A COMPARISON OF OXIDIZED CARBON ABUNDANCES AMONG COMETS. M. A. DiSanti¹, M. J. Mumma¹, B. P. Bonev¹², G. L. Villanueva¹², Y. L. Radeva¹³, K. Magee-Sauer¹, E. L. Gibbs¹ NASA-Goddard Space Flight Center, Greenbelt, MD USA (michael.a.disanti@nasa.gov), Catholic University of America, Washington, DC USA, University of Maryland, College Park, MD USA, Rowan University, Glassboro, NJ USA, U. Missouri-St. Louis, St. Louis, MO USA.

Comets contain relatively well preserved icy material remaining from the epoch of Solar System formation, however the extent to which these ices are modified from their initial state remains a fundamental question in cometary science. As a comet approaches the Sun, sublimation of the ices contained in its nucleus (termed "native ices") releases parent volatiles into the coma, where they can be measured spectroscopically.

One means of assessing the degree to which interstellar ices were processed prior to their incorporation into cometary nuclei is to measure the relative abundances of chemically-related parent volatiles. For example, formation of C₂H₆ by hydrogen atom addition (e.g., to C₂H₅) on surfaces of ice-mantled grains was proposed to explain the high C₂H₆/CH₄ abundance observed in C/1996 B2 (Hyakutake) [1]. The large C₂H₆/CH₄ abundance ratios measured universally in comets, compared with those predicted by gas phase production of C₂H₆, establishes H-atom addition as an important and likely ubiquitous process.

CO should also be hydrogenated on grain surfaces. Laboratory irradiation experiments on interstellar ice analogs indicate this to require very low temperatures (T ~ 10-25 K), the resulting yields of H₂CO and CH₂OH being highly dependent both on hydrogen density (i.e., fluence) and on temperature (12,13). This relatively narrow range in temperature reflects a lack of mobility below 8-10 K on the one hand, and reduced sticking times for H-atoms as grain surfaces are warmed above 20 K on the other. The relative abundances of these three chemically-related molecules in comets provides one measure of the efficiency of H-atom addition to CO on pre-cometary grains (Fig. 1).

Here, we compare the oxidation sequence of carbon in several comets observed with modern near-infrared echelle spectrometers, which favor the detection of native ices. We use our measured native abundances of CO, H₂CO, and CH₂OH (Fig. 1, top panel) to assess the efficiency of H-atom addition. Our results show their relative abundances to vary considerably among comets (Fig. 1, bottom panel), suggesting a range of conditions experienced by their nascent ices. Comparison with laboratory yields can provide information on conditions in the formative environment.

Fig. 1. A. Abundances (relative to H₂O) of oxidized carbon in several comets. Error bars show 1σ uncertainties, and upper limits (↑) are 3σ. The small beams afforded by modern IR spectrometers favor detection of native ices. B. Corresponding CO conversion efficiencies. These are expressed in terms of the ratio ([H₂CO]+[CH₂OH])/([CO]+[H₂CO]+[CH₂OH]), where [] indicates mixing ratio (abundance relative to H₂O). This approach assumes that formaldehyde and methanol are produced solely from CO, and that they are the only products. It does not include potential loss of CO (e.g., in the subsequent proto-solar environment) or its incorporation into more complex entities (e.g., polymers such as polyoxymethylene, POM). Note that, by the above definition, for comets in which H₂CO was not measured (e.g., C/2006 M4), or for which only upper limits for H₂CO or CH₂OH were obtained (8P/Tuttle, C/2007 W₃, C/2001 S₄), the conversion efficiency could be higher than shown. Conversely, loss of CO by other means (such as incorporation into POM) would lower the efficiency.