

## CORRECTION

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## Correction: Ethanolic gasoline, a lignocellulosic advanced biofuel†

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Correction for 'Ethanolic gasoline, a lignocellulosic advanced biofuel' by Mícheál Séamus Howard *et al.*, *Sustainable Energy Fuels*, 2019, 3, 409–421, <https://doi.org/10.1039/C8SE00378E>.

The ignition delay time (IDT) data reported in this article<sup>1</sup> were recently found to be erroneous. Two issues caused errors in the previous data, (i) a fraction of the volatile fuel, diethyl ether (DEE), was lost during mixture preparation, and (ii) a fraction of the high boiling point fuel, ethyl levulinate (EL), was lost due to condensation on the walls of the combustion chamber. These issues were realized after the publication of the original paper while working on another project. After fixing these problems, we obtained new IDT results which deviated significantly from the previous data at low temperatures. New IDT data of the EtOH/DEE/EL blend showed NTC behavior, in contrast to previous data which primarily exhibited Arrhenius-like behavior. The revised Fig. 2 below plots the updated data, and these data are provided in tabular form in the ESI.<sup>†</sup>

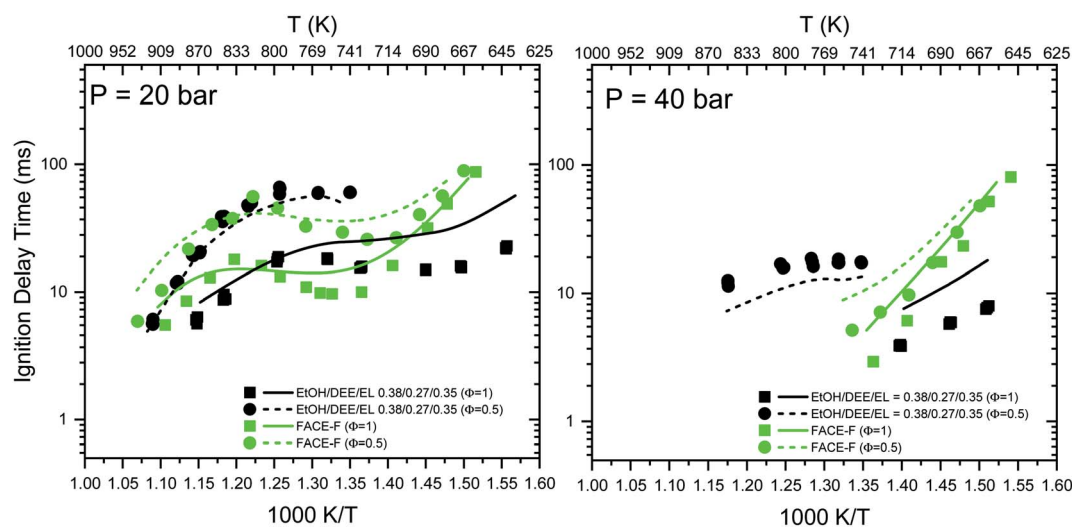


Fig. 2 RCM ignition delay times for the EL/DEE/EtOH gasoline and FACE-F.<sup>‡</sup> Symbols correspond to measurements. Lines correspond to present model predictions. FACE-F experimental data are taken from Sarathy *et al.*<sup>2</sup>

The chemical kinetic model reported in the original paper was developed and tuned to match the erroneous experimental data originally reported. The new experimental data showed that our original model needed further improvements. Therefore, we have now revised the model based on the newly acquired IDT data. The predictions of the updated model are shown in the updated Fig. 2. The following modifications are made to the previous model:

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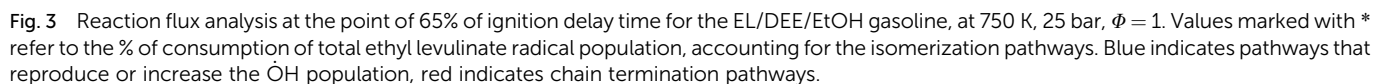
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† Electronic supplementary information (ESI) available. See <https://doi.org/10.1039/d2se90024f>

‡ Mícheál Séamus Howard and Stephen Dooley could not be contacted prior to publication of this Correction.

- Ignition delay times of conventional hydrocarbon-based gasoline (FACE-F) and oxygenated gasoline (EL/DEE/EtOH) exhibit negative temperature coefficient (NTC) behavior at intermediate temperatures (see the updated Fig. 2). However, FACE-F shows steeper NTC compared to the oxygenated gasoline. Both fuels have similar high-temperature reactivity, but the oxygenated gasoline shows lower reactivity (longer ignition delays) than FACE-F at low temperatures.



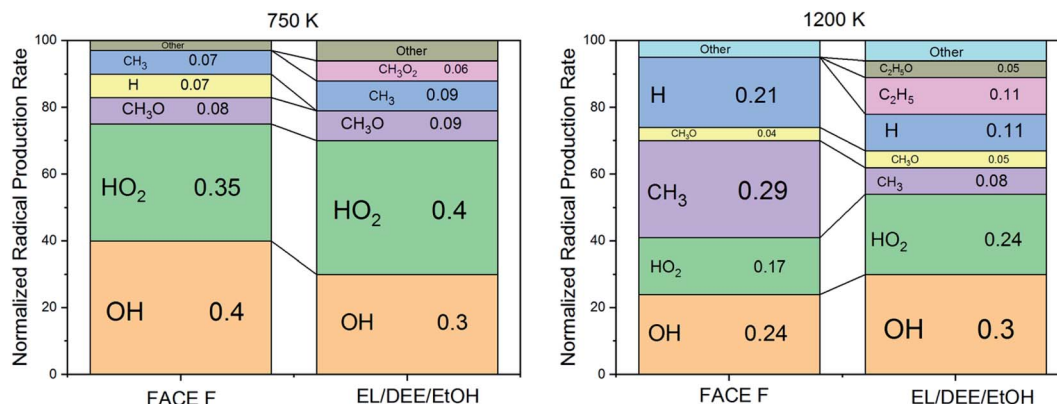


Fig. 4 Normalized small radical rate of production for EL/DEE/EtOH gasoline and FACE-F<sup>2</sup> after 65% of the computed ignition delay time at 750 K and 1200 K, 25 bar, and  $\Phi = 1$ .

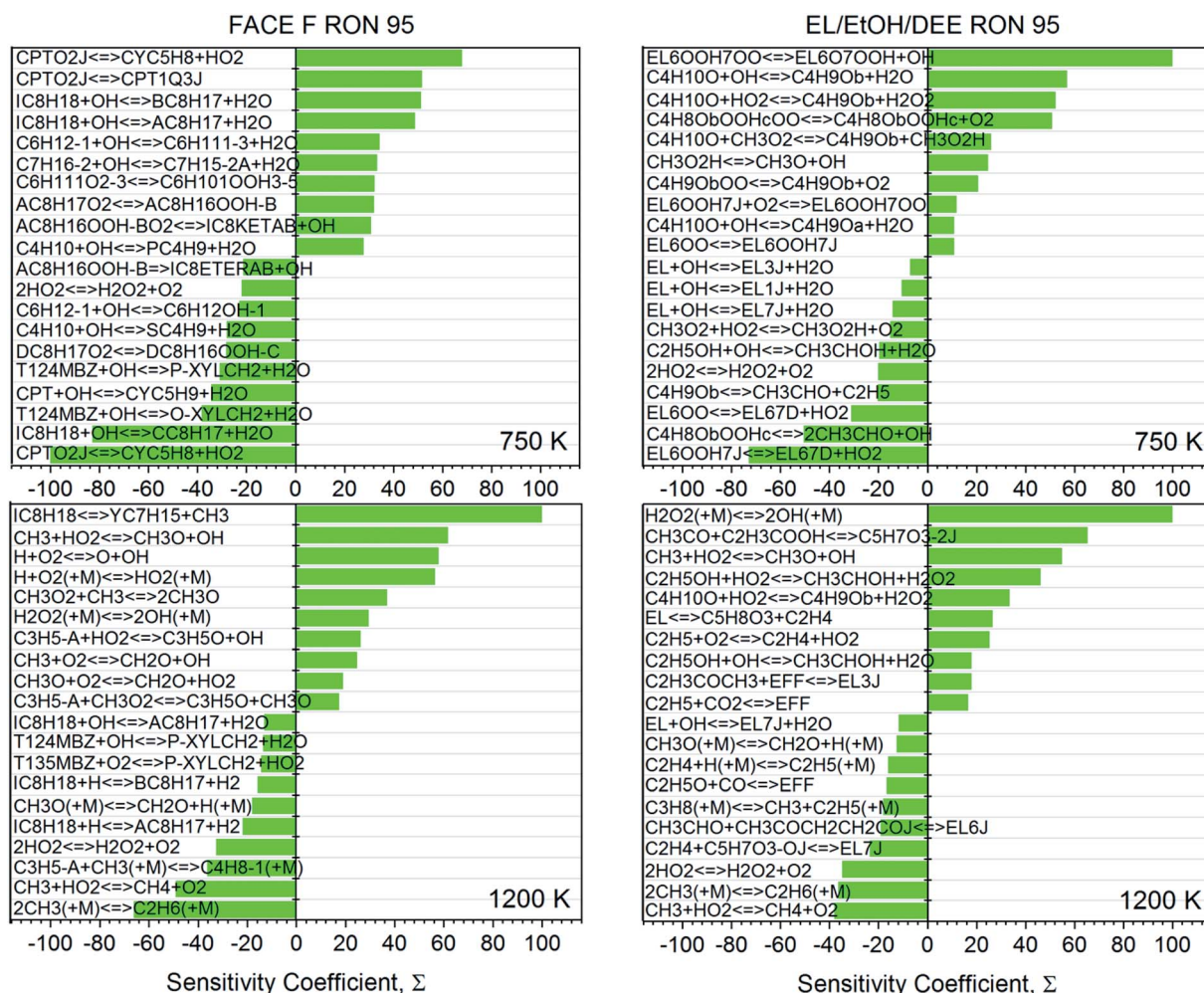


Fig. 5 Sensitivity analysis to  $\Delta T$  for EL/DEE/EtOH gasoline and FACE-F<sup>2</sup> at 750 K and 1200 K, 25 bar, and  $\Phi = 1$ . Top 20 sensitivity coefficients are shown, normalized to the absolute most sensitive reaction. Reactions are described in the direction of their net flux from left to right.

• The nominally similar reactivity behavior of the two gasolines is in-line with their similar octane numbers. However, the flatter ignition delay profile of the oxygenated gasoline indicates that it should possess a larger octane sensitivity ( $S = \text{RON} - \text{MON}$ ). This anomaly could result from the limitations of octane rating tests, as discussed in the original paper. Octane measurements of



oxygenated fuels are quite complicated and may suffer from larger uncertainties compared to the well-established octane measurements of conventional gasoline.

- Reaction flux analysis (Fig. 3) shows that the OH radical leads the consumption of the three fuel components, consuming 73%, 59% and 53% of EL, DEE and EtOH, respectively. On the other hand, HO<sub>2</sub> consumption of these fuels is 15%, 21%, and 37%, respectively. Therefore, modelling analysis (750 K, 25 bar) shows that EL is primarily consumed by OH radicals, while the HO<sub>2</sub> radical plays a larger role in the consumption of EtOH. For this tri-blend, a large pool of reactive radicals is formed from DEE, which then promotes the oxidation of EL.

- The radical rate of production analysis (Fig. 4) indicates that HO<sub>2</sub> formation is more significant for EL/DEE/EtOH compared to FACE-F at both low (750 K) and high (1200 K) temperatures. This augmented rate of production of HO<sub>2</sub> for EL/DEE/EtOH stems from the dominance of RO<sub>2</sub> and the alkylhydroperoxide radical elimination pathways.

- Sensitivity analyses (Fig. 5) show that OH consumption of the fuel components dictates the system reactivity at 750 K. In the case of FACE-F, for example, reactions of OH with iso-octane and *n*-heptane promote reactivity, while those with *n*-butane, cyclopentane, trimethylbenzene and the tertiary site of iso-octane inhibit reactivity. For EL/DEE/EtOH, the reaction of OH with DEE promotes reactivity, while those with EL and EtOH inhibit reactivity. At 1200 K, small radical chemistry dominates for both gasolines but an enhanced role of HO<sub>2</sub> reactions and H<sub>2</sub>O<sub>2</sub> decomposition is observed for EL/DEE/EtOH. Thus, the ignition modes of the two gasolines are somewhat different even though they have very similar reactivity at high temperatures.

- In conclusion, the petroleum gasoline (FACE-F) and ethanolic gasoline (EL/DEE/EtOH) with almost identical octane ratings have very similar high-temperature reactivity and both fuels show NTC behavior at intermediate temperatures. The ethanolic gasoline shows a flatter NTC profile and reduced reactivity at low temperatures, which are likely due to the increased role of HO<sub>2</sub>-related inhibition pathways.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

## References

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- 3 G. Issayev, S. Mani Sarathy and A. Farooq, Autoignition of diethyl ether and a diethyl ether/ethanol blend, *Fuel*, 2020, **279**, 118553.

