Advanced Plasma Pyrolysis Assembly (PPA) Reactor and Process Development

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Design and development of a second generation Plasma Pyrolysis Assembly (PPA) reactor is currently underway as part of NASA’s Atmosphere Revitalization Resource Recovery effort. By recovering up to 75% of the hydrogen currently lost as methane in the Sabatier reactor effluent, the PPA helps to minimize life support resupply costs for extended duration missions. To date, second generation PPA development has demonstrated significant technology advancements over the first generation device by doubling the methane processing rate while, at the same time, more than halving the required power. One development area of particular interest to NASA system engineers is fouling of the PPA reactor with carbonaceous products. As a mitigation plan, NASA MSFC has explored the feasibility of using an oxidative plasma based upon metabolic CO₂ to regenerate the reactor window and gas inlet ports. The results and implications of this testing are addressed along with the advanced PPA reactor development.

**Nomenclature**

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Definition</th>
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<tbody>
<tr>
<td>1stGen</td>
<td>first generation</td>
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<tr>
<td>AR</td>
<td>atmosphere revitalization</td>
</tr>
<tr>
<td>C</td>
<td>carbon</td>
</tr>
<tr>
<td>CH₄</td>
<td>methane</td>
</tr>
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<td>CM</td>
<td>crew member</td>
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<tr>
<td>CRA</td>
<td>Carbon Dioxide Reduction Assembly</td>
</tr>
<tr>
<td>H₂</td>
<td>hydrogen</td>
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<td>H₂O</td>
<td>water</td>
</tr>
<tr>
<td>MSFC</td>
<td>Marshall Space Flight Center</td>
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**O₂** = oxygen  
**OGA** = Oxygen Generation Assembly  
**PPA** = Plasma Pyrolysis Assembly  
**SBIR** = Small Business Innovative Research  
**sccm** = standard cubic centimeters per minute  
**SDU** = Sabatier Development Unit  
**SOW** = statement-of-work

### I. Introduction

Effective methods for recovery and regeneration of valuable resources are needed to facilitate extended duration manned missions such as a return to the lunar surface or an expedition to Mars. Current technology for recovery of oxygen (O₂) from carbon dioxide (CO₂) is embodied in the Carbon Dioxide Reduction Assembly (CRA).\(^1\)\(^\text{17}\) The CRA employs a fixed-bed ruthenium-alumina catalytic reactor and molecular hydrogen (H₂) to reduce CO₂ to methane (CH₄) and water (H₂O) via the Sabatier reaction as shown in Eq. (1), where O₂ is ultimately recovered via water electrolysis in the Oxygen Generation Assembly (OGA).

\[
\text{CO}_2 + 4\text{H}_2 \rightarrow 2\text{H}_2\text{O} + \text{CH}_4 \tag{1}
\]

In the Sabatier reaction, half of the H₂ is lost as CH₄. Without additional processing, this CH₄ becomes a waste product because no equivalent amount of oxidant is produced, hence CH₄ cannot be used as a fuel or propellant unless additional oxygen is provided. Under NASA Phase I and Phase II Small Business Innovative Research (SBIR) contracts, development efforts using microwave plasma pyrolysis techniques were pursued.\(^\text{18-21}\) This approach allows for the recovery of up 75% of the H₂ currently lost as CH₄ per the acetylene formation reaction shown in Eq. 2.

\[
\text{2CH}_4 \rightarrow \text{C}_2\text{H}_2 + 3\text{H}_2 \tag{2}
\]

This technology was embodied in a first generation Plasma Pyrolysis Assembly (PPA), shown in Figure 1, and subsequently delivered to NASA-Marshall Space Flight Center (MSFC) where it has undergone extensive independent testing as both a stand-alone device and integrated with the Sabatier Development Unit (SDU).\(^\text{22-24}\) The 1st generation (1stGen) PPA, while proving adequate to demonstrate the potential of plasma pyrolysis techniques, falls short of the processing throughput and process efficiencies needed for effective loop closure in application as a H₂ recovery, post-processor for the Sabatier reactor. Recent advanced PPA development, however, has led to a second generation evolution of the PPA technology which addresses these shortcomings.

Specific advances in second generation PPA technology include: attaining a full one crew-member (CM) sized CH₄ processing capacity (compared to less than ½-CM scale in the first generation device) while maintaining 90% CH₄ conversion in a single pass configuration; incorporation of a more robust quartz window; addition of a redundant double seal design to promote safe operation; reduction in required amount of microwave power required to drive the process as well as reduction in the amount of microwave power wasted in reflections from the reactor; and a corresponding improvement in the specific energy efficiency required for chemical conversion. All this was achieved while maintaining low amounts of carbon production with improved selectivity for acetylene formation. Finally, performance for each reactor design was measured over several hours of operation with successive long duration tests and the relative advantages of each reactor design were identified.

It is believed that the greatest possible risk to long-term efficiency of the PPA is carbonaceous material formation on the quartz window or in the inlet/outlet gas ports. Carbonaceous build-up on the quartz window will eventually inhibit microwave transmission into the reactor, thereby reducing reactor performance. In addition, partial blockage of the gas ports will cause changes to flow dynamics inside the reactor, causing changes to the flow

![Figure 1. First Generation Plasma Pyrolysis Assembly.](image-url)
paths and corresponding reactor conversion efficiency. For this reason, MSFC performed initial PPA reactor regeneration testing, concurrent with advanced PPA reactor development work, to determine the feasibility of using CO₂, a gas already readily available in the Atmosphere Revitalization (AR) loop, to remove accumulated carbon and carbonaceous build-up via the Reverse-Boudouard reaction:

\[ \text{CO}_2 + \text{C} \rightarrow 2\text{CO} \tag{3} \]

Classically, this reaction only occurs at elevated temperature (>700 °C). The method developed here, however, utilizes the microwave chamber to create a CO₂ plasma in situ (in place of the H₂/CH₄ plasma present during methane plasma pyrolysis). Highly reactive species produced in the CO₂ plasma are then available to oxidize carbon compounds as they impinge upon coated surfaces via both forced and natural convection along with slower molecular diffusion. PPA Regeneration testing at MSFC included testing to prove the feasibility of igniting a CO₂ plasma, the feasibility of the CO₂ plasma resulting in the Reverse-Boudouard reaction, and the effect of reactor operating parameters on the relative level of regeneration.

II. Advanced PPA Reactor Design

The design effort began by reviewing the 1stGen reactor development work and performance results. Specific performance improvement goals were outlined in the Statement Of Work (SOW) for the Advanced PPA reactor. Targeted improvements included greater quartz window stability, more robust vacuum seal integrity, reduced carbon formation, increased energy efficiency, higher methane conversion, and improved acetylene selectivity all at a full 1-CM scale (400 sccm) CH₄ processing rate. Conceptual designs for two 1-CM scale advanced plasma reaction chambers were developed to achieve these objectives. Detailed designs were prepared for a Low-Risk reactor, which built upon the core design of the first generation device, and a Higher-Risk reactor, which employed novel features offering potentially much greater performance. The two reactors were subsequently fabricated using CNC milling techniques.

By critically considering both the positive and negative aspects of the various design iterations evaluated during development of the 1stGen PPA, key design aspects were identified. Along with the internal dimensions of the reactor, the presence and location of a plasma locating stub were determined to arguably be the most important design elements that impact process throughput, methane conversion efficiency, and energy efficiency. Factors impacting the nature and quantity of carbonaceous compounds formed in the reactor during testing were also identified. The H₂ to CH₄ feed ratio entering the PPA has the most significant impact on the amount and nature of condensed carbon compounds formed in the reactor over time. High H₂ to CH₄ ratios yield the least solids buildup, but the carbonaceous coatings that do form are of a brittle nature and difficult to remove from the reactor walls by mechanical action. Conversely, low H₂ to CH₄ ratios produce more solids, but this material is light and powdery in nature, making it relatively manageable via manual cleaning. Long duration testing at MSFC (>45 hours) using a nominal 4:1, H₂:CH₄ molar feed ratio demonstrated low solids production for the 1stGen PPA.

A. Design Goals

Specific process performance improvement targets were identified in the Phase 3 SOW for the Advanced PPA reactor technology. These performance goals are presented in Table 1 and compared to the demonstrated performance of the 1stGen PPA.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>1st Generation Reactor Performance</th>
<th>Advanced Reactor Targets</th>
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<tbody>
<tr>
<td>Microwave Power</td>
<td>700 W</td>
<td>≤ 350 W</td>
</tr>
<tr>
<td>Energy Efficiency @ 400sccm</td>
<td>6.20%</td>
<td>≥ 25%</td>
</tr>
<tr>
<td>% CH₄ Conversion @ 400sccm</td>
<td>80%</td>
<td>≥ 90%</td>
</tr>
<tr>
<td>CH₄ Throughput @ 90% CH₄ Conversion</td>
<td>160 sccm</td>
<td>≥ 400 sccm (1CM)</td>
</tr>
<tr>
<td>Acetylene Selectivity (% of carbon from converted CH₄) @ 400sccm</td>
<td>62%</td>
<td>≥ 75%</td>
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In addition to modifications that yield greater reactor performance, design alterations that make an advanced reactor safer and more robust were also sought out. Periodic failure of the quartz window (cracking along centerline through the narrow dimension) proved to be the single most important factor limiting test duration. This phenomenon is most likely related to the thermal stresses created within the quartz by the close proximity of the hot reactive plasma volume, which heats the center of the quartz window via both conductive and radiative heat transfer, and the relatively cold contact boundary between the window and the actively cooled metal walls. As such, thicker (∆z), more robust quartz windows that better withstand the temperature and pressure extremes experienced in a PPA reactor, were utilized in all designs considered during this advance development effort. In addition, these windows were also made larger in their x and y dimensions so as to support a double seal configuration for multilayer leak protection. While a rare event, the 1stGen PPA reactor with a single o-ring design did experience a minor air leak during at least one test at MSFC. By adding a second concentric seal, inherent system safety is significantly improved.

B. Low-Risk Reactor

The concept for the Low-Risk reactor consisted of key improvements to the 1stGen PPA reactor. This reactor would not only facilitate evaluation and verification testing of next generation design improvement concepts such as number and location of methane feed jets, removal of the window sweep plate, and reduction or elimination of the hydrogen feed stub/plasma locator nozzle, but also potentially serve as the deliverable reactor if sufficient performance gains were made. This reactor is shown in Figure 2.

C. Higher-Risk Reactor

For the Higher-Risk reactor, more risky design concepts were envisioned, which also offered potentially greater reactor performance gains. Firstly, the quartz window was moved farther from the hot plasma, thereby protecting it from thermal degradation. Secondly, a smaller reactor cross-section was utilized which served to concentrate microwave power density and the corresponding plasma density thus requiring less power to attain pyrolysis reaction conditions. This reactor is shown in Figure 3.

Figure 2. Low-risk advanced PPA reactor design.

Figure 3. Higher-Risk advanced PPA reactor design.
III. Advanced Reactor Performance Testing

Each of the advanced PPA reactors was challenged with a variety of performance tests. These tests were designed to evaluate each reactor’s process performance and help identify their unique operational characteristics. Performance testing for each advanced reactor culminated in back-to-back nominal 8-hour tests demonstrating feasibility for long-term operational stability. Testing with each reactor is discussed below.

A. Low-Risk Reactor

Photos of the advanced PPA Low-Risk reactor taken during plasma pyrolysis and between tests are shown in Figure 5. The Low-Risk reactor integrated into the PPA test stand is shown in Figure 4.

Using an 11/32 inch stub located on the reactor outlet port, two successive, long-duration tests were performed with a 26% CH\textsubscript{4} feed concentration at microwave power and process pressure levels of 315 W and 50 Torr, respectively. The result of the second back-to-back run is shown in Figure 6. Methane conversions over the course of these long duration tests were just over the targeted 90% level with a H\textsubscript{2} recovery just under 70%. This level of H\textsubscript{2} recovery corresponds directly to stoichiometric acetylene production as per Eq. (2) above, since 67.5% recovery is expected at 90% conversion. In addition, note that microwave power absorbed by the plasma remained above 98% for the duration of these tests. This high degree of microwave power to plasma heating conversion efficiency is the direct result of the well-tuned single mode cavity design. Microwave energy to chemical conversion energy efficiency was 17% for each test.

Figure 5. Low-Risk reactor. Reactor plasma during methane pyrolysis (left photo) and with the chamber end-plate removed prior to test (right photo).

Figure 4. Advanced PPA Low-Risk reactor and test stand. Reactor integrated into test stand and ready for pyrolysis testing.

Figure 6. Second Advanced PPA Low-Risk Reactor long-duration performance test. Test conducted at 400 sccm, 26 mol% methane, 50 Torr, and 315W for 7 hrs.
B. Higher-Risk Reactor

The Higher-Risk advanced PPA reactor is shown in Figure 7 while integrated into the PPA test stand. Forced air cooling of the reactor was found to be inadequate so copper cooling plates were attached to the outside of the reactor. The perforated plate viewport permits observation of the 3-dimensional behavior of the plasma within the reactor as seen in Figure 8. This viewing capability was found to be critical for developmental testing, allowing rapid process trouble-shooting and immediate feedback on the effects of changing process variables such as power, pressure and flow rate.

Two successive, long-duration tests were performed using the Higher-Risk reactor operating at a 26% methane feed at 400 sccm, a microwave power of 320 W, and a process pressure of 70 Torr. The result of the second of the back-to-back runs is shown in Figure 9. Methane conversions for both runs were over 90% with energy conversion efficiency at 17%. Hydrogen recovery for the Higher-Risk reactor was typically just under 70% for the majority of each test which was similar to that observed for the Low-Risk reactor (compare plot in Figure 9 to that in Figure 6). Absorbed microwave power was somewhat less than the 98% observed for the Low-Risk reactor, but nevertheless remained over 95%, which is still indicative of a well-tuned reactor. Both the Low-Risk reactor and the Higher-Risk reactor appear to have very similar performances with the obvious operational difference between the two being the required operating pressure (50 Torr for the Low-Risk reactor and 70 Torr for the Higher-Risk reactor).
C. Performance Summary

Second generation PPA reactor development has thus far demonstrated significant technology advancements over the 1stGen reactor as summarized in Table 2. Both advanced reactors have more than doubled the methane processing rate to a full 400 sccm, 1-CM scale while, at the same time, more than halving the required microwave power, going from 700 W to ≤320 W. Importantly, these performance gains have been attained while also improving the single-pass CH₄ conversion efficiency from 80% to over 90%. Despite near optimal utilization (>98%) of the microwave power by the plasma to create excited plasma species, the net microwave energy to chemical energy efficiency for conversion of CH₄ to acetylene and H₂ nevertheless remained around 17% during performance testing with either of the advanced reactors. While significantly better than the 6% demonstrated by the 1stGen PPA reactor, 17% remains somewhat below the 25% level targeted in our advanced PPA development work and well below the best value of 62% reported in the scientific literature.³⁵ While somewhat disappointing, this shortfall may well be an inherit design limitation that is related to the geometric aspects of these advanced PPA reactor designs (i.e., 17% may be the performance wall for these types of designs at 50 Torr). Alternatively, since energy efficiency has been observed to be dependent on operating pressure and CH₄ flow (reaching as high as 22% during Low-Risk reactor performance testing at 77 Torr, 800 sccm and 25% CH₄), the 25% first year goal or even the 62% best value goal may be attainable under the right pressure, total flow, and CH₄ conversion conditions. The final performance parameter listed in Table 2, acetylene selectivity, was improved from 62% for the first generation PPA well beyond the targeted 75% to ≥86% for the advanced PPA reactors.

Table 2. Performance improvements for advanced PPA reactors.

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<tr>
<td>Energy Efficiency @ 400sccm</td>
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<td>% CH₄ Conversion @ 400sccm</td>
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<td>91%</td>
<td>92%</td>
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<tr>
<td>CH₄ Throughput @ 90% CH₄ Conversion</td>
<td>160 sccm</td>
<td>≥400 sccm (1CM)</td>
<td>400 sccm</td>
<td>400 sccm</td>
</tr>
<tr>
<td>Acetylene Selectivity (% of carbon from converted CH₄) @ 400sccm</td>
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<td>≥75%</td>
<td>&gt;86%</td>
<td>&gt;87%</td>
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IV. Advanced PPA Process Development

Long-term operation of a PPA in flight application is expected to be limited by the effects of carbonaceous build-up, as mentioned previously. To mitigate this concern, PPA Regeneration testing was completed on the PPA Test Stand at MSFC’s Environmental Control and Life Support Systems Development Facility. For this effort, the 1stGen reactor was used as integrated into the PPA at MSFC, as shown in Figure 1. The PPA was developed at UMPQUA Research Co. and delivered to MSFC in 2009 as part of a Phase II SBIR contract. Regeneration testing of the 1stGen reactor involved three testing phases at MSFC and an implementation phase at UMPQUA as described below.

Phase I was completed to determine the parameters necessary for igniting a CO₂ plasma in the PPA. A full factorial parametric test was completed with CO₂ feed rates of 50, 200, and 500 sccm and reactor pressures of 15, 30, or 45 Torr. This test was replicated three times.

Phase II testing was completed to determine the feasibility of removing accumulated PPA carbon with a CO₂ plasma, thereby forming CO. Testing was completed in a series of two-day trials. The first day of each trial was dedicated to accumulating carbon for 6 hours with 200 sccm of CH₄ and a total of 800 sccm of H₂ continuously fed to the reactor. The system was operated at total reactor pressure of 45 Torr and a microwave power setting of 600 W. The second day of each trial was dedicated to regeneration for 15 minutes, 30 minutes, or 2 hours at 100 sccm CO₂ flow rate, a reactor pressure of 45 Torr and 600 W microwave power. At the end of each regeneration cycle, the reactor was photographed before the remaining carbonaceous product was mechanically removed from the reactor surfaces and weighed. During mechanical removal of residual carbon, the quartz window was left untouched to protect the surface.
Phase III testing was completed to determine the effects of pressure, microwave power, and CO$_2$ flow rate on PPA regeneration. For each trial, carbon was accumulated for 6 hours as described in Phase II testing. Each regeneration trial was conducted for 20 minutes from the moment of CO$_2$ plasma ignition. Tested reactor pressures included 20, 35, and 50 Torr. Tested CO$_2$ flow rates included 100, 200, and 500 sccm. Tested microwave powers included 400, 500, and 600W. A single replicate was completed for each data point.

Test results from MSFC’s development work of the CO$_2$ plasma cleaning step were applied to cleaning the advanced PPA reactors between selected discrete performance evaluation tests. In particular, for the Higher-Risk reactor, a regularly scheduled CO$_2$ plasma cleaning step was used after each nominal 8 hours of methane pyrolysis.

A. Phase I PPA Regeneration Test Results
Using the 1stGen PPA reactor, a CO$_2$ plasma was reliably ignited at a reactor pressure of 15 Torr regardless of the CO$_2$ flow rate, as shown in Figure 10. However, the power required to ignite the plasma varied dramatically from run to run. There was no significant difference in power required for ignition with respect to CO$_2$ flow rate. At 30 Torr, ignition of the CO$_2$ plasma was shown to be possible; however, ignition was unreliable and often required multiple attempts. At the 50 sccm and 200 sccm CO$_2$ flow rates, the plasma was only ignited in two of the three replicates. At the 500 sccm CO$_2$ flow rate, the plasma was only ignited in one of the three replicates. A CO$_2$ plasma could not be ignited at 45 Torr.

Based on these findings, it is clear that the reactor pressure will need to be below 30 Torr to reliably and repeatedly ignite a CO$_2$ plasma. For long duration missions, PPA regeneration capability may be necessary. However, constraining the design of the system vacuum pump to accommodate these low pressures may not be as advantageous as simply replacing the quartz window or entire reactor chamber. Future trade analyses are planned to explore these options.

B. Phase II PPA Regeneration Test Results
Following a six hour carbon accumulation, the first generation PPA reactor was regenerated for 15 minutes, 30 minutes, or 2 hours. As shown in Figure 11, the majority of the carbonaceous product was removed in the first 15 minutes of the regeneration cycle. Due to these findings, a second test was run in which the reactor accumulated carbon for 18 hours, followed by a 15 minute CO$_2$ regeneration cycle. Photos a, b, c, and d in Figure 12 show the carbon accumulation on the quartz window over 6, 12, and 18 hours (photos a, b, and c, respectively) and the effect of the relatively short regeneration cycle (photo d). This data suggest that as long as the inlet/outlet ports are not blocked, very short regeneration cycles will be sufficient to maintain a clean quartz window. It should also be noted that no decrease in performance was observed.

Figure 10. Average Input Power vs. Pressure for CO$_2$ plasma ignition.

Figure 11. PPA Carbon Removal via CO$_2$ Plasma.
after 18 hours of carbon accumulation suggesting that regeneration cycles may be performed at much longer intervals than originally believed. Another important consideration for advanced PPA reactor development is that placement of the plasma inside the reactor may well have a significant influence on the quartz window regeneration. As such, redesign of the reactor chamber may alter the location of this plasma, thus changing the localized effect of CO$_2$ regeneration on the quartz window.

C. Phase III PPA Regeneration Test Results

Phase III testing was completed to investigate the influence of reactor pressure, microwave power, and CO$_2$ flow rate on the relative reactor regeneration performance. Although only one replicate was completed for this testing, images of the inside of the reactor, particularly of the plasma stub, strongly suggest that pressure is the factor most influencing regeneration performance. This can be seen in the examples in Figure 14 at 20 and 50 Torr, each operating at 400 W and 100 sccm CO$_2$ flow. Over the ranges tested, microwave power appeared also to have an effect on regeneration performance, though less influential than pressure. This is shown in the examples in Figure 13 at 400 and 600 W, each operating at 50 Torr and 500 sccm CO$_2$ flow. CO$_2$ flow rate variations appeared to have no effect.
D. Advanced PPA Reactor Regeneration Implementation Results

Test results from MSFC’s development work of the CO$_2$ plasma cleaning step were applied to cleaning the advanced PPA reactors between selected discrete performance evaluation tests. In particular, for the Higher-Risk reactor, a regularly scheduled CO$_2$ plasma cleaning step was used after each nominal 8 hours of methane pyrolysis. Before, during (plasma), and after photos of the reactor are shown in Figure 15. Note the color difference between this CO$_2$ plasma and the H$_2$/CH$_4$ plasma seen in Figure 8; the CO$_2$ plasma appearing as a light blue. The before and after chamber photos located in Figure 16 and Figure 17, respectively, show that most of the carbon build-up around the outlet stub is removed during the CO$_2$ plasma cleaning step. Preferential cleaning of the upper half compared to the lower half is clearly evident. This is presumably due to gravity-driven natural convective flow of hot reactive gases produced during cleaning (in the presence of colder feed gas flow). As such, in the absence of gravity, more symmetrical cleaning about the outlet stub would be expected, except as moderated by gas flow dynamics within the reactor, which are related to both geometry and magnitude of flow. Clearly, while not removing all carbon buildup in the reactor, significant cleaning was observed. A thorough evaluation of this periodic cleaning technique would require several weeks or months’ worth of PPA testing where the impact of any long term persistent (un-removed) carbon deposition on reactor performance would be determined.

V. Conclusion

In summary, second generation PPA development has demonstrated significant technology advancements over the 1stGen device by more than doubling the methane processing rate to 1-CM scale while, at the same time, requiring less than half the power. Advanced PPA process development work was performed by personnel at NASA MSFC concurrent with the advanced PPA reactor development effort. Here, an in-situ technique that utilized a CO$_2$ plasma was used to periodically remove carbonaceous material byproducts that naturally accumulate in the reactor over time as a result of non-equilibrium methane pyrolysis. This work represents interim results of an on-going Phase 3 development project. Second year work will focus on scaling to a 4-CM design while further improving PPA performance.

Acknowledgments

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