The Thermal Electrical Conductivity Probe (TECP) for Phoenix

Aaron P. Zent
Michael H. Hecht
Doug R. Cobos
Gaylon S. Campbell
Colin S. Campbell
Greg Cardell
Marc C. Foote
Stephen E. Wood
Manish Mehta

a, MS 245-3, NASA Ames Research Center, Moffett Field, CA 94035
b, Jet Propulsion Lab, 4800 Oak Grove Drive, Pasadena, CA 91109
c, Decagon Devices, 2345 NE Hopkins Court, Pullman, WA 99163
d, University of Washington, 367 Johnson Hall, Seattle WA 98195
e, University of Michigan, Department of Atmospheric, Oceanic and Space Sciences, Ann Arbor, MI 48109

Revised: July 30, 2008

Running Title: Phoenix TECP
Email: Aaron.P.Zent@nasa.gov
ABSTRACT

The Thermal and Electrical Conductivity Probe (TECP) is a component of the Microscopy, Electrochemistry, and Conductivity Analyzer (MECA) payload on the Phoenix Lander. TECP will measure the temperature, thermal conductivity and volumetric heat capacity of the regolith. It will also detect and quantify the population of mobile H₂O molecules in the regolith, if any, throughout the polar summer, by measuring the electrical conductivity of the regolith, as well as the dielectric permittivity. In the vapor phase, TECP is capable of measuring the atmospheric H₂O vapor abundance, as well as augment the wind velocity measurements from the meteorology instrumentation. TECP is mounted near the end of the 2.3 m Robotic Arm, and can be placed either in the regolith material or held aloft in the atmosphere.

This paper describes the development and calibration of the TECP. In addition, substantial characterization of the instrument has been conducted to identify behavioral characteristics that might affect landed surface operations. The greatest potential issue identified in characterization tests is the extraordinary sensitivity of the TECP to placement. Small gaps alter the contact between the TECP and regolith, complicating data interpretation. Testing with the Phoenix Robotic Arm identified mitigation techniques that will be implemented during flight. A flight model of the instrument was also field tested in the Antarctic Dry Valleys during the 2007-2008 International Polar year.
I Introduction

Phoenix will be the first Martian lander to operate at polar latitudes, affording a unique opportunity to study the current climate, the role of surface-atmosphere exchange, and the polar climate record in the structure and distribution of ground ice. It will make the first-ever measurements to constrain volatile and energy fluxes in a region that is key to Martian climatic evolution, and the first physical and chemical analyses of Martian ground ice. By combining observations of present-day processes, and the geologic record of past processes, Phoenix will provide fundamental insights into Martian history, and its astrobiological potential.

The Thermal and Electrical Conductivity Probe (TECP), a component of the Microscopy, Electrochemistry, and Conductivity Analyzer (MECA) on the Phoenix Lander, will address these topics. Based on the small, dual-probe sensors that are routinely used to monitor soil thermal properties and water content, TECP performs 6 distinct measurements: dielectric permittivity, electrical conductivity, temperature, thermal conductivity, volumetric heat capacity, and relative humidity. In addition, application of the thermal properties protocol when the TECP is held aloft by the Robotic Arm (RA) allows constraints to be placed on the average wind speed.

This paper describes the TECP, its measurements, its design and implementation, its calibration, characterization and field testing, as well as some of the anticipated science results. The importance of the TECP dataset is enhanced by the uniqueness of the Phoenix landing site in the Martian polar regions.
**A. The High Martian Latitudes**

The Phoenix lander touched down on May 25, 2008 at 68.16°N 233.35°E, (Figure 1 and Arvidson et al., *This Issue*), an area that is fairly typical of the Vastitas Borealis plains. Investigation of a near-polar landing site gives us the opportunity to characterize a part of Mars that is more dynamic, and arguably, over the past several $10^7$ years at least, more globally significant, than previous landing sites. Many components of the Martian climate are effectively controlled, or play out most dramatically, at these latitudes.

**H$_2$O Reservoir**

The high latitude surface is underlain by probably the largest H$_2$O reservoir on the planet. Substantial permafrost deposits were first predicted on energy balance considerations (Leighton and Murray, 1966). Post-Viking, there was supporting geomorphic evidence for widespread ground ice at high latitudes (Rossbacher and Judson, 1981; Squyres 1986). The presence of high-latitude ice deposits were confirmed by Mars Odyssey (Boynton et al., 2002).

Among the criteria for choosing the Phoenix site was that the modeled Odyssey GRS data suggests that the dry overburden thins somewhat at these longitudes; access to both the ground ice and its dry overburden were considered fundamental objectives.

**Seasonal Cycle**

The Phoenix landing site is also part of the circumpolar terrain covered annually by the seasonal CO$_2$ cap. Within the seasonal cap perimeter, the surface energy balance controls the condensation of H$_2$O and CO$_2$. Cap condensation and sublimation are the dominant climatic processes in the current epoch. Phoenix is well placed to observe and begin characterization of cap condensation processes.
Once the seasonal cap has retreated poleward of the Phoenix site, the high-latitude regolith is likely a source of H₂O to the atmosphere, which along with cap, buffers planetary atmospheric H₂O (Jakosky, 1983). Depending on the thermal and diffusive properties of the regolith, H₂O vapor exchange with the ground ice could have a major impact on the annual atmospheric water budget (Schorghofer and Aharonson, 2005).

Results from the Planetary Fourier Spectrometer (PFS) suggest the possibility that atmospheric H₂O is preferentially concentrated in the lowest few kilometers of the surface at high northern latitudes during summer (Tschimmel et al., 2008). If true, this would affect the upper boundary condition in atmospheric H₂O, and hence subsurface ice stability models.

**Obliquity Effects**

In addition to the important role the polar regolith plays in determining seasonal cycles, the high latitudes experience the most extreme variations over orbital oscillations. In particular, variations in orbital obliquity have a pronounced effect on polar temperatures. At low obliquity, variations in polar insolation over the year are minimized. It was formerly believed that semi-permanent CO₂ caps would form at both poles at minimum obliquity (Toon et al. 1980, Pollack and Toon, 1982, Fanale et al, 1982), cold-trapping regolith ice away. More recent modeling however, calls into question the nature of the cap at low obliquity (Haberle et al., 2003; Armstrong et al., 2004). When obliquities are high, the polar regions experience extreme seasonal insolation variations. The north polar H₂O cap becomes unstable at some obliquity between 35° and 45° (Mischna et al., 2003). The stable location for ice at obliquities > 45° is the low-latitude highlands, as first suggested by Jakosky and Carr, (1985), but more recently
supported by Head et al. (2003).

Because it is at the poles that obliquity variations have the most pronounced effect, the subsurface at the Phoenix site may provide a record of the processes and mechanisms that drive long term climate change.

**Periglacial Processes**
There are other similarities between the Martian high latitude surface, and some of Earth’s periglacial terrain. Understanding their true origin and nature is essential if we are to read the geologic record, both at the surface, and in the subsurface.

**Polygonal Ground**
Much of the Northern Plains, including the Phoenix landing site, are covered with patterned ground that is remarkably similar in appearance to examples that are commonly seen at terrestrial polar latitudes. (Figure 1; Arvidson et al., *This Issue*; McEwen et al. 2007)

Polygon formation begins with formation of a network of vertically-propagating cracks due to the thermal contraction of ground ice during autumn and winter. Mellon (1997) modeled the process for the Martian climate, and found the yield strength of frozen ground is exceeded during autumnal cooling, leading to mechanical failure and crack formation (Lachenbruch, 1962; French, 1996). The cracks then accumulate either clastic particles, or, if it is available, liquid water. It is also possible to condense additional H$_2$O vapor in these cracks, a process that could likewise contribute to the accumulation of excess ground ice (Fisher, 2005).

At current pressures and temperatures, liquid water is not stable on Mars long enough to fill the cracks, but past conditions may have permitted this process (Costard et al., 2002).
Whether these features are relict of freeze-thaw cycles, due solely to thermal contraction, or are the product of other processes is a key question that Phoenix must resolve.

**Excess Ground Ice**
The Gamma Ray Spectrometer on Mars Odyssey shows the presence of an ice-rich layers with unexplained abundances of more than 60% of ice by volume (Boynton et al., 2002; Feldman et al., 2002). Buried H₂O deposits, such as snow, layered with dust (Mischna et al., 2003) are a possible explanation for the large amount of interstitial ice, but periglacial freeze-thaw cycles could produce the same result.

During freezing events, as the freezing front penetrates downward into the surface, unfrozen, interfacial water is removed to the ice phase, creating a hydraulic gradient along which additional water migrates. As long as the total (sensible + latent) heat flux carried by unfrozen water from below can balance the diffusion of heat upwards to the surface, the freezing front will remain stationary, and ice lenses will develop in freezing soil. Once the regolith beneath the freezing front is depleted of H₂O, the sensible + latent heat flux to the freezing front is no longer sufficient to balance conduction, and the freezing front again begins to propagate to depth. This continues until another wet zone is reached, and another ice lens begins to form. Terrestrial periglacial terrain often exhibits numerous ice lenses in the shallow subsurface.

**Habitability**
The thermal physics of Martian permafrost are also of interest because Earth’s high-latitude permafrost, which bears many morphological similarities to the Phoenix landing site, is a rich habitat for psychrophiles.

Rivkina et al., (2000) argue, based on growth metabolism in the laboratory, that if the
temperature is stable, as in deep permafrost, the minimum temperature for growth is perhaps 253 K. Growth metabolism is easily detectable at 263 K (Finegold, 1996).

Price and Sowers (2004) reviewed the literature on psychrophilic life, and could find no detectable lower temperature limit for metabolism. They concluded that the extremely low energy costs of survival energy enable microbial communities in extreme environments to survive indefinitely. They also reported evidence for survival-level metabolism all the way down to 233K, meaning that much of the Martian regolith and ground ice qualifies as “survivable” if only thermal limits are considered.

Jakosky et al (2003), examined the minimum temperatures at which interfacial unfrozen water can exist, the minimum subfreezing temperatures at which terrestrial organisms can grow and multiply, and the maximum temperatures that can occur in Martian high-latitude and polar regions, to see if there is overlap. They concluded that high-latitude Martian ground ice was just at the limit of being habitable.

The most probable lifetime limitation is exposure to galactic cosmic rays (Kminek et al., 2003); an accumulation of damage eventually overwhelms the cell’s repair mechanisms. Kminek et al (2003) predicted that the time required to reduce a viable spore population to $10^{-6}$ would be $< 6 \times 10^5$ years in the top meter of the regolith, and $< 7 \times 10^4$ years in the otherwise-favorable upper 10 cm.

Zent (2008) used a time-resolved thermal model of the subsurface, tuned to the Phoenix landing site, along with the 10-My orbital calculations of Laskar et al. (2004), to calculate several possible 10 My thermal histories of Phoenix’s target ice. In all cases, there were many clement periods when growth, even rapid growth, was possible. Also however, in
every case, the maximum dormancy period that must be survived was at least an order of magnitude greater than the $\sim 7 \times 10^4$ years quoted by Kminek et al. (2003)

**B. Science and Measurement Objectives**

The science objectives of the Phoenix Mission are:

1. To study the history of Martian water in all its phases.
2. To search for evidence of habitable zone and assess the biological potential of the ice-soil boundary.

The polar regolith holds a substantial fraction of the remaining Martian H$_2$O inventory, and is a dynamic reservoir for H$_2$O over both seasonal and astronomically-driven climate cycles. The abundance, distribution, and structure of polar ground ice may also have preserved a geologic record of the recent climate and subsurface conditions. Finally, as we have already discussed, polar ground ice may be just at the margin of habitability.

The TECP was included on Phoenix specifically to determine the regolith thermophysical properties that control energy fluxes between the atmosphere and subsurface, to characterize the processes that control the distribution and exchange of H$_2$O between the atmosphere and subsurface in the current climate, and to document the occurrence of any unfrozen water, both as thin films in the regolith, and as atmospheric vapor.

**1. Science Objectives**

**a. Energy Fluxes**

Regolith thermal properties, particularly the thermal conductivity ($\kappa$) and the volumetric heat capacity ($C_v$) are key, and currently unknown, parameters in any model that purports to address the distribution of Martian ground ice, the climatic cycles associated with
variations in orbital parameters, or the annual cycles of H_2O and CO_2. These properties
determine how effectively heat conducts from the illuminated surface to depth, and fix
the thermal profile from the surface down at least as far as the ice table.

Based upon these measurements, estimates of the thermal inertia of the surface can be
made, and compared with thermal inertias that are derived from measurement of the
surface temperature throughout a thermal cycle, whether by TECP itself, or with thermal
inertias derived from orbital data (e.g. Mellon et al., 2000).

b. Regolith H_2O

Another set of objectives involves understanding the abundance, state, and mobility of
H_2O in the regolith. H_2O redistributes itself continuously through the regolith,
continually accumulating in the lowest-energy reservoir that is thermodynamically
available. Because soil and ice co-exist in the regolith, it is necessarily true that there is
a finite and variable population of un-frozen H_2O molecules; if sufficiently thick films of
unfrozen water occur, surface diffusion in the film can outstrip diffusion in the vapor
phase. These mobile H_2O molecules are most easily detected in measurements of soil
electrical properties, which are otherwise dominated by highly insulating silicates and
ice.

Because TECP can measure both unfrozen water and temperature over a diurnal cycle, it
may be possible to characterize both the soil water characteristic curve (the relationship
between soil water content and soil water pressure potential) as well as the hysteresis
curve of H_2O adsorption in the regolith materials. These are key parameters in
determining the flow and transport of unfrozen water in the regolith.
c. Atmosphere

The atmosphere serves as the conduit through which H$_2$O is distributed around the planet, and as the upper boundary condition controlling the state and distribution of H$_2$O throughout the regolith. Exchange between the atmosphere and regolith is controlled substantially by the absolute humidity of the atmosphere, and somewhat to a lesser extent by the wind fields. Therefore, TECP will also measure atmospheric relative humidity, and, by applying the same protocol used to determine regolith thermal properties, can constrain average wind speeds averaged over the duration of the measurement.

2. Measurement Objectives

The measurement objectives for TECP are:

• Measure the temperature of the Martian regolith, both at the surface, and within the excavation trench, as a function of Local Solar Time (LST), and the sub-solar angle ($L_s$) which is used to express Martian seasons. If different types of materials can be accessed, these measurements will be repeated over the same timescales for each material. Based upon surface temperature observations (Kieffer and Titus, 2001), measurement requirements on the temperature range from 165K to 323 K, with 2K precision, and an accuracy of ±10%.

• Measure the thermal conductivity and heat capacity of accessible Martian regolith materials. The TECP thermal conductivity requirement is to measure soil thermal conductivity ($\kappa$) with 10% accuracy over the range 0.03 – 2.5 [W m$^{-1}$ K$^{-1}$]. A typical value for thermal conductivity in frozen soils or solid ice is $\kappa = 2.0 \pm 1.0$ [W m$^{-1}$ K$^{-1}$] at $T = 223$ K. This value is from many earth observations over many
types of soil, including pure ice. We concluded that we would not require TECP
to measure pathologically low thermal conductivity, since the landing event and
the measurement attempt itself would probably destroy any such fluffy surface;
simply confirming an off-scale low $\kappa$ would be useful. Note that for such a
surface the thermal inertia, $I = \sqrt{\kappa cp} = (0.03*800 \text{ [J/kg K]} * 800 \text{ [kg/m}^3]\right) = 138,$
which is extremely low. The upper limit on the thermal conductivity requirement
was a result of heritage design constraints. Thermal conductivities above 2.5 [W
m$^{-1}$ K$^{-1}$] result in very small temperature increases at the heated needle, which are
difficult to resolve with the 12 bit ADC.

- The volumetric heat capacity ($C_V$) of a particulate mixture is the heat capacity of
each component, weighted by its volumetric relative abundance; or approximately
the specific heat of the regolith times its density. The requirement for the
instrument is to measure the volumetric heat capacity of the regolith with 10%
accuracy over range 0.4 to 4 [MJ/m$^3$ K], which approximately spans the range
from dry frozen ground to nearly the heat capacity of liquid H$_2$O (4.19 [MJ/m$^3$
K]) (Hinkel et al., 2001; Kane et al., 2001).

- Measure the electrical conductivity (inverse of the electrical resistivity) of the
regolith materials, particularly as a function of measured regolith temperature.
The requirements for electrical conductivity are to measure soil and ice electrical
conductivity from $10^7$ [nS/cm] to 1 [nS/cm] with 10% accuracy. Conductivity, ($\sigma$)
is a property of a material (e.g., copper), as opposed to conductance, which is a
property of a specific conductor (e.g. a wire made of copper). Conductivity has
units of Siemens per meter [S/m]; conductivity will typically be referred to in terms of [nS/cm]

$$\sigma \left[ \frac{S}{m} \right] = \frac{f \left[ \frac{m}{s} \right]}{R \left[ \Omega \right] A \left[ m^2 \right]} = 10^7 \left[ \frac{nS}{cm} \right]$$  \[1\]

De-ionized water has a conductivity of $5.5 \times 10^{-6}$ [S/m] (55 [nS/cm]); Siberian permafrost has been measured with electrical resistivity, $R$, as high as $1.4 \times 10^4$ [$\Omega$-m] ($\sigma = 715$ [nS/cm]) at depth $>2.2$ m just before warming begins (Olhoeft, 1978; Pozdnyakov and Chan, 1979).

- Measure the bulk dielectric permittivity of the regolith materials over the range of 1 to 20, with resolution of 0.005, as a function of regolith temperature. The dielectric permittivity of any gas is $\sim 1$, and $\varepsilon$ for water is 78.54. We chose not to measure the full range because high sensitivity (hence high resolution) is desired at the low end of the range to maximize sensitivity to small changes in the abundance of unfrozen water. The upper limit of 20 is imposed because MECA data words are only 12 bits, so achieving sensitivity of 0.005 at the low end of the range limited the upper limit of the dynamic range to 20.

The relationship between the water content of a soil and dielectric permittivity is complex and continues to be studied (Topp et al. 1980; Jones and Or, 2003; Lebron et al., 2004; Sweeney et al. 2007). The overall dielectric is the sum of the dielectrics of air, mineral, ice and water. In a non-frozen system, we typically assume unfrozen water is the only constituent changing in the medium; all other dielectrics stay the same. The change in dielectric permittivity can then be interpreted solely in terms of the change in unfrozen water. However, $T$ changes
will affect permittivity as well; therefore all $\varepsilon$ measurements are carried out with simultaneous $T$ measurements of the medium.

- Measure the $\text{H}_2\text{O}$ vapor density of the Martian atmosphere as a function of LST, $L_s$, and height above the surface, including measurements made within the trench and particularly just above the ice layer if an ice layer is detected. TECP also includes a GE Panametrics MiniCap 2 polymer relative humidity ($R_H$) sensor, a commercial capacitive polymer-based sensor whose performance at temperatures and frost-points below 233 K was not well known initially (see the Calibration Discussion, below). The requirement is to measure the $\text{H}_2\text{O}$ vapor pressure from 0 – 0.5 kPa over the temperature range of 195 K – 270 K, with accuracy of ±10% of reading. Because the sensor is sensitive to $R_H$, the absolute partial pressure can only be determined with a measurement of air temperature. Strategies for determining absolute $\text{H}_2\text{O}$ partial pressure are discussed below. An additional consequence of this approach is that the resolution in absolute pressure is a function of the sensor temperature, which is related to, but not identical to, the air temperature.

- Measure, or characterize the wind speed as a function of LST, $L_s$, and height, including within the trench.

II Technical Approach

In order to meet the measurement requirements, TECP was developed as a single electronics box, (Figure 2) integrated with four needle-like “probes”, each of which has a role as a component in a unique set of circuits (Figure 3).
The TECP needles are fitted into a plastic housing, which in turn is attached to the instrument electronics box. The plastic interface is machined from polyethylethylketone (PEEK), which has low thermal and electrical conductivity ($\kappa = 0.249 \text{ W m}^{-1} \text{ K}^{-1}$; $C_V = 2.177 \text{ kJ kg}^{-1} \text{ K}^{-1}$), and provides insulation between the thermally and electrically activated needles and the electronics box. The aluminum electronics housing is anodized to increase emissivity, enhancing thermal radiation, and minimizing heating when TECP is operating in direct sunlight. A port in the side of the instrument housing allows the relative humidity sensor to access the Martian atmosphere. The sensor is shielded behind a Teflon membrane that is porous to atmospheric $\text{H}_2\text{O}$ but prevents dust from fouling the humidity sensor.

TECP has a single rotational degree of freedom relative to the RA, (Figure 4), which in turn has 3 degrees of freedom relative to the lander and the environment.

**A. Temperature**

The temperature of three of the needles (1, 2, and 4) is measured as the difference between the sense and reference junctions of a Type E (chromel-constantan) thermocouple. The thermocouple was fabricated by soldering the ends of a 7.5 cm piece of constantan thermocouple wire to the ends of two 75 cm pieces of chromel thermocouple with Kester Sn63Pb37 rosin core solder and Indium Flux #1. The reference junction of the thermocouple is on the TECP analog electronics board, adjacent to a National Semiconductor LM234DT temperature-sensing current source. The temperature derived from this element is referred to as the board temperature ($T_b$); the output current is directly proportional to the absolute temperature, and extrapolates to zero at 0 K.
**B. Thermal Properties**

The *in situ* thermal properties of the Martian regolith are determined via the transient heated needle technique (de Vries, 1952). In this approach, a heat pulse is applied via a heater that approximates a line source, and the thermal response of the medium is measured at some radial distance from the heat source.

In addition to thermocouples, needles 1, 2 and 4 are also equipped with resistance heaters. The heater is Evanohm, an alloy used extensively for precision wound resistors. Its composition is nominally 0.75 Ni, 0.20 Cr, 0.025 Al, and 0.025 Cu. The Evanohm is dipped in lacquer because the thermal grease used to ensure good thermal contact to the needles has a metallic component, which would cause the heater to short to the needles.

In typical operations, needle 1 is heated, and the temperature change of both needle 1 and needle 2 is monitored. The temperature change of needle $i$, between time 0 and time $t$ is determined from

$$\Delta T_{i,t} = (T_{i,t} - T_{i,0}) - (T_{4,0} + T_{4,t})$$  \[2\]

Needle four (farthest from the heated needle) is used as a temperature reference, as it is unaffected by the heat pulse from the needle for heat pulses of normal duration. This method also has the advantage of removing the effects of thermal drift of the bulk material or the TECP electronics board during the measurement.

Carslaw and Jaeger, (1959) modeled the temperature surrounding an infinite line heat source with constant heat output and zero mass, in an infinite medium. When a constant amount of heat is emitted from unit length of a zero-mass heater over a period of time, the temperature response at radial distance $r$, is
where $q$ is the rate of heat dissipation [W m$^{-1}$], $t_1$ is the heating time, and $Ei$ is the exponential integral. The temperature change after the heat is turned off is given by:

$$
\Delta T_{i,t} = -\frac{q}{4\pi \kappa} Ei\left(\frac{-r^2 C_V}{4\kappa t}\right) 0 < t < t_1 \quad [3]
$$

Material thermal properties are determined by fitting the time series temperature data during heating to Eq. (3), and during cooling to Eq. (4); note that the absolute temperature $T$ doesn’t enter into the calculation, only $\Delta T_i$. Thermal conductivity and heat capacity can be obtained from the temperature of the heated needle (single needle), with $r$ taken as the radius of the needle and independently by fitting equations 3 and 4 to the temperatures measured at the adjacent needle, 7 mm from the heated needle.

For the fitting procedure, a multi-stage Monte Carlo method described by Conley (1985) is used to obtain values close to the global minimum, and then a Marquardt method is used to find the final values. The exponential integral function was approximated using formulae from Abramowitz and Stegun (1972).

The outputs of the solver function are $\kappa_{f1}$ and $C_{f1}$, which are the thermal conductivity and volumetric heat capacity measured at the heated needle (needle 1), and $\kappa_{f2}$ and $C_{f2}$, which are thermal conductivity and volumetric heat capacity measured at the adjacent needle (needle 2). The values for $\kappa_f$ and $C_f$ found via fitting to Equations 3 and 4 are inherently inaccurate due to inevitable discrepancies between the TECP and the mathematically idealized model. However, the true thermal properties can be determined.
if functional calibration is performed against materials of known thermal properties as described below.

**C. Dielectric**

To measure the dielectric permittivity of the regolith, two of the needles (3 and 4) are used as the plates of a capacitor; with the regolith serving as the dielectric. The measurement involves putting a microwave frequency (~6.25 MHz) voltage square wave on the capacitor, and measuring the time history of the charge on the plates.

The time, \( t \), required to charge a capacitor from an initial voltage \( (V_i) \) to a final voltage \( (V_f) \), with an applied voltage \( (V) \) is

\[
t = -RC \ln \left( \frac{V_f - V}{V_i - V} \right)
\]

where \( R \) is the series resistance and \( C \) is the capacitance. The capacitance, in turn, is a function of the dielectric permittivity of the medium between the plates (or in this case, the regolith between the needles)

\[
C = \varepsilon L_C
\]

where \( L_C \) is a geometric factor that encompasses the area of the capacitive plates and their separation. \( L_C \) has units of length [m]. The relationship between the time history of the plate charge and the dielectric permittivity is then

\[
\frac{1}{\varepsilon} = \frac{1}{t} \left[ -RL_C \ln \left( \frac{V_f - V}{V_i - V} \right) \right]
\]

Because we are measuring the effective permittivity of a mixture of phases, rather than a pure phase, we refer to the measured quantity as the bulk dielectric permittivity (\( \varepsilon_b \)).
**D. Electrical Conductivity**

The electrical conductivity ($\sigma$) of the soil is measured between needles 1 and 2. The regolith conductance ($G_R$), the inverse of the regolith resistance ($R_R$), is measured via a simple voltage divider (cf. Figure 3). Three fixed resistors ($R_F = 1\, \text{k}\Omega; 100\, \text{k}\Omega; 3\, \text{M}\Omega$) are used to allow the 12-bit data word to cover the entire dynamic range specified in the requirements with sufficient precision. This results in three distinct measurement channels, each with its own effective range, which are measured sequentially (Table 4).

The input voltage ($V_i$) is 2.5V, and the resistance of the regolith is found from the output voltage ($V_o$) via:

$$R_R = \frac{V_o}{V_i} \left( \frac{R_F}{V_i - V_o - 1} \right)$$

The conductivity of the regolith is found by multiplying conductance ($1/R_R$) by the probe constant ($C_P$), which in effect normalizes for the geometry of the electric field that permeates the regolith between the needles. The determination of the probe constant is described below.

**E. H$_2$O Vapor**

The TECP H$_2$O measurement is based on a GE Panametrics MiniCap 2 polymer relative humidity sensor. It is a capacitance-based instrument that measures the permittivity of a polymer film which adsorbs H$_2$O. The circuit in the TECP is read out in a manner similar to the capacitance measurement already described for the soil measurement.

Solid-state humidity sensors of this type are sensitive to relative humidity, almost independent of temperature. Specifically, the $R_H$ is measured relative to the equilibrium
pressure of liquid H₂O, rather than ice, regardless of temperature. At $T < 273$ K, the saturation vapor pressure of supercooled water is used.

The physical basis for this behavior is likely that atmospheric H₂O forms an adsorbed film on the sensing polymer in equilibrium with the atmosphere, and it is the thickness of this adsorbate that is actually sensed as a change in permittivity. It can be shown that, for many adsorbents, the mass (or thickness) of this adsorbed layer is nearly the same for a given RH, regardless of temperature (Anderson, 1995). Above some limiting film thickness, corresponding to $R_H = 100\%$, the thickness of the film increases without limit, and bulk H₂O condenses.

Unfortunately, there is a low-temperature limit in humidity measurements of this type. To substantially affect the bulk permittivity of the sensing film, the H₂O molecules must be free to align with the applied electric field; that is they must remain “unfrozen”.

Should some of the water adsorbed in the sensing film undergo spontaneous crystallization to ice, the device would instantly lose sensitivity to that portion of H₂O, resulting in a discontinuity in sensor output.

One may think of the interfacial, unfrozen water in the sensor as being in dynamic equilibrium with the competing forces of the H₂O hydrogen bonds in ice, and the van der Walls bonds of adsorbing polymer. As $T$ decreases below 273 K, the hydrogen bonds between H₂O molecules increasingly overcome the sum of the H₂O kinetic energy + the energy of the H₂O-polymer bond, and the maximum possible thickness of the unfrozen water layer decreases. At whatever temperature this ever-decreasing maximum reaches the film thickness corresponding to $R_H = 100\%$, ice will condense, and the instrument will begin to lose sensitivity, beginning at the highest humidity. Experience suggests that
in the TECP, this lower limit is reached ~ 205 K. Additional characterization work, and examination of the flight data will determine this limit more precisely.

In TECP, the humidity sensor is located adjacent to the board temperature sensor on the analog electronics board, and $T_b$ is used as a measure of the $R_H$ sensor. Due to power dissipation on the electronics board, $T_b$ is typically slightly higher than the ambient air temperature during operation. This works to our advantage, in that it maintains the sensor above the frost point when the atmospheric $R_H$ is high. However, it also means the $R_H$ of the sensor is lower than that of the atmosphere. In practice however, we can use the measured $R_H$ along with $T_b$ to calculate the absolute humidity of the atmosphere. If an estimate of atmospheric $R_H$ is desired, it can be calculated from absolute humidity along with an independent atmospheric temperature measurement, such as those made by the meteorology payload.

III Calibration

A. Temperature

1. Board Temperature Sensor

The board temperature sensors were calibrated to NIST-traceable temperature standards in a two part process. Opening the flight units and attaching temperature sensors directly to the boards was not an acceptable option, so an Engineering Model (EM) unit was used to characterize the temperature difference between $T_b$ and the temperature at the outside of the TECP enclosure (on the aluminum cover directly above the board temperature sensor). NIST-traceable temperature references were then attached to the flight TECP
units above the board temperature sensors and measured while the TECP was brought to
temperatures over its full expected temperature range. The correction determined on the
EM unit was then applied and the board temperature calibration function was determined.
The temperature difference between the PCB and the analog board cover varied smoothly
as a function of temperature according to

$$\Delta T = -4.3917 \times 10^{-6} T^2 + 2.8044 \times 10^{-4} T + 0.64736$$  \[9\]

with $R^2 = 0.996$. This polynomial was applied to the calibration data.

Diode type, NIST-traceable temperature sensors (Lake Shore Cryotronics model DI-470)
were attached with kapton tape to the TECP aluminum cover directly over the board
temperature sensors on the TECP flight units. The TECP units were placed in an
aluminum thermal enclosure in a Delta Design environmental test chamber. The
chamber was used to bring the thermal enclosure and TECP to equilibrium at eight
post-processing, the difference between $T_b$ and the analog cover temperature (Eq. 9) was
applied to the temperatures measured by the diode sensors. Figure 5 shows the calibration
function for $T_b$ on the TECP flight unit.

2. Thermocouple Calibration

Thermocouple calibration involved determining the Seebeck coefficient ($C_S$) vs.
temperature for the thermocouple alloys used in the TECP needles. This allows accurate
determination of the temperature difference between the TECP needles and $T_b$. The
Seebeck coefficient measured here was from the same spool used to fabricate the flight
model thermocouples.
Two cylindrical pieces of copper 1.3 cm long x 4.6 cm diameter were bolted 6.5 cm apart on a piece of acrylic, which was set on top of a block of open cell foam in a Delta Design temperature chamber. The two blocks each had a 3.2 mm diameter hole bored to their center from the rounded surface which housed pre-calibrated temperature sensors and thermocouple junctions. The thermocouple junctions and NIST traceable temperature sensors were co-located and potted in the borehole in the copper blocks with Arctic Silver high thermal conductivity grease.

In one of the blocks, a 0.4 cm hole was bored completely through the block to facilitate a 180 Ω electrical heating element, allowing the block temperature to be controlled independently, and thus achieve a temperature difference between the two thermocouple junctions. (Figure 6).

The Delta Design temperature chamber was used to control the temperature of the atmosphere around the thermocouples. Seebeck coefficients were determined for the thermocouple at approximately 30 K increments from 150 to 330 K. The blocks were allowed to equilibrate at each ambient temperature setting for at least 30 minutes. At each ambient temperature, a voltage between 0 and 32 V was applied to the heater in the copper block producing a temperature difference (ΔT) between the two blocks. At least 60 seconds were allowed to pass between the final voltage adjustment and the recording of data to ensure that the system was in thermal equilibrium.

Figure 7 shows the Seebeck coefficient vs. temperature data collected here, which closely follow a third order polynomial function. The Seebeck coefficient function shown here is used to calculate the temperature difference between the needles and $T_b$ on the flight unit.
**B. Thermal Properties**

The TECP thermal properties measurement functions were calibrated at the system level. Measurements in materials of known thermal properties were used to construct a calibration function for the thermal conductivity measurement and volumetric heat capacity measurement for TECP in a manner described in Cobos et al., (2006) and summarized here.

TECP thermal properties time series data sets were collected in five materials with known thermal properties (Table 1). The Macor ceramic, high density polyethylene (HDPE), and glycerol were obtained from commercial vendors. The expanded polystyrene (EPS) was obtained from a larger block of certified material used as a thermal conductivity calibration standard at Decagon Devices. The water was stabilized with agar powder at a mixing ratio of 5 [g/l] to prevent convective heat loss from the heated needle.

In the case of the stabilized water and glycerol, thermal properties values were taken from published reference literature. The thermal properties of the Macor and HDPE samples were determined with a commercial thermal properties meter (model KD2 Pro, Decagon Devices), which was calibrated immediately prior to testing. The EPS material is a commercially available, NIST traceable thermal conductivity standard material obtained from LaserComp, Inc. The volumetric specific heat of the foam standard was calculated calorimetrically from published values of the specific heat and density of polystyrene and the measured density of the EPS.

All measurements were taken in a drybox using a vertical stage to insert the TECP needles into the standard materials. An N₂ purge was used during measurement of the
glycerol, which is hygroscopic. The stabilized water and glycerol were contained in 600 ml beakers, into which the TECP was lowered until the metal sensing needles were fully immersed in the fluid, but the PEEK shoulders were not. The TECP sensing needles were inserted into the EPS block using the vertical stage, and were again inserted until the needles were fully encased, and the PEEK shoulders were not. The HDPE and Macor samples were machined to accommodate the four TECP needles. A total of six needle holes were machined into the HDPE and Macor blocks, with the center two holes purposely cut slightly under-sized to ensure that the cones of needles 1 and 2 made solid thermal contact with the material. This resulted in a small gap between the PEEK shoulder and the calibration block. Before inserting the TECP into the Macor and HDPE blocks, all of the holes were completely filled with DI water to promote thermal contact between the needles and the material. The excess water that was forced from the holes during the insertion of the TECP was removed from the surface of the material block with a pipette. After insertion into each material, the system was allowed to come into thermal equilibrium before data were collected. Additionally, between repeated measurements on the same sample, the system was also allowed to come back to thermal equilibrium after introduction of the preceding heat pulse.

For each thermal properties measurement, the TECP control software was configured to record the output of the board temperature sensor, the three needle thermocouples, and the heater current sense resistor at 1 Hz for 160 s. A 60 second heat period was applied to needle 1 after an initial 10 second buffer period with no heat. After cessation of the heat pulse, 90 s of cooling data were recorded. Each thermal properties measurement was repeated at least twice in each material. For the glycerol, the needles were removed
and re-inserted into the same sample between measurements. For the stabilized water, the needles were removed from the first sample and inserted into an identical, virgin sample between measurements. For the EPS, the needles were removed and inserted into a virgin section of the same block of the material. For the HDPE sample, the needles were removed from the sample, and the orientation of the sample was reversed so that needle 1 was in the hole formerly occupied by needle two. For the Macor sample, two measurements were made in the original orientation, and two in the reversed orientation for a total of four measurements.

The measured thermal conductivity ($\kappa_m$) and heat capacity ($C_m$) are related to the values derived from the fitting process by the following calibration functions:

$$\kappa_m = C_1 f_1 + C_2 f_2 + C_3 C_f 1 + C_4 C_f 2 + C_5$$  

[10]

and

$$C_m = C_6 f_1 + C_7 f_2 + C_8 C_f 1 + C_9 C_f 2 + C_{10}$$  

[11]

The calibration coefficients $C_1$….$C_{10}$ were determined during calibration by minimizing the error between actual values ($\kappa_a$ and $C_a$), and the measured values ($\kappa_m$ and $C_m$) determined in Eqs. 10 and 11. Values for $f_1$, $f_2$, $C_f 1$ and $C_f 2$, generated from the MSMCO-Marquardt fitting process for the 5 materials tested are shown in Table 2. The relationship between the measured values, derived from equations [10] and [11], and the actual values for the 5 known calibration materials results in the relations shown in Figures 8 and 9, which are the final conversion from the measured to actual $\kappa$ and $C_V$ values.

26
C. Dielectric

TECP dielectric measurements were taken in a series of four fluids with relative dielectric permittivities from 1 to 20 (Table 3). Each of the fluids was housed in a 30.4 cm diameter by 17.5 cm depth glass bowl, filled to within 1 cm of the top. Earlier finite element analysis and experimental testing showed that >95% of the electromagnetic field generated by the TECP dielectric measurement was contained within this volume for all calibration fluids. All measurements were performed in a drybox under dry N₂ purge; the TECP was lowered into the fluids with a translation stage. The TECP units were inserted into the fluid until the needles were submerged. For the air, isopropyl, and 1-decanol samples, the fluids were allowed to equilibrate to the temperature of the drybox for at least 1 hour prior to data collection. The tert-butanol sample was held at a steady, above-ambient temperature with an electrical heating pad due to its high melting temperature (301 K). In each fluid, data were collected from the TECP unit for 25 minutes to assure thermal equilibrium between the needles and the fluid. Thermal drift was not identified in any of the samples after one minute of equilibration time, so the TECP output values presented are the averages of approximately the final 24 minutes of data collection.

Prior to making TECP measurements in the fluids, six sub samples of each of the three measured alcohols were collected. The fluids were transferred to 100 ml, septa-equipped, airtight glass containers in the drybox under N₂ purge. The relative dielectric permittivity of the standards was independently determined at the National Institute of Standards and Technology’s Dielectric and Magnetic Material Laboratory. The NIST measurements were performed with two different parallel plate devices at the same measurement
frequency (6.25 MHz) and temperature as the TECP calibrations. Using the data in Table 3, the TECP calibration function was determined (Figure 10).

**D. Electrical Conductivity**

The electrical conductivity calibration was a two-step process. First, the TECP conductivity function was characterized with a set of fixed resistors. Second, aqueous standard solutions were used to find the probe constant vs. resistance. By combining these two functions, the electrical conductivity of the bulk medium surrounding the needles can be accurately determined for each of the three resistance channels (Table 4). The TECP flight unit was placed in the Delta Design chamber. Copper blocks machined at the same cone angle as the TECP needles were used to make electrical contact between the needles and two clip-type leads that were routed through mouseholes to the exterior of the chamber. The resistance of the leads was measured at 0.06 Ω, and has been neglected in the following analysis. The clip leads were used to sequentially connect the TECP needles across a series of 24 axial lead resistors ranging from 50 Ω to 250 MΩ. The resistance of each of the fixed resistors was measured just prior to each TECP measurement with a calibrated Agilent multimeter. To allow circuit settling, data were collected across each resistor for 120 seconds for each of the three measurement ranges. The average of the last 10 seconds of TECP output for each measurement range has been used in all subsequent analyses. This process was repeated at five different temperatures, spanning the range 160 K to 323 K.

Measurements were then taken in a series of six aqueous standard solutions ranging from 1 to 10,000 μS/cm. The standards were poured to 500 ml in a 600 ml Pyrex beaker. All experiments were performed in the dry box with the TECP lowered into the solutions
with the translation stage until the metal needles were fully submerged. The PEEK shoulders were not submerged in the solution. Measurements began with the 1μS/cm sample (DDI water), and progressed to the highest salt concentration sample. The TECP needles were wiped dry and wiped with an isopropyl alcohol clean room wipe between samples. Before measuring with the TECP, the conductivity of each solution was measured with a calibrated Orion multi meter. In the special case of the 1μS/cm solution, where a very small amount of salt contamination would affect the conductivity of the standard significantly, the TECP was first dipped into the sample and removed, then the conductivity was measured with the Orion meter, and then the TECP was placed back into the sample and a measurement was taken. The calibration again consisted of a 120 second measurement in each of the three measurement channels. Again, the average of the last 10 seconds of TECP output for each measurement range has been used in all subsequent analyses.

The data collected across the 24 resistors were used to construct a matrix of ADC output as a function of resistance and temperature for each channel. After each data set was abbreviated to its useful range (cf. Table 4) it was modeled as a 4\textsuperscript{th} order polynomial function of TECP output vs. ln resistance (Table 5).

Preliminary characterizations indicated that the TECP probe constant ($C_P$) varies as a function of the measured resistance at least for the highest conductivity channel. The known resistance vs. output functions determined above were used in conjunction with the measurements in standard solutions to determine the probe constant vs. resistance relationship. The probe constant is calculated by:
\[ C_P = \frac{10^6}{R\sigma} \]  

where \( C_P \) is the probe constant [cm], \( R \) is the measured resistance [\( \Omega \)], and \( \sigma \) is the electrical conductivity of the sample [\( \mu S/cm \)]. The probe constant vs. resistance functions are shown in Figure 11. Only the lowest conductivity solution (DDI water) fell within the useful resistance range of the low \( \sigma \) channel, so a cell constant vs. measured resistance relationship could not be determined for the low \( \sigma \) function. The single point cell constant has been applied to all low \( \sigma \) function data described here.

**E. \( H_2O \) Vapor**

Calibration of the TECP flight units for humidity was done at the University of Washington Mars Atmospheric Simulation Chamber (MASC) facility in March 2006. Over 50,000 humidity measurements at 5 to 30 second intervals were collected over a period of a week. The measurements covered a wide range of frost point temperatures (193 K – 263 K) and instrument temperatures (\( T_b \), 208 K to 303 K). Most of the data was collected under 500 [Pa] of CO\(_2\) (7 [mb]), flowing through the chamber at \(~200\) cm\(^3\) min\(^{-1}\) (STP). The absolute humidity was measured by two different NIST-traceable chilled mirror hygrometers (an EdgeTech DewPrime I and a Buck Research CR-1).

The air temperature in the chamber was typically a few degrees colder than \( T_b \), due to self-heating of the circuit board. This temperature difference, combined with the small anisothermality of the environmental chamber, prevented us from reaching higher relative humidities due to ice formation in the chamber. For example, the vapor pressure corresponding to \( R_H = 0.60 \) at -55°C is equal to the saturation vapor pressure at -58.9°C.
The ADC counts are related to $T_b$ and the relative humidity, $R_H$, via a quadratic equation

$$ADC = -1.5332R_H^2 + (963.34 + 14.705T_b)R_H + \left(2804.7 - 1.0373T_b - 6.9089 \times 10^{-3}T_b^2\right)$$  \[13\]

This relationship was obtained by a least-squares fit of nearly 12,000 data points using a 6 parameter quadratic function of relative humidity and temperature, and has a standard deviation of 4.5 ADC counts (Figure 12). Applying the calibration involves plugging the board temperature into equation 13, and solving for the root of the quadratic.

The sensor output voltage is measured with a 12-bit ADC, and has a minimum value (at $R_H = 0$) of around 2800 digital number (DN), and an apparently temperature-dependent maximum (at $R_H = 100\%$) of at least 3200 DN (the highest value we observed). At 213K the maximum value of the response function is only 2900, so this standard deviation represents a 5\% error. However at 233 K the increased range of response reduces the standard deviation to about 2\%.

**IV. Laboratory and Field Characterization**

Although the TECP meets the measurement requirements as verified in the lab, practicalities of field and spacecraft operation will affect the accuracy of the various measurements on the Martian surface. This section outlines some of those factors, and presents preliminary characterization data from the laboratory, spacecraft testbed activities, and field testing in the Antarctic Dry Valleys. Additional characterization work is underway to further improve science return from TECP.
A. Characterization

1. Needle temperature vs. regolith temperature

TECP can accurately measure the temperature of its individual needles. However, needle temperature may not be identical to the regolith or air temperature. If, for example, the thermal conductivity of the regolith is extremely low, the TECP needles may be equally coupled to the regolith and the TECP body.

To characterize this effect, an experiment was conducted using three different test materials: water stabilized with agar, dry silt loam soil, and EPS. In each instance, the material was placed in a thermal chamber and brought to a temperature below ambient. The sample was then removed from the temperature chamber, placed in an insulating container at ambient pressure, and the TECP needles were inserted into the sample. The needle temperature, $T_b$, and sample temperature were monitored until they equilibrated, and then recorded. This process was repeated at different sample temperatures ranging from well below to well above the TECP temperature.

It is apparent from the data that the error is linearly related to the temperature difference between the TECP body ($T_b$) and the needles – both measurable quantities. The slope of the linear relationship is, to first order, linearly related to the thermal conductivity of the medium in which the TECP needles are inserted. Fortunately, thermal properties are based on $\Delta T$ measurements, and are independent of sample temperature. So, the preliminary characterization data presented here indicate that regolith temperature can be derived from measured needle temperature, $T_b$, and the independently measured regolith thermal conductivity.
2. Imperfect needle insertion
In an idealized experiment, the TECP needles are fully seated in an infinite homogeneous half space, with no voids either at the sides or at the end. However, in practice, the TECP will be put into use at the end of the 2.3 m Robotic Arm, and the possibility of gaps exists. Can measurements be made with imperfect placement?

A series of experiments were conducted to understand the errors introduced into the various TECP measurements from: a) incomplete needle insertion; b) complete insertion with partial retraction, and; c) complete insertion with lateral movement. The objectives were accomplished by using a manual mill to precisely control the position of TECP in relation to the sample, and create voids of known dimension. The materials tested during the incomplete insertion portion of this exercise were glycerol, stabilized water, AETB-8, a thermal protection foam, dry clay soil, dry silt loam soil, and dry quartz sand. All of these samples with the exception of the glycerol and stabilized water were tested in the partial retraction and lateral void exercises. The results of these experiments are summarized in Figure 13 for the three soils, and the salient points are described below. The description of mitigation approaches against non-ideal insertions is deferred until the discussion of TECP-Robotic Arm Interoperability tests (below).

Electrical Properties

1. One can think of the overdrive data as an investigation of the effects of compression on the sample, the partial retraction as the effects of air gaps, and the lateral movement experiment as their combined effects.
2. Progressive linear immersion of the needles in the soil particles has a more or less linear effect on measured permittivity, but an exponential effect on electrical conductivity.

3. Compaction increases the measured permittivity linearly, and the measured conductivity exponentially.

4. The creation of voids around the needles decreases the permittivity linearly, but decreases the measured conductivity exponentially.

5. In lateral motions, the effects of compaction initially dominate the electrical properties, but beyond 0.25mm of lateral movement, the effects of the voids appears to dominate.

6. The sensitivity of the measured electrical properties to the placement of the needle is proportional to their permittivity and conductivity. It's useful to look at the actual measured quantities, to see how the effects of insertion gaps are proportional to the permittivity and conductivity.

**Thermal Properties**

1. Although the heat capacity measurement is nicely behaved, varying linearly with insertion, and being relatively insensitive to compaction, the thermal conductivity results appear to be complicated by the data reduction technique.

2. Compression of the soil by the TECP results in linear increases in thermal properties, at least up to 3 mm, due to the loss of pore voids in the vicinity of the needles.

3. Voids result in linear decrease in the measured heat capacity, but the effects on the derived thermal conductivity are difficult to deconvolute from the idiosyncracies of the mathematical modeling that must be done to extract $\kappa$. 
4. In lateral motions, the effects of compaction initially dominate both the electrical and thermal properties, but beyond 0.25 mm of lateral movement, the effects of the growing voids become increasingly important.

3. Wind

There are no requirements for TECP to measure wind velocity. However, application of a protocol similar to the thermal properties experiments has shown some promise in characterizing wind speeds, particularly in high-wind regimes where the wind-sock associated with the Meteorology package is not capable of measuring wind as well.

The procedure involves heating needle one for a period up to 70 seconds, and then following the cooling curve for another 90 seconds. Preliminary analysis of the heating and cooling data acquired at Mars-like pressures in a wind tunnel at the University of Michigan suggest that both peak temperatures and the cooling time constant

$$\tau_c = \frac{1}{\frac{dT}{dt}}$$

are sensitive to wind velocity. Characterization of the TECP response to wind velocity is ongoing, and the determination of whether or not to try wind velocity measurements on the Martian surface will be made once those analyses are complete.

B. Antarctic Dry Valley Field Testing

Field testing of a flight model unit, TECP010, were carried out in Beacon Valley, McMurdo Dry Valleys, Antarctica, during the 2007-2008 International Polar Year deployment.
1. Field Site and Data Collection

The test site in Beacon Valley (77° 50’ 24” S; 160° 39’ 27”E) is at an elevation of 1400 m. The annual average temperature is -22 °C. Snowmelt is confined to a layer ≤ 5 cm in depth, based upon observations of oxidized soil horizons and salt concentrations. Below this, H2O transport is restricted to thin films or vapor. At the time of the field observations, there was no snowcover, and the surface was visibly dry.

The surface materials are silica-rich sands from the surrounding Beacon Sandstone, along with clastic fragments of dolerite, which occurs as intrusive sills throughout the Beacon Sandstone. The near surface is very rich in salts, predominantly mirabilite (Na2SO4•10H2O), halite, and a number of minor phases. Rocks and boulders are glacial erratics with a wide variety of compositions. Additional analyses of the sampled material is pending.

During the austral summer, the sun does not set, so all measurements were conducted with the TECP, and the soil into which it was inserted, either in full sunlight, or shadow. Shadowing resulted from one of three sources. The TECP was placed near the north (afternoon sunlit) side of a pyramidal 30 x 19 x 17 cm rock. A Vaisala HM 34 Humidity and Temperature Sensor was placed next to it, and used periodically throughout data collection for comparison to the TECP humidity data.

During the “nighttime” hours, (2100h – 0400h), the sun illuminates the soil and face of the rock opposite the TECP (Figure 14, left). Between 0400h and 0800h, the entire site was shadowed by the steep valley walls. The sun emerged from behind the valley wall at ~0800h, and for several hours, fully illuminated the TECP, the rock face, and the soil being sampled. Between 1700 and 1730h, the shadow of the Science Tent, from which
the TECP was operated, covered first the soil, then the TECP body itself, and finally the rock.

Data collection spanned two periods, January 2, 2008 from ~2100h to 0400h, and the following day from 1540h to 1940h. Raw DN counts for each channel were converted to physical units via the functions published for TECP010 in the TECP Calibration Report. For the most part, the data set was nominal, and captured expected patterns in thermal and electrical properties. However, at least one anomalous and interesting event was recorded that merits explicit discussion.

2. Selected Results and Discussion

Throughout the initial period of data acquisition, the TECP and soil were shadowed, although the rock, and the soil opposite the TECP were continuously illuminated.

Needle temperatures decreased steadily until ~0300h. Beyond that, needle and board temperatures appear to be approximately stable, suggesting that the power dissipated by the TECP electronics is approximately balanced the heat loss of the TECP to the environment.

The most interesting part of the data are the electrical properties. The Bulk Relative Permittivity ($\varepsilon_b$) decreases along with temperature until ~0215h, and then begins a rapid increase (Figure 14, right). The total change from one time series to the next is greater than the variation within time series, suggesting that the effect is not due to noisy readings. The Electrical Conductivity data exhibit a somewhat different, but correlated response (not shown). Both the EC Low and EC Medium channels exhibit a sawtooth pattern; an abrupt drop in $\sigma$ correlates with the minimum in the $\varepsilon_b$ data, after which, $\sigma$ and $\varepsilon_b$ increase sharply throughout the remainder of the night.
This is a puzzling observation. At the temperature (265 K – 270 K) and the frequencies of the TECP measurement (1 kHz for \( \sigma \); 8 MHz for \( \varepsilon_b \)), both \( \sigma \) and \( \varepsilon_b \) are positively correlated with temperature in dry, rock-forming minerals (Oheloft, 1981, Buehler et al. 2003; Najipolo et al., 1998). The electrical properties change slope simultaneously, and approximately an hour before the cooling rate of the soil changed.

Since the measurements were acquired in sublimation till above ice-cemented permafrost (depth to ground ice at the site was ~25 cm), and the \( R_H \) sensor registered a non-zero and stable atmospheric relative humidity in spite of ambient cooling, it is reasonable to suppose that there existed a population of unfrozen water on the surfaces of the soil particles. Normally, however, the volumetric unfrozen water content (\( \theta \)) of soil in a closed system decreases as the temperature decreases, which would also be inconsistent with these observations.

It is possible however, to increase \( \varepsilon_b \) and \( \sigma \) in a soil by adding unfrozen H\(_2\)O molecules, (open system). The correlation between the unfrozen water content and dielectric is a well-known phenomenon; there is even a "universal" relationship, which describes the relationship between \( \varepsilon_b \) and \( \theta \) for a remarkably large variety of soil types (Topp et al. 1980)

\[
\varepsilon_b = 3.03 + 9.30 \theta + 1460 \theta^2 - 76.7 \theta^3
\]

This equation assumes that a change in \( \theta \) is the only variable affecting \( \varepsilon_b \). If applied to the data recorded here, it would suggest an increase in unfrozen water of 0.25% between 0215h and 0345h (from 4.15% to 4.4%).
The humidity data are stable, suggesting the atmosphere is not the source of an intriguing possibility, which may be consistent with the large $d^2 \varepsilon / dt^2$ at 0215h, is the arrival at the needles of additional H$_2$O molecules, migrating in thin films from the continuously warming far side of the rock. While this hypothesis requires further analysis, the unexpected relationships between thermal and electrical properties is suggestive of the type of events that might prove particularly exciting in data from the Martian surface. **C. PIT Testing**

Since the measured electrical and thermal properties are so sensitive to the details of needle insertion, characterization of the interoperability of the Robotic Arm with the TECP were conducted in the Payload Interoperability Testbed (PIT) at the University of Arizona, to assess the likely scope of gaps associated with non-ideal insertion.

The original goal was to conduct all TECP insertions along an angle of attack of $<0.08$ radians, which is the taper angle of the needles. However, this proved impracticable. Movement of the RA in the direction of the TECP needles ($z$-axis) results in bouncing by several mm during start and stop transients, with the principle motions directed along the $x$ and $z$ axes (see Figure 15 for an example, and the co-ordinate axes referenced in the discussion). In the cohesive Baghouse Dust used in the PIT, this motion leaves needle tracks in the soil, and effects the quality of the TECP insertion.

To establish an operational baseline for the performance of the RA/TECP operations, we first conducted a manual insertion of the TECP into the Baghouse Dust, in a manner as near to ideal as possible. Results for various mitigation strategies were compared against the results of the ideal insertion to assess possible strategies for mitigating gap formation.
The mitigating approach that proved most successful was to insert the TECP to the best of the RA capabilities, and then run the RASP for 15 seconds. The RASP is a small drill-like sampling device, included in the Robotic Arm Scoop in order to improve the chances of collecting very cold, and very hard, subsurface ice. In highly cohesive soils, the high-frequency, small-amplitude vibration caused by the RASP motor, coupled to the needles through the RA and TECP body, results in the collapse of any substantial voids surrounding the needles.

Of course, this technique does produce a alteration of the soil particulate structure from its original state, but the introduction of any measurement device will have the effect of disturbing the medium. Continued characterization efforts are underway to quantitatively assess the errors introduced by the non-ideal insertions, combined with effects of the RASP to collapse the resulting voids.

**D. RAC Images**

One means of characterizing the cleanliness of any given insertion is by imaging the TECP with the Robotic Arm Camera (RAC). In Figure 16, the RAC has imaged the TECP during insertion tests, as a function of the rotation angle of the TECP relative to the RA. The insertion angles range from 30° to 77° (nearly orthogonal to the RA). In some instances, the entry point of at least one needle can be seen. As the rotational angle gets smaller, it is harder to verify insertion and possible gaps with the RAC. Other things being equal, we prefer to make measurements in an orientation that permits verification of TECP placement with the RAC. In any event, the RAC will image any area sampled by TECP both prior to insertion and immediately after retraction.
V Surface Operations
Operations Planning for TECP is somewhat unique for the Phoenix mission, because TECP is an *in situ* measurement of the regolith properties, which change with the environmental variables. In addition, by trenching in the regolith, we will alter the physical structure of the regolith, and disturb some of the properties we intend to measure. Because we do not yet know the rate at which trenching operations will occur, nor the total depth to ground ice, nor the physical configuration of the trench, nor the availability of undisturbed materials in the workspace accessible to the RA, actual scheduling and targeting decisions will have to be made as part of the strategic and tactical processes. However, some considerations are described below that should guide planning for TECP measurements during the expected mission.

The physical and chemical activity of water is dependent upon temperature, and therefore, expected mission planning for characterization of regolith H_2O activity should focus on the local warmest period during operations.

Landing occurred May 25, 2008 which corresponds to L_s=76°, in the late northern spring. We can do the arithmetic very crudely and say there are 0.53842° L_s/Sol, so the approximate mapping of Mission Sol against L_s would indicate that summer solstice would occur around Sol 26 (Figure 17). This will be earlier than the local thermal maximum, and subsequent observations should be made approximately every 5° of L_s until after the local thermal maximum. Therefore, roughly, H_2O observation campaigns would also occur as follows:

L_s 95° = Sol 35

L_s 100° = Sol 44
The ideal would be to do an insertion into the soil during the warmest part of the sol, and monitor the temperature, electrical conductivity, dielectric permittivity and $R_{th}$, but not thermal properties (heating during thermal properties measurements will alter, and perhaps redistribute $H_2O$) on or about sols 26, 35, 44 and 55.

The desired insertion point, or points, cannot be determined in advance, because we do not know the configuration of the trench at each date, the specific shadowing associated with the lander, the depth to the ice table and so on. However, the $H_2O$ mobility campaigns should be conducted for an hour or two around the thermal maximum of a target selected during the tactical process in the sols leading up to the thermal campaign sols.

The temperature and thermal properties should be measured in at least one material near the beginning ($L_s \sim 80^\circ$), middle ($L_s \sim 105^\circ$) and near the end ($L_s \sim 125^\circ$) of the mission. These correspond roughly to sols 10, 50 and 90. The diurnal maximum surface temperature will change relatively little over the course of the mission, but the nighttime minima will vary considerably more (Figure 14). By targeting the beginning, middle, and end of the mission, we will be able to characterize the thermal properties of the accessible materials, as a function of temperature, over the greatest possible range.

The temperature and thermal properties measurements should be conducted in material that is minimally disturbed. If horizontal compositional layering is evident in the dry overburden, each layer should be characterized.
At least one of these thermal properties campaign days should be correlated with THEMIS observations of the Phoenix landing site, to characterize the relationship between in situ thermal properties measurements and those derived from remote sensing. Thus, a reasonable but modest regolith characterization campaign could be conducted by prioritizing TECP measurements on approximately 7 sols throughout the mission. The specific sols on which the H$_2$O and thermal properties are characterized can be decided tactically, but the rough guidelines given above will provide good coverage of temperature, and maximize the probability of catching any translational or rotational mobility in regolith H$_2$O molecules.

**VI Summary**

The TECP will provide an exciting opportunity to measure the thermal properties of the Martian regolith in situ, as well as to characterize the response of the regolith in the high latitudes to the extreme variations in boundary conditions, both for temperature and H$_2$O abundance, that are characteristic of the polar summer. In combination with other Phoenix instruments, it is hoped that the physical and chemical processes associated with energy and H$_2$O fluxes through the regolith can be quantitatively understood in the present climate, and that this knowledge will give us the capability to decode the geologic history of the Martian polar latitudes as uncovered by the Phoenix mission.

TECP is a thoroughly calibrated instrument, but there are practicalities of field and spacecraft measurements that must be taken into account. Mitigating procedures, such as using the Phoenix Robotic Arm RASP to collapse voids around the TECP needles are
described. TECP was field tested in the Antarctic Dry Valleys, and showed promise in
detecting small changes in unfrozen water under Antarctic conditions.
Acknowledgements
The authors would like to thank John Michael Morookian, Troy Hudson, Mike Janeczic of the NIST Dielectric and Magnetic Material Laboratory, Roger Tanner, Tom Squire, Matt Switzer, Matt Siegler, Richard Quinn, Martin Buehler, Marsha Presley, Lynne Cooper, Nilton Renno, Bob Bonitz, Mike Shirbacheh, Matt Schneider, Adam Bruckner, Carl Knowlen, and Troy Hudson.
References


Conley, W., Linear systems revisited, *Simulation,* 45, 15-18, 1985


Pozdnyakov, A.I., K.Y. Chan, The technique of the electrical sounding and profiling in soil investigations. (In Russian.) Vestnik, 1, 46-54, 1979


Table 1. Thermal properties standard materials used in the TECP calibration.

<table>
<thead>
<tr>
<th>Material</th>
<th>Thermal conductivity [W m(^{-1}) K(^{-1})]</th>
<th>Source</th>
<th>Heat capacity [kJ m(^{-3}) K(^{-1})]</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Macor (glass ceramic, 25 C)</td>
<td>1.402</td>
<td>Decagon</td>
<td>2010</td>
<td>Decagon</td>
</tr>
<tr>
<td>Water (stabilized, 25 C)</td>
<td>.608</td>
<td>CRC</td>
<td>4174</td>
<td>CRC</td>
</tr>
<tr>
<td>HDPE (23 C)</td>
<td>0.513</td>
<td>Decagon</td>
<td>1690</td>
<td>Decagon</td>
</tr>
<tr>
<td>Glycerol (20 C)</td>
<td>0.285</td>
<td>Ganic &amp; Hicks</td>
<td>2963</td>
<td>Ganic &amp; Hicks</td>
</tr>
<tr>
<td>Expanded polystyrene (EPS, 20 C)</td>
<td>0.033</td>
<td>Laser Comp</td>
<td>63</td>
<td>calculation</td>
</tr>
</tbody>
</table>

Table 2. Thermal conductivity (\(\kappa\)) and heat capacity (\(C_V\)) values measured at needle 1 and 2 with the TECP flight unit. The values of \(\kappa_m\) and \(C_{V,m}\) are generated from equations 13 and 14, and the known values of \(\kappa_a\) and \(C_{V,a}\) are taken from literature or other sources.

<table>
<thead>
<tr>
<th>Material</th>
<th>(\kappa_1)</th>
<th>(\kappa_2)</th>
<th>(C_V)</th>
<th>(C_m)</th>
<th>(C_{V,m})</th>
<th>(\kappa_a)</th>
<th>(C_{V,a})</th>
</tr>
</thead>
<tbody>
<tr>
<td>EPS</td>
<td>0.110</td>
<td>0.199</td>
<td>1.741</td>
<td>1.539</td>
<td>0.033</td>
<td>0.065</td>
<td>0.033</td>
</tr>
<tr>
<td>EPS 2</td>
<td>0.112</td>
<td>0.202</td>
<td>1.727</td>
<td>1.545</td>
<td>0.034</td>
<td>0.061</td>
<td>0.033</td>
</tr>
<tr>
<td>Glycerol</td>
<td>0.498</td>
<td>0.526</td>
<td>1.815</td>
<td>5.149</td>
<td>0.283</td>
<td>2.913</td>
<td>0.285</td>
</tr>
<tr>
<td>Glycerol 2</td>
<td>0.517</td>
<td>0.534</td>
<td>1.669</td>
<td>5.097</td>
<td>0.291</td>
<td>2.823</td>
<td>0.285</td>
</tr>
<tr>
<td>Water</td>
<td>0.936</td>
<td>1.071</td>
<td>1.077</td>
<td>7.232</td>
<td>0.595</td>
<td>4.296</td>
<td>0.606</td>
</tr>
<tr>
<td>Water 2</td>
<td>0.951</td>
<td>1.125</td>
<td>1.086</td>
<td>7.348</td>
<td>0.614</td>
<td>4.367</td>
<td>0.606</td>
</tr>
<tr>
<td>Water 3</td>
<td>0.942</td>
<td>1.035</td>
<td>1.131</td>
<td>6.728</td>
<td>0.602</td>
<td>4.097</td>
<td>0.606</td>
</tr>
<tr>
<td>Macor</td>
<td>1.629</td>
<td>2.846</td>
<td>0.409</td>
<td>3.095</td>
<td>1.391</td>
<td>2.008</td>
<td>1.401</td>
</tr>
<tr>
<td>Macor 2</td>
<td>1.636</td>
<td>2.832</td>
<td>0.401</td>
<td>3.149</td>
<td>1.392</td>
<td>2.059</td>
<td>1.401</td>
</tr>
<tr>
<td>Macor 3</td>
<td>1.682</td>
<td>2.947</td>
<td>0.351</td>
<td>3.085</td>
<td>1.441</td>
<td>2.024</td>
<td>1.401</td>
</tr>
<tr>
<td>Macor 4</td>
<td>1.634</td>
<td>2.904</td>
<td>0.377</td>
<td>3.151</td>
<td>1.404</td>
<td>1.986</td>
<td>1.401</td>
</tr>
<tr>
<td>HDPE</td>
<td>0.683</td>
<td>1.082</td>
<td>1.268</td>
<td>3.484</td>
<td>0.501</td>
<td>1.679</td>
<td>0.513</td>
</tr>
<tr>
<td>HDPE2</td>
<td>0.694</td>
<td>1.122</td>
<td>1.199</td>
<td>3.627</td>
<td>0.514</td>
<td>1.693</td>
<td>0.513</td>
</tr>
</tbody>
</table>
Table 3. The materials and results used in the TECP dielectric calibration

<table>
<thead>
<tr>
<th>Material</th>
<th>Nominal dielectric</th>
<th>Temperature (°C)</th>
<th>NIST Dielectric</th>
<th>Output TECP FM (ADC)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Air</td>
<td>1</td>
<td>23</td>
<td>1</td>
<td>642</td>
</tr>
<tr>
<td>1-decanol</td>
<td>7.5</td>
<td>24</td>
<td>8.07</td>
<td>1410</td>
</tr>
<tr>
<td>Tert butanol</td>
<td>10.9</td>
<td>32</td>
<td>11.22</td>
<td>2300</td>
</tr>
<tr>
<td>Isopropyl</td>
<td>18.3</td>
<td>23</td>
<td>19.76</td>
<td>4016</td>
</tr>
</tbody>
</table>

Table 4. Useful resistance ranges for each TECP measurement range

<table>
<thead>
<tr>
<th>Channel</th>
<th>minimum resistance</th>
<th>maximum resistance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hi Conductivity (ECH)</td>
<td>--</td>
<td>6 kΩ</td>
</tr>
<tr>
<td>Medium Conductivity</td>
<td>6 kΩ</td>
<td>1MΩ</td>
</tr>
<tr>
<td>Low Conductivity</td>
<td>156kΩ</td>
<td>--</td>
</tr>
</tbody>
</table>

Table 5. The coefficients for the 4th order polynomial fit of $\ln(R)$ to the ADC counts as a function of temperature.

<table>
<thead>
<tr>
<th>$T$ [K]</th>
<th>Hi range</th>
<th>Mid range</th>
<th>Low range</th>
</tr>
</thead>
<tbody>
<tr>
<td>160</td>
<td>$\ln (R) = -8.341 \times 10^{-14} ADC^4 + 8.237 \times 10^{-10} ADC^3 - 2.751 \times 10^{-06} ADC^2 + 4.743 \times 10^{-03} ADC + 3.136$</td>
<td>$\ln (R) = -3.248 \times 10^{-14} ADC^4 + 4.654 \times 10^{-10} ADC^3 - 1.922 \times 10^{-06} ADC^2 + 4.037 \times 10^{-03} ADC + 7.918$</td>
<td>$\ln (R) = 1.3236 \times 10^8 ADC^3 - 1.1648 \times 10^4 ADC^2 + 3.4326 \times 10^1 ADC + 323.41$</td>
</tr>
<tr>
<td>200</td>
<td>$\ln (R) = -7.892 \times 10^{-14} ADC^4 + 7.927 \times 10^{-10} ADC^3 - 2.691 \times 10^{-06} ADC^2 + 4.718 \times 10^{-03} ADC + 3.128$</td>
<td>$\ln (R) = -3.263 \times 10^{-14} ADC^4 + 4.605 \times 10^{-10} ADC^3 - 1.903 \times 10^{-06} ADC^2 + 4.019 \times 10^{-03} ADC + 7.905$</td>
<td>$\ln (R) = 1.2165 \times 10^8 ADC^3 - 1.0593 \times 10^4 ADC^2 + 3.0932 \times 10^1 ADC + 286.57$</td>
</tr>
<tr>
<td>240</td>
<td>$\ln (R) = -7.809 \times 10^{-14} ADC^4 + 7.858 \times 10^{-10} ADC^3 - 2.673 \times 10^{-06} ADC^2 + 4.699 \times 10^{-03} ADC + 3.136$</td>
<td>$\ln (R) = -3.128 \times 10^{-14} ADC^4 + 4.498 \times 10^{-10} ADC^3 - 1.877 \times 10^{-06} ADC^2 + 4.000 \times 10^{-03} ADC + 7.901$</td>
<td>$\ln (R) = -3.100 \times 10^{-14} ADC^4 + 4.478 \times 10^{-10} ADC^3 - 1.873 \times 10^{-06} ADC^2 + 3.998 \times 10^{-03} ADC + 7.901$</td>
</tr>
<tr>
<td>280</td>
<td>$\ln (R) = -7.742 \times 10^{-14} ADC^4 + 7.807 \times 10^{-10} ADC^3 - 2.661 \times 10^{-06} ADC^2 + 4.690 \times 10^{-03} ADC + 3.139$</td>
<td>$\ln (R) = -3.100 \times 10^{-14} ADC^4 + 4.478 \times 10^{-10} ADC^3 - 1.873 \times 10^{-06} ADC^2 + 3.998 \times 10^{-03} ADC + 7.901$</td>
<td>$\ln (R) = 1.9291 \times 10^8 ADC^3 - 1.7078 \times 10^4 ADC^2 + 5.0553 \times 10^1 ADC + 483.93$</td>
</tr>
<tr>
<td>323</td>
<td>$\ln (R) = -7.769 \times 10^{-14} ADC^4 + 7.820 \times 10^{-10} ADC^3 - 2.662 \times 10^{-06} ADC^2 + 4.689 \times 10^{-03} ADC + 3.141$</td>
<td>$\ln (R) = -3.100 \times 10^{-14} ADC^4 + 4.478 \times 10^{-10} ADC^3 - 1.873 \times 10^{-06} ADC^2 + 3.998 \times 10^{-03} ADC + 7.901$</td>
<td>$\ln (R) = 1.9291 \times 10^8 ADC^3 - 1.7078 \times 10^4 ADC^2 + 5.0553 \times 10^1 ADC + 483.93$</td>
</tr>
</tbody>
</table>
Figure 1. Geologic Map of the Phoenix Landing site, by K. D. Seelos (see Arvidson et al., This Issue), with examples of patterned ground seen in Hi-Rise imagery (all examples ~ 50 cm/pixel).
Figure 2. Photograph of the TECP instrument (top) and with the external cover removed to allow access to the electronics board (bottom). For each needle, the numerical designation and functionality is identified at right. The TECP is 118.76 mm in length. Its needles are 15 mm long.
**Figure 3**

The electrical circuits employed by TECP to measure electrical and thermal properties of the regolith and atmosphere.

**Figure 3.** The electrical circuits employed by TECP to measure electrical and thermal properties of the regolith and atmosphere.
Figure 4. The TECP mounted on the Robotic Arm. The Robotic Arm Camera (RAC) is at right, and can image either the scoop contents, or the TECP when it is rotated counterclockwise for use.
Figure 5. TECP FM board temperature sensor calibration.

\[ T_B = 0.082607\text{ADC} - 3.045 \]
\[ R = 0.99995 \]
Figure 6. The calibration setup for the thermocouple calibration.
Figure 7. The Seebeck Coefficient of the thermocouple wire used for the TECP FM.

\[ C_s = 1.052 \times 10^{-6} T^3 - 1.045 \times 10^{-3} T^2 + 0.4317T - 1.6204 \]

\[ R = 0.9999 \]
Figure 8. The volumetric heat capacity calibration curve. $C_{v,a}$ is the volumetric heat capacity of the laboratory standard materials, and $C_{v,m}$ is the measured heat capacity – which is actually derived not through direct measurement, but through fitting Eqs. 10 – 11 to the measured heating and cooling curves associated with heating one of the needles.
Figure 9. The thermal conductivity calibration curve. $\kappa_a$ is the thermal conductivity of the laboratory standard materials, and $\kappa_m$ is the measured thermal conductivity – which is actually derived not through direct measurement, but through fitting Eqs. 10-11 to the measured heating and cooling curves associated with heating one of the needles.
Figure 10. The dielectric permittivity calibration curve for the TECP flight unit.

\[ \varepsilon = 1.172 \times 10^9 \text{ADC}^3 - 8.528 \times 10^6 \text{ADC}^2 + 2.289 \times 10^2 \text{ADC} - 10.58 \]
Figure 11. The Probe Constant for the TECP as a function of the resistance, for each of the 3 channels of the EC function.
Figure 12. (A) Contours of the calibration function for the TECP humidity sensor at four different values of $T_b$. The dashed portions of each line indicate an extrapolation beyond the range of data used to derive the calibration function. (B) Scatterplot of the environmental conditions corresponding to each calibration data point.
Figure 13. Errors introduced by incomplete and excess insertion, partial withdrawal, and lateral gaps in percent error deviation from an ideal insertion (marked with crosshairs).
Figure 14. The TECP deployed in Beacon Valley (left). Needle 3 temperature and bulk dielectric permittivity recorded during the overnight hours of 2 January, 2008.
Figure 15. The TECP during interoperability testing with the Robotic Arm in the Payload Interoperability testbed. The axes refer to the TECP-centered coordinate system used by the RA. The needle holes are evidence of overshooting in the $z$ direction during a commanded move of $+1$ cm in the $z$ direction. Their shape reflects uncommanded movement of $\sim3.5$ mm along the $x$ axis.
Figure 16

Figure 16. The RAC view of the TECP at insertion angles of 30°, 47° and 77° (top to bottom).
Figure 17. The predicted surface temperature at 70°N (top). The arrow and vertical red lines indicate the nominal Phoenix mission duration. For comparison the lower plot is $T_{18}$, indicating approximate surface kinetic temperature. (From Kieffer and Titus, 2001) from TES. The horizontal white line indicates 70°N.