

# Conversion of CH<sub>4</sub> / CO<sub>2</sub> by a Nanosecond Repetitively Pulsed Discharge

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**Abstract.** A possible way to store both renewable energy and CO<sub>2</sub> in chemical energy is to produce value-added chemicals and fuels starting from CO<sub>2</sub> and green electricity. This can be done by exploiting the non-equilibrium properties of gaseous electrical discharges. Discharges, in addition, can be switched on and off quickly, thus being suitable to be coupled with an intermittent energy source. In this study, we have used a nanosecond pulsed discharge to dissociate CO<sub>2</sub> and CH<sub>4</sub> in a 1:1 mixture at atmospheric pressure, and compared our results with literature data obtained by other discharges. The main products are CO, H<sub>2</sub>, C<sub>2</sub>H<sub>2</sub>, water and solid carbon. We estimate an energy efficiency of 40% for syngas (CO and H<sub>2</sub>) production, higher if also other products are considered. Such values are among the highest compared to other discharges, and, although not very high on an absolute scale, are likely improvable along possible routes discussed in the paper and by coupling to the discharge a heterogeneous catalysis stage.

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## 1. Introduction

Satisfying the increasing energy demand while reducing negative impacts on the Earth climate is a paramount challenge in the present days. In the past 10 years, the energy scenario has changed radically due to the increased availability of natural gas, not to say coal. The present established reserves of natural gas of about 186 trillion cubic meters compare with an annual (2013) world consumption of about 3348 billion cubic meters, which means they will meet the gas demand for at least 55 years [1]. In addition, unconventional gas sources, thought currently not exploited, might play an increasing

role in the future. However, a main problem with natural gas is its low energy density in comparison with a liquid fuel. As an example, in order to be shipped overseas, natural gas must be cryogenically liquefied, with obvious costs. Thus finding new techniques to convert gas into a liquid product is a valuable option.

Nowadays the main concern is the increasing atmospheric concentration of  $CO_2$ . In the recent past, it remained almost constant at about 280 ppm until about two hundred years ago, when it started to grow in conjunction with the mounting exploitation of fossil fuels. At present,  $CO_2$  concentration stands at 400 ppm, and consequently the power of the Earth's greenhouse effect has been enhanced.

To cut down the emission of  $CO_2$ , the obvious way is to substitute fossil fuels with renewable energy sources, which are  $CO_2$  neutral but frequently intermittent, and to recycle  $CO_2$  to produce value-added chemicals and fuels [2]. In fact, solar energy is quite abundant, about four orders of magnitude the current energy consumption, but needs to be stored and transported.

In this framework, the production of syngas ( $CO + H_2$ ) from  $CO_2$  by using solar energy can be a way to store intermittent renewable energy into the chemical production chain. Converting  $CO_2$  into chemical fuels requires a source of hydrogen. If  $CH_4$  is used, as in the so called dry reforming reaction  $CH_4 + CO_2 \rightarrow 2CO + 2H_2$ , two main greenhouse gases are used. In addition,  $CH_4$  can be eventually converted via gas-to-liquid chemistry into a high energy-density fuel. Biogas valorization is an ideal application of such a technology, since renewable sources of methane, carbon dioxide and energy can be used to produce value-added chemicals and eventually liquid fuels by the Fischer-Tropsch process.

The dry reforming is a very endothermic reaction ( $H_{298K} = 247 \text{ kJ mol}^{-1}$ ), since both  $CO_2$  and  $CH_4$  are quite stable molecules. Thermodynamics calculations [3] indicate that a temperature of at least  $1000^\circ\text{C}$  is needed to get appreciable conversions. Low temperature values can be reached by using catalysts, but then the problem is their deactivation by carbon deposition.

Therefore, an attractive possibility is to consider the non-equilibrium properties of gaseous electrical discharges for channeling energy in the molecular dissociation rather than in heating the gas [4]. A further advantage of using plasma techniques to dissociate  $CO_2$  and  $CH_4$  is directly related to the storage of green electricity. Since discharges can be switched on and off quickly, plasma systems are suitable to be coupled with an intermittent electric power source, thus providing a flexible and scalable technology for storing renewable energy into chemical energy.

By far the most studied discharge is the Dielectric Barrier Discharge (DBD), due to its simplicity and reliability [5]. When  $CO_2$  and  $CH_4$  are the feed gases, a DBD produces syngas and light hydrocarbons, but also liquid oxygenates [6, 7] and liquid hydrocarbons [8, 9]. Unfortunately, both the conversion rate and the global energy efficiency are low. To overcome these limitations, the synergy between plasma and heterogeneous catalysis is currently actively investigated [10]. A complementary route is to attempt different discharge configurations, such as corona [11], microwave [12], spark

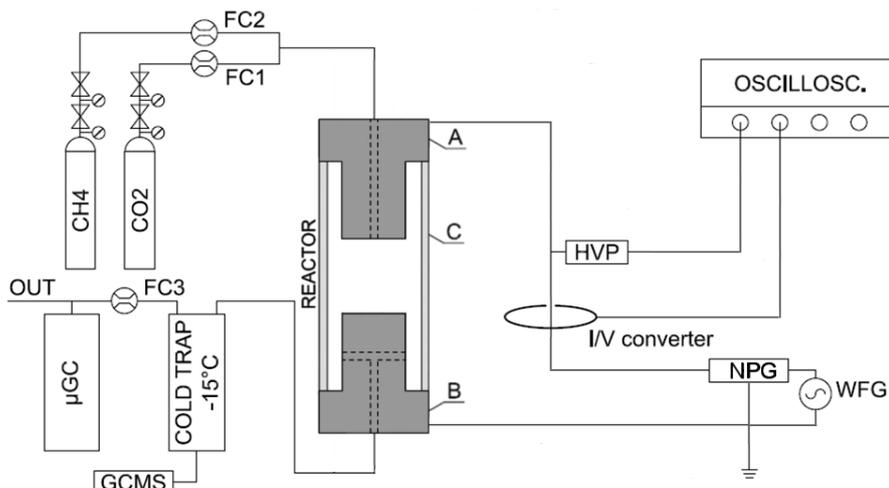


Figure 1: Schematic representation of the experimental setup. A HV electrode; B grounded electrode; C glass tube reactor; HPV High Voltage Probe; NPG Nanosecond Pulsed Generator; WFG wave-form generator; FC mass flow controller.

[13] and gliding-arc [14]. In this context, a repetitive pulsed excitation with ns scale pulse rise time and duration (NRP) appears to be a promising candidate due to its highly non-equilibrium nature [15]. In this paper we report on experiments carried out by an atmospheric pressure NRP discharge in a  $\text{CH}_4$ - $\text{CO}_2$  mixture, with a detailed analysis of the products, not limited to syngas only, which is very important to define correctly the energy efficiency. We present results on reactant conversion, product selectivity and energy efficiency, and compare them with available literature data relevant to other discharge types.

We find that the NRP energy efficiency is larger than the values of DBD, corona and microwave discharges, and comparable to that of gliding arcs. Possible limiting factors to the efficiency increase are discussed.

## 2. Experimental Methods

The experimental setup is shown in figure 1. The reactor is made of a quartz tube (internal diameter 10 mm, external diameter 13 mm), containing two brass discs (diameter 8 mm). The discharge in plane-to-plane configuration occurs in an inter-electrode gap that can vary up to 10 mm.

The discharge is produced by a nanosecond-scale pulsed power supply (NPG 18/3500, Megaimpulse Ltd.), triggered by a wave-form generator (WFG, 33220A, Agilent Technologies Inc.). Discharge current and voltage values are measured by a I/V converter (CT-D-1.0, Magnelab) and a high-voltage probe (P6015A, Tektronix), respectively. V/I signals are recorded by a digital oscilloscope (WaveSurfer 104MXs-A,

LeCroy). The pulse repetition rate is controlled by the WFG and ranges from 100 up to 3000 Hz.

Two mass flow controllers set the input flow; a third one, placed at the exit of a cold trap at  $-15^\circ\text{C}$ , records the output flow. The input flow is changed between 200 sccm and 600 sccm, which translates in a variation of the residence time in the discharge between 0.012 s and 0.036 s for a gap of 2.5 mm.

On-line gas detection is performed by using a gas-chromatography (3000 Micro GC, Agilent Technologies Inc.). Helium, hydrogen, methane and carbon monoxide were measured by a Molesieve column with back-flush; carbon dioxide, ethane, ethylene, acetylene, propane and propyne by using a Plot U column. Standard compounds were used to calibrate the instrument. A small amount ( $< 0.5\%$ ) of pure helium was added to the reactant flux as internal standard to accurately take into account possible variations of the response factor of the third mass flow meter. Compounds condensed in the cold trap were dissolved in acetonitrile and analyzed using a GC-MS (Trace GC Ultra, Finningham). Water, the main liquid by-product, was estimated by the standard addition method with a Carbowax column. The quantitative measurement of water requires very long discharge runs in order to collect a sufficient amount. For such a reason, we did not carry it out routinely. Solid carbon was collected from the reactor wall and characterized by scanning electron microscopy, FTIR and XPS spectroscopy. Carbon quantitative measurement is prevented by the impossibility to completely collect it.

### 2.1. Discharge power measurement

The discharge power measurement is a delicate issue when dealing with short voltage and current pulses. The time delay introduced by the acquisition system introduces a spurious phase shift between  $I$  and  $V$ , that in turn affects the power estimation. We measured this time delay, with the discharge off, by reducing to zero the time integral of the  $I \times V$  product [16]. To this purpose, we maintained the same circuit configuration and reactor geometry, by filling the reactor with Freon-113 vapor to prevent breakdown. For a 2.5 mm gap, the time delay was estimated ( $2.15 \pm 0.09$ ) ns. The instantaneous power is then calculated as the  $I \times V$  product accounting for the spurious delay. The pulse energy is the time integral of the instantaneous power. Figure 2 shows an example of the recorded voltage and current signals. The instantaneous power and pulse energy are also shown.

### 2.2. Process characterization

Quantities for process characterization must be chosen accurately. Many parameters have been introduced in the literature to compare different discharges. Some authors consider only the conversion of reactants, defining the conversion ability [12] or the energy efficiency [17] or the energy cost for converting  $\text{CH}_4$  and  $\text{CO}_2$  [18]. These choices are limiting, since not all the products are energetically useful, and therefore

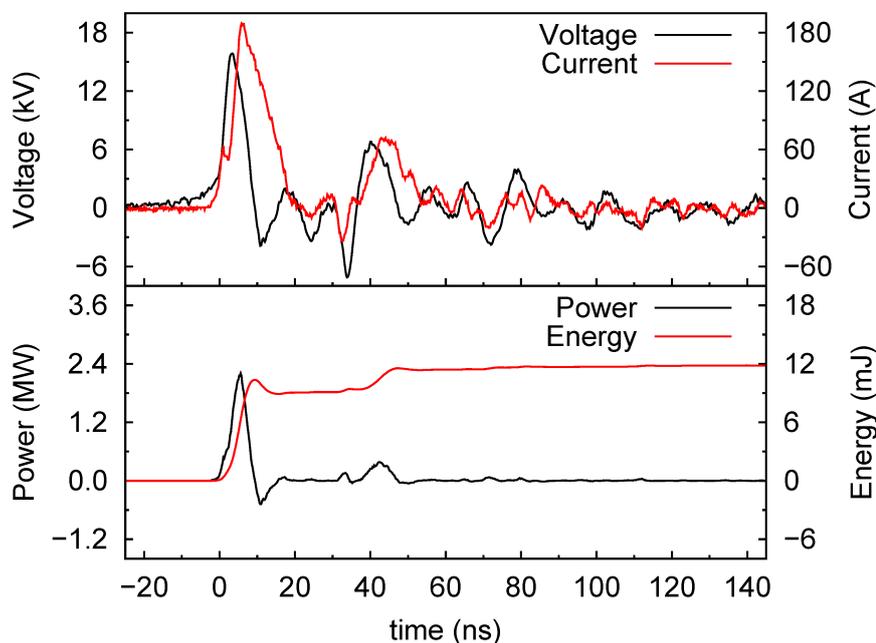


Figure 2: Voltage, current, instantaneous power, and energy for a 6 ns FWHM, 15.8 kV pulse.

efficiency calculations based only on conversion substantially overestimate the efficiency and mask its trends as a function of discharge parameters. Tao *et al* [19] introduce specific energy and energy conversion efficiency, considering only  $\text{H}_2$  and  $\text{CO}$  as products, while Zhu *et al* [18] calculate the energy cost for  $\text{H}_2$  production. Also these quantities are incomplete, since other hydrocarbon products should be considered in the energy evaluation, especially if their selectivity is larger than 10%.

Following [20], we consider the energy conversion efficiency, ECE, defined as the ratio between the energy contained in the products and the sum of the energy of the converted reactants with the energy injected in the plasma. In the following we give the formulas for the quantities used in the paper. Setting:

- $n_r^{\text{in}}$  and  $n_r^{\text{con}}$  the input and converted moles of reactants  $r$ ;
- $P$  and  $E$  the power and energy of the discharge;
- $\Phi_r$  the total flux of reactants;
- $n_x$  the number of moles of specie  $x$ ;
- $\text{LHV}_r$  and  $\text{LHV}_{\text{pr}}$  the Lower Heating Value of reactants and products.

We define the conversion of reactants (methane or carbon dioxide):

$$C_r = \frac{n_r^{\text{con}}}{n_r^{\text{in}}} \times 100. \quad (1)$$

The specific energy input – SEI – i.e. the average amount of energy injected in the plasma per unit volume:

$$SEI = \frac{P}{\Phi_r} \quad (\text{kJ dm}^{-3}). \quad (2)$$

The gaseous products selectivity:

$$\begin{aligned} S_{H_2} &= \frac{n_{H_2}}{2 n_{CH_4}^{con}} \times 100; \\ S_{CO}^C &= \frac{n_{CO}}{n_{CH_4}^{con} + n_{CO_2}^{con}} \times 100; \\ S_{CO}^O &= \frac{n_{CO}}{2 n_{CO_2}^{con}} \times 100; \\ S_{C_xH_y}^C &= \sum_{x,y} \frac{x n_{C_xH_y}}{n_{CH_4}^{con} + n_{CO_2}^{con}} \times 100; \\ S_{C_xH_y}^H &= \sum_{x,y} \frac{y n_{C_xH_y}}{4 n_{CH_4}^{con}} \times 100. \end{aligned} \quad (3)$$

In which the hydrocarbon selectivity is the sum over all the measured hydrocarbon products (total hydrocarbon). The Energy Conversion Efficiency – ECE:

$$ECE = \frac{LHV_{pr}}{LHV_r + E} \times 100. \quad (4)$$

And, finally, the Energy Storage Efficiency – ESE:

$$ESE = \frac{LHV_{pr} - LHV_r^{con}}{E} \times 100. \quad (5)$$

### 3. Results and Discussion

The discharge has generally a glow structure, but its nature shows differences depending on the gap between the electrodes. For a gap of 2 mm, the presence of few filaments was observed, while at a higher gap (8 mm), the filaments disappear and the discharge extends up to the quartz tube walls. For the latter configuration, we guess that carbon powder formation on the glass might increase the surface conduction and promote the surface discharge, thus changing the nature of the discharge itself. For this reason, we report only the results for a gap of 2.5 mm, where the higher stability allows to work in a wider range of pulse repetition rates. Conversion, selectivity and ECE/ESE have been calculated from measured products and are reported as a function of SEI. The SEI is varied up to  $10 \text{ kJ dm}^{-3}$ , by varying both the HV pulse repetition rate and the reactant flux. By changing  $\Phi_r$  and  $P$ , such as SEI remains constant, we got similar results. The SEI upper limit is set by the difficulty of handling the experiment at low flux values. For the highest SEI of  $10 \text{ kJ dm}^{-3}$ , we estimated a temperature of the reactor exhaust around  $170^\circ\text{C}$  by using a thermocouple.

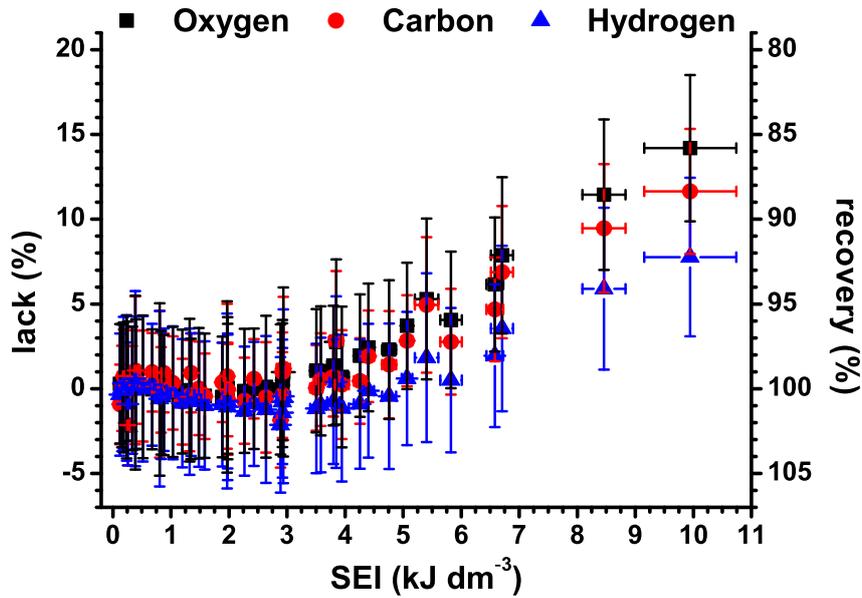


Figure 3: Mass balance as a function of SEI. Lack is the missing percentage of a given element in the mass balance. Recovery is the percentage of a detected element with respect to the amount contained in the reactant.

### 3.1. Mass balance

The mass balance, calculated from gaseous products data, is shown in figure 3. At SEI lower than  $3 \text{ kJ dm}^{-3}$ , the recovery is total. At higher SEI, the balance indicates that a significant amount of mass is not detected, and this behavior increases with SEI.

Oxygen is the specie with the lower recovery, we guess due to water formation. Indeed, in a long discharge run at  $\text{SEI}=6.5 \text{ kJ dm}^{-3}$ , we have measured about 500 mg of water that fits quite well with the 600 mg calculated from the oxygen lack.

Carbon lack is due to the formation of carbon powder, which is dispersed throughout the reactor. Its analysis confirms the presence of C–C and C–H bonds, both aliphatic and aromatic.

### 3.2. Reactants conversion

The reactant conversion is shown in figure 4a and figure 4b. Methane conversion is roughly linear with the SEI variation, and its value is higher than for  $\text{CO}_2$  conversion, similarly to what observed in other cold plasmas [21] and model calculations [22].  $\text{CO}_2$ , instead, shows a rather parabolic behavior. The rationalization of these results requires complex model calculations. We just note that, if we calculate the total conversion,  $0.5 \text{ C}_{\text{CH}_4} + 0.5 \text{ C}_{\text{CO}_2}$ , we find a non-linear behavior that looks similar, although in a different SEI range, to that measured in a coaxial DBD reactor and calculated by a 0 dimensional kinetic model [23].

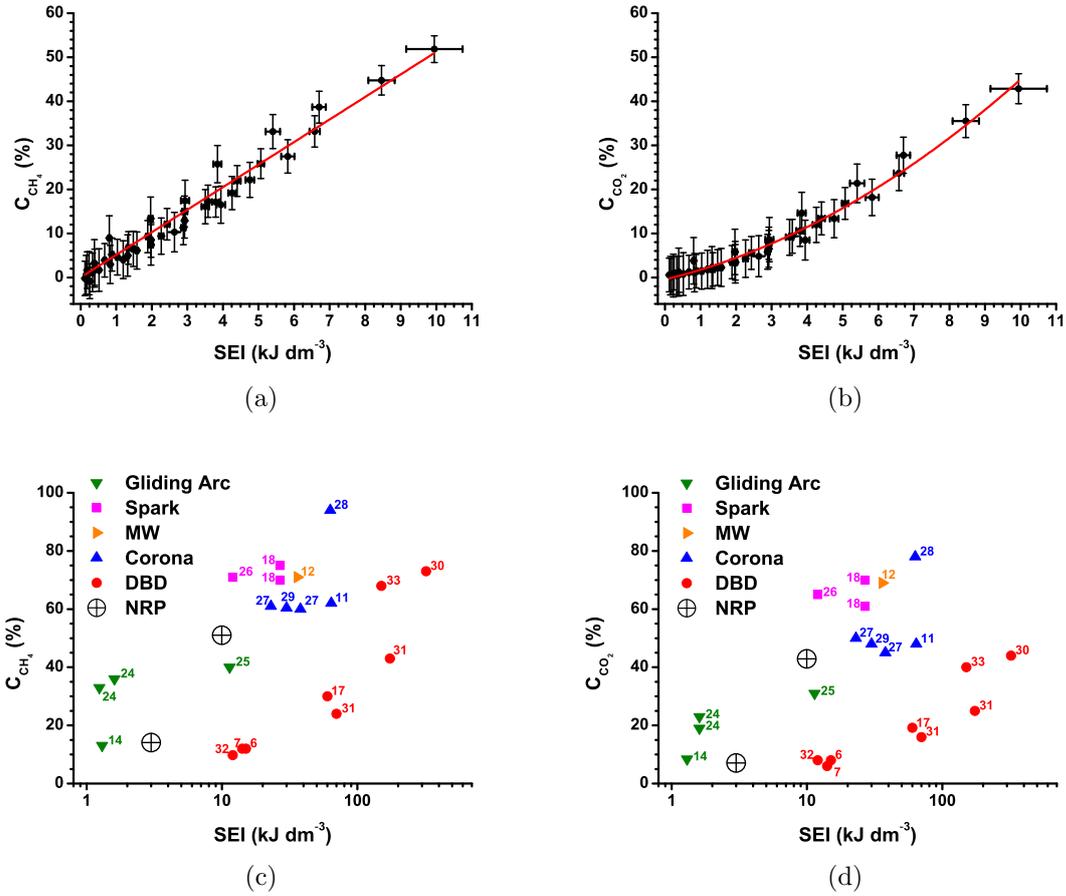


Figure 4: Conversion rates as function of the SEI. Present data are reported in (a) and (b), while in (c) and (d) literature data for various discharges are compared. For simplicity, we show our results for two SEI values only. Numbers correspond to references.

In figure 4c and figure 4d we compare conversion values for different plasmas, whose values are calculated from recent literature. At higher SEI the conversion increases for all the discharges. DBDs are easy to develop and well-studied, but they show low performance. Corona, another well-known discharge, has better performance, but the discharge volume is very small in comparison to others. Microwave discharges have good performance and large volumes, but require complex systems. Gliding, spark and NRP have similar performances.

### 3.3. Product selectivity

The selectivity, as defined in (3), is reported in figure 5. Analysis of the discharge effluents shows that the main products are  $\text{H}_2$  and  $\text{CO}$ . Their selectivity with respect to hydrogen and carbon are almost constant with SEI variation. The selectivity of  $\text{CO}$  respect to oxygen shows an higher decrease, which we attribute to water formation.

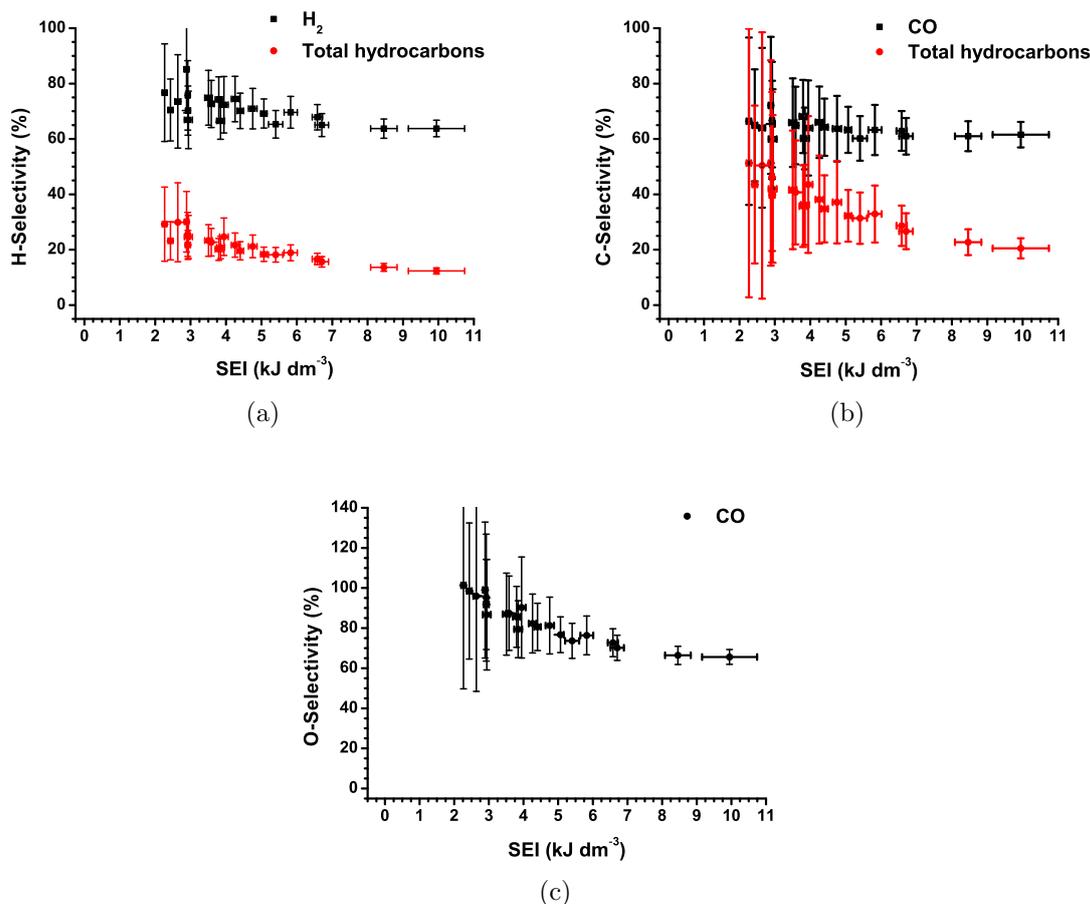


Figure 5: Selectivity. (a): respect to hydrogen; (b): respect to carbon; (c): respect to oxygen.

For hydrocarbons, the selectivity with respect to both hydrogen and carbon slightly decreases with SEI.

The most abundant hydrocarbon product is acetylene, but other alkynes, as methylacetylene, are also detected. Since in DBD reforming the main product is ethane [6], this difference suggests a different initial radical abundance, i.e. in NRP  $\text{CH}$  prevails on  $\text{CH}_3$ , while in a DBD the reverse occurs. This hydrocarbon distribution is more similar to that found in gliding arc or spark discharges, with a prevalence of unsaturated byproducts [17, 26].

At low SEI, hydrogen, CO and light hydrocarbons are the main products. Increasing the discharge power, water and carbon powder become important products.

### 3.4. Energy efficiency

We have selected ECE to evaluate the process efficiency, since this quantity correctly accounts for the amount of chemical energy stored in the products. In figure 6a, ECE is plotted vs SEI, both for syngas only and for all the gaseous products (carbon powder

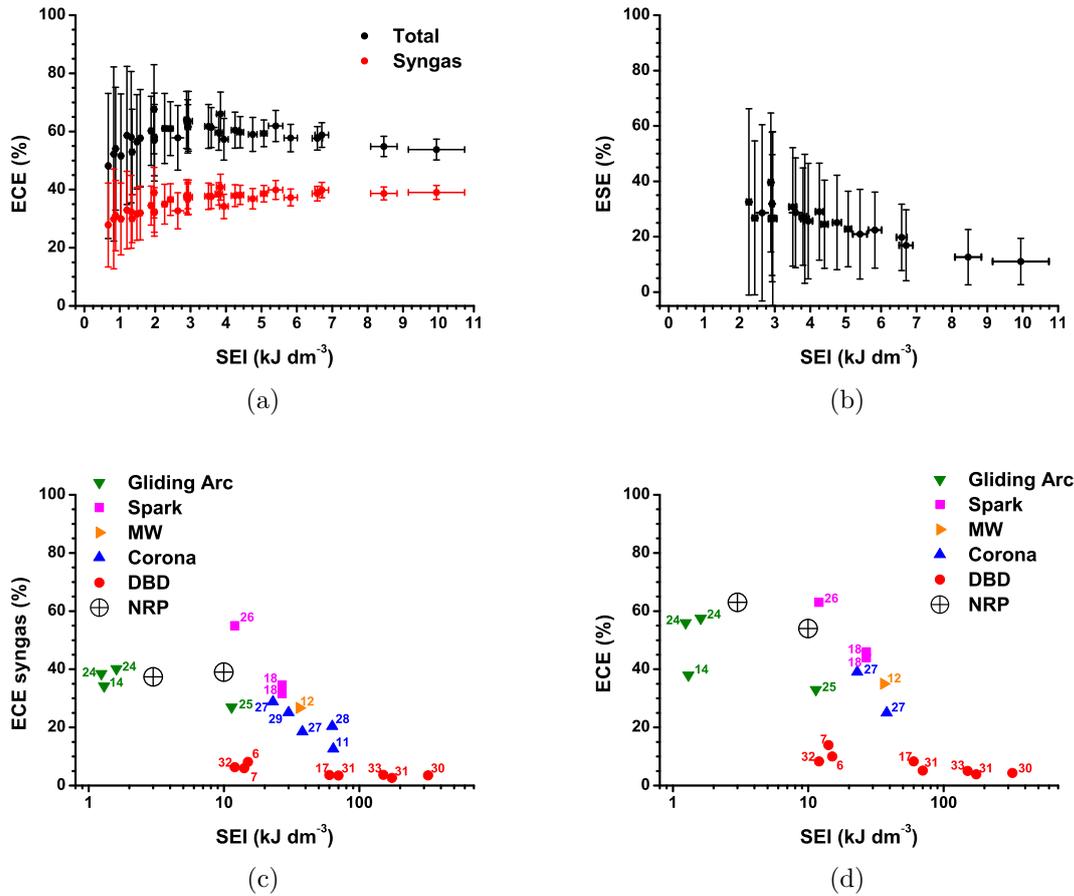


Figure 6: ECE and ESE as function of SEI. Present data are shown (a) and (b), while in (c) and (d) a comparison with literature data for various discharge types is reported. For simplicity, we show our results for two SEI values only. Numbers correspond to references.

is still excluded). The difference between the two ECE values suggests that the energy stored in products other than syngas is not negligible. The energy efficiency reaches a plateau at about  $3 \text{ kJ dm}^{-3}$  and then starts to decrease slowly, likely due to a competing production of water and carbon. This appears to be a limiting factor that prevents the efficiency to increase as a function of SEI.

Considering other cold plasma techniques, NRP has better performance with respect to DBDs, MW and corona discharges, and similar to gliding and spark discharges. We finally observe that the best efficiency at low SEI values, and the decrease of efficiency at high SEI, is a general feature of all the kinds of discharges. Therefore we guess that water and carbon formation might be a general limiting factor of the plasma treatment, occurring in different SEI ranges according to the energy deposition mechanisms of the various discharge technologies.

ECE expresses the efficiency of total process, but it does not give information on

the efficiency of energy storing. The energy efficiency in energy storage can be calculated as the difference of LHV between products and reactants divided by discharge energy (see (5)). It is shown in figure 6b. The best performance we achieve is about 30%.

In figure 6c and figure 6d we compare our ECE values with literature data. The latter have been calculated by using the products reported in each reference.

In addition to energy, also  $CO_2$  can be stored, since a fraction of carbon contained in the products derives from  $CO_2$ .

#### 4. Conclusions

The main conclusions we can draw are:

- (i) The energy efficiency of NRP treatment is among the highest compared to other kinds of discharge. Its value is sufficiently high as to justify further research to improve it.
- (ii) Both  $CH_4$  and  $CO_2$  conversions increase as a function of SEI. However, the selectivity towards syngas decreases, likely due to a competing production of water and carbon powder.
- (iii) The bare discharge is not totally selective towards syngas.

To improve the process one should first of all have clear in mind the final target: for example, syngas production, liquid fuels production, energy storage, and  $CO_2$  disposal.

The step forward must seek for selectivity improvements, playing with discharge parameters [6], or with heterogeneous catalysis process integration [17].

From the point of view of energy efficiency, a key point seems to be finding a way to inhibit water formation (note that the energy content of carbon powder can be in principle re-utilized, then increasing the global energy efficiency). For this purpose, we should first of all understand water formation mechanisms. From present results, however, we already understand that working at low SEI values is advantageous, and we might envisage a cascade of low-SEI discharge stages as a possible way to pursue.

As a pure energy storage technique, the efficiency appears to be still too low as compared, for example, with storage in  $H_2$  by electrolysis, that reaches efficiencies of the order of 80-90%.

The advantages of plasma reforming of  $CH_4 + CO_2$  must then be looked for in a multi-target application concept, in which value-added chemicals conversion, energy storage and  $CO_2$  disposal are simultaneously achieved. Biogas treatment appears to be an ideal application of plasma reforming.

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