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In Situ Solid–Gas Reactivity of Nanoscaled Metal Borides from Molten Salt Synthesis

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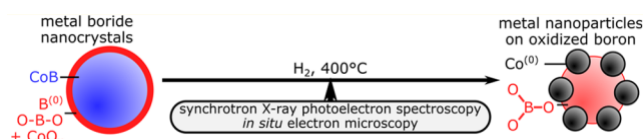
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ABSTRACT: Metal borides have mostly been studied as bulk materials. The nanoscale provides new opportunities to investigate the properties of these materials, e.g., nanoscale hardening and surface reactivity. Metal borides are often considered stable solids because of their covalent character, but little is known on their behavior under a reactive atmosphere,

especially reductive gases. We use molten salt synthesis at 750 °C to provide cobalt monoboride (CoB) nanocrystals embedded in an amorphous layer of cobalt(II) and partially oxidized boron as a model platform to study morphological, chemical, and structural evolutions of the boride and the superficial layer exposed to argon, dihydrogen (H₂), and a mixture of H₂ and carbon dioxide (CO₂) through a multiscale in situ approach: environmental transmission electron microscopy, synchrotron-based near-ambient-pressure X-ray photoelectron spectroscopy, and near-edge X-ray absorption spectroscopy. Although the material is stable under argon, H₂ triggers at 400 °C decomposition of CoB, leading to cobalt(0) nanoparticles. We then show that H₂ activates CoB for the catalysis of CO₂ methanation. A similar decomposition process is also observed on NiB nanocrystals under oxidizing conditions at 300 °C. Our work highlights the instability under reactive atmospheres of nanocrystalline cobalt and nickel borides obtained from molten salt synthesis. Therefore, we question the general stability of metal borides with distinct compositions under such conditions. These results shed light on the actual species in metal boride catalysis and provide the framework for future applications of metal borides in their stability domains.



INTRODUCTION

Synthetic routes toward nanocrystalline metal borides are drawing increasing interest^{1–8} because of the properties of these materials, which differ from those of other solids, including magnetism, plasmonics, thermoelectricity, superconductivity, and superhardness.^{2,9–18} Nanoscaled metal borides are also attractive as heterogeneous catalysts of the selective hydrogenation of various functional organic groups.^{8,19,20} Moreover, cobalt, nickel, iron, and palladium borides⁸ are currently considered for the production of hydrogen, by catalysis of the hydrolysis of sodium borohydride (NaBH₄)^{21,22} and electrocatalysis of the hydrogen evolution reaction.^{23–26} In all cases, the nanoscale provides a parameter to influence or enhance properties, but it has been only scarcely explored because of the difficulty in synthesizing nanoscaled

metal borides. Assessing the chemical stability and structural evolution of these nanomaterials is of utmost importance to identify suitable temperature ranges for operation as thermo-electric materials, hard coatings, or heterogeneous catalysts, for instance.^{22,27} Such applications require chemical and thermal stability that is not met by readily available amorphous boride nanoparticles but should be strongly improved in crystalline nanoparticles. Although the stability of borides under air and inert atmospheres has been examined,^{1,13,28–31} compositional, structural, and morphological evolutions of nanocrystalline metal borides in contact with other gases, especially the reductive atmosphere encountered in catalytic processes,

are unknown because of the scarcity of crystalline boride nanomaterials.^{1–8}

Herein we use inorganic molten salts as media to synthesize crystalline cobalt and nickel monoboride (CoB and NiB) nanoparticles embedded in an amorphous layer comprising cobalt(II) species and partially oxidized boron.^{5,6} We then explore the reactivity of these nanomaterials, exposing high surface areas. Our multiscale in situ analytical methodology combines two techniques enabling studies under reactive atmospheres. First, environmental transmission electron microscopy (TEM) enables one to investigate the nanoscale morphology evolution at pressures of up to 1 bar.^{32,33} Second, ensembles of nanoparticles are analyzed by synchrotron-based near-ambient-pressure X-ray photoelectron spectroscopy (NAP-XPS)³⁴ and near-edge X-ray absorption spectroscopy (NEXAFS) to provide information on chemical and structural evolutions occurring at the surface and in the core of the material, respectively, under a few millibars of several gases, especially hydrogen. On CoB, we demonstrate that the boride is surprisingly decomposed under reducing conditions, yielding cobalt particles at a relatively low temperature of 400 °C. We then show that hydrogen treatment activates CoB for the catalysis of carbon dioxide (CO₂) methanation, through these cobalt(0) species. Similar decomposition occurs in NiB nanocrystals exposed to oxidizing conditions, suggesting that this behavior may be generalized to many other metal boride nanomaterials under a reactive atmosphere, thus providing the frame for their future use as catalysts and functional coatings.

RESULTS AND DISCUSSION

CoB Nanoparticles Synthesized in Molten Salts. CoB nanocrystals were synthesized under argon at 750 °C from the reaction between cobalt(II) chloride (CoCl₂) and NaBH₄ in a lithium chloride (LiCl)/potassium chloride (KCl) mixture (45/ 55 wt %; melting point of 353 °C). After cooling to room temperature (RT), the frozen salt medium was washed with deionized water. A black powder was recovered and stored in air.

X-ray diffraction (XRD; Figure 1a) shows that CoB is the only crystalline phase. The apparent size is 14 ± 2 nm according to the Scherrer formula. TEM and high-resolution TEM (HRTEM; Figure 1b,c) reveal that the sample is composed of single-crystal CoB nanoparticles with diameters ranging from 4 to 20 nm, in agreement with XRD. These particles are smaller than any other CoB nanoparticles reported in the literature to date (18–22 nm).³⁵ The particles are dispersed in an amorphous matrix exhibiting a lower contrast and a 2–5 nm thickness. High-contrast inclusions of ca. 1 nm (black arrows in Figure 1b) are detected in the amorphous matrix. Z-contrast pictures acquired by high-annular-angle dark-field detection in scanning TEM mode (STEM-HAADF; Figure 1d) highlight cobalt-rich regions as bright areas corresponding to the CoB nanoparticles and to subnanometric cobalt-rich inclusions in the amorphous layer. These latter species are not detected by XRD. Energy-dispersive X-ray analysis indicates that cobalt, boron, and oxygen are the only elements present in the material (data not shown). The measured composition of Co_{0.30}B_{0.53}O_{0.17} is in accordance with the Co/B ratio of 0.6 measured by inductively coupled plasma (ICP). On the basis of a previous study, oxygen is expected to be located in the amorphous matrix.⁵

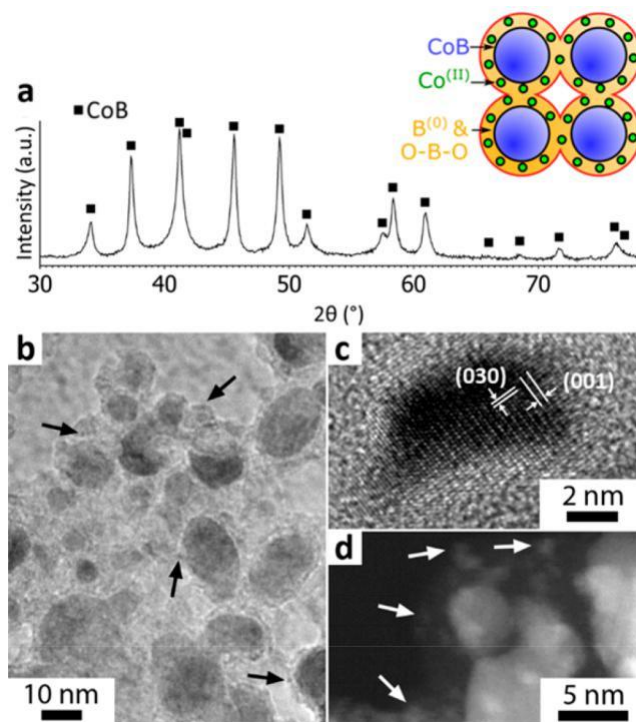


Figure 1. (a) Powder XRD pattern of a CoB sample synthesized in inorganic molten salts at 750 °C for 1.5 h. Scheme of the sample nanostructure deduced from XRD, TEM, and XPS. (b) TEM picture of CoB nanoparticles embedded in an amorphous matrix. Black arrows highlight subnanometer inclusions in the matrix. (c) HRTEM picture of a CoB nanoparticle. (d) Z-contrast picture from STEM-HAADF. White arrows indicate bright cobalt-rich inclusions in the superficial layer.

Brunauer–Emmett–Teller analysis of the dinitrogen sorption isotherms indicates a specific surface area of 71 m² g⁻¹, more than twice the highest value reported in the literature (32 m² g⁻¹) for CoB materials.³⁶ Compositional information on the superficial amorphous layer was further provided by XPS under ultrahigh vacuum (UHV; Figure 2). Using synchrotron-based radiation (TEMPO beamline at Synchrotron SOLEIL, Gif sur Yvette Cedex, France) enabled changing of the inelastic mean-free path of the photoelectron, and, hence, the depth of analysis,^{37–39} by varying the incident photon energy (IPE):^{40–42} 1.7 nm for O 1s and B 1s photoelectrons (IPE 1050 and 725 eV, respectively); 1.1 and 2.1 nm for Co 2p_{3/2} photoelectrons (IPE 1050 and 1500 eV, respectively). Under IPE 1500 eV, the Co 2p_{3/2} region shows a peak at 778.4 eV attributed to CoB⁴³ and accompanied by a broad signal between 776 and 785 eV characteristic of cobalt(II).⁴⁴ The O 1s contribution at 532.0 eV is consistent with oxidized cobalt species, confirming the oxygen environment surrounding cobalt(II). The O 1s contribution at 533.0 eV corresponds to oxygen species linked to boron in agreement with the B 1s contribution at 193.0 eV,⁴⁵ attributed to partially oxidized boron(II). The two other B 1s contributions at 189.2 and 187.9 eV correspond to elemental boron (B⁰) and CoB,⁴³ respectively. The high binding energy (BE) for B⁰ compared to the reported values^{22,45} is due to the nanostructure that yields charge effects.⁴⁶ CoB is detected in the B 1s region but also in the Co 2p_{3/2} region for a photon energy of 1500 eV. At a lower photon energy of 1050 eV probing a thinner layer of ca. 1.1 nm, cobalt(II) is the only species detected in the Co 2p_{3/2} region, with peaks more resolved than those at a photon energy of 1500 eV: 779.3 eV and a broader satellite at 782.5 eV (Figure 2).⁴⁴

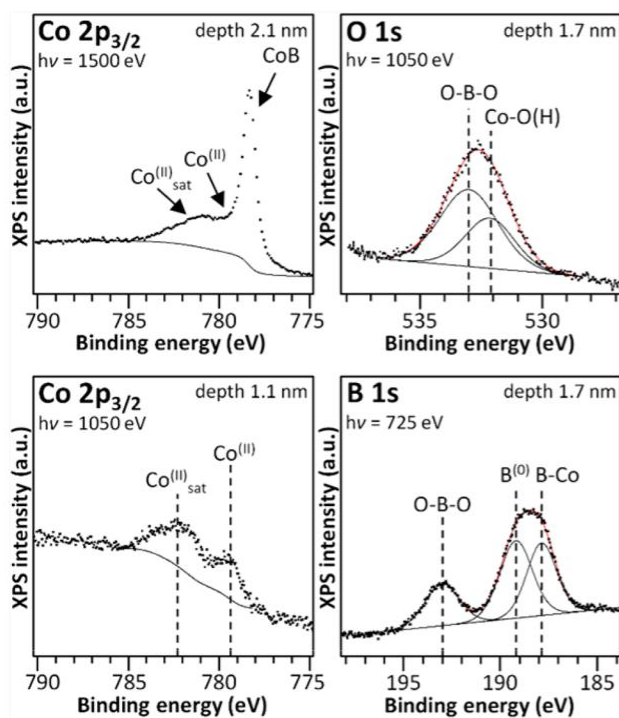


Figure 2. UHV-XPS spectra of the CoB nanocomposite in the Co $2p_{3/2}$, O $1s$, and B $1s$ regions in the initial state. $h\nu$ = photon energy. The approximate depths of analysis are also given.

These XPS data provide strong support to XRD and STEM-HAADF analyses, showing that molten-salt-derived CoB is a nanocomposite made of CoB nanocrystals buried below a ca. 2-nm-thick amorphous layer, composed of partially oxidized amorphous boron (B^0 and BO_2 species) and cobalt oxide and/or (oxyhydr)oxide inclusions (scheme in Figure 1).

Because the synthesis was performed in oxygen-free conditions, the oxidized boron and cobalt(II) species in the amorphous layer originate from exposure of the material to water during the washing step and air during storage. Interestingly, the cobalt(II) species may be interesting precursors to cobalt(0) catalysts, especially for carbon monoxide (CO) or CO_2 methanation, like cobalt oxides.⁴⁷ We thus explore in the following parts how a catalyst activation procedure, namely, reduction under hydrogen gas, impacts the nanostructure and local structure of the material. Then, as an example, we examine how a typical gas reagent mixture used for methanation [dihydrogen (H_2)/ CO_2] can impact the activated material.

Both morphological and compositional evolutions were characterized using (i) environmental STEM with Z-contrast HAADF imaging, (ii) XPS under UHV and NAP conditions, which is sensitive only to modifications of the superficial layer (1–2 nm), and (iii) NEXAFS for a complementary spectroscopic characterization to a depth of approximately 10 nm, i.e., comprising the CoB nanocrystals. Although there is a pressure gap between environmental TEM (1 bar) and NAP-XPS/NEXAFS (a few millibars), in the latter case, the material surface is fully saturated with adsorbates, thus providing relevant information on the actual state under higher pressure.^{48–51} Morphology Evolution of Nanoscaled Cobalt Boride under H_2 . STEM (Figure 3, left) and STEM-HAADF (Figure 3, right) images of the nanocomposite powder were recorded under environmental conditions.

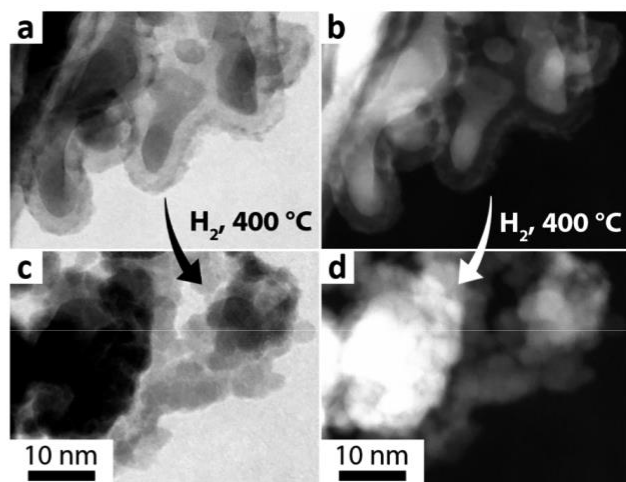


Figure 3. In situ STEM (left) and STEM-HAADF (right) images of the nanocomposite under different conditions: (a and b) under 0.3 bar of argon at 150 °C; (c and d) under 0.9 bar of H_2 at 400 °C. The four micrographs are taken from the same region with an identical scale.

Under 0.3 bar of argon at 150 °C (Figure 3a,b), the morphological characteristics of the powder are similar to those of the pristine sample (Figure 1): CoB nanoparticles are surrounded by the amorphous layer containing cobalt-rich inclusions. The same region is observed under reducing conditions (Figure 3c,d). Upon exposure to 0.9 bar of H_2 at 400 °C, core-shell particles reorganize into nanoparticles with homogeneous contrast, thus demonstrating strong morphological reconstruction of the initial crystalline particles (Figure S1), with fragmentation into smaller nano-particles. Chemical and structural evolutions accompanying these morphological changes under reactive gases were then investigated using XPS.

Surface Reactivity of Nanoscaled Cobalt Boride under Hydrogen. XPS under UHV conditions was performed on the CoB powder after exposure to a static atmosphere of 1 bar of pure H_2 at 400 °C (Figure 4). The spectra in the Co $2p_{3/2}$, O $1s$, and B $1s$ regions were recorded with photon energies of 1050, 725, and 725 eV, respectively, corresponding to approximate depths of analysis of 1.1, 0.9, and 1.7 nm. The Co $2p_{3/2}$ region shows one peak at ~ 779.0 eV corresponding to reduced cobalt, in accordance with the absence of (H)O–Co species in the O $1s$ region.⁵² One single peak is detected in the B $1s$ region at 195.4 eV. Similarly, in the O $1s$ region, one single peak is detected at 535.0 eV. Both signals correspond to fully oxidized boron. Compared to the usual BEs assigned to oxidized boron in previous literature data, they are shifted by 1.8 eV to higher BEs.⁵³ This discrepancy may originate from charge effects due to the nanostructure. Compared to the initial material in the B $1s$ region, no B^0 or B^{II} is observed after H_2 treatment.

Surface Reactivity of Nanoscaled Cobalt Boride under a Mixture of H_2 and CO_2 . The nanomaterial treated under 1 bar of H_2 at 400 °C and then analyzed by XPS under UHV conditions was exposed to a H_2/CO_2 gas mixture (4:1 volume ratio) at 400 °C during analysis. To enable NAP-XPS acquisition (Figure 4), the pressure in the chamber was kept at 2 mbar. Boron remains fully oxidized, as shown by the B $1s$ and O $1s$ contributions at 195.4 and at 534.8 eV, respectively. The absence of a peak at lower BEs in the B $1s$ region indicates that neither CoB nor elemental and partially oxidized boron is present on the surface of the material (depth of analysis of 1.7 nm). In the O $1s$ region,

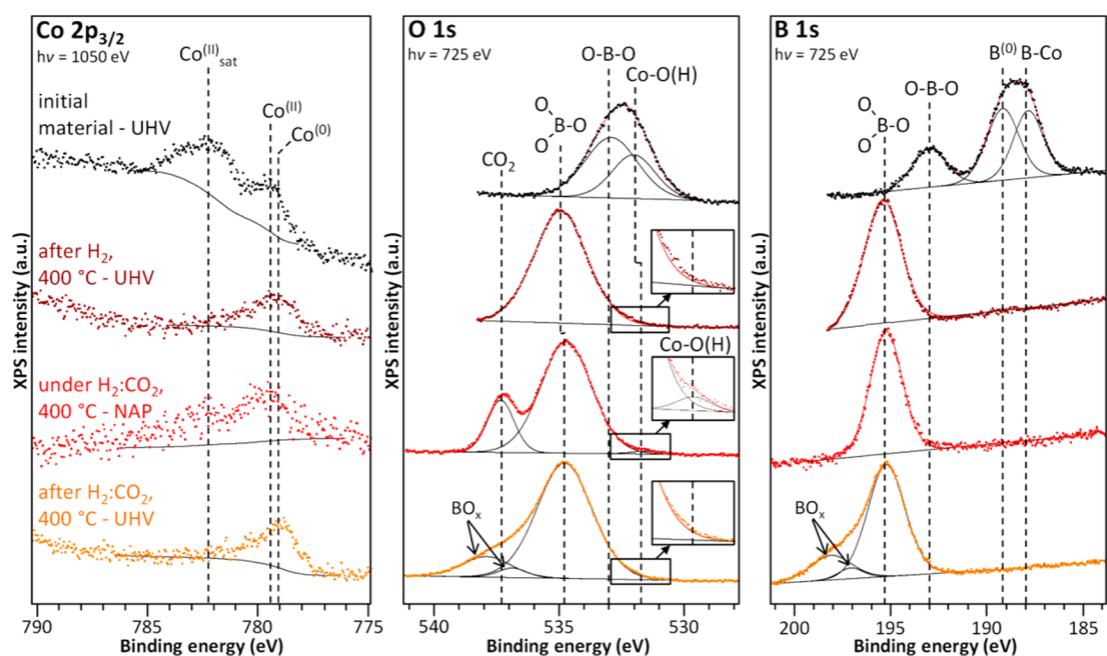


Figure 4. XPS spectra in the Co $2p_{3/2}$, O $1s$, and B $1s$ regions of the CoB nanocomposite. From top to bottom: initial state under UHV; after exposure to 1 bar of H_2 at 400 °C under UHV; near-ambient-pressure conditions of 2 mbar of H_2/CO_2 (4:1 volume ratio) at 400 °C; under UHV at RT after exposure to H_2/CO_2 (4:1 volume ratio) at 400 °C. $h\nu$ = photon energy.

the contribution at 537.4 eV corresponds to gas-phase CO_2 . The shoulder of weak intensity at 532.0 eV is ascribed to (H)O–Co species (inset). In the Co $2p_{3/2}$ region, the broad satellite of oxidized cobalt is detected between 781 and 785 eV, despite a poor signal-to-noise ratio due to the gas phase absorbing most of the emitted photoelectrons. Oxidized cobalt species thus form on the surface of the material when exposed to H_2/CO_2 at 400 °C.

Back to RT and under UHV conditions (Figure 4), the signal in the Co $2p_{3/2}$ region shows the exclusive presence of metallic cobalt, while the shoulder at 532.0 eV in the O $1s$ region disappears. Thus, the (H)O–Co species formed under NAP conditions are not stable under UHV conditions, and fully reduced cobalt species are restored, as observed after exposure to 1 atm of pure H_2 at 400 °C (Figure 4). Note that two additional species (“ BO_x ” in Figure 4) are detected. They are attributed to UHV-specific species and not representative of the material exposed to environmental conditions (discussion in the Supporting Information and Figure S2). All in all, zerovalent cobalt species are formed on the surface of the material initially composed of partially oxidized boron and Co^{II} species. The contribution of deeper CoB in the B $1s$ region also disappears, but the changes of the monoboride core cannot be fully assessed using XPS. Indeed, XPS is only able to probe the first 1–2 nm beneath the surface. Hence, to probe the core of the material, especially the CoB nanoparticles buried under the superficial layer, we have performed NEXAFS on the same synchrotron beamline and the same materials as XPS measurements, also in NAP conditions. The total electron yield mode of NEXAFS was used to provide a depth of analysis of ca. 10 nm, 1 order of magnitude deeper than that of XPS.^{54,55}

Core Reactivity of Nanoscaled Cobalt Boride under H_2 and a H_2/CO_2 Mixture. NEXAFS spectra have been recorded to probe both crystalline nanoparticles and the

amorphous component (analysis depth ~ 10 nm) at the Co L-edge, O K-edge, and B K-edge (Figure 5). The spectra for the initial state show contributions of CoB, oxidized and nonoxidized (broad signal) of amorphous boron, and oxidized cobalt inclusions. Hence, NEXAFS confirms the conclusions obtained from XRD, TEM, and XPS: the initial material is constituted of CoB nanocrystals embedded in a ~ 2 -nm-thick amorphous layer, containing cobalt(II) in the form of oxides or (oxyhydr)oxides (scheme in Figure 1), dispersed in partially oxidized boron, with BO_2 , BO_3 , and B^0 species. Between the initial state and the sample exposed to H_2 at 400 °C, the contributions of oxidized cobalt⁵⁶ and elemental boron⁵⁵ have disappeared, in agreement with UHV-XPS. NEXAFS thus confirms that the whole CoB core is converted into oxidized boron and metallic cobalt after exposure to the reducing atmosphere. NEXAFS spectra of the sample under H_2/CO_2 (NAP conditions) at 400 °C are also displayed. Although poorer signal-to-noise ratios are obtained, they are similar to those in the spectra after the reducing treatment. This indicates that no further change happens in the core of the material upon exposure to the H_2/CO_2 mixture, in agreement with XPS.

In brief, combined TEM, XPS, and NEXAFS studies demonstrate that the oxidized cobalt inclusions in the superficial layer, as well as the CoB nanoparticles, are converted into metallic cobalt nanoparticles during a H_2 pretreatment, while boron is further oxidized by surrounding oxygen species initially bound to cobalt and possibly adsorbed water molecules. After the reducing treatment, the compositional homogeneity between the initial core and superficial layer observed by XPS and NEXAFS is in accordance with the TEM observations: core-shell nanoparticles turning to homogeneously contrasted ones.

Reactivity of Nanoscaled Nickel Boride under Hydrogen and a Mixture of H_2 and Dioxygen (O_2). To assess the generality of the structural modifications evidenced on CoB upon exposure to a reactive gas atmosphere, NiB was investigated. The nanoparticles were synthesized using the

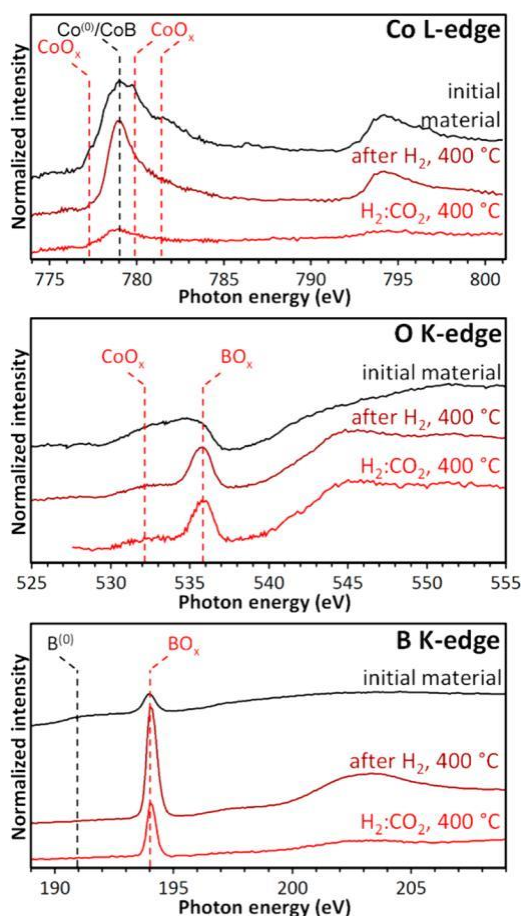


Figure 5. NEXAFS spectra at the Co L-edge, O K-edge, and B K-edge. From top to bottom: the initial CoB nanocomposite under UHV; after exposure to 1 bar of H₂ at 400 °C under UHV; near-ambient-pressure conditions with 2 mbar of H₂/CO₂ (4:1 volume ratio) at 400 °C.

same inorganic molten salt route. The NiB nanocomposite is described in detail elsewhere.⁶ This material is made of NiB nanocrystals surrounded by an amorphous layer. The XRD pattern of the as-obtained powder highlights the presence of NiB crystallites with apparent sizes of 9 ± 2 nm⁶ together with traces of Ni₄B₃ (Figure 6b, top). In contrast to CoB, it has been shown recently by TEM that no

nickel atom resides in the superficial amorphous layer⁶ and that all the nickel atoms are involved in NiB. The NiB nanoparticles are exposed consecutively to reducing and oxidizing conditions at 1 atm. The different steps are schematized in Figure 6a. Chromatography analysis of the exhaust gas does not show any H₂ consumption during all reducing steps (not shown). Oxygen depletion from 320 to 420 °C during the first oxidation step (Figure 6c) indicates oxidation of the nanocomposite. No further O₂ consumption is observed in the subsequent oxidizing steps (Figure 6d), despite the intermediate reducing treatments. The first oxidation process is thus irreversible. After subsequent cycles, the NiB black powder turns into a black monolith and a white powder deposits onto the reactor walls. The XRD pattern of the monolith highlights the presence of metallic nickel and traces of Ni₃B instead of the parent NiB and Ni₄B₃ (minor amount) material (Figure 6b, bottom). The resulting nickel particles exhibit a crystallite size of ca. 30 nm. Segregation of nickel and boron can then be correlated to boron oxidation during the first oxidation treatment. The O/B molar ratio is 1.5 according to O₂ consumption in the first oxidation step, in good agreement with the expected stoichiometry of B₂O₃. The white powder deposited on the reactor walls is therefore B₂O₃, which is known to melt at 450 °C and shows high volatility in this temperature range.⁵⁷

Catalysis of CO₂ Methanation over CoB-Derived Cobalt Nanoparticles. In situ studies shown above demonstrate that exposure to H₂ at 400 °C triggers phase segregation of CoB into oxidized boron and cobalt(0), a typical catalyst of carbon (di)oxide methanation. Hence, we assessed the catalytic activity of our CoB material toward CO₂ methanation. The material was deposited on TiO₂-P25 Degussa (mixture of 20–50 nm rutile/anatase titania nanoparticles in a 20/80 mass ratio; specific surface area of about 60 m² g⁻¹) as a support already used for CO₂ methanation.^{58,59} The supported material (10 wt %) was treated in situ under H₂ (1 atm) at 400 °C for 2 h and then exposed to a H₂/CO₂/helium mixture with a 4:1:5 volume ratio at 1 atm. The gas outlet was analyzed by gas-phase chromatography. No species other than methane (CH₄) and

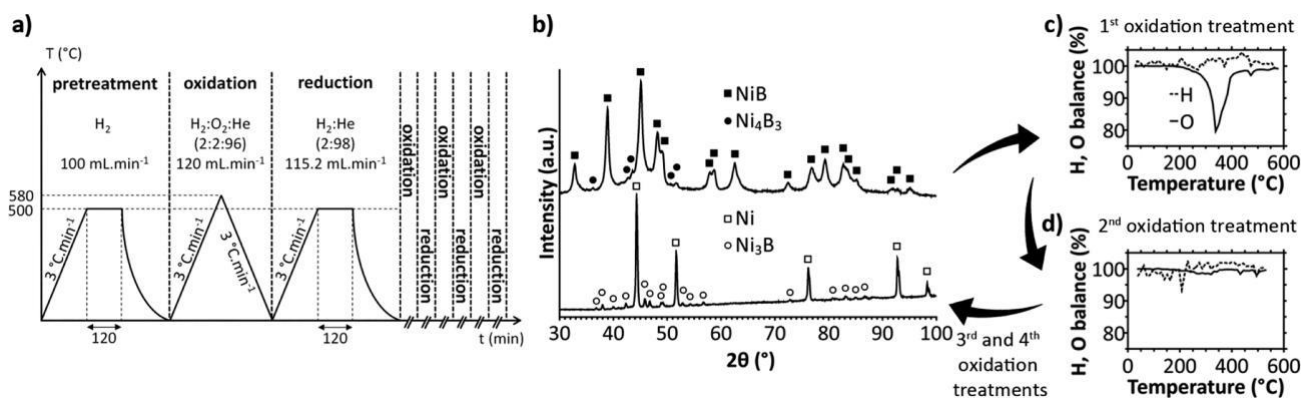


Figure 6. Destabilization of NiB nanoparticles during reduction and oxidation treatments. (a) Consecutive treatments under gases. The oxidation–reduction treatment was repeated four times. (b) XRD patterns of a sample composed of NiB nanoparticles (top) synthesized in inorganic molten salts at 750 °C for 1.5 h and the sample after the reducing–oxidizing treatments (bottom). (c and d) Elemental hydrogen and oxygen balances during the two first consecutive oxidizing heat treatments of NiB nanoparticles: H₂/O₂/helium mixture (2:2:96 volume ratio) to 580 °C at 3 °C min⁻¹. Two oxidizing treatments are separated by exposure to H₂/helium (2:98 volume ratio) at 500 °C for 2 h, not showing any consumption of hydrogen (not shown). Elemental balances during the third and fourth oxidation treatments are similar to that shown in part d.

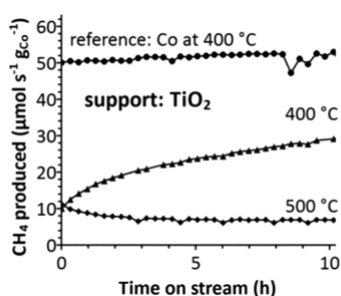


Figure 7. CH₄ production rate as a function of the time on stream for TiO₂-supported CoB at 400 and 500 °C. The behavior of a Co/TiO₂ reference sample is also shown for comparison. Inner gas mixture: H₂/CO₂/helium (4:1:5 volume ratio), 1 atm, 20 mL min⁻¹. Pretreatment: H₂, 1 atm, 30 mL min⁻¹ at 400 °C for 2 h.

production rate measured after 20 min on stream strongly depends on the temperature (Figure S3). It is maximized between 400 and 500 °C. Figure 7 shows the CH₄ production rate as a function of the time on stream at 400 and 500 °C. Deactivation occurs at 500 °C, but the behavior at 400 °C is drastically different: the CH₄ production rate continuously increases during the whole experiment. Increases of 2- and 3-fold are observed after 2 and 10 h on stream, respectively. Note that, after 10 h on stream, the production rate of CH₄ has not yet reached a steady state, but it is still increasing over ca. 30 mol s⁻¹ gCo⁻¹. In comparison, a reference Co/TiO₂-P25 sample with the same cobalt content, formed by wet impregnation of Co₃O₄ supported on titania,⁶⁰ shows a constant CH₄ production rate of 50 mol s⁻¹ gCo⁻¹. Activation of the CoB sample with time on stream at 400 °C therefore indicates that significant modifications of the catalyst occur, with the activity trend toward the behavior of cobalt, in accordance with the structural evolutions described in the previous sections. To confirm such changes, TiO₂-supported CoB materials have been extracted after each step of the catalysis test (Figure S4). Indeed, powder XRD patterns show phase segregation from CoB to cubic cobalt. The generality of the decomposition process has been confirmed by using silica as an alternative support (Figures S5 and S6).

Instability of Metal Borides Exposed to Reactive Atmospheres. The experiments reported above clearly demonstrate both chemical and morphological changes of the CoB-based nanocomposite due to exposure to H₂. These changes are schematized in Figure 8. In the superficial amorphous layer, the formation of cobalt nanoparticles occurs through borothermal reduction: reduction of cobalt(II) oxides

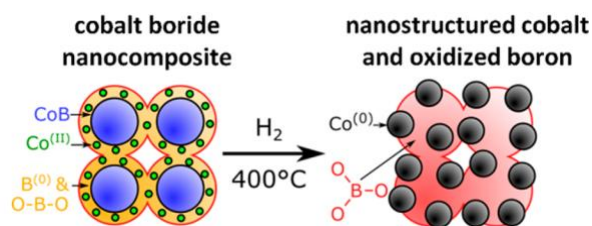


Figure 8. Chemical and morphological modifications of the CoB nanocomposite upon exposure to H₂.

and/or (oxyhydr)oxides by the initially present elemental boron and partially oxidized boron species.

In the core, elemental segregation leads to metallic cobalt, while boron is oxidized either in boron oxide or in boric acid. Heat treatments of amorphous cobalt–boron thin films under various atmospheres were shown by Patel et al. to lead to metallic cobalt only under pure O₂.⁶¹ In our case, treatment under H₂ diluted in argon (5 vol %) does not destabilize CoB up to 800 °C (Figure S7). Hence, temperature is not the only parameter explaining decomposition of CoB nanocrystals, but the atmosphere is important: strong reducing conditions are also required to form cobalt nanoparticles. On the other side, NiB undergoes decomposition when exposed to oxidizing conditions, through the formation of boron oxide. Metallic nickel is the major phase obtained and boron is irreversibly oxidized, as was already seen for CoB.^{19,61} Our original observation is the generality of the decomposition of metal boride nanomaterials previously handled under air and moisture, when they are exposed to gas-phase treatment, in strongly oxidizing and reducing conditions.

In the catalytic hydrolysis of NaBH₄ by cobalt–boron materials, the nature of the active site has been extensively discussed,^{21,22,27,36,61} whether it is composed of cobalt only, cobalt and boron in an amorphous alloy, or one of the known crystalline cobalt–boron phases. Besides, boron (0.5 wt %) is known to act as a promoter of cobalt catalysts of the Fischer–Tropsch process by hindering coking.⁶² Our results provide evidence for the instability of crystalline cobalt boride under a H₂ atmosphere and at relatively low temperatures (400 °C) compared to the temperature of synthesis (750 °C), yielding metallic cobalt as the only cobalt species. Our results put also into perspective that the intrinsic activity of cobalt–boron materials in the hydrolysis of NaBH₄ may result from its segregation to metallic cobalt nanoparticles. Indeed, Arzac et al.²⁷ highlighted the presence of cobalt particles through ex situ characterization in such systems. In the present study, NAP-XPS experiments unravel the true nature of the active surface composed of elemental cobalt, Co–O(H) in small proportions, and oxidized boron. Similar in situ studies applied to amorphous cobalt boride should provide useful information on the real active site in these systems, for which no consensus has been reached so far.²² In the mechanism of CO₂ methanation, the detection of Co–O(H) during in situ NAP-XPS is in favor of a steady state comprising partially oxidized cobalt. This is in accordance with previous works describing the dissociation of CO₂ by cobalt through the formation of CoO.⁶³ Under the catalytic conditions of methanation, the surface concentration of oxidized cobalt might be even higher than that during NAP-XPS experiments (2 mbar) because the CO₂ partial pressure is much higher (0.1 bar).

Catalytic Activity of Metal Cobalt Nanoparticles Surrounded by Oxidized Boron. Cobalt nanoparticles obtained from CoB decomposition during the H₂ pretreatment are the actual active species catalyzing CO₂ methanation. After H₂ pretreatment, the size of the cobalt nanoparticles varies from 15 to 20 nm when supported on silicon dioxide (SiO₂), according to the XRD data (Figure S6), in agreement with the particle sizes reported for cobalt nanoparticles supported on SiO₂ and titanium dioxide (TiO₂).⁶⁴ Interestingly, NAP-XPS in the B 1s region during exposure to H₂/CO₂ indicates no specific species involving boron in the steady state.

The nature of the support onto which CoB was deposited has a drastic influence on the CO₂ methanation reaction.

CoB/TiO₂ indeed shows a CH₄ production rate up to 5 times higher than that found on CoB/SiO₂ at a reaction temperature of 400 °C. The CH₄ production rates for the cobalt reference and the CoB-based/TiO₂ samples are of the same order of magnitude after almost 14 h on stream at 400 °C, in agreement with the formation of dispersed cobalt nanoparticles from segregation in the CoB precursor upon in situ exposure to H₂. The continuous activity increase with the time on stream for TiO₂-supported CoB at 400 °C should be related to reorganization of boron (hydr)oxide, leading to a better exposure of cobalt nanoparticles. Such a rearrangement could facilitate interactions between cobalt particles and the TiO₂ support. For cobalt-based catalytic materials, Riva et al. showed that TiO₂ favored the dispersion of cobalt nanoparticles during the reduction treatment thanks to metal-support interactions, while the cobalt particles sinter on SiO₂ due to the lack of significant interactions.⁶⁰ In the present case, TiO₂ might also favor dispersion of cobalt nanoparticles after elemental segregation occurred.

CONCLUSION

To conclude, we have exploited in situ ensemble and local analyses to unravel the formation of metal nanoparticles upon exposure of nanocrystalline CoB to a reducing gas phase. Similarly, nickel borides decompose under oxidizing conditions. Such a decomposition at relatively low temperatures (400 °C) under a reactive atmosphere appears as a general process for metal borides obtained from the molten salt synthesis. In this process, boron is fully and irreversibly oxidized by oxygen species from the gas phase or from exposure to water and air prior to thermal treatment. Accordingly, our study points out strong morphological, chemical, and structural evolutions, which stress the special care that must be taken to attribute the measured properties, especially catalytic properties, to these crystalline materials. From these observations, the general instability of metal boride nanomaterials at relatively low temperatures (400 °C) under reactive atmospheres is questioned. By exploring with this study the stability ranges of boride nanomaterials, we delineate their acceptable ranges of study and application and we uncover the possible use of reducing atmospheres to exploit decomposition in boron compounds to yield complex materials.⁶⁵

EXPERIMENTAL SECTION

Synthesis. The reaction medium was prepared in an argon-filled glovebox (M-Braun). A total of 4 mmol of NaBH₄ (min 98%, Alfa Aesar) and 2 mmol of anhydrous CoCl₂ (99.99%, Alfa Aesar) were mixed with 5 g of a vacuum-dried LiCl/KCl mixture at the eutectic composition (45/55 wt %). The powders were finely ground in a ballmill (Retsch MM400) for 2 min at 20 Hz and then transferred into a glassy-carbon crucible. The crucible was maintained in a quartz tube, which was connected to an argon ramp. The medium was heated in a vertical oven (Carbolite) from RT to 750 °C (10 °C min⁻¹) and maintained at 750 °C for 1.5 h. The medium was stirred with a glassy-carbon rod rotating at 100 rpm. After the apparatus cooled to RT, the frozen salt medium was washed with deionized water, the suspension was centrifuged for 15 min at 20 krpm (Nalgene), and the powder was separated from the supernatant. This washing procedure was repeated three times. The black powder was finally evacuated at 40 °C overnight.

H₂ Reduction and H₂/O₂ Oxidation Treatments. A total of 50.6 mg of NiB powder (synthesis described in ref 6) was introduced into a continuous-flow quartz reactor. First, the powder was treated under 100 mL min⁻¹ of H₂ (1 atm) for 2 h at 500 °C. After cooling to RT, the sample was exposed four times to oxidation (120 mL min⁻¹ of a H₂/O₂/helium mixture; volume ratio of 2:2:96; heating at 3 °C min⁻¹ from RT to 580 °C) and reduction treatments (115.2 mL min⁻¹ of a H₂/helium mixture, volume ratio of 2:98, 3 °C min⁻¹ from RT to 500 °C and 2 h at 500 °C). H₂, O₂, and H₂O concentrations were measured using a CP4900 microchromatograph (Varian).

CO₂ Methanation. Supported CoB Materials. A total of 40.0 mg of the cobalt boride powder was dispersed in deionized water and added dropwise to 360.0 mg of TiO₂-P25 (Degussa) or SiO₂-Aerosil 200 (Degussa; 10 nm nanoparticles; specific surface area of 200 m² g⁻¹) suspended in deionized water during stirring. The resulting mixture was then sonicated in a USC100TH bath (VWR) for 5 min and stirred for 5 min three times. The dispersion was dried using a rotary evaporator (Büchi) at 60 °C. The reference-supported cobalt material was obtained from Co₃O₄ supported on TiO₂. This precursor was synthesized by the following procedure: 341.3 mg of Co(NO₃)₂·6H₂O was dissolved in deionized water and added dropwise to 735.8 mg of TiO₂-P25 (Degussa) suspended in deionized water during stirring. The suspension was stirred for 48 h at RT and then dry-evaporated at 95 °C. The powder was then annealed at 500 °C during 20 h in air.

Methanation Experiments. A total of 200 mg of precatalyst with particle sizes between 100 and 315 µm was loaded into a continuous-flow fixed-bed reactor and reduced in situ at 400 °C for 2 h under 30 mL min⁻¹ of H₂ (1 atm) prior to the catalytic reaction. The reaction was carried out at 1 atm from 200 to 500 °C under a reaction mixture composed of CO₂/H₂/helium (1:4:5 volume ratio) at 20 mL min⁻¹. The exhaust gases were quantified using a gas chromatograph (Varian CP3800), equipped with Hayesep Q, Molsieve 5 Å, and CP-Sil-5CB columns. The reactants and products were detected with a flame ionization detector (CH₄) and a thermal conductivity detector (CO and CO₂). The exhaust gases were analyzed every 19 min with ~1% (relative) accuracy for the CH₄ production rate (moles of CH₄ produced per gram of cobalt per second). All transfer lines were maintained at 110 °C to avoid water condensation.

In Situ TEM. STEM-HAADF and environmental TEM were performed at the Institut de Physique et de Chimie de Strasbourg on a 200 kV JEOL 2100 F instrument [METSA (Microscopie Electronique et Sonde Atomique) national network] equipped with a Cs probe corrector, allowing one to achieve spatial resolutions down to 0.11 nm. For environmental TEM, the sample was exposed to a continuous 0.01 mL min⁻¹ gas flow. The sample holder was an Atmosphere model from Protochips able to withstand pressures of up to 1 atm and temperatures of up to 1000 °C (±3 °C and with high heating/cooling speeds of 50 °C s⁻¹) under gas flow. These severe conditions do not alter the microscope performances in terms of spatial and energy resolution. The main drawback of the environmental TEM is electron beam damage, which may impact the microstructural changes within the region of interest during the treatment. To overcome this, specimen exposure to the electron beam was minimized, by carrying out all preliminary alignments away from the areas examined during the tests.

XPS. Most XPS experiments were performed on the NAP-XPS end station (SPECS) on the TEMPO beamline of the Synchrotron SOLEIL (Gif sur Yvette Cedex, France). The hemispheric electron analyzer was a Phoibos 150 NAP apparatus (SPECS).

The samples were prepared by the dip-coating technique using a dip-coater (SolGelWay). Nanoparticles suspended in ethanol (30 mg mL⁻¹) were deposited on a silicon wafer coated with a 50-nm-thick gold layer. The XPS spectra were calibrated according to the Au 4f_{7/2} signal at 84.0 eV. Data analysis was performed on the CasaXPS software. Peaks were fitted using GL(30) component line shapes with linear (B 1s and O1s) and Shirley (Co 2p_{3/2} and Au 4f_{7/2}) background types. The BE and full width at half-maximum (BE, eV; fwhm, eV) of the contributions are as follows: (i) in the B 1s region, CoB (187.9, 1.8), B⁰ (189.2, 2.0), BO₂ (193.0, 2.1), BO₃ (195.4, 2.2), and the high-BE BO₃ species (197.1, 1.5) and (198.1, 2.5); (ii) in the O 1s region, Co–O(H)

(532.0, 2.3), BO₂ (533.0, 2.4), BO₃ (534.9, 2.4), CO₂ (537.4, 1.3), and the high-BE BO₃ species (536.9, 1.4) and (537.9, 2.3). The spectrum recorded at 1500 eV was obtained on an Omicron Technology apparatus at the IMPC. It was equipped with a hemispheric analyzer (SPECS) and an aluminum cathode with an Al K α monochromator. In this case, the XPS spectrum was calibrated according to the C 1s signal at 285.0 eV. The depth of XPS analysis in each region was evaluated by calculating the inelastic mean-free path of electrons extracted with a given kinetic energy from a given material.⁶⁶ Because the superficial layer is mainly composed of boron and oxygen, it is approximated to boron oxide for the calculations (considering that only boron yields similar values).

NEXAFS. NEXAFS spectra were recorded on the TEMPO beamline at Synchrotron SOLEIL (see above) in the total electron yield mode. The studied energy windows were as follows: (i) B K-edge, 185–205 eV with 0.1 eV steps and 205–220 eV with 0.2 eV steps; (ii) O K-edge, 525–530 eV with 0.2 eV steps, 530–540 eV with 0.1 eV steps, and 540–555 eV with 0.2 eV steps; (iii) Co L-edge, 770–774 eV with 0.2 eV steps, 774–785 eV with 0.1 eV steps, and 785–810 eV with 0.2 eV steps.

ASSOCIATED CONTENT

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Characterization methods and Figures S1–S7 (PDF)

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Notes

The authors declare no competing financial interest.

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Supplementary information

In situ solid-gas reactivity of nanoscaled metal borides from molten salt synthesis

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Methods

Figure S1. *In situ* TEM analysis of the CoB nanocomposite under (a) 0.3 bar of argon at 150 °C, and (b) 0.9 bar of H₂ at 400 °C.

Figure S2. XPS surveys of the CoB nanocomposite at various steps of the *in situ* study.

Figure S3. Methane production rate as a function of temperature for various samples after 20 min on stream.

Figure S4. XRD diagrams of CoB supported on TiO₂ at different steps of the methanation.

Figure S5. CH₄ production rate as a function of time on stream for SiO₂- and TiO₂-supported CoB at 400 and 500 °C.

Figure S6. Powder XRD patterns of the CoB nanocomposite supported on SiO₂ before exposure to the reducing treatment and after different steps of the catalytic process.

Figure S7. Powder XRD patterns of CoB nanoparticles before and after heat treatment under H₂:Ar (5:95 volume ratio).

Techniques

Powder X-ray diffraction (XRD). XRD measurements were performed on a D8 Bruker apparatus operating at the Cu-K α radiation. Patterns were indexed according to ICDD files: CoB (04-003-2122), cubic Co (04-001-2681), hexagonal Co (04-01-3273), Co₃O₄ (04-05-4386), NiB (01-074-1207), Ni₄B₃ (01-073-2551), Ni₃B (04-014-0853) and Ni (04-010-6148). Apparent crystallite sizes were calculated using the Scherrer formula and pseudo -Voigt functions to fit peaks. The samples after the pretreatments and the catalytic reactions were extracted and analyzed by XRD in inert atmosphere using a glovebag filled with nitrogen and a protective sample holder.

Transmission electron microscopy (TEM). A 120 kV Tecnai Spirit microscope was used for TEM. High Resolution-TEM (HRTEM) was also operated on a 200 kV JEOL JEM 2011 apparatus, at the Institut des Matériaux de Paris Centre (IMPC). Samples were prepared by evaporating a drop of suspension in ethanol on a carbon-coated copper grid. SAED pictures were calibrated with gold nanoparticles.

Energy Dispersive X-Ray spectrometry (EDX). Elemental EDX analyses were obtained on a Scanning Electron Microscope (SEM) S-3400-N (Hitachi) equipped with an EDX detector (Oxford) calibrated prior measurements to allow retrieving quantitative data. 100 x 50 μm^2 areas were probed. Acceleration tension of the source was 10 kV. A titanium grid was used for quantitative calibration. The samples were fixed to a conductive carbon tape.

Inductive coupled plasma-Optical Emission Spectrometry. ICP-OES measurements were acquired on an iCAP 6000 apparatus (ThermoFischer). The sample was prepared by dissolving the cobalt boride powder in a 5 mol L⁻¹ nitric acid solution followed by a 10 times dilution in deionized water. Each concentration is the average of three measurements.

Nitrogen sorption. Samples were degassed at 150 °C overnight before measuring on an ASAP 2000 apparatus (Micromeritics).

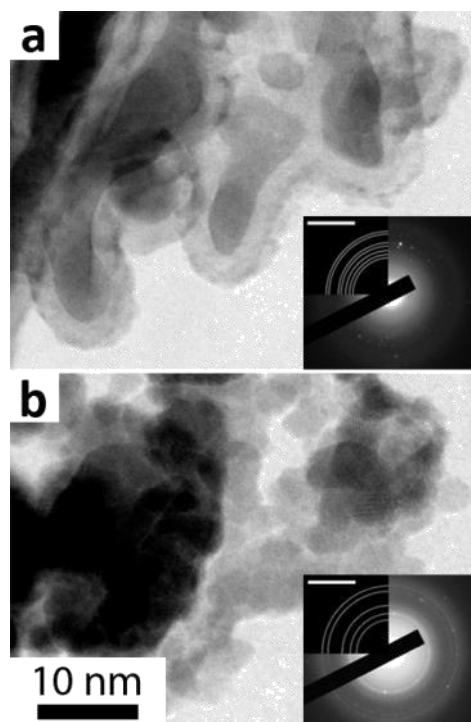


Figure S1. *In situ* TEM analysis of the CoB nanocomposite under (a) 0.3 bar of argon at 150 °C, and (b) 0.9 bar of H₂ at 400 °C. The SAED patterns are indexed (a) along the CoB structure and (b) along the Co₃O₄ structure. Note that the surprising occurrence of Co₃O₄, contrary to UHV-XPS, NAP-XPS and *ex situ* XRD, arises from the very small gas flow and non-optimized gas inlet used in this apparatus, that likely resulted in traces of oxygen impurities in the flow.

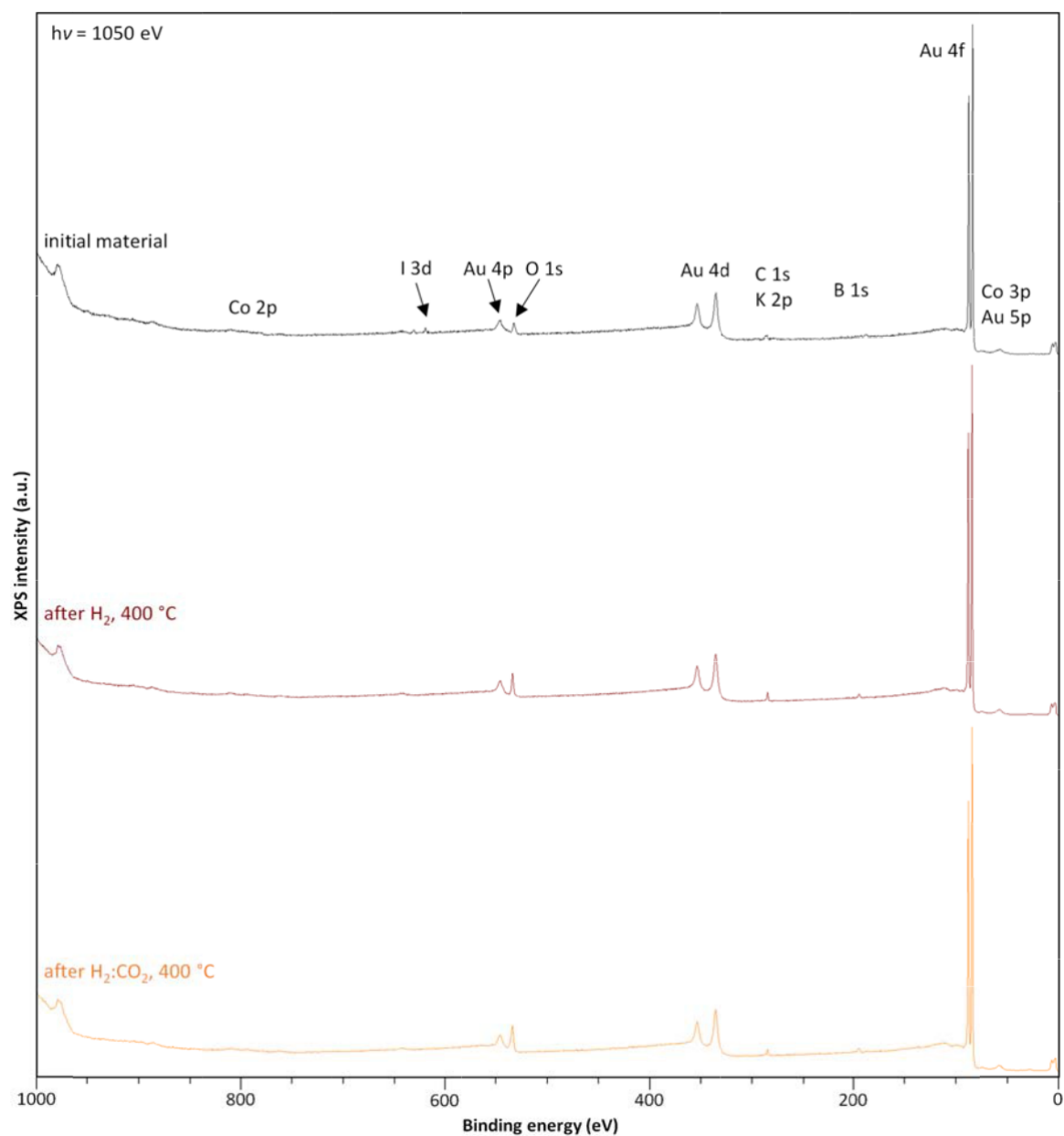


Figure S2. UHV-XPS surveys of the initial CoB nanocomposite (black), after exposure to 1 bar of H₂ at 400 °C in UHV (maroon) and after exposition to H₂:CO₂ (4:1 vol.) at 400 °C. hv: photon energy. Potassium and iodine are detected only in the initial material. These impurities come from the inorganic salt medium in which the nanocomposite is synthesized. These elements were not detected by MEB-EDX. They are thus present in very small proportion, are located only at the surface and eliminated during the first heat treatment.

Note that after NAP-XPS experiments and return to room temperature and UHV conditions, two other species are detected by XPS (“BO_x” on Figure 4 in the main text). Indeed, two oxygen contributions appear at 536.9 and 537.9 eV. In the B 1s region under UHV conditions, two peaks at 197.1 and 198.1 eV appear in addition to the BO₃ peak at 195.4 eV. The area ratios between the two oxygen peaks and

between the two boron peaks are identical. The four contributions are thus interlinked, with couples (O 1s, B 1s) attributed to two unusual fully oxidized boron species with binding energies that are much higher than any other reported previously.(M. M. Ennaceur, B. Terreault, XPS Study of the Process of Oxygen Gettering by Thin Films of PACVD Boron. *J. Nucl. Mater.* 2000, **280**, 33–38; B. R. Strohmeier, Surface Characterization Of Ammonium Fluoborate. 1989, **40**, 249–263)

When the material is heated again under NAP conditions (data not shown), these two species disappear and the NAP-XPS spectra are found similar to those shown in Figure 4, red spectra, thus indicating the reversibility of such structural changes under UHV conditions. These high-energy peaks may originate from locally distorted boron oxides only stable under UHV conditions. Similar phenomena have been described during the reduction of cerium oxides.(J. Holgado, G. Munuera, J. Espinós, A. González-Elipe, XPS Study of Oxidation Processes of CeO_x Defective Layers. *Appl. Surf. Sci.* 2000, **158**, 164–171) These UHV metastable species are not representative of the material exposed to environmental conditions and have not been further investigated.

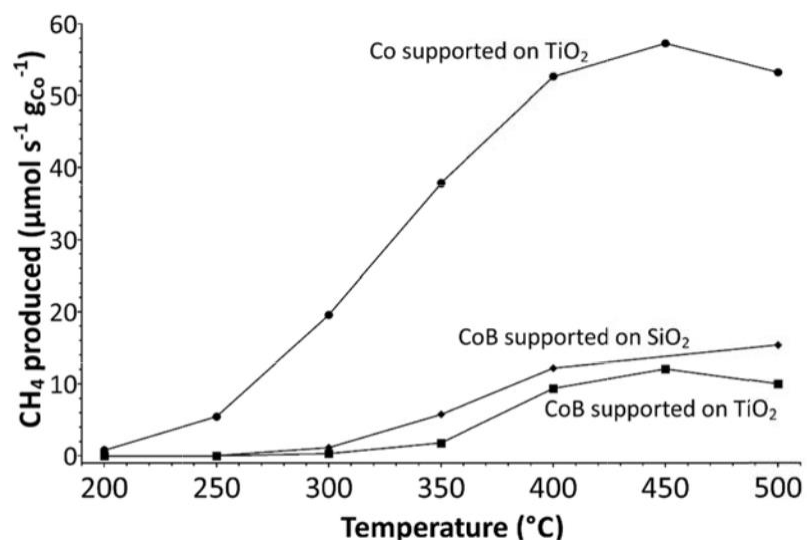


Figure S3. Methane production rate as a function of temperature for various samples after 20 min on stream. Supported CoB were treated under H₂ (1 atm, 30 mL min⁻¹) at 400 °C. The reactive gas mixture during the methanation reaction was CO₂:H₂:He (vol. ratio 1:4:5, 1 atm, 20 mL min⁻¹).

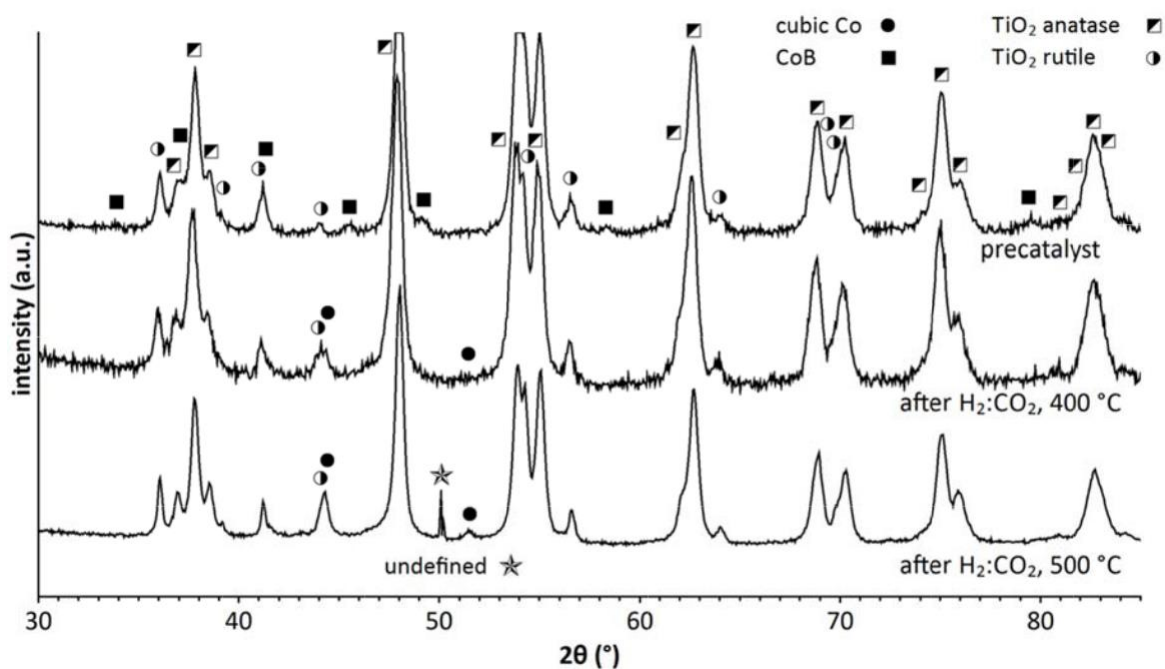


Figure S4. XRD patterns of CoB supported on TiO₂ at different steps. The anatase and rutile phases of TiO₂ are indexed according to the ICDD files 00-021-1272 and 00-021-1276, respectively.

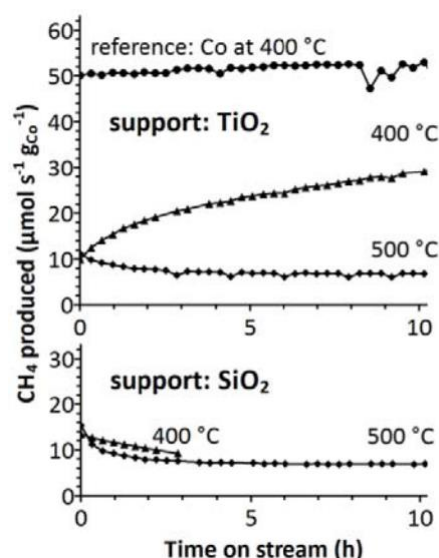


Figure S5. CH₄ production rate as a function of time on stream for SiO₂- and TiO₂-supported CoB at 400 and 500 °C. The behavior of a Co/TiO₂ reference sample is also shown for comparison. Inner gas mix: H₂:CO₂:He (4:1:5 vo l.), 1 atm, 20 mL min⁻¹. P retreatment: H₂, 1 atm, 30 mL min⁻¹ at 400 °C for 2 h. For CoB supported on SiO₂, the CH₄ production rate decreases upon time on stream, leading to a 2-fold decrease at 500 °C within 2 h on stream. Then the rate reaches steady state at around 7 mol s⁻¹ gCo⁻¹.

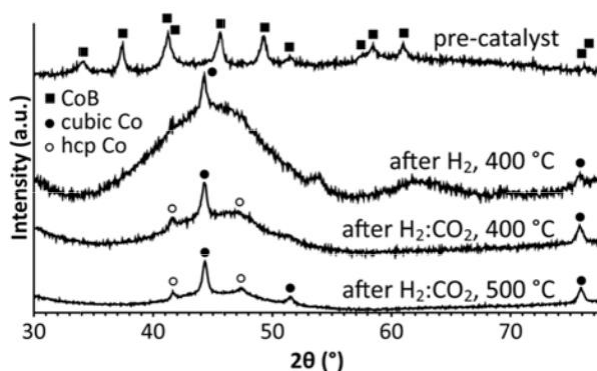


Figure S 6. Powder XRD patterns of the CoB nanocomposite supported on SiO₂ before exposure to the reducing treatment and after different steps of the catalytic process. In the initial state, the XRD pattern is similar to that recorded for the unsupported material (Figure 1a in the main text), with broad peaks corresponding to CoB nanoparticles and crystallite sizes of 14±3 nm according to the Scherrer formula. After exposure to H₂ at 400 °C for 2 h, CoB reflections disappear, whereas peaks corresponding to cubic Co are detected, in agreement with UHV-XPS and NEXA FS data, with an apparent diameter of 18±4 nm. A broad contribution to the baseline is observed in addition to crystalline Co, indicating that the amorphous part of the material has evolved compared to the pre-catalyst. After exposure to the methanation conditions at 400 °C for 3 h, metallic cobalt is also observed in the cubic and hexagonal structures with a crystallite size of 15±2 nm. After methanation at 500 °C, the presence of cubic and hexagonal Co crystallites is also identified with the same apparent size.

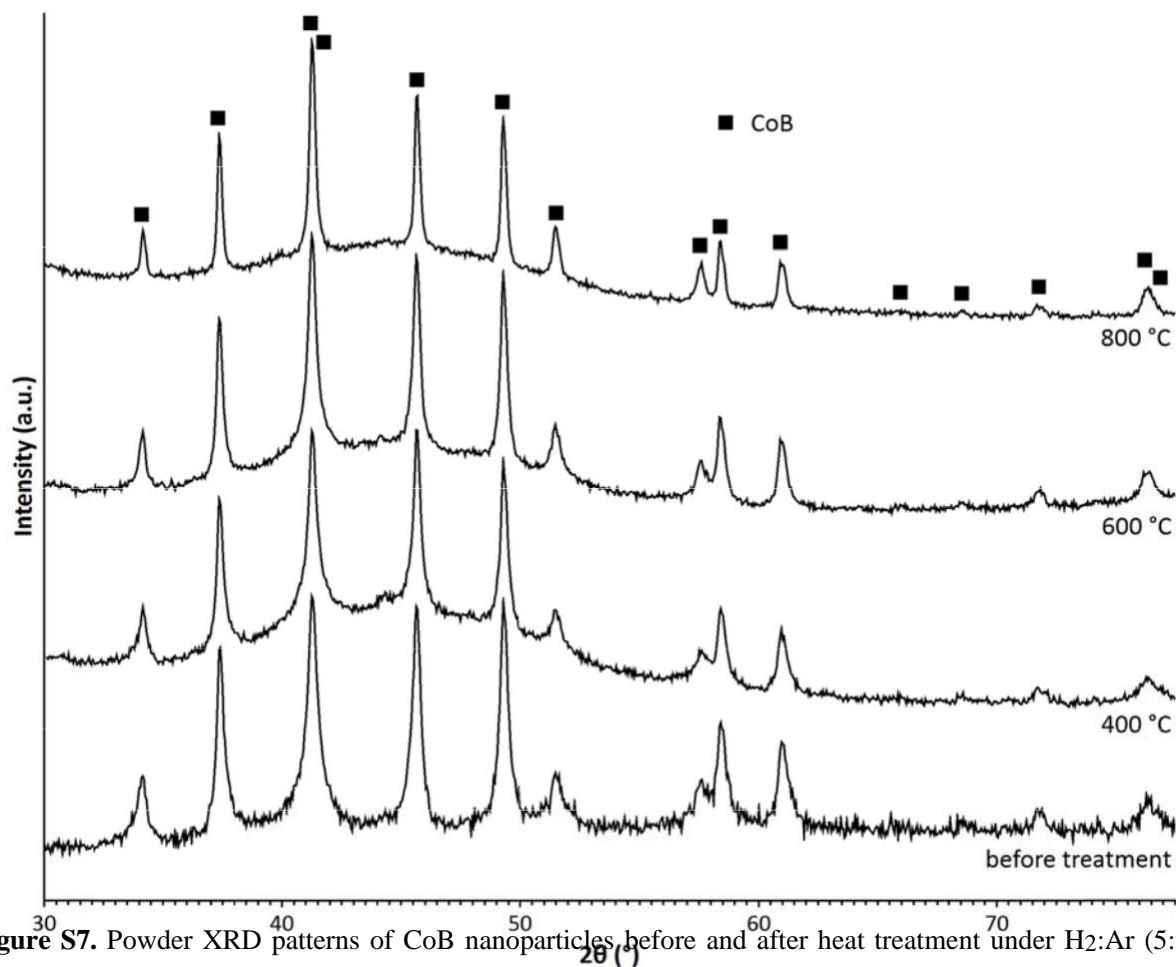


Figure S7. Powder XRD patterns of CoB nanoparticles before and after heat treatment under H₂:Ar (5:95 volume ratio) at various temperatures for 2 h. Heat treatments in a mix of argon and dihydrogen (5 %) do not lead to cobalt boride decompositions. CoB remains the only crystalline phase detected, even after heating at 800 °C. Reflections sharpen with increasing temperature due to nanocrystal growth.