

METHODS

Mars Subsurface Ice Model (MSIM)

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Contents

1	1D Thermal Model for Mars	3
1.1	Semi-Implicit Scheme on Irregular Grid	3
1.1.1	Upper boundary condition: prescribed T	4
1.1.2	Upper boundary condition: Stefan-Boltzmann radiation law	5
1.1.3	Lower boundary condition	6
1.1.4	Validations	6
1.2	Other Model Components	8
1.2.1	Seasonal frost cover	8
1.2.2	Thermal properties of the ground	9
1.2.3	Incidence angle and orbit	10
1.2.4	Mars atmospheric extinction and sky irradiance	10
1.3	Surface Energy Balance for Planar Slope on Mars	12
1.3.1	Direct solar irradiance on slope	12
1.3.2	Terrain and sky irradiance for planar slope	12
	Bibliography	13
2	Diffusion of Water Vapor with Phase Transitions	15
2.1	Governing Equations	15
2.2	Discretizations	16
2.2.1	Possible discretizations of spatial derivatives	16
2.2.2	Discretization of time derivative	17
2.2.3	Complete scheme	17
2.2.4	Upper boundary condition	18
2.2.5	Lower boundary condition	18
2.3	Discussion	19
	Bibliography	19
3	Long-Term Ice Evolution: Equilibrium and Non-equilibrium Ice Tables	20
3.1	Equilibrium Ice Table on Mars	20
3.2	Asynchronous Model for Near-Surface Ice on Mars	21
	Bibliography	22

Preface

Companion to <https://github.com/nschorgh/MSIM/>

Cite source code or the description of methods as:

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The most recent release is usually behind the most recent version available on GitHub, but releases have DOIs that can be cited.

Part 1

1D Thermal Model for Mars

1-Dimensional Numerical Model of Thermal Conduction and Surface Energy Balance

The heat flow in the shallow subsurface is described by the heat equation:

$$\rho c \frac{\partial T}{\partial t} = \frac{\partial}{\partial z} \left(k \frac{\partial T}{\partial z} \right) \quad (1.1)$$

where $T(z, t)$ is temperature, t time, z depth, ρc the volumetric heat capacity, and k thermal conductivity. The (negative of the) heat flux is $F = k \frac{\partial T}{\partial z}$. Boundary conditions are specified below.

1.1 Semi-Implicit Scheme on Irregular Grid

Authors & History: originally implemented by Samar Khatiwala in 2001 (including upper radiation boundary condition for semi-implicit scheme); extended to variable thermal properties and irregular grid by Norbert Schörghofer 2002–2003; added predictor-corrector step in 2019

Consider grid points at depths z_1, \dots, z_N in a direction normal to the surface, with z_1 the first point below the surface. A flux-conservative discretization on an irregularly-spaced grid is given by

$$\frac{\partial}{\partial z} F_j = \frac{F_{j+\frac{1}{2}} - F_{j-\frac{1}{2}}}{(z_{j+1} - z_{j-1})/2} = 2 \frac{k_{j+\frac{1}{2}} \frac{T_{j+1} - T_j}{z_{j+1} - z_j} - k_{j-\frac{1}{2}} \frac{T_j - T_{j-1}}{z_j - z_{j-1}}}{z_{j+1} - z_{j-1}}$$

Subscript j refers to position z_j . The spatial discretization of the heat equation (1.1) then becomes

$$\begin{aligned} (\rho c)_j \frac{\partial T_j}{\partial t} = & \frac{2k_{j+\frac{1}{2}}}{(z_{j+1} - z_j)(z_{j+1} - z_{j-1})} T_{j+1} - \frac{2}{z_{j+1} - z_{j-1}} \left(\frac{k_{j+\frac{1}{2}}}{z_{j+1} - z_j} + \frac{k_{j-\frac{1}{2}}}{z_j - z_{j-1}} \right) T_j + \\ & + \frac{2k_{j-\frac{1}{2}}}{(z_j - z_{j-1})(z_{j+1} - z_{j-1})} T_{j-1} \end{aligned}$$

Introduce the coefficients

$$\alpha_j = \frac{\Delta t}{(\rho c)_j} \frac{k_{j+\frac{1}{2}}}{(z_{j+1} - z_j)(z_{j+1} - z_{j-1})} \quad \text{and} \quad \gamma_j = \frac{\Delta t}{(\rho c)_j} \frac{k_{j-\frac{1}{2}}}{(z_j - z_{j-1})(z_{j+1} - z_{j-1})} \quad (1.2)$$

The discretized system of equations then becomes

$$\Delta t \frac{\partial T_j}{\partial t} = 2\alpha_j T_{j+1} - 2(\alpha_j + \gamma_j) T_j + 2\gamma_j T_{j-1}$$

A semi-implicit time discretization of (1.1) is of the form (Crank and Nicolson, 1947; Press et al., 1992)

$$(\rho c)_j \frac{T_j^{n+1} - T_j^n}{\Delta t} = \frac{1}{2} \left(\frac{\partial}{\partial z} F_j^{n+1} + \frac{\partial}{\partial z} F_j^n \right)$$

where superscript n refers to the time step. Hence,

$$T_j^{n+1} - T_j^n = \alpha_j T_{j+1}^{n+1} - (\alpha_j + \gamma_j) T_j^{n+1} + \gamma_j T_{j-1}^{n+1} + \alpha_j T_{j+1}^n - (\alpha_j + \gamma_j) T_j^n + \gamma_j T_{j-1}^n$$

which leads to the system of equations

$$\boxed{-\alpha_j T_{j+1}^{n+1} + (1 + \alpha_j + \gamma_j) T_j^{n+1} - \gamma_j T_{j-1}^{n+1} = \alpha_j T_{j+1}^n + (1 - \alpha_j - \gamma_j) T_j^n + \gamma_j T_{j-1}^n} \quad 1 < j < N \quad (1.3)$$

This tridiagonal linear system can be solved in $O(N)$ steps.

Whereas the temperature T_j is defined on grid point z_j , the conductivity k is defined in between points. In the equations above, $(\rho c)_j$ is defined on z_j , but in the program implementations, $2(\rho c)_j = (\rho c)_{j+\frac{1}{2}} + (\rho c)_{j-\frac{1}{2}}$. In this way, the thermal properties k and ρc are defined on the same points. (In the case of an interface between two layers with greatly different thermal properties, a grid point can be placed on the interface whereas the thermal properties do not need to be defined on the interface.) Since array indices must be integers, we choose $\mathbf{k}[j] = k_{j-\frac{1}{2}}$, and the same for ρc .

Although the derivation was made with time-constant thermal parameters k and ρc , it remains applicable if these parameters change slowly with time.

1.1.1 Upper boundary condition: prescribed T

Take as boundary condition a prescribed surface temperature $T_s = T(0, t)$. The general formulas (1.2) and (1.3) with $T_0 = T_s$ and $z_0 = 0$ yield

$$\alpha_1 = \frac{\Delta t}{(\rho c)_1} \frac{k_{3/2}}{(z_2 - z_1)z_2} \quad \text{and} \quad \gamma_1 = \frac{\Delta t}{(\rho c)_1} \frac{k_{1/2}}{z_1 z_2}$$

$$-\alpha_1 T_2^{n+1} + (1 + \alpha_1 + \gamma_1) T_1^{n+1} = \alpha_1 T_2^n + (1 - \alpha_1 - \gamma_1) T_1^n + \gamma_1 (T_s^n + T_s^{n+1}) \quad (1.4)$$

which is implemented in `conductionT.f90`. This is a standard Crank-Nicolson solver for an irregular spaced grid.

1.1.2 Upper boundary condition: Stefan-Boltzmann radiation law

The surface energy balance on an airless body is given by

$$Q + k \left. \frac{\partial T}{\partial z} \right|_{z=0} = \epsilon \sigma T^4 \Big|_{z=0} \quad (1.5)$$

$Q(t)$ is the absorbed solar flux. On the right-hand side, ϵ is the (infrared) emissivity of the surface and σ is the Stefan-Boltzmann constant. To use an implicit method, the nonlinear boundary condition needs to be linearized.

Introduce the auxiliary quantity T_0 , such that the surface temperature $T_s = (T_0 + T_1)/2$. On the surface,

$$\left. \frac{\partial T}{\partial z} \right|_{z=0} = \frac{T_1 - T_0}{\Delta z} \quad \text{and} \quad T^4 \Big|_{z=0} = \left(\frac{T_0 + T_1}{2} \right)^4 \quad \text{with} \quad \Delta z = 2z_1$$

The temperature is linearized around a reference temperature T_r , $T = T_r + T'$. Equation (1.5) becomes

$$\begin{aligned} Q + k_{1/2} \frac{T_1 - T_0}{\Delta z} &= \epsilon \sigma \left(\frac{2T_r + T'_0 + T'_1}{2} \right)^4 \\ &\approx \epsilon \sigma T_r^4 + 2\epsilon \sigma T_r^3 (T'_0 + T'_1) = -3\epsilon \sigma T_r^4 + 2\epsilon \sigma T_r^3 (T_0 + T_1) \end{aligned}$$

This results in the following expression for T_0 :

$$T_0 \left(\frac{k_{1/2}}{\Delta z} + B(T_r) \right) = Q + 3\epsilon \sigma T_r^4 + T_1 \left(\frac{k_{1/2}}{\Delta z} - B(T_r) \right) \quad \text{where} \quad B(T_r) = 2\epsilon \sigma T_r^3$$

Introduce $a = (Q + 3\epsilon \sigma T_r^4) / \left(\frac{k}{\Delta z} + B \right)$ and $b = \left(\frac{k_{1/2}}{\Delta z} - B \right) / \left(\frac{k_{1/2}}{\Delta z} + B \right)$. The relation for $j = 1$ in (1.3) becomes

$$-\alpha_1 T_2^{n+1} + (1 + \alpha_1 + \gamma_1 - \gamma_1 b^{n+1}) T_1^{n+1} = \alpha_1 T_2^n + (1 - \alpha_1 - \gamma_1 + \gamma_1 b^n) T_1^n + \gamma_1 (a^n + a^{n+1}) \quad (1.6)$$

Define $\beta = \frac{\Delta t}{(\rho c)_1} \frac{1}{2\Delta z^2}$, then $\alpha_1 = \beta k_{3/2}$ and $\gamma_1 = \beta k_{1/2}$. The surface temperature is computed as

$$T_s = \frac{1}{2}(T_0 + T_1) = \frac{1}{2}(a + bT_1 + T_1)$$

As reference temperature choose $T_r = T_s^n$. The semi-implicit solver with this boundary condition is implemented in `conductionQ.f90`.

The uppermost layer is a half-layer, and the first few grid points must be chosen as $z_0 = 0$ and $z_2 = 3z_1$ (in other words $z_1 = \Delta z/2$, $z_2 = z_1 + \Delta z$). The coefficients for the nonlinear upper boundary condition are designed for that. Subroutine `setgrid` can be used to generate a suitable grid.

Optional treatment of the upper boundary helpful in some situations: The above linearization of the Stefan-Boltzmann law, σT^4 , works well as long as the surface temperature changes slowly, such as for Mars orbit and a horizontal surface. A temporary instability,

where the surface temperature overshoots, can occur when a sloped, shadowed surface suddenly emerges into sunlight. In this case the energy input changes abruptly.

Above, the thermal emission is linearized around the reference temperature $T_r = T_s^n$. If T_s^{n+1} is far from T_s^n , a significant error was incurred in the evaluation of the emitted energy. This can be addressed by repeating the calculation with a new reference temperature T_r somewhere in between T_s^n and T_s^{n+1} . An empirical choice is the geometric mean of the previous reference temperature and the new surface temperature. This predictor-corrector step is iteratively applied until T_r is within 20% of T_s^{n+1} .

Another approach is “artificial flux smoothing” where the time step is subdivided into many substeps, using linear interpolation of the incoming flux from Q^n to Q^{n+1} . It turns a discontinuous change in Q into a continuous change. However, this approach does not identify a nascent instability or answer how many substeps are required to cure it.

1.1.3 Lower boundary condition

At the lower boundary of the domain, a known value is imposed on the heat flux. For the chosen spatial discretization, the flux is given by

$$F_{N+\frac{1}{2}} = k_{N+\frac{1}{2}} \frac{T_{N+1} - T_N}{z_{N+1} - z_N} = F_{\text{geothermal}}$$

The flux boundary condition is offset by half a layer, but considering how slowly the geothermal heat flux varies with depth, this is accurate. The condition amounts to

$$T_{N+1} = T_N + F_{\text{geothermal}} \frac{z_{N+1} - z_N}{k_{N+\frac{1}{2}}}$$

Assume the position of the hypothetical next grid point is set by $z_{N+1} - z_N = z_N - z_{N-1}$ and $k_{N+\frac{1}{2}} = k_{N-\frac{1}{2}}$. In the system (1.3) the equation for $j = N$ becomes

$$(1 + \gamma_N)T_N^{n+1} - \gamma_N T_{N-1}^{n+1} = (1 - \gamma_N)T_N^n + \gamma_N T_{N-1}^n + \frac{\Delta t}{(\rho c)_N} \frac{F_{\text{geothermal}}}{z_N - z_{N-1}} \quad (1.7)$$

with

$$\gamma_N = \frac{\Delta t}{(\rho c)_N} \frac{k_{N-\frac{1}{2}}}{2(z_N - z_{N-1})^2}$$

1.1.4 Validations

The following tests were performed for these solvers:

a) For a sinusoidally varying surface temperature, the solution to the heat equation is known analytically:

$$T = T_m + T_a e^{-z/\delta} \sin\left(\frac{z}{\delta} - \frac{2\pi t}{P}\right) \quad (1.8)$$

where

$$\delta = \frac{\Gamma}{\rho c} \sqrt{\frac{P}{\pi}} \quad (1.9)$$

is the thermal skin depth, Γ the thermal inertia, and P the period. This expression can be used to validate `conductionT` for uniform thermal properties (Figure 1.1). The heat flux is given by

$$F = -k \frac{\partial T}{\partial z} = -\sqrt{2} k \frac{T_a}{\delta} e^{-z/\delta} \cos\left(\frac{z}{\delta} - \frac{2\pi t}{P} + \frac{\pi}{4}\right)$$

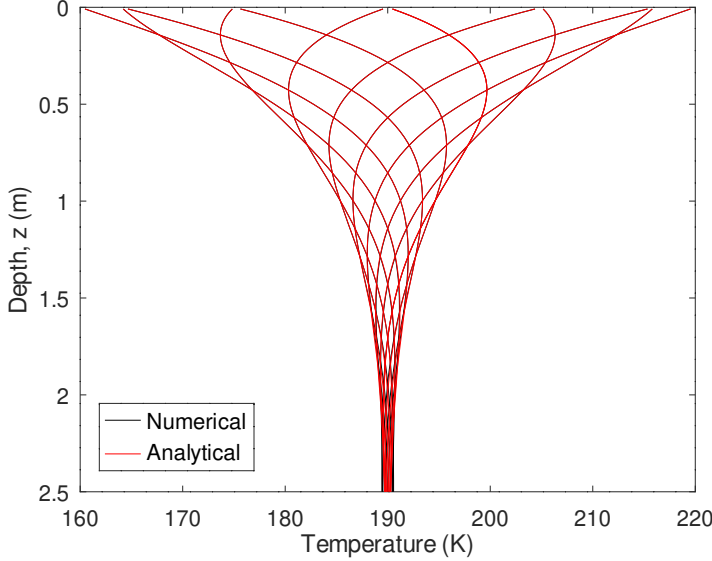


Figure 1.1: Comparison of numerical with analytical solution for Crank-Nicolson solver with periodic surface boundary condition. The deviations at the bottom are justified because the analytical solution (1.8) is for an infinitely deep domain. Non-equidistant grid points were used in this example.

b) Convergence of some solutions with Δt and Δz has been verified. For example, Figure 1.2 demonstrates that the error decreases proportionally to Δt^2 , as expected for a semi-implicit scheme. (An explicit or fully implicit scheme would only converge with Δt to the first power.)

c) For periodic solutions (with boundary condition 1.1.1 or 1.1.2) the heat flux $F = -k\partial T/\partial z$, time-averaged over one period, must be the same at all depths and equal to the heat flux imposed at the bottom boundary. Consider the time average of eq. (1.1) over one period. After the solution has equilibrated (has become periodic) and as long as the heat capacity does not vary with time, the time average of F must be constant with depth, even if the thermal properties vary with depth. Figure 1.3 shows a flux conservation test.

d) A short-term solution for the heat equation with Stefan-Boltzmann radiation surface boundary condition was derived by Handelsman and Olmstead (1972). Their non-dimensional equations are $T_t = T_{zz}$, $T_z(0, t) = T^n(0, t) - f(t)$, $T(z, 0) = 0$, $\lim_{z \rightarrow \infty} T(z, t) = 0$. In our case $n = 4$ and $f(t) = T_e^4$, where T_e is an ambient temperature. In this case, their solution is $T(0, t) = \frac{2}{\sqrt{\pi}} T_e^4 \sqrt{t}$, for small t . After re-dimensionalizing, the surface temperature is found to change as

$$T(0, t) = T_0 + \frac{2}{\sqrt{\pi}} \frac{\epsilon \sigma}{\Gamma} (T_e^4 - T_0^4) \sqrt{t} \quad \text{for small } t \geq 0 \quad (1.10)$$

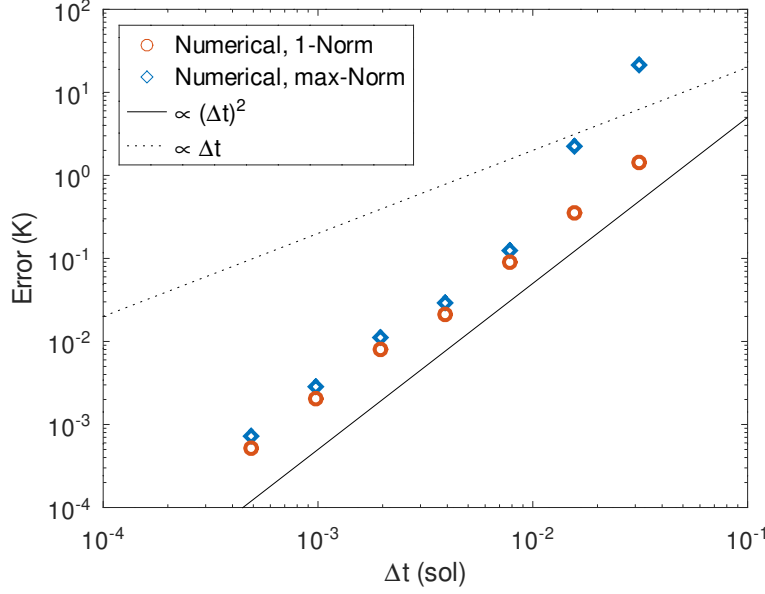


Figure 1.2: Convergence with time step Δt for the Crank-Nicolson method with nonlinear boundary condition. Errors are evaluated as $\|T_{\Delta t}(z, t) - T_{\Delta t/2}(z, t)\|$, where the subscript indicates the time step. The rate of convergence is second order. The black lines have slopes 1 and 2, respectively, and arbitrary prefactors.

where $T_0 = T(0, 0)$ is the initial surface temperature. Figure 1.4 shows that the numerical solver reproduces the expected behavior for this discontinuous change in incoming flux.

1.2 Other Model Components

1.2.1 Seasonal frost cover

On Mars, atmospheric CO_2 can condense (desublimates) seasonally. The surface energy balance with the latent heat of CO_2 sublimation added is

$$Q + k \left. \frac{\partial T}{\partial z} \right|_{z=0} = \epsilon \sigma T^4 \Big|_{z=0} + L \frac{dm_{\text{CO}_2}}{dt} \quad \text{with} \quad m_{\text{CO}_2} \geq 0 \quad (1.11)$$

where L is the specific latent heat and m_{CO_2} is the areal density of CO_2 ice. Time integration has to switch between boundary conditions (1.1.1) and (1.1.2). Call `conductionQ` if the surface temperature T_s is above the CO_2 frost point temperature or if $m_{\text{CO}_2} = 0$. Call `conductionT` if T_s is below the CO_2 frost point or if $m_{\text{CO}_2} > 0$. In the latter case, calculate the energy difference and update m_{CO_2} . Adjust the surface albedo and infrared emissivity. Repeat this at every time step. An implementation of the 1D thermal model for Mars is provided by `mars_thermal1d.f`.

Modeled surface temperatures have been compared to TES (Thermal Emission Spectrometer) surface temperatures.

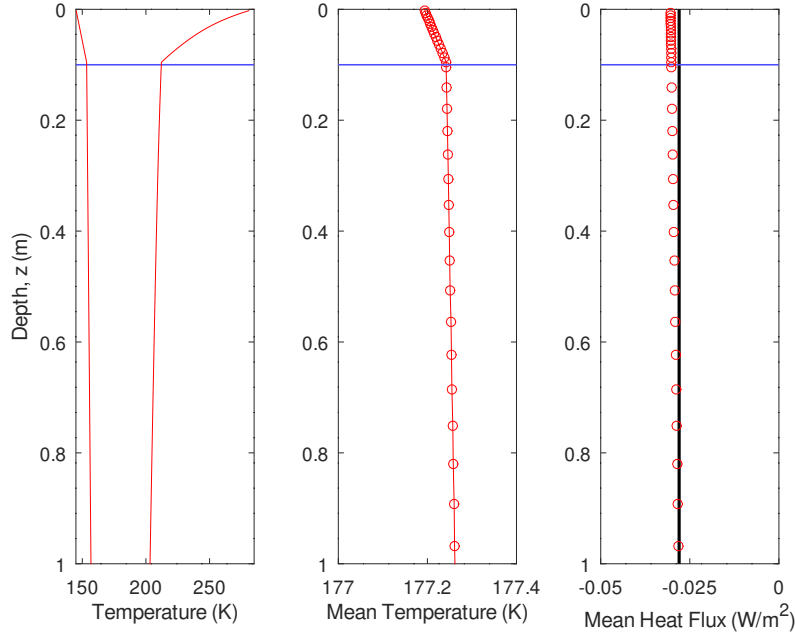


Figure 1.3: Validation of the conservation of the heat flux (flux-conservative discretization). The ice table at 10 cm depth causes dramatic changes in thermal properties. Left panel: Minimum and maximum subsurface temperatures over one Mars year. Middle panel: Temperatures averaged over one Mars year, which change linearly as the thermal conductivity is constant within each of the two layers. Right panel: Heat flux averaged over one Mars year, which is preserved across changes in thermal properties and equals to the heat flux imposed at the bottom boundary of 0.028 W/m^2 .

The Mars thermal model was used in Schorghofer and Aharonson (2005) and many subsequent papers, including Schorghofer (2008).

1.2.2 Thermal properties of the ground

See Winter and Saari (1969) for heat capacity of silicates as a function of temperature. Notably, many silicates have about the same specific heat capacity. See Handbook of Chemistry and Physics (Lide, 2003) for the temperature dependence of ice. At around 200 Kelvin: $c_{\text{ice}} \approx 1540 \text{ J/(kg K)}$, $\rho_{\text{ice}} \approx 927 \text{ kg/m}^3$, $k_{\text{ice}} \approx 3.2 \text{ W/(m K)}$, which results in a thermal inertia of $2137 \text{ J m}^{-2} \text{ K}^{-1} \text{ s}^{-1/2}$. The thermal conductivity of granular media varies by four orders of magnitudes on planetary surfaces, primarily due to the dependence on grain size.

Influence of ice on thermal properties: Ice can greatly change the thermal properties of porous ground. The following is one possible parametrization. In retrospective, it agrees well with the laboratory measurements by Siegler et al. (2012) for vapor-deposited ice.

$$\begin{aligned} \rho c &= (1 - \epsilon) \rho_{\text{regolith}} c_{\text{regolith}} + \epsilon f \rho_{\text{ice}} c_{\text{ice}} \\ k &= (1 - \epsilon) k_{\text{regolith}} + \epsilon f k_{\text{ice}} + (1 - f) \epsilon k_{\text{air}} \end{aligned}$$

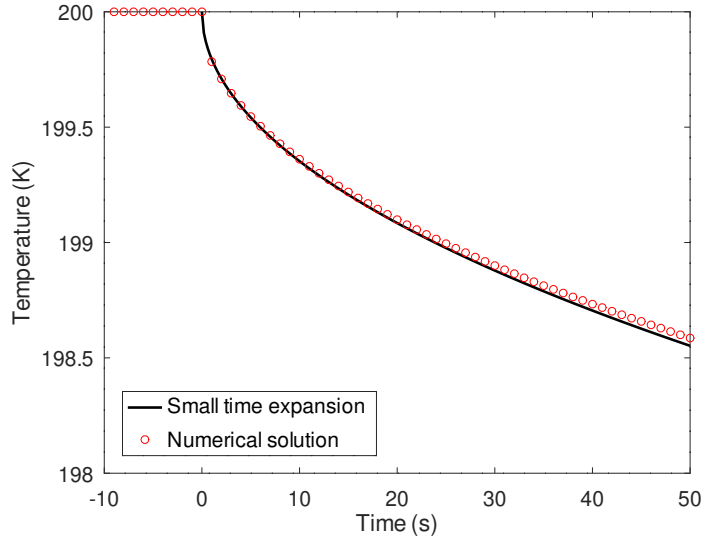


Figure 1.4: Response of numerical solution to a sudden change in incoming flux compared to the analytically obtained expansion for small times, eq. (1.10).

where ρ is density, c heat capacity, k thermal conductivity, ϵ porosity (void space / total volume), and f the ice filling fraction (between 0 and 1, but it can be larger than 1 for excess ice).

In the program, k and ρc are defined halfway between grid points, whereas f and T are defined on grid points.

The thermal inertia of ice-cemented soil is not drastically different from pure ice, which lends some justification to calculating an equilibrium ice table agnostic to the lithic content of the ice.

1.2.3 Incidence angle and orbit

The elevation β of the sun above an horizontal horizon is given in terms of geographic latitude λ , declination δ of the sun, and the hour angle h by

$$\sin \beta = \cos \lambda \cos \delta \cos h + \sin \lambda \sin \delta. \quad (1.12)$$

Equations for the ephemerides of present-day Mars are implented in `marsorbit.f90`. For past orbital elements, `generalorbit.f` can be used, which solves Kepler's equation with the Newton method.

1.2.4 Mars atmospheric extinction and sky irradiance

The absorbed direct solar irradiance is approximately

$$Q_{\text{solar}} = \frac{S_0}{R^2} (1 - A)(1 - f)^{1/\max(\sin \beta, 0.04)} \sin \beta, \quad (1.13)$$

where S_0 is the solar constant, R the distance from the sun in AU, A the albedo, and f due to the extinction in the atmosphere. The length of the path through the atmosphere is approximately proportional to $1/\sin \beta$ and the transmission is taken to be exponential in this path length. The nadir optical depth of the atmosphere is $-\ln(1-f) \approx f$. For small extinction and away from the horizon, $(1-f)^{1/\sin \beta} \approx 1-f/\sin \beta$. The maximum atmospheric path length ℓ_{\max} is limited due to the curvature of the planet, $H/\ell_{\max} \approx \sqrt{H/2R} \approx 0.04$, where H is the scale height of the atmosphere and R the radius of the planet.

In addition to the direct insolation, the diffuse irradiance from the sky contributes to the surface energy balance (sky irradiance). Atmospheric emission is approximated by a fraction f_{IR} (typically 2–4%) of noontime insolation and is kept constant throughout a solar day (Kieffer et al., 1977):

$$Q_{\text{atm,IR}} = f_{\text{IR}} \frac{S_0}{R^2} \sin \beta_{\text{noon}} \quad (\text{all day}) \quad (1.14)$$

This approximation fails during polar winter; in this case, Kieffer et al. (1977) replaces the noontime insolation with the surface frost emission.

In addition, there is scattered light when $\sin \beta > 0$, which is approximated by

$$Q_{\text{atm,scat}} = \frac{1}{2} f_{\text{scat}} \frac{S_0}{R^2}. \quad (1.15)$$

Half of the scattered light is assumed to be lost to space.

The total absorbed flux is

$$Q = Q_{\text{solar}} + (1 - A)Q_{\text{atm,scat}} + \epsilon Q_{\text{atm,IR}}$$

For the purpose of discussion, we determine the total energy budget of the atmosphere for a horizontal land mass. To first order $(1-f)^{1/\sin \beta} \sin \beta \approx \sin \beta - f$, in eq. (1.13), so at any time the sun is above the horizon the energy absorbed and scattered in the atmosphere is approximately $(S_0/R^2)f$. Over a solar day

$$\frac{S_0}{R^2} f \int_{\text{daytime}} dh \approx \frac{S_0}{R^2} \pi f. \quad (1.16)$$

The infrared emission from the atmosphere over the same time period is

$$2\pi f_{\text{IR}} \frac{S_0}{R^2} \sin \beta_{\text{noon}} \quad (1.17)$$

and the scattered energy

$$\pi f_{\text{scat}} \frac{S_0}{R^2}. \quad (1.18)$$

The globally averaged $\sin \beta_{\text{noon}}$ is

$$\frac{1}{\pi} \int_{-\pi/2}^{\pi/2} d\lambda \cos \lambda \sin \beta_{\text{noon}} \approx \frac{1}{\pi} \int_{-\pi/2}^{\pi/2} d\lambda \cos^2 \lambda = \frac{1}{2} \quad (1.19)$$

Global balance, (1.16)=(1.17)+(1.18), is achieved with $f = f_{\text{IR}} + f_{\text{scat}}$. This relation does not hold at an individual latitude, but it does hold globally.

1.3 Surface Energy Balance for Planar Slope on Mars

Planar slopes are much simpler than the general 3D problem. This section describes a model for the thermal balance on a tilted plane in the form of two coupled 1D thermal models plus a quasi-0D atmosphere. The model is for Mars, but easily simplified to airless bodies. It was used in Aharonson and Schorghofer (2006) and Schorghofer and Edgett (2006).

History: developed 2002–2005

1.3.1 Direct solar irradiance on slope

The elevation β of the sun above a horizontal horizon is given by (1.12). The angle θ of the sun above a sloped surface is

$$\sin \theta = \cos \alpha \sin \beta - \sin \alpha \cos \beta \cos(\Delta a) \quad (1.20)$$

where α is the slope angle and Δa is the difference between the azimuth of the sun and the azimuth of the topographic gradient. The sun is assumed to be below the horizon if either $\sin \beta < 0$ (horizontal horizon at infinity) or $\sin \theta < 0$ (self shadowing of slope).

On an airless body the direct insolation is

$$Q_{\text{direct}} = \frac{S_0}{R^2} \sin \theta \quad (1.21)$$

where S_0 is the solar constant and R the distance from the sun in AU.

For Mars, the direct solar insolation is

$$Q_{\text{direct}} = \frac{S_0}{R^2} (1 - f)^{1/\max(\sin \beta, 0.04)} \sin \theta \quad (1.22)$$

where f arises from the extinction in the atmosphere. This is a generalization of eq. (1.13) in subsection 1.2.4, where this approximate expression is further justified.

1.3.2 Terrain and sky irradiance for planar slope

Let α denote the slope angle. The heat balance on the surface is given by (1.11) with

$$Q = (1 - A)[Q_{\text{direct}}(\alpha) + Q_{\text{atm,scat}}(\alpha) + Q_{\text{land,vis}}(\alpha)] + \epsilon[Q_{\text{atm,IR}}(\alpha) + Q_{\text{land,IR}}(\alpha)] \quad (1.23)$$

Q is the absorbed irradiance from the sun, atmosphere, and surfaces within field of view. Figure 1.5 illustrates the contributions.

Terrain irradiance: The surface reemits radiation in all directions, but receives additional energy from surfaces in its field of view (terrain irradiance). This emission is weighted according to the incidence angle ι (Greek letter iota) and integrated over the spherical angle Ω subtended by the visible land surfaces. If we consider a horizontal surface at uniform temperature T_1 (Kreslavsky and Head, 2005):

$$Q_{\text{land,IR}} = \epsilon_1 \sigma T_1^4 \int \cos \iota d\Omega = \sin^2 \left(\frac{\alpha}{2} \right) \epsilon_1 \sigma T_1^4 \quad (1.24)$$

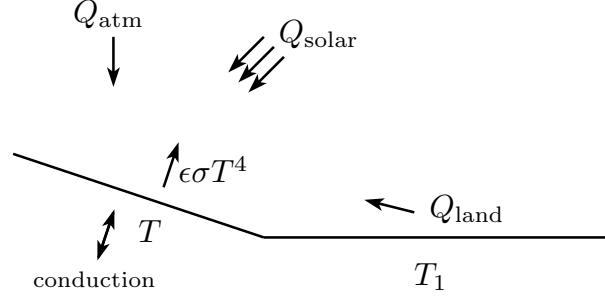


Figure 1.5: Contributions to the energy balance on a slope with surface temperature T .

If one assumes $T_1 = T$ and $\epsilon_1 = \epsilon$, then this term can be brought to the right-hand side of eq. (1.11), leading to an effective emissivity of $\epsilon \cos^2(\alpha/2)$. However, this is often not a good approximation, as demonstrated in Aharonson and Schorghofer (2006), Fig. 2b. It is more accurate to base T_1 on a separate 1D model for a flat unobstructed surface.

The terrain irradiance also has a short-wavelength component, usually much smaller than the long-wavelength (infrared) component. It uses the same geometric factor. The short-wavelength terrain irradiance onto a planar slope is

$$Q_{\text{land,vis}} = \sin^2\left(\frac{\alpha}{2}\right) A_1 Q_{\text{direct},1} \quad (1.25)$$

where subscript 1 denotes quantities on the flat surface.

Sky irradiance for planar slope. The diffuse irradiance from the sky adds a small amount of energy to the surface. In the spirit of the Kieffer approximation (subsection 1.2.4),

$$Q_{\text{atm,IR}} = \frac{S_0}{R^2} F_{\text{sky}} f_{\text{IR}} \sin \beta_{\text{noon}} \quad (\text{all day}) \quad (1.26)$$

and if the sun is up, then

$$Q_{\text{atm,scat}} = \frac{S_0}{2R^2} F_{\text{sky}} f_{\text{scat}} \quad \text{when } \sin \beta > 0 \quad (1.27)$$

otherwise $Q_{\text{atm,scat}} = 0$.

More than one definition of (sky) view factor F_{sky} and various approximations for the sky irradiance are used in the literature; see Flo Heggem et al. (2001) and Rakovec and Zakšek (2012). For isotropic (Lambertian) irradiance from the sky, the diffuse irradiance from the atmosphere is weighed by the cosine of the incidence angle. For a planar slope this results in $F_{\text{sky}} = \cos^2(\alpha/2)$. Spiga and Forget (2008) have derived a more detailed parameterization for $Q_{\text{atm,scat}}$ on a planar slope on Mars.

For a 3D horizon, the sky view factor can be calculated from horizon elevations. The detailed expression is given in the User Guide of the Planetary-Code-Collection, section 2.5 (“Diffuse sky irradiance in the presence of horizons”).

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Part 2

Diffusion of Water Vapor with Phase Transitions

1-Dimensional Diffusion of Water Vapor in Porous Medium with Phase Transitions;
variable diffusivity; irregular grid

3 phases: vapor, free (macroscopic) H₂O ice, H₂O adsorbate
implemented in `vapordiffusioni.f`

History: developed 2003–2004

2.1 Governing Equations

indices: v ... gas (vapor), f ... free ice (solid), a ... adsorbed water
 $\bar{\rho}$... mass per total volume, \bar{J} ... vapor flux per total area

conservation of mass:

$$\frac{\partial}{\partial t}(\bar{\rho}_v + \bar{\rho}_f + \bar{\rho}_a) + \nabla \cdot \bar{J} = 0 \quad (2.1)$$

vapor transport: (Landau and Lifshitz, 1987, Vol. VI, §57, §58)

$$J = -D\rho_0\nabla c \quad (2.2)$$

c ... concentration $c = \rho_v/\rho_0$

ρ_0 ... total density of air, including water vapor

ρ_v ... density of vapor

$$p_v = nkT = \rho_v \frac{k}{m_v} T \quad (2.3)$$

m_v ... mass of water molecule; k ... Boltzmann constant

ϵ ... porosity (= void space / total volume)

$\epsilon(1 - \rho_f/\rho_{\text{ice}})$... fraction of space available to gas

$\bar{\rho}_v = \rho_v\epsilon(1 - \rho_f/\rho_{\text{ice}})$ ρ_v ... vapor density in void space

$\bar{\rho}_f = \rho_f\epsilon$ ρ_f ... ice density in volume not occupied by regolith

$\bar{J} = J\epsilon(1 - \rho_f/\rho_{\text{ice}})$ J ... vapor flux through void area

$\rho_{\text{ice}} \approx 926 \text{ kg/m}^3$... density of ice when it's really cold

adsorption: $\bar{\rho}_a = A(p, T)$

reversible and not kinetically-limited

Conservation of mass becomes

$$\begin{aligned} \frac{\partial}{\partial t} \left(\rho_v \left(1 - \frac{\rho_f}{\rho_{\text{ice}}} \right) + \rho_f + \frac{1}{\epsilon} \bar{\rho}_a \right) + \partial_z \left(1 - \frac{\rho_f}{\rho_{\text{ice}}} \right) J &= 0 \\ \frac{\partial}{\partial t} \left[\rho_v \left(1 - \frac{\rho_f}{\rho_{\text{ice}}} \right) + \rho_f + \frac{1}{\epsilon} \bar{\rho}_a \right] &= \partial_z \left[\left(1 - \frac{\rho_f}{\rho_{\text{ice}}} \right) D \partial_z \rho_v \right] \end{aligned}$$

introduce $\varphi = 1 - \frac{\rho_f}{\rho_{\text{ice}}}$ and $\gamma = \frac{k}{m} \frac{1}{\epsilon}$

$$\partial_t \left(\frac{p}{T} \varphi + \frac{k}{m_v} \rho_f \right) + \gamma \left(\frac{\partial \bar{\rho}_a}{\partial p} \partial_t p + \frac{\partial \bar{\rho}_a}{\partial T} \partial_t T \right) = \partial_z \left[D \varphi \left(\partial_z \frac{p}{T} \right) \right] \quad (2.4)$$

This is an equation for p or ρ_f , depending on whether ice is present or not. If ice is present, p is determined by the saturation (frost point) temperature. If there is no ice, then

$$\left(\frac{1}{T} + \gamma \frac{\partial \bar{\rho}_a}{\partial p} \right) \partial_t p + \left(-\frac{p}{T^2} + \gamma \frac{\partial \bar{\rho}_a}{\partial T} \right) \partial_t T = \partial_z \left(D \partial_z \frac{p}{T} \right)$$

2.2 Discretizations

2.2.1 Possible discretizations of spatial derivatives

Note: These spatial discretizations are not necessarily optimal in terms of discretization error.

$$\partial_z(a \partial_z b)|_j = \frac{1}{\Delta z^2} \left(a_{j+1/2}(b_{j+1} - b_j) - a_{j-1/2}(b_j - b_{j-1}) \right) + O(\Delta z^2) \quad (2.5)$$

or

$$\partial_z(a \partial_z b)|_j = \frac{1}{2\Delta z^2} \left((a_{j+1} + a_j)(b_{j+1} - b_j) - (a_j + a_{j-1})(b_j - b_{j-1}) \right) + O(\Delta z^2) \quad (2.6)$$

or

$$\begin{aligned} \partial_z(a \partial_z b)|_j &= a \partial_{zz} b + (\partial_z a) \partial_z b \\ &= \frac{1}{\Delta z^2} \left[a_j(b_{j+1} - 2b_j + b_{j-1}) + \frac{1}{4}(a_{j+1} - a_{j-1})(b_{j+1} - b_{j-1}) \right] + O(\Delta z^2) \end{aligned} \quad (2.7)$$

The most general discretization which is accurate to $O(\Delta z^2)$, rather than just $O(\Delta z)$, is of the following form

$$\begin{aligned} \partial_z(a \partial_z b)|_j &= \frac{1}{\Delta z^2} \left(c a_j b_j + \left(-1 - \frac{c}{2} \right) a_{j-1} b_j + \left(-1 - \frac{c}{2} \right) a_{j+1} b_j \right. \\ &\quad - \frac{c}{2} a_j b_{j-1} + \frac{3+c}{4} a_{j-1} b_{j-1} + \frac{1+c}{4} a_{j+1} b_{j-1} \\ &\quad \left. - \frac{c}{2} a_j b_{j+1} + \frac{1+c}{4} a_{j-1} b_{j+1} + \frac{3+c}{4} a_{j+1} b_{j+1} \right) + O(\Delta z^2) \end{aligned} \quad (2.8)$$

Choices (2.6) and (2.7) above correspond to $c = -1$ and $c = -2$, respectively.

Another set of schemes does not involve the cross-terms $a_{j+1}b_{j-1}$ and $a_{j-1}b_{j+1}$. They are of the following form

$$\begin{aligned}\partial_z(a\partial_z b)|_j &= \frac{1}{\Delta z^2}(-a_j b_j - c a_{j-1} b_j + (-1+c)a_{j+1} b_j + \\ &\quad (1-c)a_j b_{j-1} + c a_{j-1} b_{j-1} + c a_j b_{j+1} + (1-c)a_{j+1} b_{j+1}) + \\ &\quad \left(c - \frac{1}{2}\right) O(\Delta z) + O(\Delta z^2) \\ &= \frac{1}{\Delta z^2}[(1-c)a_{j+1}(b_{j+1} - b_j) + c a_{j-1}(b_{j-1} - b_j) + \\ &\quad + a_j(c b_{j+1} - b_j + (1-c)b_{j-1})] + O(\Delta z)\end{aligned}\tag{2.9}$$

For $c = 1/2$ this reduces to scheme (2.6) above

If starting with complete pore filling, $c > 0$ is required for downward motion of ice table.

On irregular grid: General scheme without cross-terms

$$\begin{aligned}\partial_z(a\partial_z b)|_j &= -\frac{2c + (1-2c)h_+/h_-}{h_- h_+} a_j b_j + \frac{-1 + (1-2c)h_+/h_-}{h_-(h_- + h_+)} a_{j-1} b_j + \frac{2c-2}{h_+(h_- + h_+)} a_{j+1} b_j + \\ &\quad + \frac{1 + (1-2c)h_+/h_-}{h_-(h_- + h_+)} a_j b_{j-1} + \frac{1 + (2c-1)h_+/h_-}{h_-(h_- + h_+)} a_{j-1} b_{j-1} + \frac{2c}{h_+(h_- + h_+)} a_j b_{j+1} \\ &\quad + \frac{2-2c}{h_+(h_- + h_+)} a_{j+1} b_{j+1} + O(h_+ + h_-)\end{aligned}\tag{2.10}$$

where $h_+ = z_{j+1} - z_j$ and $h_- = z_j - z_{j-1}$. For $h_+ = h_- = h$ this reduces to (2.9)

2.2.2 Discretization of time derivative

use eq. (2.4), $A \equiv f$

$$\begin{aligned}\frac{p_j^{n+1}}{T_j^{n+1}} \varphi_j^{n+1} - \frac{p_j^n}{T_j^n} \varphi_j^n + \frac{k}{\mu} (\rho_{f_j}^{n+1} - \rho_{f_j}^n) + \gamma \left. \frac{\partial f}{\partial p} \right|_j^n (p_j^{n+1} - p_j^n) + \\ + \gamma \left. \frac{\partial f}{\partial T} \right|_j^n (T_j^{n+1} - T_j^n) = \Delta t \left(\partial_z D \varphi \partial_z \frac{p}{T} \right)_j^n\end{aligned}\tag{2.11}$$

derivatives of the isotherm are not expanded to keep it linear

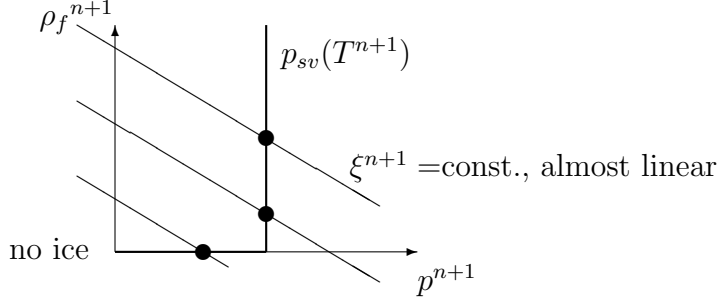
2.2.3 Complete scheme

using (2.11) and (2.10)

$$\begin{aligned}\xi_j^{n+1} &= \frac{p_j^n}{T_j^n} \varphi_j^n + \frac{k}{\mu} \rho_{f_j}^n + \gamma \left. \frac{\partial f}{\partial p} \right|_j^n p_j^n - \gamma \left. \frac{\partial f}{\partial T} \right|_j^n (T_j^{n+1} - T_j^n) + \\ &\quad \frac{\Delta t}{\Delta z^2} \left[D_j \varphi_j^n \left(\frac{p_{j+1}^n}{T_{j+1}^n} - 2 \frac{p_j^n}{T_j^n} + \frac{p_{j-1}^n}{T_{j-1}^n} \right) + \frac{1}{4} (D_{j+1} \varphi_{j+1}^n - D_{j-1} \varphi_{j-1}^n) \left(\frac{p_{j+1}^n}{T_{j+1}^n} - \frac{p_{j-1}^n}{T_{j-1}^n} \right) \right]\end{aligned}$$

where $\xi^{n+1} = \frac{p^{n+1}}{T^{n+1}} \left(1 - \frac{\rho_f^{n+1}}{\rho_{\text{ice}}} \right) + \frac{k}{\mu} \rho_f^{n+1} + \gamma \left| \frac{\partial f}{\partial p} \right|^n p^{n+1}$

$p \leq p_{sv}(T)$ and $0 \leq \rho_f \leq \rho_{\text{ice}}$



p_{sv} ... saturation vapor pressure

Try $\rho_f^{n+1} = 0 \Rightarrow p^{n+1} = \frac{T^{n+1} \cdot \xi^{n+1}}{1 + T^{n+1} \gamma \left| \frac{\partial f}{\partial p} \right|^n}$ and $\rho_f^{n+1} = 0$

If $p^{n+1} > p_{sv}(T^{n+1})$ then $p^{n+1} = p_{sv}(T^{n+1})$ and

$$\rho_f^{n+1} = \frac{\xi^{n+1} - \frac{p_{sv}(T^{n+1})}{T^{n+1}} - \gamma \left| \frac{\partial f}{\partial p} \right|^n p_{sv}(T^{n+1})}{\frac{k}{\mu} - \frac{p_{sv}(T^{n+1})}{T^{n+1} \rho_{\text{ice}}}}$$

introduce $p_{\text{frost}}^{n+1} = p_{sv}(T^{n+1})$

2.2.4 Upper boundary condition

- 1) $p(z=0, t) = p_{\text{atm.}}(t)$
- 2) $D(z=0) = D_0$
- 3) $\varphi_0 = 1$

$$\partial_z \left(D \varphi \partial_z \frac{p}{T} \right) \Big|_{j=0} = \frac{1}{\Delta z^2} \left[D_1 \varphi_1 \left(\frac{p_2}{T_2} - 2 \frac{p_1}{T_1} + \frac{p_{\text{atm}}}{T_{\text{surf}}} \right) + \frac{1}{4} (D_2 \varphi_2 - D_0 \varphi_0) \left(\frac{p_2}{T_2} - \frac{p_{\text{atm}}}{T_{\text{surf}}} \right) \right] \quad (2.12)$$

for half-shifted grid ($z_2 = 3z_1$):

$$a \partial_{zz} b + (\partial_z a) \partial_z b = \frac{1}{\Delta z^2} \left[a_1 \left(\frac{8}{3} b_s - 4b_1 + \frac{4}{3} b_2 \right) + \left(-\frac{4}{3} a_s + a_1 + \frac{1}{3} a_2 \right) \left(-\frac{4}{3} b_s + b_1 + \frac{1}{3} b_2 \right) \right] \quad (2.13)$$

2.2.5 Lower boundary condition

no vapor flux (impermeable) $J = 0 \Rightarrow \partial_z \rho_v = 0 \Rightarrow \partial_z \frac{p}{T} = 0 \Rightarrow \frac{p_{N+1}}{T_{N+1}} = \frac{p_{N-1}}{T_{N-1}}$

$$\partial_z \left(D \varphi \partial_z \frac{p}{T} \right) \Big|_{j=N} = \frac{1}{\Delta z^2} 2D_N \varphi_N \left(\frac{p_{N-1}}{T_{N-1}} - \frac{p_N}{T_N} \right) \quad (2.14)$$

2.3 Discussion

For 1D diffusion and advection with multiple gases, the governing equations are more complex, but the numerical implementation is not. For the governing equations in non-isothermal environments see works such as Cunningham and Williams (1980); Bouziani and Fanale (1998); Hudson et al. (2007).

Water vapor diffusion calculations for Mars are carried out in Schorghofer and Aharonson (2005).

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Part 3

Long-Term Ice Evolution: Equilibrium and Non-equilibrium Ice Tables

Long-term evolution of sub-surface ice due to vapor exchange with the atmosphere using diurnally- and seasonally-resolved temperatures.

History:

2002–2004 equilibrium ice table on Mars

2006–2011 asynchronous model for ice on Mars

3.1 Equilibrium Ice Table on Mars

The equilibrium depth to the ice table is defined by a balance between the vapor pressure at the ice table and the atmosphere (Fig. 3.1). It is the end result of atmosphere-subsurface vapor exchange after an asymptotically long time. If no equilibrium is possible, then subsurface ice is “unstable”. Using the thermal model described in Part 1, this model calculates the equilibrium depth based on matching the vapor density at the ice table with that in the atmosphere. The thermal model is run over a number of Mars years to equilibrate, and annual means are calculated for the last orbit. The ice content of the subsurface changes the thermal properties, and the thermal model is repeatedly equilibrated.

Figure 3.2 shows the result of thermal model calculations (using the methods described

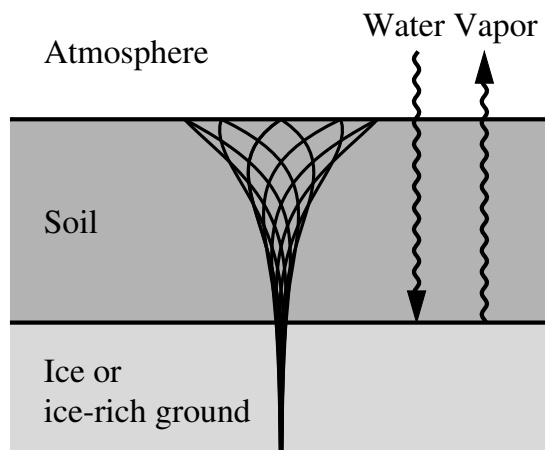


Figure 3.1: Subsurface ice exchanges water vapor with the atmosphere through a layer of porous soil. Temperature oscillations decay with depth, as illustrated by a set of instantaneous temperature profiles.

in Part 1) and vapor diffusion calculations (using the methods described in Part 2). The gradient in temperature abruptly changes at the ice table due to the change in thermal properties (Fig. 3.2a). The annual mean of the vapor density profile can be determined from the annual mean of the vapor density on the surface and at the ice table (Fig. 3.2c). The time-averaging can be justified mathematically by swapping the time integral with the gradient in Fick’s diffusion law. These boundary values can be calculated without solving the vapor diffusion equation, which provides a major computational advantage.

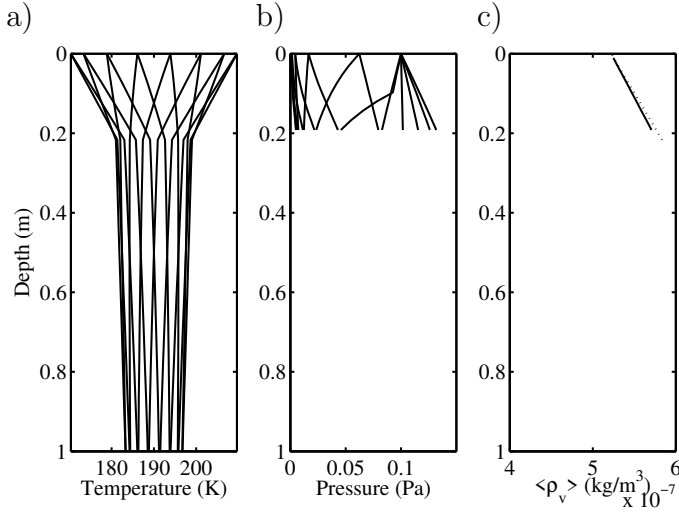


Figure 3.2: Temperature, partial pressure of H₂O, and annual mean vapor density $\langle \rho_v \rangle$ when the ice table is above equilibrium (and therefore retreats). In this conceptional example, a sinusoidal surface temperature dependence is assumed. The vapor density profiles vary with time, but the annual mean of ρ_v simply increases linearly with depth. The dotted line is based on the boundary values of the mean annual vapor density and closely approximates the annual mean from the microphysical vapor diffusion calculations.

The equilibrium ice table calculations determine at what depth the annual mean vapor density at the ice table matches that on the surface. Because the ice content changes the thermal properties, multiple thermal model runs need to be carried out to arrive at the equilibrium depth. A root-finding procedure is deployed to find the equilibrium depth. In `mars_mapi.f` the root-finding procedure is a bisection method. The more recent but simpler program `mars_mapii.f` instead iteratively updates the instantaneous equilibrium depth to arrive at the unique final equilibrium depth. Then `mars_mapt.f` can be used to output additional variables for a given ice table depth.

The model was extensively used in Schorghofer and Aharonson (2005), where further description is available. With extensions described in section 1.3.2 for planar slopes, it also forms the core of the model used in Aharonson and Schorghofer (2006).

3.2 Asynchronous Model for Near-Surface Ice on Mars

The model couples a diurnally-resolved thermal model with a long-term ice evolution model. Ice (massive or interstitial) can be lost to the atmosphere and, vice-versa, pore spaces can

Table 3.1: Programs that calculate the equilibrium ice table on Mars.

Main program	Task
<code>mars_mapt</code>	temperatures for prescribed ice table depth
<code>mars_mapi</code>	finds equilibrium ice table with bisection method
<code>mars_mapii</code>	finds equilibrium ice table with iteration method
<code>mars_mapi2p</code>	finds equilibrium ice table for planar slopes with bisection method

be recharged with interstitial ice. In difference to the model for the equilibrium ice table (section 3.1), this *dynamical* model calculates changes in ice volume and the interstitial spaces may only be partially filled with ice.

Schorghofer (2010) provides a description of this rather complex model, which is not repeated here. In brief, the time-averaged vapor transport equations are solved, which avoids having to solve the microphysical vapor transport equations. The numerical method involves a one-sided derivative at the moving ice table, otherwise a numerical instability occurs due to the strong contrast in thermal properties at the ice table. The model allows for up to three layers: ice-free, soil with interstitial ice (plus void spaces), and massive ice with dust. In the current implementation, one of the two interfaces is tracked explicitly. The model is extensively used in Schorghofer (2007) and Schorghofer and Forget (2012) for studies of the Martian ice age cycle.

Table 3.2: Overview of current implementations of asynchronous dynamic models for Mars

Main program	Task
<code>mars_fast</code>	ice evolution on Mars (massive and interstitial ice)
<code>exper_fast</code>	ice evolution in lab experiment
<code>stabgrow_fast</code>	growth of pore ice

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