

Identifying Different Types of Catalysts for CO₂ Reduction by Ethane through Dry Reforming and Oxidative Dehydrogenation

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Abstract: The recent shale gas boom combined with the requirement to reduce atmospheric CO₂ have created an opportunity for using both raw materials (shale gas and CO₂) in a single process. Shale gas is primarily made up of methane, but ethane comprises about 10 % and reserves are underutilized. Two routes have been investigated by combining ethane decomposition with CO₂ reduction to produce products of higher value. The first reaction is ethane dry reforming which produces synthesis gas (CO + H₂). The second route is oxidative dehydrogenation which produces ethylene using CO₂ as a soft oxidant. The results of this study indicate that the Pt/CeO₂ catalyst shows promise for the production of synthesis gas, while Mo₂C-based materials preserve the C–C bond of ethane to produce ethylene. These findings are supported by density functional theory (DFT) calculations and X-ray absorption near-edge spectroscopy (XANES) characterization of the catalysts under in situ reaction conditions.

Recent exploitation of shale gas reserves has increased the importance of using methane and the heavier gas fractions as raw materials for producing chemicals and fuels. Methane is the primary component of shale gas, but ethane comprises approximately 10 %, with the exact concentration depending on the particular source.^[1] Most ethane is converted by thermal dehydrogenation into ethylene, a valuable commodity chemical, at high temperatures, between 750–950 °C, in the presence of steam,^[2] which is energy-intensive and leads to coke formation on the reactor walls and catalyst bed.^[3] Catalysts that can effectively convert ethane to useful products at lower temperatures, without coke formation, are very desirable. Ideally, a catalyst that can use CO₂ as the oxidant is particularly useful because the reduction of CO₂ can help mitigate ocean acidification and climate change,^[4] while unreacted CO₂ can also reduce coke formation through the reverse Boudouard reaction (CO₂ + C → 2CO), which has

been shown to occur as low as 500 °C,^[5] thus increasing conversion and catalyst stability.^[3]

CO₂ reduction by ethane can proceed through two distinct pathways: 1) reforming to produce synthesis gas (2CO₂ + C₂H₆ → 4CO + 3H₂) and 2) oxidative dehydrogenation (ODH) to produce ethylene (CO₂ + C₂H₆ → C₂H₄ + CO + H₂O). For decades, researchers have attempted dry reforming of methane and CO₂, but breakthroughs have been difficult because of the high reaction temperatures (ca. 700 °C) and deactivation of catalysts.^[6] Dry reforming of ethane, however, becomes thermodynamically favorable at a lower temperature than for methane (ca. 600 °C), making the process more feasible under milder conditions.^[7] The difference in reaction temperatures between ethane and methane dry reforming also provides a potential opportunity for reactively separating shale gas, thereby directly producing synthesis gas from ethane and CO₂, while leaving the methane unreacted.

ODH of ethane by CO₂ is another important pathway because of the high energy requirement for the current ethane dehydrogenation processes.^[8] Several groups have studied ODH of ethane with pure O₂,^[9] which is more reactive but does not have the added benefit of reducing CO₂ emissions. ODH of ethane with CO₂ as a mild oxidant is proposed to occur through a Mars-van-Krevelen-type mechanism,^[10] where the catalyst first dehydrogenates ethane into ethylene and H₂, then H₂ reduces CO₂ to CO through the reverse water-gas shift (RWGS) reaction, producing H₂O as a byproduct.^[11] At 600 °C, both the reforming and ODH reactions occur simultaneously, thus it is necessary to identify classes of catalysts to kinetically control the extent of reforming versus ODH. It is highly desirable to identify distinct types of catalysts that can 1) efficiently break the C–C bond to produce synthesis gas, or 2) selectively break C–H bonds of ethane while preserving the C–C bond to produce ethylene.

Previous research indicates that typical catalysts used for CO₂ reduction by ethane are Cr-based oxide catalysts. Cr₂O₃ catalysts convert high amounts of ethane, but overall CO₂ conversion is low with short lifetimes, likely because of coke formation.^[12] There is a support effect, with SiO₂ displaying the highest activity,^[13] and doping the catalyst with small amounts of Fe, Co, and Mn further improves ethylene selectivity.^[14] The high activity of Cr-based catalysts could stem from their ability to exist in several oxidation states, similar to Mn-based molecular sieve catalysts.^[12] Ga₂O₃ and Ca-doped ThO₂ are reducible materials that are also active and selective for this reaction,^[15] while a nanostructured CeO₂ catalyst doped with CaO oxidizes ethane into CO and H₂O, with ODH as the side reaction. For carbide catalysts, Mo₂C/SiO₂ has about 10 % conversion with high ethylene selectiv-

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ity; however, pure Mo₂C and MoO₃/SiO₂ are not reported as active.^[16]

Although there have been studies into CO₂ reduction by ethane, details are lacking on how to control the extent of dry reforming versus ODH. In the current study, two types of catalysts, metals (Pt-based) and metal carbides (Mo₂C), are investigated under identical reaction conditions to differentiate the two reaction pathways. It is hypothesized that Pt and Mo₂C-based systems are promising catalysts to study reforming and ODH during CO₂ reduction by ethane because Pt catalysts supported on CeO₂^[17] and Mo₂C^[18] are both active for RWGS from their high oxygen mobility, a critical feature required for CO₂ reduction by ethane. The findings of this study confirm that although there is deactivation, Pt/CeO₂ favors the reforming reaction and Mo₂C-based catalysts are more selective at producing the ODH product, ethylene. These novel findings are supported by DFT calculations and in situ X-ray absorption near-edge spectroscopy (XANES) measurements.

To help identify reaction pathways over the two types of catalysts, density functional theory (DFT) calculations were performed to compare differences in binding energies of potential reaction intermediates. The results in Table 1 show that all intermediates bind more strongly on Mo₂C(001) than on Pt(111). It is noted that, for adsorbates with similar binding configurations, the binding energy difference between Pt(111) and Mo₂C(001) is more pronounced for species adsorbed by the oxygen atom than those using the carbon atom. This is due to the much higher affinity of the Mo₂C(001) surface toward O relative to Pt(111). The DFT calculated BE_{diff} is 3.53 eV for atomic oxygen, while it is only 1.60 eV for atomic carbon.

Table 1: DFT-calculated binding energies (in eV) of ethane dehydrogenation intermediates and atomic O and C on Pt(111), Ce₃O₆/Pt(111), and Mo₂C(001) surfaces.

Species	Pt(111)	Ce ₃ O ₆ /Pt(111)	Mo ₂ C(001)	$BE_{\text{diff}} = BE_{\text{Pt(111)}} - BE_{\text{Mo}_2\text{C(001)}}$
CH ₃ CH ₂ O	-1.62	-2.42	-4.22	2.60
CH ₃ CHO	-0.31	-0.95	-2.30	1.99
CH ₃ CO	-2.36	-2.99	-3.81	1.45
CH ₃ CH ₂	-1.97	-1.97	-2.98	1.01
CH ₂ CH ₂	-1.09	-1.24	-2.23	1.14
CH ₂ CH	-3.18	-3.42	-4.64	1.46
CH ₃	-2.11	-2.26	-3.14	1.03
CO	-1.76	-2.19	-2.72	0.96
H	-2.75	-2.80	-3.37	0.62
O	-4.11	-4.33	-7.64	3.53
C	-7.17	-5.55	-8.77	1.60

DFT was also used to calculate the enthalpy change for the oxidative C–C and C–H bond cleavage of ethane on Pt(111) and Mo₂C(001) surfaces. The C–C bond cleavage of ethane produces CH₃*, CO*, and H₂O(g), while C–H bond scission results in the formation of CH₂CH₂* and H₂(g). For the case of Pt(111) in Figure 1a, atomic hydrogen is assumed to form H₂O(g) since both OH (O* + H* → OH*) and H₂O formation (H* + OH* → H₂O*) are exothermic processes on the Pt(111) surface.^[19] This is not the case, however, for

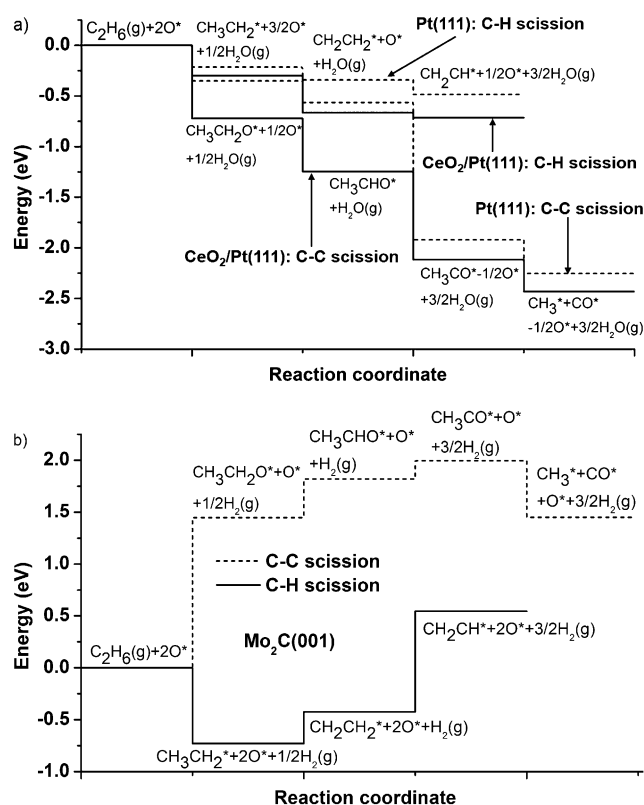


Figure 1. DFT-calculated energy profile of reforming and oxidative dehydrogenation of ethane on a) Pt(111) and CeO₂/Pt(111) and b) the Mo₂C(001) surface.

Mo₂C(001) in Figure 1b, where both OH (O* + H* → OH*) and H₂O formation (H* + OH* → H₂O*) are highly endothermic processes.^[20] Over Mo₂C(001), the atomic hydrogen prefers to recombine and desorb from the surface in the form of hydrogen gas.

Furthermore, Figure 1a shows that on Pt(111), the C–C bond cleavage of ethane to form CH₃*, CO*, and H₂O(g) is not only exothermic, but also more favorable compared to the selective C–H bond cleavage to form CH₂CH₂*. More specifically, oxidation of ethane to form CH₃CH₂O* is energetically more favorable than the dehydrogenation of ethane to form CH₃CH₂* in the presence of oxygen on the surface. On Pt(111), CH₃CH₂O* undergoes two successive dehydrogenation reactions to form CH₃CO*, which then undergoes C–C bond cleavage to form CH₃* and CO*. Thus, the Pt(111) surface promotes reforming of ethane to form CO.

A different reaction scheme from that on Pt(111) is observed on Mo₂C(001). The stronger affinity of Mo₂C(001) for oxygen makes C–O bond formation less favorable on Mo₂C(001) than on Pt(111), which agrees well with previous studies showing that Mo₂C(001) binds carbon and oxygen very strongly, leading to spontaneous cleavage of the C–O bond.^[21] Therefore, as shown in Figure 1b, the Mo₂C(001) surface preferentially dehydrogenates ethane to form CH₂CH₂*. Further discussion of the possible origins of the selective adsorption of different intermediate species over Pt(111) and Mo₂C(001) is provided in the Supporting Information.

Further DFT calculations were performed on $\text{CeO}_2/\text{Pt}(111)$ to investigate the role of the Pt– CeO_2 interface in the energy profile of reforming and oxidative dehydrogenation of ethane. The O-containing species bind at the Pt– CeO_2 interface and most of the hydrocarbon species bind at the Pt sites, with detailed figures available in the Supporting Information. More importantly, the calculated energy profile for the reforming of ethane on $\text{CeO}_2/\text{Pt}(111)$ and $\text{Pt}(111)$ in Figure 1 shows a similar trend; however, the reaction is more thermodynamically favorable on $\text{CeO}_2/\text{Pt}(111)$, which is associated with the strong CeO_2 –Pt(111) interaction^[22] and therefore strengthened binding of O-containing species at the interface, as shown in Table 1. Thus, the DFT results suggest that similar to Pt(111), $\text{CeO}_2/\text{Pt}(111)$ also enhances the reforming of ethane.

To verify the DFT predictions, reactions of ethane and CO_2 were evaluated in a quartz reactor under atmospheric pressure. The activity and selectivity as a function of time on stream for the catalysts, Pt/CeO₂ and Mo₂C/γ-Al₂O₃, are compared in Figure 2. Furthermore, alloying Co with the Pt/CeO₂ catalyst improves the catalytic stability. The comparison between the two catalysts clearly reveals differences in performance. Pt/CeO₂ shows higher activity and selectivity toward reforming products, but Mo₂C/γ-Al₂O₃ has a lower activity with increased ethylene selectivity, confirming the trend seen in the DFT calculations.

The difference in activity between the two classes of catalysts can be further explained through measuring the apparent activation barrier of both CO_2 and ethane over Pt/CeO₂ and Mo₂C/γ-Al₂O₃, as shown in Figure 3. Over Pt/CeO₂ the activation barriers are 16.0 kcal mol^{−1} and 21.6 kcal mol^{−1} for CO_2 and ethane, respectively, and on Mo₂C/γ-Al₂O₃ the values are 19.5 kcal mol^{−1} and 17.5 kcal mol^{−1}. The lower activation barrier of CO_2 over Pt/CeO₂ confirms the relatively high activity of CeO₂-based materials toward CO_2 .^[23]

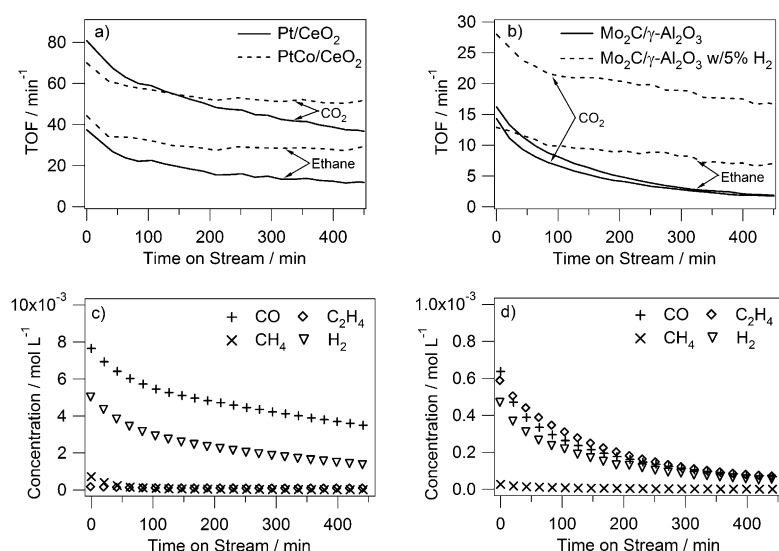


Figure 2. Turnover frequency (left column) and concentration of products (right column) for reactions of 10 mL min^{−1} ethane and CO_2 10 mL min^{−1} diluted in 20 mL min^{−1} He at 600 °C for a,c) Pt/CeO₂ and b,d) 8% Mo₂C/γ-Al₂O₃ plotted versus time on stream. PtCo/CeO₂ and Mo₂C/γ-Al₂O₃ with 5% H₂ (dashed) are included in (a) and (c), respectively as references.

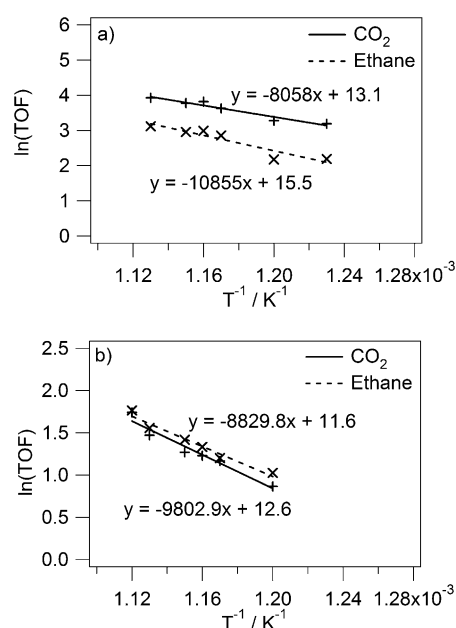


Figure 3. Activation barrier plots represented as ln(TOF) versus 1/T for a) Pt/CeO₂ and b) Mo₂C/γ-Al₂O₃ at six temperatures. Values of TOF calculated by averaging data points between 2–4.5 h on stream.

Because CeO₂ has been shown to be an excellent support for reactions involving CO_2 , an additional test was performed by supporting Mo₂C on CeO₂. As seen in Table 2, comparisons of conversion indicate there is a clear support effect for Mo₂C, where γ-Al₂O₃ slightly improves the activity over the pure carbide, but CeO₂ appears to diminish the activity. This support effect likely suggests that the greater acidity of γ-Al₂O₃ promotes the binding of ethane, which is consistent with previous work indicating that oxidative dehydrogenation of alkanes is strongly dependent on the acidity of support materials.^[24] However, future studies will be required to determine the effect of acidity on the reaction pathways of CO_2 and ethane over Mo₂C/γ-Al₂O₃.

Furthermore, the behavior of Mo₂C/γ-Al₂O₃ is more similar to that of pure Mo₂C than Mo₂C/CeO₂, which is likely from the low surface area and CO uptake of Mo₂C/CeO₂ (0.8 μmol g^{−1}). It is also possible that the high intrinsic activity of both CeO₂ and Mo₂C^[21b] towards CO_2 bond scission creates an excess of adsorbed O on the catalyst surface, which quickly deactivates Mo₂C/CeO₂ through oxidation.

Deactivation is not only limited to Mo₂C/CeO₂, but is clearly seen for both Pt/CeO₂ and Mo₂C/γ-Al₂O₃ in Figure 2. To better understand the mechanism of rapid deactivation over Mo₂C/γ-Al₂O₃, reaction order measurements were performed for ethane. Clearly, as shown in the Supporting Information, the reaction order is negative with respect to ethane, meaning increasing concentrations of ethane decreases the catalytic activity. One explanation, obtained through DFT calculations, indicates the higher

Table 2: Summary of flow reactor data for CO₂ reduction by ethane experiments in 10 mL min⁻¹ ethane and CO₂ diluted in 20 mL min⁻¹ He at 600 °C. Values of conversion and selectivity calculated by averaging data points between 2–4.5 h on stream.

Catalyst	Conversion [%]		Carbon Selectivity [%]			
	CO ₂	C ₂ H ₆	CO	CH ₄	C ₂ H ₄	C ₃ +
Mo ₂ C	1.0	2.0	38.6	1.6	59.5	0.3
Mo ₂ C/γ-Al ₂ O ₃	1.8	2.1	47.0	1.2	51.6	0.2
Mo ₂ C/CeO ₂	0.3	0.1	88.1	2.1	9.8	0.0
Mo ₂ C/γ-Al ₂ O ₃ w/H ₂	8.8	3.8	73.4	0.7	25.7	0.2
Pt/CeO ₂	28.0	9.5	97.8	0.8	1.4	0.0
PtCo/CeO ₂	47.2	25.7	98.7	0.6	0.6	0.0
Pt/γ-Al ₂ O ₃	12.4	1.7	95.2	0.7	4.1	0.0

binding energy of oxygen on Mo₂C destabilizes C–O bond formation and leads to ethylene as the primary product. The high selectivity towards ethylene, combined with the high binding energy on Mo₂C(001) ($BE_{\text{diff}} = 1.14$ eV relative to Pt(111)), likely leads to coke formation through ethylene decomposition or polymerization,^[25] partially explaining the deactivation and negative order.

To help mitigate the high deactivation over Mo₂C/γ-Al₂O₃, a small amount of H₂ (5 vol%) was added to the reaction feed to remove inactive oxide species to complete the catalytic cycle and restore some of the active carbide phase, leading to higher activity (Figure 2c). This phenomenon has been seen in previous work over Mo₂C-based catalysts for CO₂ reduction by H₂.^[18b] As shown in Table 2, the addition of H₂ improves the initial catalytic activity, but the CO selectivity increases while that of ethylene decreases. The increased selectivity toward CO is likely because the addition of H₂ promotes RWGS. Additionally, the Mo₂C/γ-Al₂O₃ catalyst can be regenerated through a post-reaction treatment cycle as shown in the Supporting Information.

To further understand the mechanism of deactivation over Mo₂C, in situ XANES measurements were performed over pure Mo₂C. Full details of the XANES experiments including spectra for pure Mo₂C in various reaction environments as well as the difference spectra (ΔXANES) can be found in Figure 4. The XANES measurements of Mo₂C show clear

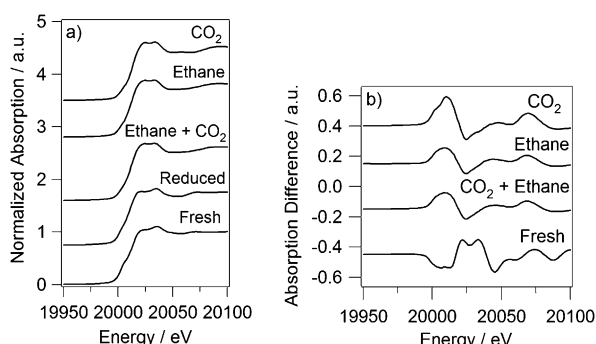


Figure 4: X-ray absorption near-edge structure measurements of Mo₂C in different gas environments, with a) raw spectra and b) ΔXANES subtracted from the “reduced sample”. All spectra measured at room temperature after treatment at 450 °C for reduction and 600 °C for CO₂ and ethane environments.

Table 3: Results from linear combination fitting of Mo₂C XANES spectra.

Condition	Mo ₂ C	MoO ₃	MoO ₂
fresh	64.1	35.9	0
reduced	75.7	6.7	17.6
C ₂ H ₆ + CO ₂	23.5	0	76.5
C ₂ H ₆	26.6	0	73.4
CO ₂	0	9.6	90.4

changes in the oxidation state of Mo during all gas treatments, except for pure ethane. To quantify these changes, each Mo spectrum was fitted by a linear combination of Mo₂C, MoO₃, and MoO₂ standards, as described previously.^[18a] The results of the linear combination fitting analysis are shown in Table 3.

By comparing the XANES results in Table 3, it is clear that treatment in H₂ removes a significant amount of the oxide species formed during passivation of the carbide during synthesis. Upon reaction of ethane and CO₂, Mo₂C is oxidized to a greater degree. Treatment in pure CO₂, however, completely oxidizes Mo₂C. Therefore, the adsorbed O from CO₂ is necessary to form an oxycarbide phase, which dehydrogenates ethane, but also likely contributes, at least partially, to the deactivation seen over Mo₂C-based catalysts.

In summary, Pt/CeO₂ and Mo₂C-based catalysts have been evaluated to catalytically control the products of CO₂ reduction by ethane to produce either synthesis gas or ethylene, respectively. Results of this study indicate that Pt/CeO₂ is a promising catalyst to produce synthesis gas, while Mo₂C/γ-Al₂O₃ preserves the C–C bond of ethane to produce ethylene, although the catalyst deactivates. These findings agree with DFT predictions that the Pt(111) surface preferentially forms synthesis gas, while Mo₂C(001) leads to ethylene formation. Future studies should focus on designing materials that selectively produce ethylene with high stability and minimal deactivation.

Experimental Section

Spin-polarized DFT^[26] calculations were performed using plane wave basis set Vienna Ab-Initio Simulation Package (VASP) code.^[27] A plane wave cut-off energy of 400 eV and 3 × 3 × 1 Monkhorst-Pack^[28] grid were used for total energy calculations. Details of the DFT calculations are provided in the Supporting Information.

The synthesis procedures of Pt-based and Mo₂C-based catalysts are provided in the Supporting Information. Reactions of ethane and CO₂ were carried out in a quartz reactor under atmospheric pressure. In each experiment, approximately 100 mg catalyst (60–80 mesh) was loaded into the flow reactor. Before the reaction, the catalyst was reduced under a 1:1 hydrogen and helium mixture (40 mL min⁻¹ total flow) at 450 °C for 1 h. To start the reaction, the flow of CO₂ and ethane were each set at 10 mL min⁻¹ with 20 mL min⁻¹ of He as a diluent. For each experiment, the temperature was raised to 600 °C and the reaction was run for approximately 6 h.

X-ray absorption near-edge spectroscopy (XANES) measurements were used to confirm the oxidation state of Mo₂C under reaction conditions. Catalyst samples were diluted with boron nitride, pressed into a pellet, then grinded and sieved (100–150 mesh). The mass of catalyst and dilution were determined so the sample amount was two absorption lengths to maximize the signal-to-noise ratio of each sample. The XAFS spectra for Mo₂C was collected in a custom designed in situ glassy carbon cell,^[32] using a 4-channel vortex fluorescence detector.

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