



On the mechanism of visible-light accelerated methane dry reforming reaction over Ni/CeO_{2-x} catalysts

Kristijan Lorber^{a,c}, Janez Zavašnik^b, Jordi Sancho-Parramon^d, Matej Bubaš^d, Matjaž Mazaj^a, Petar Djinović^{a,c,*}

^a Department of Inorganic Chemistry and Technology, National Institute of Chemistry, Hajdrihova 19, SI-1000 Ljubljana, Slovenia

^b Jožef Stefan Institute, Jamova cesta 39, SI-1000 Ljubljana, Slovenia

^c University of Nova Gorica, Vipavska 13, SI-5000 Nova Gorica, Slovenia

^d Ruđer Bošković Institute, Division of Materials Physics, Bijenička cesta 54, 10000 Zagreb, Croatia

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ABSTRACT

The methane dry reforming reaction (DRM) converts methane and CO₂ into syngas, a mixture of H₂ and CO. When illuminated by 790 mW cm⁻² of white light, the 2Ni/CeO_{2-x} catalyst converts CH₄ and CO₂ beyond thermodynamic equilibrium, while the energy efficiency reaches 33%. The DRM reaction is sustained in a purely photocatalytic mode without external heating, yielding CH₄ and CO₂ rates of 0.21 and 0.75 mmol (g_{cat} • min)⁻¹, respectively. Theoretical analysis of Ni/CeO_{2-x} optical properties agrees with *in-situ* UV-Vis DRS results and reveals partly reduced Ce³⁺ sites crucial for extending the optical absorption of Ni/CeO_{2-x} into the visible light range. Two photocatalytic mechanisms are postulated to occur: the hot charge-carrier driven photocatalytic mechanism and the near-field induced resonant energy transfer, depending on the energy of photons used to stimulate the catalyst. This work identifies sub-stoichiometric Ni/CeO_{2-x} as highly efficient for boosting methane activation by visible light under mild conditions.

1. Introduction

Methane dry reforming reaction (DRM, CH₄ + CO₂ ↔ 2CO + 2H₂) represents an attractive pathway for converting methane (an abundant and cheap hydrocarbon) and CO₂ (greenhouse gas) into a mixture of H₂ and CO (syngas). DRM is among few catalytic CO₂ reduction processes (such as CO₂ hydrogenation to methanol, reverse water gas shift and Sabatier reaction), which could be implemented on a global megaton scale to mitigate anthropogenic CO₂ emissions, and thus minimize their impact on climate change [1–3]. Syngas is widely used in the petrochemical industry for synfuel production via the Fischer-Tropsch process, as well as for hydrogenation, hydroformylation and carbonylation reactions [4]. The downsides of the DRM reaction are high endothermicity (ΔH_r = 247 kJ mol⁻¹, ΔG_{298 K} = 171 kJ mol⁻¹) and unfavorable thermodynamic equilibrium values at low reaction temperatures, which limit achievable conversion and hydrogen selectivity. The DRM reaction has been extensively studied at high temperatures (> 600 °C) over transition (Ni, Co and Fe), as well as noble metal catalysts (Ru, Rh, Pt and Pd) [5]. Nickel exhibits high activity for

methane activation [6] and when dispersed in the form of nanoparticles over CeO₂, yields an affordable and outstandingly active DRM catalyst [7,8]. Also, if the nickel cluster size is maintained below 5 nm, carbon accumulation during the reaction can be avoided [9], which overcomes a substantial hurdle in the industrial application of this reaction [5]. Abundance of oxygen vacancies on ceria, manifested through its redox activity [10], is regarded crucial for kinetic balancing of carbon accumulation and gasification reactions during DRM, preventing carbon accumulation on catalysts and consequent deactivation [11,12]. In the past years, catalysis over nano-shaped ceria revealed notable improvements of activity in several reduction and oxidation reactions [13], which appear to be correlated with ease of oxygen vacancy formation over its different terminating facets [13,14]. Activity dependence on ceria shape was also observed in DRM reaction over Ni/CeO₂ catalysts [15].

Photocatalysis is a promising pathway for converting intermittent light (photon energy) into storable chemical energy [16–18]. Illumination of the catalyst can significantly accelerate the reaction rates, and conversions beyond the thermodynamic equilibrium can be achieved

* Correspondence to: Department of Inorganic Chemistry and Technology, National Institute of Chemistry, Slovenia.

E-mail address: petar.djinovic@ki.si (P. Djinović).

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due to non-thermal, vibrational and electronic stimulation of adsorbed reactants and reaction intermediates [19,20]. Thus, (photo)catalytic reactions can be performed at lower temperatures and pressures, which is often beneficial in terms of product selectivity, catalyst stability as well as process cost and design.

Several works investigating plasmonic and non-plasmonic metals (Au, Pd, Pt and Rh) supported on black titania, Al₂O₃, TaON, Ta₃N₅ and SiO₂ report increased H₂ and CO yields during photocatalytic, compared to thermocatalytic DRM reaction [21–24]. Shoji et al. [19], report that Rh/SrTiO₃ catalyst enables methane and CO₂ conversions far beyond thermodynamic equilibrium under UV-light irradiation and without additional heating. The photocatalytically produced syngas contained an equimolar H₂ and CO content, which is not achievable during thermocatalytic DRM reaction due to the large fraction of hydrogen being converted to water by the kinetically dominant RWGS reaction [25].

Halas et al. [26], utilized plasmonic Cu antennas decorated with Ru single sites, which enabled up to a 5-fold increase of methane reaction rate compared to thermocatalysis, and 100% H₂ selectivity when the catalyst was illuminated by > 16 W cm⁻² of white light.

It is becoming clear that visible and UV light have a notable benefit on catalytic activity and selectivity in DRM reaction over noble metals (Rh [19], Au [21,23], Pt [22], Pd [23] and Ru [26]). From an academic, industrial, and practical perspective, a strong photocatalytic response in DRM reaction over transition metal/semiconductor photocatalysts that can be produced with a scalable synthesis method is highly desired.

This work investigates the applicability of Ni/CeO_{2-x} nanorod catalyst for visible-light driven DRM reaction at mild conditions and analyzes the underlying reaction mechanisms. The *in-situ* UV-Vis DRS analysis showed facile catalyst activation by visible light, confirmed by simulation of its electromagnetic properties. The consequences of catalyst illumination by 790 mW cm⁻² of white light (400 < λ < 700 nm) were up to an 8.5-fold increase in methane rate and improved H₂ selectivity (H₂/CO increased from 0.23 to 0.59) compared to thermocatalytic DRM at a constant catalyst temperature of 400 °C. Also, CH₄ and CO₂ conversions beyond the thermodynamic equilibrium were achievable at temperatures below 250 and 300 °C, respectively.

2. Experimental section

2.1. Material synthesis

The selection of ceria nanorod morphology as the catalyst support is based on our preliminary screening of nickel on ceria nanorod and nanocube catalysts (Fig. S1). The nanorod based catalyst was more active, accumulated far less carbon and maintained a higher fraction of Ce³⁺, which is important for visible light absorption [27]. The CeO₂ nanorods were synthesized according to Zabilskiy et al. [28] by dissolving 53.8 g of NaOH (99% purity, Merck) in 140 ml of ultrapure water. Then, 84 ml of the aqueous solution containing 4.9 g of Ce(NO₃)₃·6H₂O (99% purity, Sigma-Aldrich) was added under vigorous stirring. The suspension was stirred for additional 30 min and transferred into Teflon® lined stainless steel autoclaves (volume ~ 35 ml each), where it was aged for 24 h at 100 °C in a laboratory drier. After autoclave quenching, the suspension was filtered, freeze dried overnight and calcined in air (4 h at 500 °C, heating ramp of 5 °C min⁻¹, Nabertherm P330).

Different nickel loadings (0.5, 1, 2 and 4 wt% nominal) were deposited by adding 2.5% NH₄OH solution to the aqueous suspension containing CeO₂ nanorods and an appropriate amount of dissolved Ni(NO₃)₂ (Sigma Aldrich, purity 99%), until pH reached 7.7. After 2 h of stirring, the pH of the suspension was raised to 9.5 using 25% NH₄OH and stirred for an additional 30 min. The efficiency of nickel deposition was verified spectrophotometrically by analyzing the mother liquor using a Spectroquant® NOVA60 analyzer (Merck). In all cases, the efficiency of nickel deposition was above 98%, and nominal loadings were considered actual. Finally, the suspension was filtered, dried overnight

at 70 °C in a laboratory drier, and calcined in air for 4 h at 500 °C (Nabertherm P330). The samples are denoted as CeO₂-R for the bare ceria support and xNi for nickel ceria nanorod catalysts, where “x” represents the nickel loading.

2.2. Characterization

The *in-situ* UV-Vis DRS analysis was performed on the Lambda 650 apparatus (Perkin Elmer) equipped with a HVC-VUV-5 reaction chamber from Harrick. Finely powdered samples (~ 10 mg) were analyzed in air at 25 °C and in 5% H₂/N₂ (flow of 10 ml min⁻¹, purity 5.0 by Messer) after dwelling the sample at 200 and 450 °C for 30 min. Spectralon® white standard was used to record the background.

The XRD analyses were performed on a Panalytical XP PRO MPD apparatus using Cu_{Kα1} irradiation ($\lambda = 1.5406 \text{ \AA}$) in the 2theta range between 10° and 80°, step size of 0.034° and fully opened 100 channel X'Celerator detector.

The H₂ adsorption experiments were performed on a IMI-HTP manometric sorption analyzer (Hiden Isochema Inc.). Prior the isothermal measurements, the catalyst samples were reduced *in-situ* in pure H₂ flow of 10 ml min⁻¹ while ramping the reactor temperature at 5 °C min⁻¹ to 350 °C. After the reduction pretreatment lasting 16 h, the sorption isotherms were measured at 110 °C in the pressure range between 50 and 1000 mbar. The chemisorbed H₂ values were estimated from extrapolating the linear part of the H₂ desorption isotherm to zero pressure as described by Słowik et al. [29] and the average nickel particle size was adopted from Velu et al. [30].

The N₂ physisorption technique (Tristar II apparatus from Micromeritics) was used to analyze BET specific surface area, total pore volume and average pore size (BJH method) of the synthesized catalysts. Before analysis, the samples were degassed 1 h at 90 °C, followed by 4 h at 300 °C in N₂ flow (purity 6.0, Linde) on a SmartPrep accessory (Micromeritics).

Thermogravimetric analysis (TGA) was used to quantify carbon accumulated on the catalyst during the reaction. The samples were heated in airflow (25 ml min⁻¹) from 50 to 800 °C with a 10 °C min⁻¹ ramp (Perkin Elmer, model STA6000). Carbon mass was calculated from the mass difference before and after analysis.

H₂-temperature programmed reduction (H₂-TPR) was performed on an AutoChem 2920 apparatus to analyze the reduction of materials and quantify the amount of consumed H₂. Before analysis, the samples were *in-situ* pretreated in synthetic air at 300 °C for 15 min. After cooling to 10 °C, the atmosphere was changed to 5% H₂/Ar (25 ml min⁻¹), and a temperature ramp of 10 °C min⁻¹ was used to heat the sample (100 mg) to the final temperature of 550 °C. H₂ consumption was analyzed with a TCD detector, and a LN₂/isopropanol cold trap was used to remove water vapor and eliminate its contribution to H₂ quantification.

For TEM analyses, powder samples were first dispersed in absolute ethanol and sonicated to prevent agglomeration of the nanoparticles. Such prepared suspension was transferred onto commercial lacey-carbon Cu support grids. TEM analyses (conventional TEM micrographs, HR-TEM and electron diffraction patterns) were performed on a LaB₆ JEM-2100 microscope (Jeol Inc.), operated at 200 kV and additionally equipped with an energy-dispersive X-ray spectrometer (EDS, model EX-24063JGT, Jeol Inc.). TEM micrographs were recorded by Orius SCD-1000 (Gatan Inc.) slow-scan CCD camera.

Simulation of electromagnetic properties of bare CeO₂ rods, spherical Ni particles and Ni/CeO₂ nanorod catalysts was done using the MNPBEM implementation of the boundary element method [31]. Optical constants for Ni, oxidized and reduced ceria were taken from literature [32,33]. In all calculations, the CeO₂ rods are assumed to be in an air-like environment. Calculations were done averaging over different light polarization directions with respect to the rod axis to simulate a random orientation of nanorods.

2.3. Catalytic experimentation

Catalytic tests were performed in a modified reaction chamber (HVC-MRA-5, Harrick, Figs. S2 and S3) between 180 and 470 °C. During thermo-catalytic experiments, the catalyst temperature was varied in 10–50 °C increments by changing the power output of the electric heater. During light-assisted experiments, the catalysts were illuminated by 790 mW cm⁻² of white light, and the catalyst temperature was varied by changing the power output of the electric heater. Schott KL2500 LED source (400 λ <math>< 700</math> nm, Fig. S4) was used for catalyst illumination, equipped with an optic fiber having a 9 mm active diameter and light focusing lenses (Thorlabs Inc.), which concentrated the light to a spot equal to the catalyst pellet diameter (4.5 mm) with a maximum intensity of 790 mW cm⁻² (measured by Thorlabs PM100D photometer). For all tests, 2 mg of finely powdered catalyst was used, which formed a round pellet measuring 4.5 mm in diameter and 0.5 mm in thickness. The catalyst was positioned on top of a 1 mm thick layer of powdered SiC (Sicat, 30–150 μm) to improve heat transfer from the furnace to the sample and minimize radial temperature gradient in the catalyst layer. The catalyst temperature was measured with a 0.25 mm O.D. thermocouple (Omega Engineering Inc., model SCASS-010U-12) just penetrating the catalyst bed from the bottom with the tip located about 0.2 mm below the illuminated catalytic surface. Before reaction, the catalysts were activated *in-situ* in a 10 ml min⁻¹ flow of 5% H₂/N₂ (Messer, purity 5.0) at 470 °C for 30 min. For the catalytic reactions, the atmosphere was switched to equimolar flow of CH₄ and CO₂ (Linde, purity 5.0 and 5.3, respectively, 10 ml min⁻¹ each, without any dilution, WHSV = 600 L(g_{cat} • h)⁻¹). The equimolar CH₄-CO₂ feed was used in all experiments, in accordance with the stoichiometry of the DRM reaction.

All catalytic tests were performed in the kinetic regime to ensure intrinsic activities are reported. Analysis of gas leaving the photocatalytic reactor was performed by GC (model 490, equipped with MS5A and PPU columns by Agilent). Each reported activity point is a calculated average of at least five analytical repetitions.

3. Results

3.1. Catalyst characterization

The specific surface area of CeO₂-R with the value of 84 m² g⁻¹ changed negligibly after the nickel deposition, regardless of its loading (0.5–4 wt% of Ni). The average pore size determined by BJH method defining interparticle porosity was found to be 11 nm for bare CeO₂-R and increased slightly for the Ni/CeO₂ catalysts (13–16 nm, Table S1).

The XRD analysis (Fig. S5) confirmed the presence of the Face Centered Cubic (*fcc*) fluorite-type ceria phase (PDF 00-034-0394) in bare CeO₂ support and Ni/CeO₂ catalysts. The average CeO₂ crystallite size of 11 nm was calculated by the Scherrer equation for bare CeO₂-R, and this value remained identical after deposition of nickel. No diffraction lines belonging to NiO or metallic nickel were observed in any of the samples, suggesting its presence in the form of small nanoparticles.

TEM analysis of bare CeO₂-R support (Fig. 1A and B) and 2Ni catalyst (Fig. 1D and E) confirmed the presence of ceria in the shape of nanorods, which are about 8–15 nm thick and 100–200 nm long. The CeO₂ nanorod morphology is dominated by (111) facets, with (100) and (110) facets exposed only at the tips of the nanorods, contributing less than 10% to the overall surface area. Phase identification of fluorite-type

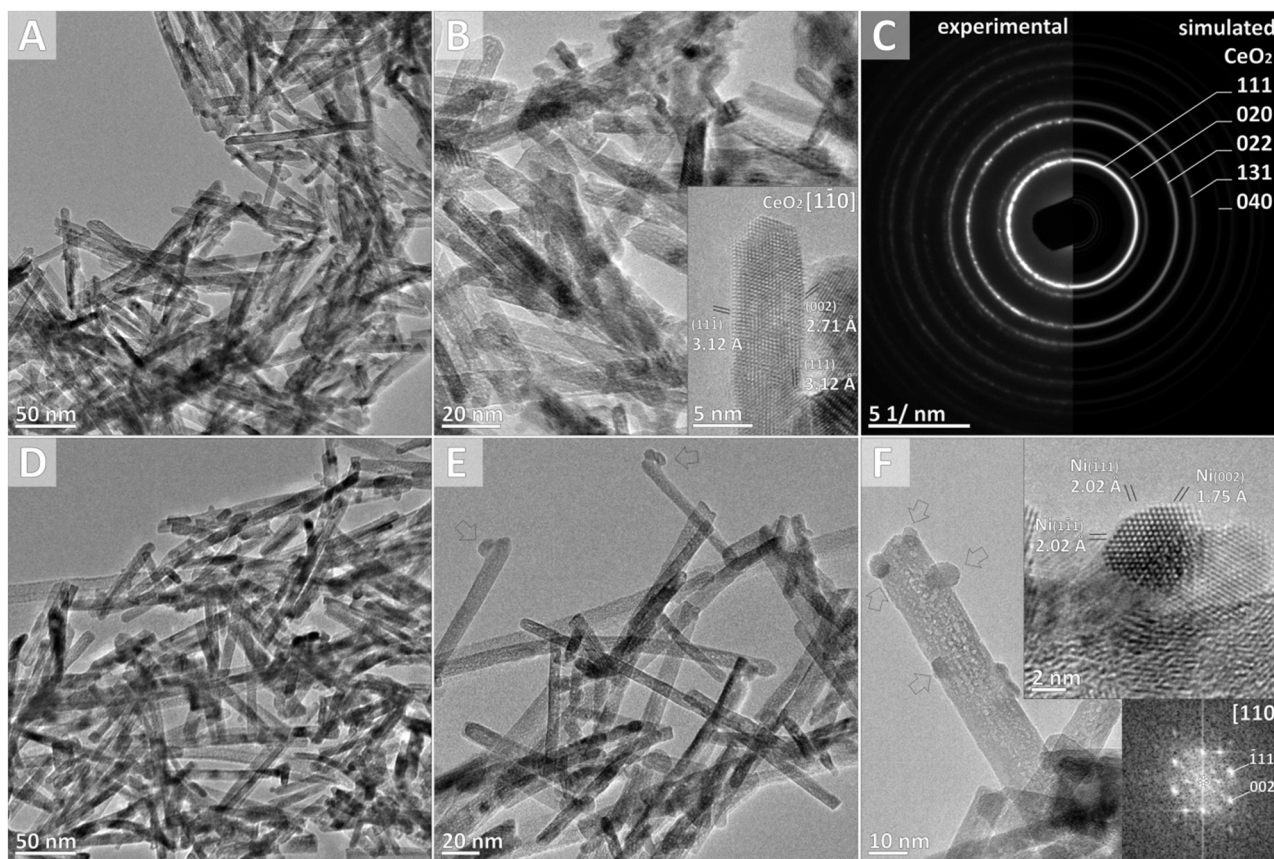


Fig. 1. TEM micrographs of bare CeO₂-R support (A, B) and 2Ni catalyst (D–F). The ceria nanorods are crystalline, showing well-developed crystal faces (B, inset). Experimental SAED pattern, recorded over multiple particles, corresponds to pure CeO₂ phase with *Fm-3m* (No. 225) space group (C). After Ni deposition, the shape and morphology of nanorods are preserved (D, E), with surface-attached nanoparticles marked by arrows (E); the fast Fourier transform (FFT), calculated from phase-contrast HR-TEM micrograph (F, inset) corresponds to cubic Ni phase with *Fm-3m* (No. 225) space group.

CeO₂ was confirmed by SAED (Fig. 1C). In the 2Ni catalyst sample, nickel was visualized as polyhedral nanoparticles decorating the surface of ceria nanorods (Fig. 1E and F). Nickel particle size, measured from the HR-TEM micrographs, range between 5.1 and 6.5 nm, matching the values obtained by the H₂ chemisorption-TPD technique (Table S2).

The redox properties of CeO₂ are indicative of a reversible generation of oxygen vacancies and Ce³⁺ sites, which serve as active sites for dissociative CO₂ adsorption [8,13]. The reducibility of CeO₂ depends on the shape of its crystals, their size and thermal history, and governs on time coke gasification during DRM, which prevents catalyst deactivation [8,12,13]. For the photocatalytic application of ceria, the existence of Ce³⁺ ensures the occupation of energy levels in the 4f orbital. The 4f orbital lies inside the bandgap of ceria, making electron promotion from the 4f orbital to the conduction band achievable by visible light photons [27].

During H₂-TPR analysis of Ni/CeO₂ catalysts (Fig. S6), the reduction started already at 30 °C, whereas for bare CeO₂ nanorods, it was initiated much later at 250 °C. The H₂ consumed during reduction increased progressively with nickel content in the catalysts (Table S2). Based on the quantification of H₂ consumed until 550 °C, 17% of Ce⁴⁺ was reduced to Ce³⁺ in bare CeO₂-R, and this value was very similar also for 1Ni and 2Ni catalysts (19% and 17%, respectively, Table S2), and decreased to 9% in 4Ni sample. To summarize, the Ni/CeO₂ catalysts contain a notable fraction of reduced Ce³⁺ sites in reducing atmosphere and in the temperature range relevant for catalytic reactions performed in this work.

The *in-situ* UV-Vis DRS was used to analyze the ability of the Ni/CeO_{2-x} catalysts to absorb visible light, which is a prerequisite for photocatalytic activity. The optical bandgap of pristine CeO₂ and Ni/CeO₂ catalysts at 25 °C in air decreased slightly, from 3.21 to 3.12 eV with increasing Ni content from 0 to 4 wt% (Figs. 2, S7 and S8). Upon heating the samples in a 5% H₂/N₂ atmosphere to 450 °C, the bandgap values of Ni/CeO_{2-x} samples decreased to values between 2.92 and 2.79 eV (Fig. 2), making them suitable for visible light harvesting. The thermal contribution to bandgap narrowing is proportional to Boltzmann constant multiplied by the temperature change (8.617×10^{-5} eV/K · ΔT) and is equal for all samples: 0.015 and 0.037 eV when heated from 25 to 200 and 450 °C, respectively. The observed bandgap narrowing is larger (0.21 eV for bare CeO₂ and 0.33 eV for 4Ni when heated in reducing atmosphere from 25 to 450 °C, Fig. 2), revealing the change is dominated by chemical alteration (reduction) of the catalysts. The partial reduction caused a gradual change of color from pale brown to dark grey, resulting in a strong absorption throughout the visible range spectrum (Fig. S7). This sub-bandgap absorption was observed as a

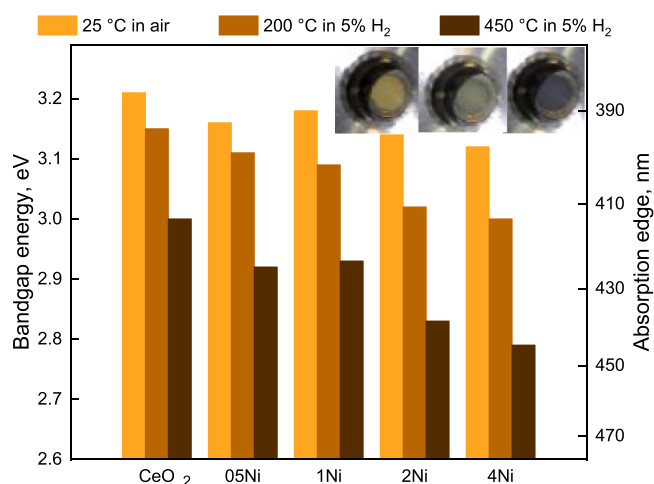


Fig. 2. Bandgap energies for bare CeO₂-R and Ni/CeO₂ catalysts containing 0.5–4 wt% Ni in air at 25 °C and 200 °C and 450 °C in 5% H₂/N₂ atmosphere. Inset shows the 2Ni catalyst at mentioned temperatures and atmospheres.

broad peak centered between 500 and 600 nm for all Ni/CeO_{2-x} catalysts in reducing atmosphere. In addition to partial reduction of ceria, the contact between Ni nanoparticles and CeO₂ likely causes formation of Ni-O-Ce bonds at their interface. As a result, trace amounts of the new mixed oxide phase could be formed, resulting in modification of the electronic band structure of CeO₂, which also contributes to narrowing of bandgap. Increasing the nickel content and temperature in the reductive atmosphere is favorable for both optical bandgap narrowing and increased sub-bandgap absorption of Ni/CeO_{2-x} catalysts.

Simulation of electromagnetic properties of bare CeO₂ nanorods (10 × 100 nm in size), bare spherical nickel nanoparticles measuring 6 nm and Ni/CeO_{2-x} nanorods is discussed in the following section to gain more insight into the occurring light-matter interactions.

The extinction cross-section as a function of wavelength for oxidized CeO₂ nanorods (Fig. S9) correlates well with the absorbance measurements of bare CeO₂ at 25 °C in air (yellow trace in Fig. S7). Over fully oxidized CeO₂, negligible absorption for wavelengths longer than 400 nm is expected. However, reduced CeO_{2-x} shows remarkable extinction in the visible part of the spectra due to the non-negligible extinction coefficient for energies below the band-gap [33], as was observed experimentally using *in-situ* UV-Vis analysis. We also computed the extinction cross-section of an oxidized CeO₂ rod covered by a thin (1 or 2 nm) external shell of partly reduced ceria (CeO_{2-x}) [34]. Radial concentration gradients of cerium oxidation state (surface enrichment with Ce³⁺) were previously experimentally observed in CuO/CeO₂ nanorod catalysts [35]. Our simulations showed a strong positive correlation between the thickness of the surface layer containing Ce³⁺ cerium sites and absorbance below the band-gap energy for CeO_{2-x}, expressed as a broad tail. It should be noted that a direct comparison between the optical response of fully reduced (Ce₂O₃) and fully oxidized (CeO₂) ceria is difficult due to the lack of studies correlating oxidation state (exact fraction of Ce³⁺ and Ce⁴⁺) and optical properties. In any case, our calculations and *in-situ* experimental data strongly indicate that partial reduction of CeO₂ and presence of Ce³⁺ leads to enhanced sub-band gap absorption.

The extinction cross-section of a bare 6 nm metallic nickel nanosphere is moderate when illuminated by wavelengths between 250 and 800 nm. (Fig. S10A.) The plasmon resonance in Ni nanoparticles strongly overlaps with the contribution of bound d electrons and can be hardly discerned in the extinction cross-section [36]. However, the plasmon-related enhancement of electromagnetic field intensity around the nanoparticle surface shows better-resolved features. The near-field response (like near field enhancement, Fig. S10B) of plasmonic nanoparticles is red-shifted with respect to the far-field response (like the extinction cross-section) for systems with significant losses [37]. However, in the case of nickel, the average and maximum near field enhancement is very moderate, making it a poor plasmonic metal.

Simulation of electromagnetic properties of partly reduced CeO_{2-x} rods decorated with one or five spherical 6 nm Ni nanoparticles (Fig. 3A) shows that the presence of Ni nanoparticles does not significantly modify the absorption properties of the partly reduced ceria.

The absorbance of CeO_{2-x} rods is 1–2 orders of magnitude larger than that of bare nickel nanoparticles (Figs. S9 and S10A). Our simulations suggest that the experimentally observed remarkable increase of absorption below the bandgap of CeO_{2-x} does not seem to be related to the presence of Ni nanoparticles. The average field enhancement over the Ni surface in the Ni/CeO_{2-x} system is comparable to the values obtained for individual Ni particles (Figs. 3B and S10B). However, the maximum field enhancement in the region around the contact point between Ni and CeO_{2-x} rod (Fig. 3C) is almost two orders of magnitude larger than the average value over the particle surfaces, thus creating 'hot spots', Fig. 4 [38]. This can have important consequences also for accelerating photocatalytic methane activation. Rodriguez et al. [39] report that at the interface between nickel and ceria, methane dissociation is strongly promoted through electronic perturbation of nickel, induced by the reduced ceria.

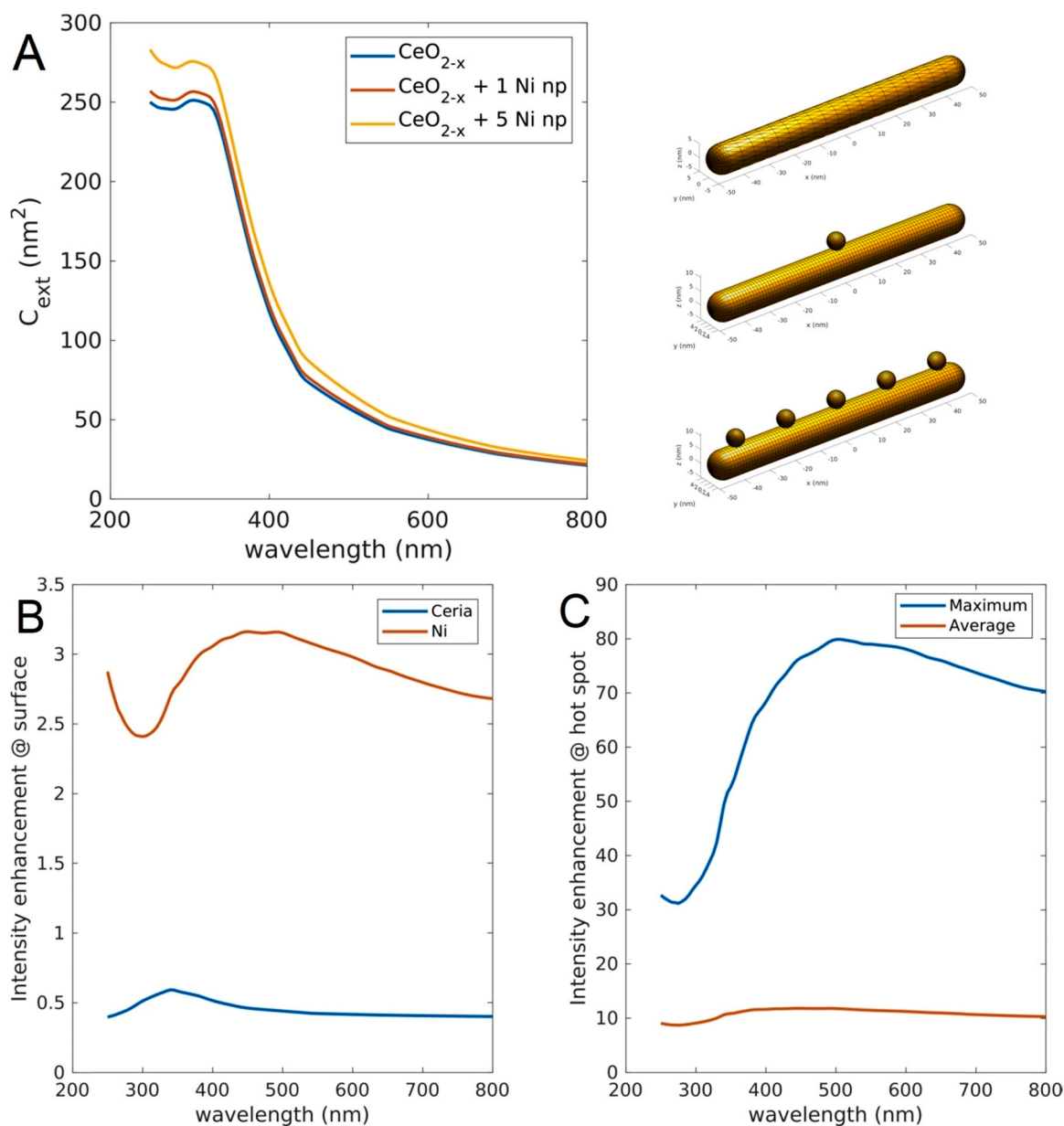


Fig. 3. (A) Extinction cross section of bare CeO_{2-x} rods and decorated with 1 or 5 spherical nickel nanoparticles measuring 6 nm in diameter. Geometric scheme of the simulated structures is shown top right. Near-field intensity enhancements of a CeO_{2-x} nanorod decorated with one spherical nickel nanoparticle: (B) average enhancement calculated over Ni and CeO_{2-x} surfaces and (C) average and maximum enhancement calculated in the hot spot region (cubic volume of 64 nm³ centered at the contact point between Ni and CeO_{2-x} nanorod).

3.2. Thermally driven and light-assisted catalytic performance

Thermally driven DRM performance of Ni/ CeO_{2-x} catalysts is shown in Figs. 5A and S11. No other reaction products were observed beside H_2 , CO and water, which is consistent with occurrence of DRM and RWGS reactions. The CH_4 and CO_2 conversion, as well as the H_2/CO ratio, increased with temperature and nickel content in the catalysts ($0.5Ni < 1Ni < 2Ni \approx 4Ni$). Increasing the nickel content accelerates the DRM reaction rate due to the higher number of active sites for methane activation, which is the rate-determining step in the DRM reaction [9, 40]. The H_2/CO ratio and consequently H_2 selectivity increased in parallel with Ni particle size, likely due to the more dominant direct methane dehydrogenation pathway over larger nickel metallic particles. Over smaller Ni clusters, the contribution of the oxygen-assisted methane activation pathway at the nickel-ceria interface, which produces water, is more dominant [41,42]. This is due to oxygen spillover

from the ceria to nickel and a relatively larger fraction of interface perimeter over smaller nickel particles. The amount of accumulated carbon on the catalysts during 6 h of the reaction was very low (0.1–0.2 wt%) for 0.5Ni, 1Ni and 2Ni samples but increased sharply to 12 wt% on 4Ni catalyst (Table S2). The carbon accumulation rate during the DRM reaction is strongly dependent on nickel particle size and accelerates quickly as nickel size exceeds 5 nm [9,12].

During the light-assisted DRM experiments, the catalysts were constantly illuminated with white light (790 mW cm⁻²), and the power output of the electric heater was adjusted to vary the catalyst temperature (Figs. 5B and S12). In the light-assisted mode, the 2Ni catalyst was most active, close followed by the 4Ni. The CH_4 and CO_2 conversions were greatly increased compared to the thermocatalytic tests at identical catalyst temperatures. Namely, the light-assisted CH_4 rates over 2Ni catalyst were 4.26 and 1.58 mmol $CH_4/g_{cat} \cdot min$ at 460 and 362 °C, respectively. During thermally-driven catalysis at these exact

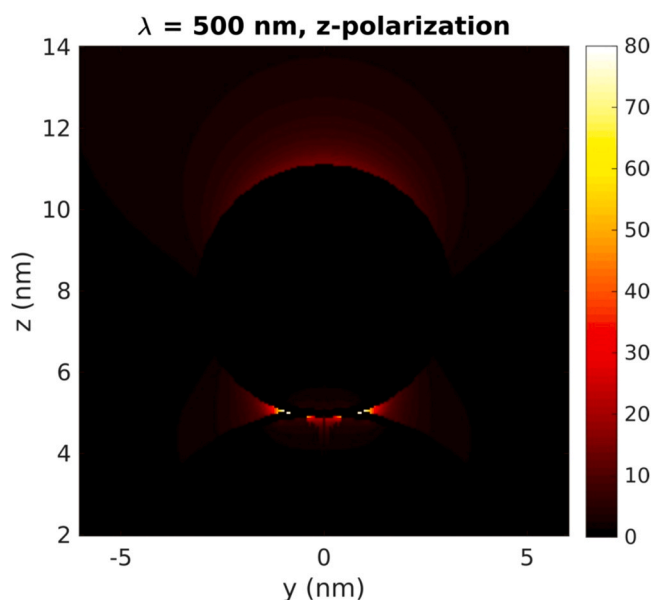


Fig. 4. Spatial distribution of near field intensity enhancement for a Ni/CeO_{2-x} catalyst calculated at the wavelength where largest field enhancement takes place for light polarization parallel to the axis joining the Ni sphere and CeO_{2-x} rod.

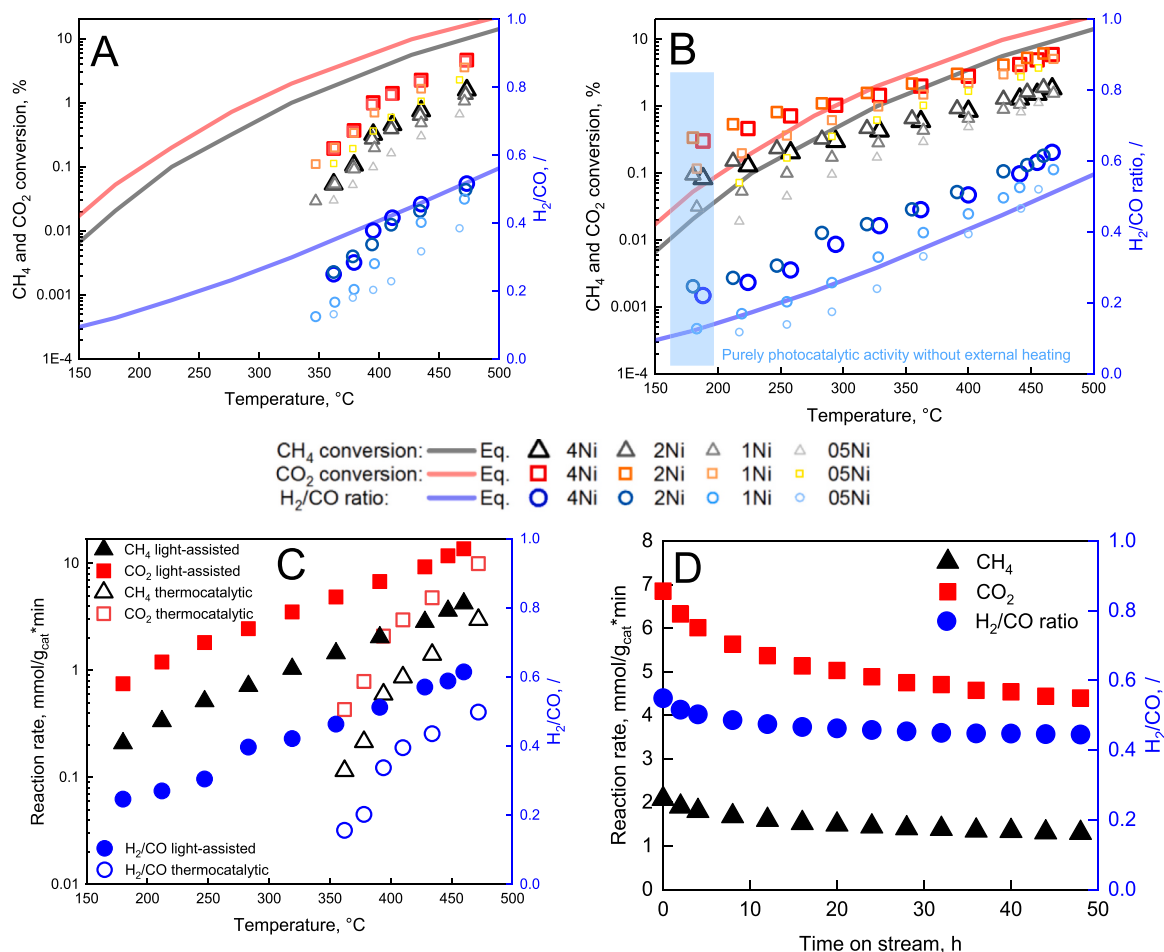


Fig. 5. (A) Conversion of CH₄ and CO₂, as well as H₂/CO ratios produced during thermocatalytic and (B) light-assisted DRM reaction over Ni/CeO_{2-x} nanorod catalysts. Thermodynamic equilibrium values of CH₄ and CO₂ conversion and H₂/CO ratios are shown by full grey, red and blue lines. (C) Comparison of CH₄ and CO₂ rates and H₂/CO ratio produced during light-assisted and thermocatalytic DRM reaction over 2Ni catalyst as a function of catalyst temperature. (D) Long-term light-assisted performance of 2Ni catalyst at 400 °C and constant irradiation by 790 mW/cm² of white light.

temperatures, the values were 2.48 and 0.097 mmol CH₄ (g_{cat} • min)⁻¹, respectively (Figs. 5C, S11A and S12A). Regardless of nickel content, no CH₄ or CO₂ conversion could be identified below 347 °C during thermally driven DRM. However, CH₄ and CO₂ rates of 0.2 and 0.8 mmol (g_{cat} • min)⁻¹ were observed over the 2Ni catalyst in purely photocatalytic mode with no external heating (highlighted by blue rectangle in Fig. 5C). Under such reaction conditions, the 2Ni catalyst temperature was 180 °C. The purely photocatalytic rates over 2Ni catalyst are comparable to those achieved by Shoji et al. [19] over Rh/SrTiO₃ catalyst, irradiated by a 150 W Hg-Xe lamp at 200 °C. A more comprehensive comparison of light-assisted DRM performance is shown in Table 1.

Also, in the light-assisted mode at temperatures up to 250 and 300 °C, respectively (Fig. 5B), the 2Ni and 4Ni catalysts enabled CH₄ and CO₂ conversions well beyond the values predicted by thermodynamic equilibrium. The photocatalytic activity is limited by the number of photons provided by the light source, whereas the thermocatalytic activity increases exponentially with increasing temperature according to the Arrhenius law [19,20,26]. As a result, the highest photocatalytic gain was observed at low temperatures, where the thermocatalytic rate is slowest.

In the light-assisted mode, the 2Ni catalyst produced syngas with the H₂/CO ratio of 0.61 and 0.47 at 460 and 362 °C, respectively. These values are substantially higher than the maximum values predicted by the thermodynamic equilibrium. During thermally-driven reaction at identical catalyst temperatures, the H₂/CO ratio was 0.48 and 0.16. Consequently, light-assisted DRM proceeds with higher H₂ selectivity

Table 1
Comparison of light-assisted DRM performance between various catalysts.

Catalyst	Catalyst temperature and feed composition	Irradiance, W/cm ²	CH ₄ rate, mmol/g _{cat} • min	H ₂ /CO ratio, /	Source
2% Ni/CeO ₂	T = 400 °C CH ₄ : CO ₂ = 1:1	0.79	2.2	0.53	This work
Pd ₉₀ Au ₁₀ /Al ₂ O ₃	T = 500 °C CH ₄ : CO ₂ = 1:1	0.21	0.9	0.9	[23]
5% Ni/Al ₂ O ₃	T = 300 °C CO ₂ : CH ₄ : Ar = 1: 1: 8	0.25	0.1	0.69	[43]
Ru _{0.2} Cu _{19.8} /MgO-Al ₂ O ₃	T = 727 °C CH ₄ : CO ₂ = 1:1	19.2	16.5	1	[26]
10% Rh/SrTiO ₃	T = ~ 120 °C CH ₄ : CO ₂ : Ar = 1: 1: 98	0.056	0.5	1	[19]
1% Pt/0.1SiO ₂ -0.9CeO ₂	T = 600 °C CO ₂ : CH ₄ : Ar = 1: 1: 8	3.6	1.5	0.58	[44]
1% Pt/Zn-CeO ₂	T = 600 °C CO ₂ : CH ₄ : Ar = 1: 1: 8	3.6	2	0.48	[45]
3% Ag-5% La-pCNNT	T = 100 °C CO ₂ : CH ₄ = 1: 1	0.15	~ 0.03	0.14 ^a	[46]
12% Ni/CeO ₂	T = ~ 25 °C CH ₄ : CO ₂ : Ar = 0.98: 1: 8.02	0.47	2.6	1	[47]

^a Methanol and C₂H₆ were identified among reaction products.

compared to thermocatalytic one at identical catalyst temperatures. The H₂ selectivity during DRM is strongly affected by the co-occurring RWGS reaction, and visible light illumination diminishes its contribution to the product selectivity over Ni/CeO_{2-x} catalysts. This is in line with previous findings of Zhou et al. [26], who attributed the 100% H₂ selectivity during DRM to visible light stimulating H₂ desorption from ruthenium active sites, thus decelerating the RWGS rate. Also, with increasing catalyst temperature, the DRM rate increases faster compared to RWGS rate (Apparent Activation Energy for DRM and RWGS were estimated at 82 and 74 kJ mol⁻¹ by Halas et al. [26]). Consequently, by increasing the temperature, a higher H₂/CO ratio is expected.

Long-term light-assisted catalytic activity was tested at 400 °C for the most active 2Ni catalyst (Fig. 5D). During the 50 h time on stream, the initial CH₄ and CO₂ rates decreased by 37% and 35%, and the initial value of H₂/CO dropped from 0.55 to 0.44, revealing catalyst deactivation takes place. Analysis of spent 2Ni catalyst identified 1 wt% of carbon on the sample, thus likely excluding active site blocking as a source of deactivation.

The following analysis was applied to exclude the localized sample heating as the source of the photocatalytic effect, observed over the Ni/CeO_{2-x} catalysts. The apparent activation energies (E_a) in the thermocatalytic mode (90–130 kJ mol⁻¹) decreased to 30–55 kJ mol⁻¹ in the light-assisted mode, where excitation by light and thermal energy were supplied in parallel to drive the reaction (inset Figs. S11A and S12A). The apparent activation energy includes the contributions of activation barriers for the rate-determining step (RDS), the enthalpy of the steps that produce species involved in the RDS and a coverage-dependent contribution related to the enthalpy required to regenerate the active sites occupied by reaction intermediates [48]. The decrease of E_a indicates a change in the reaction mechanism related to light-driven methane activation, which occurs in parallel to the thermally-driven mechanism in the light-assisted DRM mode.

Also, we estimated the local temperature increase of the 6 nm nickel particle under constant illumination, as proposed by Baffou [49], which was found to be between 1 and 2 °C. The CH₄ rate is accelerated by about 5% when the catalyst temperature increases by 2 °C. However,

over the 2Ni sample, the smallest difference between light-assisted and thermally driven CH₄ rates was 71%, in favor of the light-assisted reaction at identical catalyst temperature. Zhou et al. [26] compared the catalyst temperature upon visible light illumination in the same microreactor reactor as used in our work: The values measured by a thermocouple positioned in the catalyst bed and by the thermal camera were identical.

As a control experiment, light-assisted DRM activity was benchmarked to the thermally driven one over the 2Ni/SiO₂ catalyst (Fig. S13A). Silica is a wide bandgap insulator, and when coupled with poor visible light absorption of nickel (Fig. S10), negligible photocatalytic activity gain is expected during this experiment. Indeed, the CH₄ and CO₂ rates differ by less than 5% in thermal and light-assisted DRM experiments at identical catalyst temperatures, revealing the temperature measurement inside the catalyst layer reflects the local temperature at the metallic particles. This experiment also confirmed that the accelerated DRM rate observed over Ni/CeO₂ catalysts is related to light absorption by the CeO_{2-x}, which agrees with our analysis (Figs. 3, S9, S10 and S13B). However, during the light-assisted DRM over 2Ni/SiO₂, the H₂/CO ratio was consistently higher compared to the thermocatalytic experiment. This is likely a result of accelerated H₂ desorption from the nickel surface via desorption induced by electronic transitions (DIET) mechanism [50], which decelerated the RWGS reaction rate. Control experiments with pure CeO₂-R support and SiC confirmed their negligible contribution to photocatalytic activity and identified the presence of nickel as crucial for enabling the DRM reaction (Fig. S14). Based on the above control experiments, the observed DRM rate acceleration upon illumination has a photocatalytic, and not thermal origin.

The effect of wavelength on light-assisted DRM activity was tested on the 2Ni catalyst at a constant irradiance of 300 mW cm⁻² using different bandpass filters (Fig. 6A). The acceleration of CH₄ rate was observed for all wavelengths, compared to the thermally-driven methane rate, which equaled 0.49 mmol (g_{cat} • min)⁻¹ at identical catalyst temperature. Interestingly, the lowest light assisted CH₄ rate (0.61 mmol(g_{cat} • min)⁻¹) was observed for illumination with wavelengths shorter than 450 nm, which is the only fraction of light that enables electron promotion from the VB to the CB of CeO_{2-x} and their further migration to nickel nanoparticles, where they can assist in methane activation. The methane rate of 0.62 mmol (g_{cat} • min)⁻¹ was achieved during illumination with the lowest energy photons (λ > 600 nm), which can undergo only sub-bandgap electron transitions. The most pronounced rate acceleration was observed with wavelengths between 450 and 550 nm, which correlates with the wavelength dependence of the near-field electromagnetic intensity enhancement on nickel nanoparticles (Fig. 3C).

The effect of irradiance on the CH₄ and CO₂ rates, as well as H₂/CO ratio over the 2Ni catalyst, was probed at the constant catalyst temperature of 395 °C (Fig. 6B). To maintain the catalyst temperature constant with increasing irradiance, the output of the electric heater was decreased simultaneously. The dependence of photocatalytic rate on light intensity (I) reveals plenty about the reaction mechanisms in light-assisted reactions [51]. Four different kinetic categories are classified: sublinear (rate ∝ Iⁿ, n < 1), linear (rate ∝ I), superlinear (rate ∝ Iⁿ, n > 1) and exponential (rate ∝ e^(I)) [52]. Plasmon-induced photocatalysis has been demonstrated to exhibit each of these regimes, depending on the operating conditions, except for the sublinear dependence, which is characteristic of non-plasmonic semiconductor catalysis when charge carrier recombination is dominant [53]. The exponential dependence is characteristic for thermally driven reactions. Both linear (rate ∝ I) and superlinear (rate ∝ Iⁿ, n > 1) regimes are distinctive features of electron-driven chemical transformations on metal surfaces, but the reaction conditions are the key factor determining the prevalence of each of them [51].

The acceleration of CH₄ and CO₂ rates, as well as H₂/CO ratio, started to increase notably as irradiance exceeded 400 mW cm⁻². At

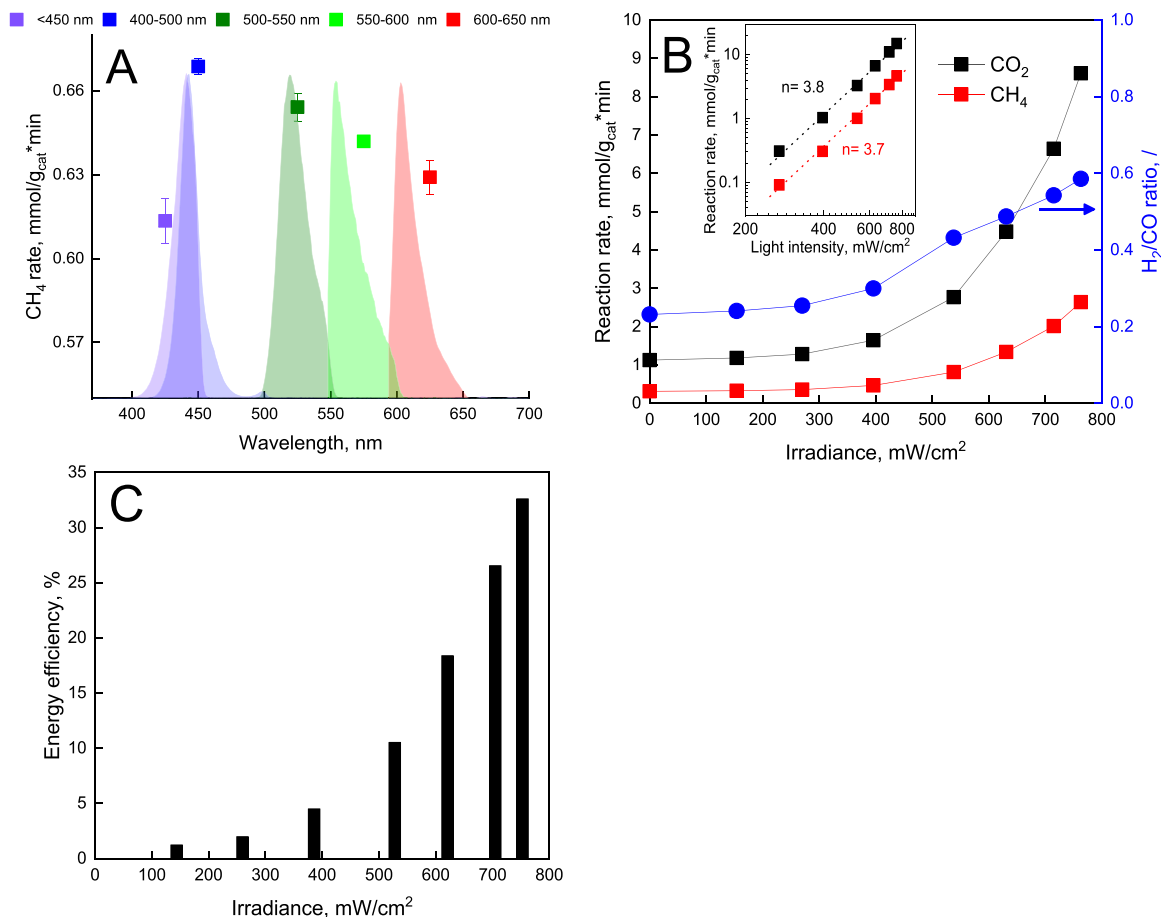


Fig. 6. (A) The CH₄ rate as a function of wavelength at a constant irradiance of 300 mW cm⁻² and constant catalyst temperature of 395 °C. Colored cones show light intensity dependence on wavelength when using different bandpass filters. (B) CH₄ and CO₂ rate with corresponding H₂/CO ratio as a function of irradiance at a constant catalyst temperature of 400 °C. (C) Energy efficiency related to the experiment shown in panel B.

790 mW cm⁻², the CH₄ and CO₂ rates were accelerated by 852% and 769% compared to the thermocatalytic experiment in the dark at identical catalyst temperature, and the H₂/CO ratio increased from 0.23 to 0.59. The inset in Fig. 6B shows the superlinear dependence of both CH₄ and CO₂ rates on irradiance over 2Ni/CeO_{2-x} catalysts. This again points to a conclusion that electron-driven chemical transformations on metal surfaces, and not thermal energy are responsible for rate acceleration.

The calculated energy efficiency (Fig. 6C) increased rapidly at irradiances above 400 mW/cm⁻² and reached 33% at 790 mW cm⁻². This revealed the 2Ni as an efficient photocatalyst for visible light utilization.

Hot carriers, generated by plasmon decay in the nickel nanoparticles were analyzed by DFT, as they could alter the reaction pathway of methane dissociation. Initial hot carrier probability distribution can be obtained from calculated density of states (DOS, see Supplementary information and Fig. S15) for photoexcitation with energy E_{ph} and Fermi distribution at room temperature. Probability distributions for hot carrier generation suggest that for photon energies corresponding to visible light (up to 3.26 eV or 380 nm), nickel should not be an efficient source of hot carriers, Fig. S16.

Cumulative hot carrier generation probability (Fig. S17) shows that the trend for hot carrier generation increases gradually and almost linearly with the increasing photon energy in the visible range, after which it sharply increases in the UV range. In contrast, the experimental data in Fig. 6A shows a sharp decrease of reaction rate below 450 nm (above 2.76 eV). Had the reaction been catalyzed by hot carriers generated in Ni nanoparticles, the reaction rate would be expected to follow the trend of hot carrier generation probability and continue increasing as the

wavelength gets shorter. It has to be noted that hot carrier generation probability is an intrinsic property of the material and the near-field intensity could change the absolute number of generated hot carriers and, consequently, the rate of reaction catalyzed by hot carriers. The aforementioned moderate hot carrier generation probability in the spectral range where the experiments were carried out, diverging trends in generation probability and reaction rate as the photon energy approaches UV range, and low near-field intensity enhancement inside Ni nanoparticles suggest that hot carriers are unlikely to be generated in Ni nanoparticles.

4. Discussion

Based on the characterization, theoretical and catalytic data, we can postulate the light-assisted DRM mechanism over the Ni/CeO_{2-x} catalyst. First, we discuss the situation where the catalyst is irradiated with photons having energy higher than its bandgap (hot-carrier mediated photocatalytic mechanism, Fig. 7A).

During DRM reaction, nickel is present as metallic nanoparticles [8] measuring about 5 nm and ceria as a partly reduced (CeO_{2-x}) semiconductor [8]. Ceria is an n-type semiconductor whose work function (5.34 eV) is larger than that of nickel (5.01 eV) [54]. After illuminating the Ni/CeO_{2-x} catalysts with photons having energy higher than 2.84 eV, (the absorption edge of 2Ni-R was estimated at ~ 440 nm, Fig. 2), the photo-excited electrons migrate from the 4f orbital of CeO_{2-x} to its conduction band, and are transferred to nickel. An upward band bending in the n-type semiconductor is created due to the accumulation of excess positive charge in the semiconductor caused by electron

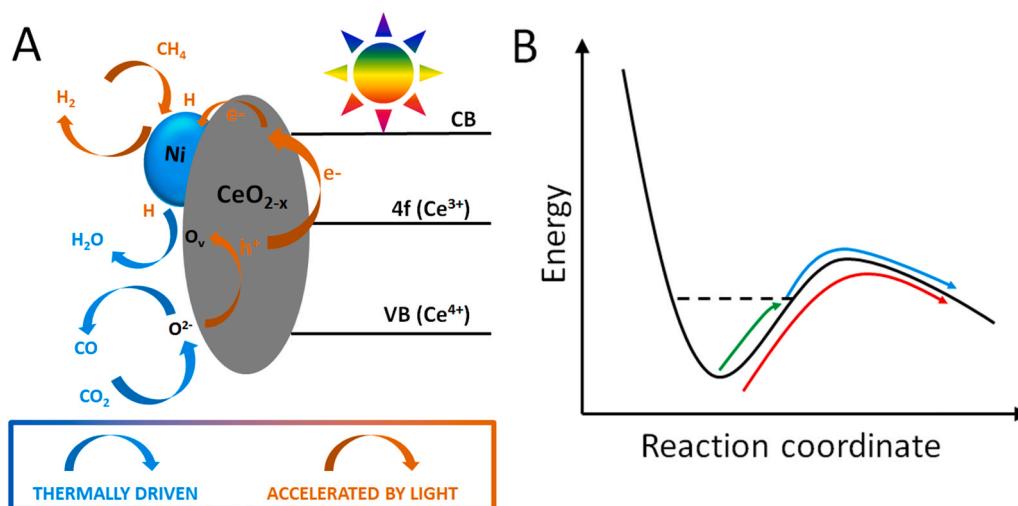


Fig. 7. A) Schematic representation of the hot-carrier driven photocatalytic mechanism on the Ni/CeO_{2-x} catalyst and B) alleviation of the energy barriers due to near field electromagnetic enhancement energy transfer.

migration. As a result, a Schottky barrier is created, which serves as an electron trap, leading to electron accumulation on the nickel and prevents their migration back to the semiconductor. The hot electrons on nickel can be transferred to the LUMO of adsorbed methane and electronically excite the CH_x species, thus accelerating C–H bond cleavage (orange arrows in Fig. 7A). Hot electrons can also participate in H⁺ and e⁻ recombination, leading to accelerated formation and desorption of molecular H₂. This should result in higher abundance of free nickel sites where activation of adsorbed methane molecules can occur, resulting in accelerated methane activation and overall increase of DRM rate (considering methane activation is the RDS of reaction).

The role of light on the photocatalytic CO₂ reaction pathway was tested with the 2Ni catalyst in the light-assisted RWGS reaction (CO₂ + H₂ ↔ CO + H₂O). The concentrations of H₂ and CO₂ were very similar to the values measured during the DRM test (Fig. S18). The RWGS reaction is rate-determined by the dissociation of CO₂, which takes place over partly reduced CeO₂ [55]. The H₂ dissociation is much faster and occurs over nickel with a low activation barrier (0.1 eV for Ni [111]). Since H₂ activation on nickel is kinetically fast, its acceleration should have a minor role on the RWGS rate, as CO₂ activation is the step which controls the overall RWGS reaction rate.

During progressively increasing irradiance from 0 to 790 mW cm⁻² (and a simultaneous decrease of thermal energy input, resulting in constant catalyst temperature of 400 °C), the CO rate increased by 11%. During the DRM reaction at identical reaction conditions, CH₄ rate increased by 852% (Fig. 6B). Thus, the very different response of the Ni/CeO_{2-x} catalyst during light-assisted RWGS and DRM reactions suggests that visible light does not play any meaningful role in accelerating CO₂ dissociation under tested reaction conditions. However, it strongly accelerates CH₄ dissociation over nickel.

Our experimental findings are in accordance with those of Zhang et al. [47], who used DFT to analyze the energetics of elementary reaction steps on nickel in ground and excited state (simulating illumination). They observed no decrease in activation barriers in the CO₂ reaction pathway, whereas activation barriers for the CH dissociation and oxidation of C and CH species, leading to accelerated DRM rates.

Based on the above, we can postulate that CO₂ activation on partly reduced ceria during DRM under reaction conditions studied is exclusively thermally driven, whereas methane activation over nickel is a light-assisted process. One has to keep in mind that methane activation is the rate determining step of the DRM reaction and the (photocatalytic) DRM rate acceleration will be noticed until the methane dissociation catches up with the CO₂ dissociation rate.

The catalytic role of hot holes (h⁺), accumulated on CeO_{2-x} is more

difficult to establish. The *in-situ* DRIFTS is a powerful surface characterization tool and was used to analyze CO₂ activation over the 2Ni catalyst in thermally-driven and light-assisted modes (see [Supplementary information](#) for more details, Fig. S19). The characteristic bands of carbonates are analyzed, which populate exclusively the ceria surface. The mono- and bidentate carbonates are the active intermediates in the CO₂ activation and dissociation pathway, whereas polydentate is an inert spectator at temperatures below 500 °C.

The surface carbonates on oxidized 2Ni sample in dark (Fig. S19A) are present mainly as polydentate (1454 cm⁻¹), accompanied by some monodentate (1588 cm⁻¹) and bidentate carbonate species (1282 cm⁻¹). Upon illumination, no change occurs in the polydentate carbonate band and a very modest change in the mono and bidentate.

Contrary, over the reduced 2Ni catalyst in dark, the carbonate distribution is very different (Fig. S19B). The monodentate and bidentate prevail over the polydentate, indicating strongly altered CO₂ adsorption sites, compared to the oxidized sample. Catalyst illumination induces a drastic decrease of the monodentate and bidentate carbonate coverage, and simultaneously increases the polydentate band intensity.

Finally, the carbonate DRIFT spectra of the reduced 2Ni sample in H₂/CO₂ atmosphere in dark (Fig. S19C) show dominant monodentate vibrations (due to more abundant Ce³⁺ at 400 °C in H₂/CO₂ flow, compared to CO₂ flow, Fig. S19B), which are attenuated upon illumination, with a concomitant rise of the polydentate band. The monodentate band decrease is less pronounced compared to the one in pure CO₂ flow (Fig. S19B and C), as H₂ atmosphere maintains a higher fraction of Ce³⁺.

Based on the similarity between carbonate bands of oxidized 2Ni, and reduced illuminated 2Ni catalysts, oxidation of ceria can be assumed to occur during illumination.

Upon illumination of the Ni/CeO_{2-x} catalyst, the electron is excited from the 4f orbital of Ce³⁺ to the conduction band and migrates to nickel, causing h⁺ accumulation on ceria. Hot holes can participate in the reaction with surface adsorbed oxide species (O²⁻) originating from CO₂, and fill the oxygen vacancy sites, causing ceria re-oxidation. This process should not be mistaken with CO₂ activation, which is driven by electron transfer [56] from electron rich Ce³⁺ sites to the adsorbed CO₂, producing CO₂⁻. The electron transfer process is the rate-limiting step for CO₂ splitting over ceria based materials at moderate temperatures [57].

The light-assisted rate acceleration does not correlate with decreasing wavelength (increasing photon energy), which is expected for the hot-carrier driven mechanism. This suggests the existence of an additional light-driven reaction mechanism, which is depicted in Fig. 7B (shown for ground state when the catalyst is illuminated by photons

having energy lower than the CeO_{2-x} bandgap). Namely, photons with a shorter wavelength can excite electrons to higher energy levels in the conduction band (CB), increasing the over-potential for charge carriers and thus increasing their reactivity. Furthermore, the methane rate, achieved by illuminating with photons having substantially lower energy than what is required for VB to CB transition in ceria, is higher compared to the methane rate achieved by photons with sufficient energy that enable bandgap transitions (Fig. 6A). The near-field electromagnetic enhancement is strongly dependent on the illumination wavelength and reaches a maximum at about 500 nm (Fig. 3C). Upon illuminating the 2Ni catalyst with 300 mW cm^{-2} of light using a band-pass filter transparent for wavelengths between 500 and 550 nm, a methane rate of $0.65 \text{ mmol (g}_{\text{cat}} \bullet \text{min})^{-1}$ was achieved which represents a 33% increase compared to the experiment in the dark. The localized near field enhancement induces resonant energy transfer into the Ni- CH_x bond (green arrow in Fig. 7B). As the vibrational energy stored in the bond increases, it gets progressively more destabilized [58]. As a result, the energy barrier which remains to be overcome by thermal energy (blue arrow in Fig. 7B) is lowered, which manifests as lower apparent activation energy (inset Figs. S11A and S12A) and accelerated DRM rate. In the thermally-driven mode (red arrow in Fig. 7B), the activation barrier remains high because thermal energy is the only means of activation of adsorbed reaction intermediates, which drives the catalytic reaction.

5. Conclusions

This work analyzes the underlying mechanism of visible light assisted DRM rate acceleration over Ni/CeO_{2-x} catalysts. The *in-situ* UV-Vis analysis and simulation of electromagnetic properties revealed that the presence of Ce³⁺ in partly reduced ceria is crucial for visible light absorption. The light-assisted DRM test showed that methane and CO₂ rates, as well as hydrogen selectivity, are greatly improved compared to thermally driven experiments at identical catalyst temperatures. Two weight percent nickel was found as optimal active metal loading, enabling high (photocatalytic) reaction rates, energy efficiency of 33% and minimal carbon accumulation. Photocatalytic DRM activity is maintained over the 2Ni catalyst in purely photocatalytic mode with 790 mW cm^{-2} of white light irradiance and without external heating. Simulation of electromagnetic properties identified the electromagnetic near field intensity enhancement occurs mainly over nickel nanoparticles, which are the active sites for methane activation. The light assisted DRM rate is wavelength dependent and reaches a maximum at about 500 nm, which also triggers the highest near field enhancement. Two mechanisms exist simultaneously during white light illumination: hot carrier driven charge transfer, employing photons with energy higher than the bandgap of partly reduced ceria and near field enhancement induced resonant energy transfer, concentrated over the nickel-ceria interphase, stimulated by sub-bandgap absorption of low energy photons. This work can act as a cornerstone for further developing Ni/CeO₂ based catalysts for visible light-driven methane activation under mild conditions.

CRedit authorship contribution statement

Kristijan Lorber: Investigation, Writing – original draft. **Janez Zavašnik:** Investigation, Formal analysis, Writing – review & editing. **Jordi Sancho-Parramon:** Formal analysis, Writing – review & editing. **Matej Bubas:** Formal analysis, Writing – review & editing. **Matjaž Mazaj:** Investigation. **Petar Djinović:** Conceptualization, Supervision, Resources, Writing – review & editing, Funding acquisition.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence

the work reported in this paper.

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at doi:10.1016/j.apcatb.2021.120745.

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