that the average sizes and the relative standard deviation (SD) values for the diameters of the spherical Fe$_3$O$_4$ MNPs was 264 ± 28 nm. Moreover, ICP analysis was performed to determine the weight of element composition of Fe$_3$O$_4$@Cu-apatite. From the ICP analysis, the Ca, P, Fe and Cu content were found to be 30.96, 18.63, 47, 1.83%, respectively.

STEM was used to analyze the microstructure of the Fe$_3$O$_4$ and Fe$_3$O$_4$@Cu-apatite materials. From Fig. 5a and b, it can be observed that Fe$_3$O$_4$ NPS show a very good dispersion of spherical particles. These Fe$_3$O$_4$ spheres were further used as cores for the growth of Cu-apatite shells to obtain the Fe$_3$O$_4$@Cu-apatite core–shell nanostructures. Compared to the Fe$_3$O$_4$ cores, the outside surfaces became coarse after the growth process, as shown in Fig. 5c and d. These results indicate that the Cu-apatite nanoparticles were successfully loaded onto the Fe$_3$O$_4$ sphere surfaces, which clearly demonstrates the formation of core–shell nanostructures. We have also noticed that Cu-apatite was still tightly anchored on the surface of Fe$_3$O$_4$ even after the severe conditions under which the sample preparation for STEM analysis were conducted (a long time of mechanical stirring and sonication), suggesting the existence of strong interactions between Cu-apatite and Fe$_3$O$_4$.

The magnetic properties of Fe$_3$O$_4$ and Fe$_3$O$_4$@Cu-apatite were investigated using SQUID from −15 000 to 15 000 Oe at room temperature. As shown in Fig. 6 the saturation
magnetization (Ms) of Fe₃O₄ is close to 57 emu g⁻¹, a higher value than that obtained for the corresponding Fe₃O₄@Cu-apatite (30.66 emu g⁻¹). These results indicate that Fe₃O₄ core was successfully coated with a Cu-apatite shell. Moreover, the field-dependent magnetization curves of Fe₃O₄ and Fe₃O₄@Cu-apatite show negligible remanence (Mr) and coercivity (Hc), indicating superparamagnetic behavior for both samples at room temperature.

This superparamagnetism can make the magnetic nanoparticles dispersible in the solution with negligible magnetic interactions between each other, which avoids magnetic clustering. As shown in the Fig. 7, Fe₃O₄@Cu-apatite can be dispersed in deionized water to form a stable brown suspension before magnetic separation. However, when a magnet was placed close to the reaction vessel for a while, it could be observed that the synthesized samples were rapidly attracted to the magnet side, and a nearly colorless solution was obtained.

### 3.2. N-Arylation of imidazole over Fe₃O₄@Cu-apatite catalyst

Firstly, the activities of some samples were screened, such as: Fe₃O₄, Cu-apatite, Fe₃O₄@Cu-apatite and Cu-apatite@Fe₃O₄ catalyzing imidazole N-arylation using phenyl boronic acid as a model reaction. The resultant obtained results are summarized in Table 2. First of all, the reaction doesn’t occur without the addition of a catalyst (Table 2, entry 1), and the iron oxide is also inactive in this reaction (Table 2, entry 2). However, the reaction of N-arylation of imidazole was occurring when Cu-apatite was used as catalyst with a yield of 98% (Table 2, entry 3). These results can be explained by the presence of the copper, which play an important role as catalytic element for enhancing the cross-coupling reaction. Additionally, coating iron oxide by copper-apatite (Fe₃O₄@Cu-apatite) in order to separate them by the simple application of a magnet has no significant effect on its catalytic activity (Table 2, entry 4) with a yield of 97%. On the other hand, coating Cu-apatite by Fe₃O₄ decrease their catalytic activity (Table 2, entry 5), this can be explain by the decrease of the contact surface of the catalytic element and the reactants. This result shows the importance of the catalytic system developed in this work (Fe₃O₄@Cu-apatite) for the N-arylation of imidazole.

| Entry | Catalyst | Yield (%) |
|-------|----------|-----------|
| 1     | —        | n.r.      |
| 2     | Fe₃O₄   | n.r.      |
| 3     | Cu-apatite | 98       |
| 4     | Fe₃O₄@Cu-apatite | 97 |
| 5     | Cu-apatite@Fe₃O₄ | 47 |

### Table 3 N-Arylation of imidazole with aryloboronic acid in various solvents and bases in the presence of Fe₃O₄@Cu-apatite

| Entry | Solvent | Base   | Catalyst (mol%) | Temp. (°C) | Yield of 3a (%) | Yield of 4a (%) |
|-------|---------|--------|-----------------|------------|----------------|----------------|
| 1     | Ethanol | K₂CO₃  | 20              | Reflux     | 87             | —              |
| 2     | Water   | K₂CO₃  | 20              | Reflux     | 66             | —              |
| 3     | Methanol | K₂CO₃ | 20              | Reflux     | 97             | —              |
| 4     | Methanol | Na₂CO₃ | 20              | Reflux     | 94             | —              |
| 5     | Methanol | KOH   | 20              | Reflux     | 88             | —              |
| 6     | Methanol | NaOH   | 20              | Reflux     | 84             | —              |
| 7     | Methanol | K₂CO₃ | 20              | 70         | 95             | —              |
| 8     | Methanol | K₂CO₃ | 20              | 60         | 93             | —              |
| 9     | Methanol | K₂CO₃ | 20              | 50         | 89             | —              |
| 10    | Methanol | K₂CO₃ | 20              | 40         | 75             | —              |
| 11    | Methanol | K₂CO₃ | 15              | 60         | 93             | —              |
| 12    | Methanol | K₂CO₃ | 10              | 60         | 84             | —              |
| 13    | Methanol | K₂CO₃ | 5               | 60         | 72             | —              |

| a Reaction conditions: aryloboronic acid (1 mmol), imidazole (0.5 mmol), base (1 mmol), solvent (2 mL) for 6 h.  b Isolated yield.
Table 4  
*N*-Arylation of (H)-heterocycles with aryl boronic acid in the presence of Fe3O4@Cu-apatite

| Samples | N(H)-heterocycle | Aryl | Yield (%) |
|---------|------------------|------|-----------|
| 1 | Imidazole | C6H5 | 85 |
| 2 | Imidazole | 4-CH3-C6H4 | 87 |
| 3 | Imidazole | 3-CH3-C6H4 | 85 |
| 4 | Imidazole | 2-CH3-C6H4 | 83 |
| 5 | Imidazole | 4-MeO-C6H4 | 92 |
| 6 | Imidazole | 4-NO2-C6H4 | 78 |
| 7 | Indole | C6H5 | 91 |
| 8 | Indole | 4-CH3-C6H4 | 86 |
| 9 | Indole | 3-CH3-C6H4 | 88 |
| 10 | Indole | 2-CH3-C6H4 | 85 |
| 11 | Indole | 4-MeO-C6H4 | 96 |
| 12 | Indole | 4-NO2-C6H4 | 72 |

* Reaction conditions: Fe3O4@Cu-apatite (15 mol%), arylboronic acid (1 mmol), N(H)-heterocycles (0.5 mmol), K2CO3 (1 mmol), methanol (2 mL), 60 °C for 6 h. * Isolated yield.

loading was investigated. The results obtained are summarized in Table 3. Initially, with 20 mol% of the catalyst and K2CO3 as a base, the screening of the effect of different solvents showed that H2O may not be the optimal solvent, when the reaction was carried out in EtOH or MeOH, the yield of the product 1-phenyl-1H-imidazole was isolated in 87 and 97%, respectively (Table 3, entry 1 and 3). After optimizing the solvent, the effect of the base was studied (Table 3, entries 4–6), and it was found that the best yields were obtained by using K2CO3 as base. Otherwise, it was noted that the presence of the base was necessary for the achievement of the reaction as reported in the literature. This is explained by the role of the base to activate the boronic acid in the mechanism of the reaction.

The study of the influence of temperature, in the case of using MeOH, indicated that this reaction is very sensitive to changes in temperature (Table 3, entries 7–10). According to these studies we can conclude that temperature is a key parameter in this type of reaction, and the optimum yield was obtained at 60 °C. Furthermore, the decreasing the catalyst loading to half results in a yield of 84% (Table 3, entry 12), which demonstrate the influence of the catalyst amount in this type of reactions. Thus, the optimum conditions for this reaction are at 60 °C in MeOH in the presence of 15 mol% of nanocatalyst with respect to imidazole and K2CO3 (1 mmol) as the base (Table 3, entry 11).

With the optimized reaction conditions in hand, a variety of arylboronic acids were chosen as the substrates in this N-imidazole cross-coupling reaction and the results are shown in Table 4. The substitution effects of arylboronic acid were investigated. It was found that the reaction with phenylboronic acids with an electron donating group afforded better yields (Table 4, entries 1–5) than with electron withdrawing groups (Table 4, entries 6). Similar observation was made when indole was used in place of imidazoles to obtain the corresponding N-arylindole (Table 4, entries 8–10).

Another benefit of this nanocatalyst system consists of its ease of recyclability. The coupling of imidazole with arylboronic acid was chosen as a model reaction for the reusability study (Fig. 8). After the cross-coupling reaction, the recovered catalyst was washed twice by dichloromethane and water, dried at room temperature before being reused under similar conditions for the next run. It has to be noted that the catalytic behavior of the Fe3O4@Cu-apatite catalyst remains nearly the same for the next run. A notable loss of the catalytic activity was observed. The N-arylation of N-arylheterocycles over Fe3O4@Cu-apatite was compared with other catalysts reported in the literature as tabulated in Table 5. From this, we can show clearly that our Fe3O4@Cu-apatite catalyst exhibited a best catalytic activity of N-arylation of imidazole comparing with other catalytic systems.

In order to investigate the behavior of the Fe3O4@Cu-apatite catalyst during recycling experiments, XRD and STEM analysis were performed (Fig. 9). It was observed from the XRD pattern of the Fe3O4@Cu-apatite catalyst that, after one catalytic cycle, the crystalline composition of the catalyst remained unchanged.

Table 5  
Comparison of activity of different heterogeneous catalysts in the N-arylation reaction of imidazole

| Catalyst               | Reaction conditions | Yield (%) | Ref.  |
|------------------------|---------------------|-----------|-------|
| CuI-zeolite            | Methanol, 65 °C, 17 h | 95        | 53    |
| Silica-supported copper(ii) | Methanol, 70 °C, 6 h | 78–98    | 54    |
| Cu-FAP                 | Methanol, K2CO3, r.t., 6 h | 88        | 55    |
| Cu(0)-cellulose        | Methanol, reflux, 2.5 h | 15        |       |
| Fe3O4@Cu-apatite       | Methanol, 60 °C, 6 h  | 96        | This work |
Also, from STEM analysis, Cu-apatite was still tightly anchored on the surface of Fe₃O₄ after one catalytic cycle. The catalytic effect of copper ions leached from the Fe₃O₄@Cu-apatite was studied in the N-arylation reaction, which was carried out under the optimum conditions. After 2 hours of heating, the catalyst was removed and the remaining reaction mixture was reheated. It was observed that the concentration of the desired product didn’t increase even after heating for an additional 8 h. To further evaluate the catalytic effect of copper ions leached from the Fe₃O₄@Cu-apatite, ICP analysis was used against the catalyst before and after reaction. The copper concentration of the catalyst was found to be 1.83 wt% for the fresh catalyst and 1.82 wt% after one catalytic cycle in N-arylation of imidazole, which confirms negligible copper leaching.

4. Conclusion

In summary, magnetic Fe₃O₄@Cu-apatite core–shell nanocatalysts have been successfully prepared using a hydrothermal method. The physico-chemical properties of these materials were evaluated by FT-IR, XRD, SEM, STEM as well as the adsorption–desorption of nitrogen. The prepared catalysts showed potential capability for the preparation of N-arylimidazoles through the N-arylation reaction under mild conditions. Thereafter, the optimal reaction conditions were explored, and methanol was identified as the optimum solvent of the reaction, which is environmentally friendly. Furthermore, the Fe₃O₄@Cu-apatite can also be easily separated by an external magnet and reused for five runs with only a slight decrease in its catalytic activity.

Conflicts of interest

The authors declare that they have no competing interests.

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