Garbage-In Garbage-Out (GIGO): The Use and Abuse of Combustion Modeling and Recent U.S. Spacelaunch Environmental Impacts

PattiMichelle Sheaffer

1Aerospace Corp (ret.)

March 21, 2023

Abstract

Although adequately detailed kerosene chemical-combustion Arrhenius reaction-rate suites were not readily available for combustion modeling until ca. the 1990’s (e.g., Marinov [1998]), it was already known from mass-spectrometer measurements during the early Apollo era that fuel-rich liquid oxygen + kerosene (RP-1) gas generators yield large quantities (e.g., several percent of total fuel flows) of complex hydrocarbons such as benzene, butadiene, toluene, anthracene, fluoranthene, etc. (Thompson [1966]), which are formed concomitantly with soot (Pugmire [2001]). By the 1960’s, virtually every fuel-oxidizer combination for liquid-fueled rocket engines had been tested, and the impact of gas phase combustion-efficiency governing the rocket-nozzle efficiency factor had been empirically well-determined (Clark [1972]). Up until relatively recently, spacelaunch and orbital-transfer engines were increasingly designed for high efficiency, to maximize orbital parameters while minimizing fuels and structural masses: Preburners and high-energy atomization have been used to pre-gasify fuels to increase (gas-phase) combustion efficiency, decreasing the yield of complex/aromatic hydrocarbons (which limit rocket-nozzle efficiency and overall engine efficiency) in hydrocarbon-fueled engine exhausts, thereby maximizing system launch and orbital-maneuver capability (Clark; Sutton; Sutton/Yang). The combustion community has been aware that the choice of Arrhenius reaction-rate suite is critical to computer engine-model outputs. Specific combustion suites are required to estimate the yield of high-molecular-weight/reactive/toxic hydrocarbons in the rocket engine combustion chamber, nonetheless such GIGO errors can be seen in recent documents. Low-efficiency launch vehicles also need larger fuels loads to achieve the same launched mass, further increasing the yield of complex hydrocarbons and radicals deposited by low-efficiency rocket engines along launch trajectories and into the stratospheric ozone layer, the mesosphere, and above. With increasing launch rates from low-efficiency systems, these persistent (Ross/Sheaffer [2014]; Sheaffer [2016]), reactive chemical species must have a growing impact on critical, poorly-understood upper-atmosphere chemistry systems.
Background: Garbage-In, Garbage-Out (GIGO) combustion calculations are defined here as the use of a detailed Arrhenius chemical-kinetic reaction/combustion suite for some of the chemical species which are present in real-world combustion, but the exclusion by choice of detailed reactions of other crucial chemical species known to be present. In the simplest instance, a detailed $H_2+O_2+CO$ reaction suite combined with a single kerosene overall- or net-reaction (pseudoreaction) can produce valid LOX/RP-1 (kerosene) combustion chamber pressures and temperatures for engine-design purposes, but the results cannot produce data on the hydrocarbons relevant to nozzle efficiency and mandated environmental assessments, because it ignores the complex hydrocarbon chemistry dependent upon trace-$O_2$ hydrocarbon cracking and oligomerization reactions (e.g., Christardo [2009], Seshardi [1982], VanKamp [1984], Speight [2003]) which ultimately produce the soot observed in real-world combustion. Non-GIGO chemistry for these key species in LOX/RP-1 combustion modeling has been available for decades (e.g., Marinov [1998]); see figure), and reproduces known LOX/RP-1 measurements: Large quantities – up to several percent of total fuels flow – of complex hydrocarbons are typical; benzene, butadiene, and polycyclic aromatic hydrocarbons (PAH) such as anthracene, fluoranthene, etc. (Thompson, Rocketdyne [1966]; figure). These hydrocarbons are the well-known soot-precursor species, and are therefore necessarily formed concomitantly with soot (e.g., Pugmire [2001]). The direct observation of large quantities of soot in a rocket plume is thus diagnostic of low combustion-efficiency and the presence of these chemical species, and low overall engine-efficiency due to rocket nozzle thermodynamics. The latter is due inefficiency in extracting molecular vibrational energy from the expansion of large polyatomic molecules such as benzene and PAH (Clark [1972]). Examples are provided. Additionally, legal limits have since been established on the production of these class-1 carcinogens from internal combustion engines (rocket engines), and appropriate employee and civil protections established, making these species important to model and measure directly in the lack of afterburning (see below). These considerations apply to both the open-cycle fuel-rich gas generators and main engine combustion chamber fuels and film-coolants in LOX/RP-1 engines since the products distribution is largely thermodynamically controlled.

During the decades of government-funded launch vehicle development, liquid fueled engines for space launch were typically designed for high efficiency to maximize, for instance, mass-to-orbit. To this end, pre-burners and high-energy fuels atomization (via high-energy swirl or impinging injectors) have been used to minimize fuel droplet starting size, maximizing gasification rate, and increasing gas-phase combustion efficiency and therefore engine/nozzle efficiency (Clark; Sutton; Sutton/Yang). Recent private LOX/RP-1 rocket engine development in the U.S. (SpaceX), and a strikingly similar Korean engine (Hanwha), favor particularly low-combustion-efficiency by eschewing fuel atomization in favor of Pintle-style injectors (Son [2016], Seedhouse; Springer [2013], Mueller [2012], Zarchan [2004]). The resulting low gasification rate of the characteristically large RP-1 fuel droplets maximizes the yield of large hydrocarbons (e.g. benzene, PAH, PM2.5, tars, and cokes), lowering gas-generator/nozzle/engine efficiency. Egregious GIGO errors can be seen, for instance, in recent U.S. Environmental Assessment/Impact documents (45" Space Wing [2007], U.S. Army [2007], Nelson Eng. [2013], F.A.A. [2014], Sierra Eng. #2003-001 [2003], Song [2017]) wherein GIGO calculations de facto instruct computer combustion models to ignore PAH formation known to be present. Examples are provided. Awareness of the above combustion science is essential knowledge for research personnel involved in understanding launch vehicle upper-atmospheric impacts and climate disruption.

Impacts: During flight in the Troposphere, a sufficient atmospheric partial pressure of $O_2$ is present to burn-off the PAH and soot generated by low-efficiency rocket engines, ignited by the hot rocket plume. This so-called “afterburning” of the excess hydrocarbons (soot, tars, PAH, etc.) gives rise to the brilliant appearance of an inefficient LOX/RP-1 plume shortly after lift-off in these engines; however, above the troposphere, and on the launch stand during water-deluge, afterburning ceases and the predominant quantities of these species are directly deposited and persist. Examples are provided. Low-efficiency launch vehicles also need significantly larger liftoff fuels loads (a factor of $>1.2$) to achieve similar orbital mass, further increasing the per-launch yield of complex hydrocarbon tars and stable PAH free radicals. These are deposited directly into launch trajectories, including the launch pad, stratospheric ozone layer, mesosphere, and above. Increasing launch rates of these new, low-efficiency engines must therefore have an increasing cumulative impact on critical and poorly-understood upper-atmosphere chemistry systems (Ross/Sheaffer [2014]; Sheaffer [2016]). For instance; the ozone reactions of gas- or condensed-phase hydrocarbon tars combined with PM2.5 have not yet been studied in detail, and potentially represent an unrecognized class of ozone-depleting compounds. The direct deposition of PAH tars during orbital maneuvers in populated orbits and orbital stations is of potential concern to the satellite remote-sensing community. The solar absorption of these species is much higher than that of the background atmosphere, and converts UV/Visible insolation into upper-atmospheric heat.