Hydrostatic, ideal-gas reference states in spherical coordinates

Loren Matilsky

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1 Basic assumptions

We wish to derive the thermodynamic state for an ideal, hydrostatic gas in spherical coordinates, assuming a known specific entropy stratification (r) and gravitational acceleration $g(r)$. In keeping with the convention used in, e.g., `Rayleigh` and `ASH`, we will denote the thermal profiles of this reference state using overbars. In mathematical terms, we have:

$$\frac{d}{dr} = -(r)g(r) \qquad \text{(hydrostatic balance)}, \tag{1.1}$$

$$\text{and} \qquad (r) = (r)\mathcal{R}(r) \qquad \text{(ideal gas law)}. \tag{1.2}$$

Here, (r) is the pressure, (r) the density, (r) the temperature, and $\mathcal{R}$ the gas constant. We further assume that the number of degrees of freedom (d.o.f.) is constant throughout, so that $\mathcal{R}$ is independent of radius. We note the relationship between the specific heats for an ideal gas:

$$c_{\mathrm{p}} = \gamma c_{\mathrm{v}} = \frac{\gamma \mathcal{R}}{\gamma - 1}, \tag{1.3}$$

where $\gamma := c_{\mathrm{p}}/c_{\mathrm{v}}$.

We note the First Law of Thermodynamics for the radial thermodynamic gradients (easiest starting point is $d = c_{\mathrm{v}}d - (/^2)d$):

$$\frac{1}{c_{\mathrm{v}}}\frac{d\overline{S}}{dr} = \frac{d\ln\overline{T}}{dr} - (\gamma - 1)\frac{d\ln\overline{\rho}}{dr}. \tag{1.4}$$

Differentiating the ideal gas law (1.2) gives

$$\frac{d\ln\overline{T}}{dr} = \frac{d\ln\rho}{dr} + \frac{d\ln T}{dr}. \tag{1.5}$$

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Combining Equations (1.4) and (1.5) yields the following useful forms of the First Law:

$$\frac{1}{c_{\mathrm{p}}}\frac{d\overline{S}}{dr} = \frac{1}{\gamma}\frac{d\ln\overline{T}}{dr} - \frac{\gamma-1}{\gamma}\frac{d\ln\overline{\rho}}{dr} \tag{1.6a}$$

$$= \frac{1}{\gamma}\frac{d\ln\overline{P}}{dr} - \frac{d\ln\overline{\rho}}{dr} \tag{1.6b}$$

$$= \frac{d\ln\overline{T}}{dr} - \frac{\gamma-1}{\gamma}\frac{d\ln\overline{P}}{dr} \tag{1.6c}$$

Using Equations (1.3) and (1.6), we can derive an ordinary differential equation for the temperature:

$$\boxed{\left[\frac{d}{dr} - \left[\frac{1}{c_{\mathrm{p}}}\frac{d\overline{S}}{dr}\right]\right](r) = -\frac{g(r)}{c_{\mathrm{p}}}.} \tag{1.7}$$

We can solve Equation (1.7) via integrating factors, yielding

$$(r) = -\frac{e^{(r)/c_{\mathrm{p}}}}{c_{\mathrm{p}}}\int_{r_0}^{r} g(x)e^{-(x)/c_{\mathrm{p}}}dx +_0 e^{[(x)-_0]/c_{\mathrm{p}}} \tag{1.8}$$

Here, we have defined r_0 as an arbitrary radius from which to start the integration. Accordingly, we define the temperature, pressure, density, and entropy at r_0 as $_0$, $_0$, $_0$, and $_0$, respectively.

Note that in equation (1.8), adding a constant σ to the entropy ($\longrightarrow +\sigma$) has no effect on the profile (r). Only the relative stratification of entropy is important. We therefore set $_0 := 0$ in what follows. Equation (1.8) thus becomes

$$\boxed{(r) = e^{(r)/c_{\mathrm{p}}}\left[_0 - \int_{r_0}^{r}\frac{g(x)}{c_{\mathrm{p}}}e^{-(x)/c_{\mathrm{p}}}dx\right].} \tag{1.9}$$

With (r) now obtained, we can obtain (r) through integrating the First Law in its form (1.6c):

$$\boxed{(r) =_0 e^{-(r)/\mathcal{R}}\left[\frac{(r)}{_0}\right]^{\gamma/(\gamma-1)}.} \tag{1.10}$$

Finally, (r) is determined from Equation (1.2):

$$\boxed{(r) =_0 e^{-(r)/\mathcal{R}}\left[\frac{(r)}{_0}\right]^{1/(\gamma-1)}.} \tag{1.11}$$

Thus, the full state of the atmosphere is purely determined by the entropy stratification $d\overline{S}/dr$, the gravitational acceleration $g(r)$, and the two constants $_0$ and $_0$ (or $_0$ and $_0$, etc.) It is well-known that with an equation of state, all the thermodynamic variables can be written as functions of the other two. Effectively, we have chosen our two "fundamental variables" to

be the entropy and pressure (because of the hydrostatic assumption, we can replace pressure by gravity).

The state specified in equations (1.9)–(1.11) is completely general, relying only on the assumptions of a hydrostatic, ideal gas with position-independent specific heats. To move forward, we must specify $d\overline{S}/dr$ and $g(r)$. For relatively thin convection zones (e.g., the Sun), it is safe to assume that the gravitational acceleration comes purely from a central spherically distributed mass, i.e.,

$$g(r) = \frac{GM_*}{r^2},\qquad(1.12)$$

where G is the Universal Gravitational Constant and M_* the stellar mass.

2 Adiabatic atmosphere

Considering the mathematical form of (1.9), the simplest atmosphere to derive is one that is adiabatic (constant-entropy):

$$(r) \equiv 0 \quad \text{and} \quad \frac{d\overline{S}}{dr} \equiv 0.\qquad(2.1)$$

In this case, all the exponentials in (1.9) cancel to 1 and we find

$$(r) = -\frac{GM_*}{c_{\mathrm{p}}} \int_{r_0}^{r} \frac{dx}{x^2} +_0$$

or

$$(r) =_0 \left[\tilde{a}\left(\frac{r_0}{r}\right) + (1 - \tilde{a})\right] \qquad \text{(adiabatic atmosphere)},\qquad(2.2)$$

$$\text{where} \quad \tilde{a} := \frac{GM_*}{c_{\mathrm{p}0}r_0}.\qquad(2.3)$$

Note that $\tilde{a}$ is (up to a factor likely of order unity, depending on convention) equal to the dissipation number, $\mathrm{Di} := \tilde{g}H/(c_{\mathrm{p}}\tilde{T})$, where H is a typical length-scale and the tildes denote typical background-state values.

Equations (1.10) and (1.11) yield

$$(r) =_0 \left[\tilde{a}\left(\frac{r_0}{r}\right) + (1 - \tilde{a})\right]^{\gamma/(\gamma-1)} \qquad \text{(adiabatic atmosphere)}\qquad(2.4)$$

$$\text{and} \quad (r) =_0 \left[\tilde{a}\left(\frac{r_0}{r}\right) + (1 - \tilde{a})\right]^{1/(\gamma-1)} \qquad \text{(adiabatic atmosphere)}.\qquad(2.5)$$

Note: to keep thermodynamic quantities positive, we must ensure that the argument

$$\tilde{\zeta}(r) := \tilde{a}\left(\frac{r_0}{r}\right) + (1 - \tilde{a}) > 0.\qquad(2.6)$$

This leads to the conditions on available radii r (for our particular choices of $\tilde{a}$—i.e., M_*, $\mathcal{R}$, and $_0$—and r_0) for which the derived atmosphere is valid:

$$0 < r < r_{\text{bound}} := \left(\frac{\tilde{a}}{\tilde{a} - 1} \right) r_0 \qquad \tilde{a} > 1 \qquad (2.7\text{a})$$

$$r > 0 \qquad\qquad\qquad \tilde{a} \leq 1. \qquad (2.7\text{b})$$

3   Polytropes

The starting assumption for a "polytrope" is that the temperature gradient is divergenceless—i.e., there is a constant flux due to radiative diffusion throughout the fluid layer. If we assume, for simplicity, a constant radiative diffusion coefficient, then $\nabla^2 = 0$, which, in spherical coordinates, is:

$$(r) = \frac{A}{r} + B.$$

The integration constants A and B are arbitrary, but one is eliminated from the condition $T(r_0) =_0$, leaving

$$(r) =_0 \left[a \left(\frac{r_0}{r} \right) + (1 - a) \right] \qquad (3.1)$$

where the new constant a is dimensionless. Mathematically, a can be any number, with $a = 0$ corresponding to an isothermal atmosphere. Physically, if we want the temperature (and as we will see shortly, the pressure and density) to decrease with radius, we restrict ourselves to

$$a \geq 0. \qquad (3.2)$$

Note the similarity of the argument

$$\zeta(r) := a \left(\frac{r_0}{r} \right) + (1 - a) \qquad (3.3)$$

in Equation (3.1) (which was derived by assuming a divergenceless temperature gradient) to the argument $\tilde{\zeta}(r)$ in Equation (2.2) (which was derived by assuming the gas was adiabatic).

Using Equation (1.2) to eliminate (r) from Equation (1.1) gives

$$(r) =_0 \exp \left[-\frac{1}{\mathcal{R}} \int_{r_0}^{r} \frac{g(x)}{(x)} dx \right],$$

from which

$$(r) =_0 \left[a \left(\frac{r_0}{r} \right) + (1 - a) \right]^{n+1},$$

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where we have defined the polytropic index n to be

$$n := \frac{b}{a} - 1 \iff a = \frac{b}{n+1}, \tag{3.4}$$

$$\text{where} \quad b := \frac{GM_*}{\mathcal{R}_0 r_0}. \tag{3.5}$$

From Equation (3.2), n is restricted to the range:

$$0 \leq n \leq \infty. \tag{3.6}$$

($n \to \infty$ corresponds to $a = 0$, the isothermal atmosphere).

Note that b is (up to a factor likely of order unity, depending on convention) equal to the dissipation number, $\mathrm{Di} := \tilde{g}H/(c_\mathrm{p}\tilde{T})$, where H is a typical length-scale and the tildes denote typical background-state values.

Putting it all together (and using the ideal gas law to derive the density), we have

$$(r) =_0 \left[a \left(\frac{r_0}{r} \right) + (1-a) \right] \quad \text{(polytropic atmosphere)}. \tag{3.7}$$

$$(r) =_0 \left[a \left(\frac{r_0}{r} \right) + (1-a) \right]^{n+1} \quad \text{(polytropic atmosphere)}. \tag{3.8}$$

$$(r) =_0 \left[a \left(\frac{r_0}{r} \right) + (1-a) \right]^{n} \quad \text{(polytropic atmosphere)} \tag{3.9}$$

We can also rewrite $\zeta(r)$ as

$$\zeta(r) = \frac{b}{n+1} \left(\frac{r_0}{r} \right) - \frac{b - (n+1)}{n+1}$$

to yield

$$(r) =_0 \left[\frac{b}{n+1} \left(\frac{r_0}{r} \right) - \frac{b - (n+1)}{n+1} \right] \quad \text{(polytropic atmosphere)}. \tag{3.10}$$

$$(r) =_0 \left[\frac{b}{n+1} \left(\frac{r_0}{r} \right) - \frac{b - (n+1)}{n+1} \right]^{n+1} \quad \text{(polytropic atmosphere)}. \tag{3.11}$$

$$(r) =_0 \left[\frac{b}{n+1} \left(\frac{r_0}{r} \right) - \frac{b - (n+1)}{n+1} \right]^{n} \quad \text{(polytropic atmosphere)}. \tag{3.12}$$

Similarly to the adiabatic atmosphere (see Equation (2.7)), the choices for a (or equivalently, b and n) and r_0 determine the range of radii for which the polytrope is valid (i.e., the range for which the thermodynamic quantities are positive):

$$0 < r < r_\mathrm{bound}(n) := \left[\frac{b}{b - (n+1)} \right] r_0 \quad b = \frac{GM_*}{\mathcal{R}_0 r_0} > n+1 \tag{3.13a}$$

$$r > 0 \quad\quad\quad b = \frac{GM_*}{\mathcal{R}_0 r_0} \leq n+1. \tag{3.13b}$$

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In general, we use a polytrope as a simplified background for a localized region of a particular star. We will thus choose representative stellar values for M_*, $\mathcal{R}$, r_0, and $_0$ (setting b), and then choose a value for n. For $n < b - 1$, higher n yields a wider range of validity, and lower n a narrower one.

To make things concrete, here we show where an adiabatic polytrope is valid for the Sun. Recall $G = 6.67 \times 10^{-8}$ cm^3 g^{-1} s^{-2}. For the Sun ($M_* = M_\odot = 1.99 \times 10^{33}$ g), we integrate from the base of the convection zone ($r_0 = r_{\mathrm{bcz}} = 5.00 \times 10^{10}$ cm; $_0 =_{\mathrm{bcz}} = 2.11 \times 10^6$ K; $c_{\mathrm{p}} = 3.50 \times 10^8$ erg g^{-1} K^{-1} or $\mathcal{R} = 1.40 \times 10^8$ erg g^{-1} K^{-1}) and assume adiabatic stratification ($n = 1.50$). Then $b = 8.98 > 2.5$ and condition (3.13a) becomes

$$0 < r < (r_{\mathrm{bound}})_\odot := 6.93 \times 10^{10} \text{ cm}. \tag{3.14}$$

The upper bound is only a little less than the solar radius, $R_\odot = 6.96 \times 10^{10}$ cm. This makes sense of course, since the solar convection zone *is* (mostly) an adiabatic polytrope, and the photosphere corresponds to (relatively) very low temperature, density, and pressure.

Note that for the polytrope, in addition to the two integration constants $_0$ and $_0$, there is another constant n (or equivalently, a). Specifying n (or a) is the same as specifying $d\bar{S}/dr$. For, plugging Equations (3.10)–(3.12) (we only need two of them) into any form of (1.6) yields

$$\frac{1}{c_{\mathrm{p}}} \frac{d\bar{S}}{dr} = \left(\frac{n - \tilde{n}}{\tilde{n} + 1} \right) \left(\frac{1}{r} \right) \left\{ \frac{1}{1 + [(1 - a)/a](r/r_0)} \right\}] \tag{3.15a}$$

$$\text{and} \quad \frac{(r)}{c_{\mathrm{p}}} = \left(\frac{n - \tilde{n}}{\tilde{n} + 1} \right) \left\{ \ln \left(\frac{r}{r_0} \right) - \ln \left[a + (1 - a) \left(\frac{r}{r_0} \right) \right] \right\} \tag{3.15b}$$

(polytropic atmosphere),

or, in terms of b,

$$\frac{1}{c_{\mathrm{p}}} \frac{d\bar{S}}{dr} = \left(\frac{n - \tilde{n}}{\tilde{n} + 1} \right) \left(\frac{1}{r} \right) \left[\frac{1}{1 + [(n + 1 - b)/b](r/r_0)} \right] \tag{3.16a}$$

$$\text{and} \quad \frac{(r)}{c_{\mathrm{p}}} = \left(\frac{n - \tilde{n}}{\tilde{n} + 1} \right) \left\{ \ln \left(\frac{r}{r_0} \right) - \ln \left[\frac{b}{n + 1} + \left(\frac{n + 1 - b}{n + 1} \right) \left(\frac{r}{r_0} \right) \right] \right\} \tag{3.16b}$$

(polytropic atmosphere),

where

$$\tilde{n} = \frac{1}{\tilde{a}} \frac{GM_*}{\mathcal{R}_0 r_0} - 1 := \frac{1}{\gamma - 1} \tag{3.17}$$

and $\tilde{a} := GM_*/c_{\mathrm{p}0} r_0$ was already defined in Equation (2.3).

From (3.16a), it is obvious that $n = \tilde{n}$ (or equivalently, $a = \tilde{a}$) corresponds to an adiabatic atmosphere, as was already shown by Equations (2.2)–(2.5)). For other values of n, we have

$$n > \tilde{n} \iff \frac{d\bar{S}}{dr} > 0 \text{ (atmosphere is stable to convection)} \tag{3.18}$$

$$\text{and} \quad n < \tilde{n} \iff \frac{d\bar{S}}{dr} < 0 \text{ (atmosphere may be unstable to convection)}. \tag{3.19}$$

Say we want to increase the stability of a solar-like atmosphere by increasing n (keeping r_0, $_0$, and $_0$ fixed). What happens? We compute, for a particular radius r:

$$\frac{\partial}{\partial n}\left(\frac{1}{c_{\mathrm{p}}}\frac{d\overline{S}}{dr}\right) = \frac{b}{\tilde{n}+1}\left(\frac{r_0}{r^2}\right)\left\{\frac{\tilde{n}+1-b+b(r_0/r)}{[n+1-b+b(r_0/r)]^2}\right\}. \tag{3.20}$$

The sign of this partial derivative is independent of n. Therefore, at any given radius r, the magnitude of the entropy gradient monotonically increases or decreases with n. Let's assume we confine ourselves to $r < (r_{\mathrm{bound}})_\odot$, even though for high n, there will technically be a wider range of validity for the polytrope. Then $\tilde{n}+1-b+b(r_0/r) \to 0$ (from above) as $r \to (r_{\mathrm{bound}})_\odot$ (from below). Thus (for $r < (r_{\mathrm{bound}})_\odot$), the partial derivative in Equation (3.20) is positive, and maximum stability is achieved for $n \to \infty$, the isothermal atmosphere.

Recalling that $\lim_{n\to\infty}(1+x/n)^n = e^x$, Equations (3.10)–(3.16a) show that the isothermal atmosphere in spherical coordinates is:

$$(r) \to_0, \tag{3.21}$$

$$(r) \to_0 \exp\left[b\left(\frac{r_0}{r}-1\right)\right], \tag{3.22}$$

$$(r) \to_0 \exp\left[b\left(\frac{r_0}{r}-1\right)\right], \tag{3.23}$$

$$\frac{1}{c_{\mathrm{p}}}\frac{d\overline{S}}{dr} \to \frac{b}{\tilde{n}+1}\left(\frac{1}{r^2}\right), \tag{3.24}$$

$$\text{and}\quad \frac{(r)}{c_{\mathrm{p}}} \to \frac{b}{\tilde{n}+1}\left(1-\frac{r_0}{r}\right) \tag{3.25}$$

(isothermal atmosphere; polytrope with $n \to \infty$).

If one tries to increase the stability a solar-like polytrope by increasing n, there is therefore a fundamental upper bound imposed by the polytropic formulation itself. Similarly, as $n \to \infty$, the total entropy contrast from top to bottom and all logarithmic derivatives of the thermodynamic variables have upper bounds, meaning that all scale heights have lower bounds. These bounds are reached for an isothermal atmosphere.

Finally, suppose we want to increase the *instability* of the system by *lowering* n to 0 (for $b = 8.98$ and $r < (r_{\mathrm{bound}})_\odot$, Equation (3.20) shows that d/dr everywhere decreases as n decreases). For $n = 0$, we find the maximally unstable (isopycnic, or constant-density) atmosphere:

$$(r) \to_0 \left[b\left(\frac{r_0}{r}\right) + (1-b)\right], \tag{3.26}$$

$$(r) \to_0 \left[b\left(\frac{r_0}{r}\right) + (1-b)\right], \tag{3.27}$$

$$(r) \to_0, \tag{3.28}$$

$$\frac{1}{c_{\mathrm{p}}}\frac{d\overline{S}}{dr} \to -\frac{\tilde{n}b}{\tilde{n}+1}\left(\frac{1}{r}\right)\left[\frac{1}{(1-b)(r/r_0)+b}\right], \tag{3.29}$$

$$\text{and}\quad \frac{(r)}{c_{\mathrm{p}}} \to -\frac{\tilde{n}}{\tilde{n}+1}\left\{\ln\left(\frac{r}{r_0}\right) - \ln\left[(1-b)\left(\frac{r}{r_0}\right)+b\right]\right\} \tag{3.30}$$

(isopycnic atmosphere; polytrope with $n = 0$).

Note that from Equation (3.13a), this $n = 0$ polytrope for the Sun (with $r_0 = r_{\text{bcz}}$ and $b = 8.98$) is only valid for $r < (r_{\text{bound}})_{\odot,n=0} := 5.63 \times 10^{10}$ cm.

3.1 Polytrope in terms of $_{\text{bcz}}$ and N_ρ

Suppose we use the polytrope to describe a convection zone (CZ) and integrate from the base of the CZ ($r_0 = r_{\text{bcz}}$). In the work of Jones et al. (2011) and in the `Rayleigh` code (Featherstone et al., 2021), the polytrope is initiated using the density at the base of the CZ ($_{\text{bcz}}$) and the number of density scale heights N_ρ across the CZ, as opposed to the two base-of-CZ values $_{\text{bcz}}$ and $_{\text{bcz}}$. Here $N_\rho := \ln\left(_{\text{bcz}}/_{\text{out}}\right)$, where $_{\text{out}}$ is the density at the top of the CZ. We can recast Equations (3.7)–(3.9) in terms of $_{\text{bcz}}$ and N_ρ by using (3.4), (3.5), and (3.9) to yield

$$a = \frac{e^{N_\rho/n} - 1}{(1 - \beta)e^{N_\rho/n}} \qquad \text{and} \qquad _{\text{bcz}} = \frac{(1 - \beta)e^{N_\rho/n}}{e^{N_\rho/n} - 1} \frac{GM_*}{(n + 1)\mathcal{R}r_{\text{bcz}}}, \tag{3.31}$$

where we have defined the aspect ratio of the CZ,

$$\beta := \frac{r_{\text{bcz}}}{r_{\text{out}}} < 1. \tag{3.32}$$

Thus, instead of specifying $_{\text{bcz}}$ and $_{\text{bcz}}$, we can specify $_{\text{bcz}}$ and N_ρ; $_{\text{bcz}}$ is then determined through (3.31) and $_{\text{bcz}}$ through the ideal gas law.

3.2 Polytrope with respect to the center of the CZ

Jones et al. (2011) define the polytrope using the variables at the center of the CZ (i.e., in our notation $r_0 = r_c := (r_{\text{bcz}} + r_{\text{out}})/2$). In that case, (3.10)–(3.12) become

$$(r) =_c \left[a_c \left(\frac{r_c}{r}\right) + (1 - a_c)\right], \tag{3.33}$$

$$(r) =_c \left[a_c \left(\frac{r_c}{r}\right) + (1 - a_c)\right]^{n+1}, \tag{3.34}$$

$$(r) =_c \left[a_c \left(\frac{r_c}{r}\right) + (1 - a_c)\right]^n, \tag{3.35}$$

where

$$a_c := \frac{GM_*}{(n + 1)\mathcal{R}_c r_c}. \tag{3.36}$$

Using a similar computation as the one leading to (3.31), one can show that

$$a_c = \frac{2\beta(e^{N_\rho/n} - 1)}{(1 - \beta)(\beta e^{N_\rho/n} + 1)}. \tag{3.37}$$

Defining

$$H := r_{\text{out}} - r_{\text{bcz}}, \tag{3.38}$$

$$c_0 := 1 - a_c = \frac{(1+\beta)(1 - \beta e^{N_\rho/n})}{(1-\beta)(\beta e^{N_\rho/n} + 1)}, \tag{3.39}$$

$$c_1 := \left(\frac{r_c}{d}\right) a_c = \frac{(1+\beta)\beta(e^{N_\rho/n} - 1)}{(1-\beta)^2(\beta e^{N_\rho/n} + 1)}, \tag{3.40}$$

$$\text{and} \quad \zeta(r) := a_c\left(\frac{r_c}{r}\right) + (1 - a_c) = c_0 + c_1\left(\frac{H}{r}\right), \tag{3.41}$$

(3.33)–(3.35) become

$$(r) =_c \zeta(r), \qquad (r) =_c [\zeta(r)]^{n+1}, \qquad (r) =_c [\zeta(r)]^n. \tag{3.42}$$

We can also see that if we define

$$\zeta_{\text{out}} := \zeta(r_{\text{out}}) = \frac{1+\beta}{\beta e^{N_\rho/n} + 1} \tag{3.43a}$$

$$\text{with} \quad e^{N_\rho/n} = \frac{1 + \beta - \zeta_{\text{out}}}{\zeta_{\text{out}}}, \tag{3.43b}$$

then

$$c_0 = \frac{2\zeta_{\text{out}} - \beta - 1}{1 - \beta} \quad \text{and} \quad c_1 = \frac{(1+\beta)(1 - \zeta_{\text{out}})}{(1-\beta)^2}, \tag{3.44a}$$

$$\text{with} \quad \zeta_{\text{bcz}} := \zeta(r_{\text{bcz}}) = c_0 + \left(\frac{1-\beta}{\beta}\right) c_1 = \frac{1 + \beta - \zeta_{\text{out}}}{\beta}, \tag{3.44b}$$

yielding exactly the formulation for the polytrope in Jones et al. (2011) (see their Equations (18) and (19)).

Also note that combining Equations (3.36) and (3.40) yields

$$c_1 = \frac{GM_*}{(n+1)\mathcal{R}T_c H} \tag{3.45a}$$

$$= \frac{GM_*}{c_{\text{p}}T_c H} \quad \text{if gas is adiabatic } (n = \tilde{n}). \tag{3.45b}$$

3.3   Non-Dimensional Polytrope

Here we show that as far as the non-dimensional system is concerned, any polytrope in a spherical shell is fully described by three numbers: the shell's aspect ratio, the number of density scale heights across the shell, and the polytropic index. We consider a spherical shell with (dimensional) inner and outer radii r_{in} and r_{out}, respectively. Then

$$\beta := \frac{r_{\text{in}}}{r_{\text{out}}} \qquad \text{(aspect ratio)} \tag{3.46a}$$

$$\text{and} \quad H := r_{\text{out}} - r_{\text{in}} \quad \text{(shell depth)}. \tag{3.46b}$$

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We choose H as our length-scale and thus consider the shell to occupy the range

$$\frac{\beta}{1-\beta} \leq \hat{r} \leq \frac{1}{1-\beta}, \tag{3.47}$$

where $\hat{r} = r/H$ refers to the non-dimensional radius.

We define the number of density scale heights across the shell as

$$N_\rho := \ln\left[\frac{\text{in}}{\text{out}}\right], \tag{3.48}$$

where the subscripts "in" and "out" mean "evaluated at r_{in} or r_{out}, respectively".

We choose $r_0 = r_{\text{in}}$, so

$$\zeta(\hat{r}) = a\left(\frac{\beta}{1-\beta}\right)\left(\frac{1}{\hat{r}}\right) + (1-a). \tag{3.49}$$

From Equation (3.9), we then find

$$e^{-N_\rho/n} = a(\beta - 1) + 1$$
$$\text{or} \quad a = \frac{b}{n+1} = \frac{1 - e^{-N_\rho/n}}{1-\beta}. \tag{3.50}$$

Each thermal profile is non-dimensionalized using a representative value from the polytrope. This could be the volume-average of the profile (Rayleigh's `reference_type = 2`; e.g., Hindman et al. 2020) or the value of the profile at the outer boundary (Rayleigh's `reference_type = 3`; e.g Heimpel et al. 2022). Here, we instead choose the value at the inner boundary. Plugging everything into Equations (3.7)–(3.15b), we find

$$\hat{T}(\hat{r}) = \left[\frac{\beta(1 - e^{-N_\rho/n})}{(1-\beta)^2}\right]\left(\frac{1}{\hat{r}}\right) - \left(\frac{\beta - e^{-N_\rho/n}}{1-\beta}\right) \tag{3.51}$$

$$\hat{P}(\hat{r}) = \left\{\left[\frac{\beta(1 - e^{-N_\rho/n})}{(1-\beta)^2}\right]\left(\frac{1}{\hat{r}}\right) - \left(\frac{\beta - e^{-N_\rho/n}}{1-\beta}\right)\right\}^{n+1}, \tag{3.52}$$

$$\hat{\rho}(\hat{r}) = \left\{\left[\frac{\beta(1 - e^{-N_\rho/n})}{(1-\beta)^2}\right]\left(\frac{1}{\hat{r}}\right) - \left(\frac{\beta - e^{-N_\rho/n}}{1-\beta}\right)\right\}^{n}, \tag{3.53}$$

$$\frac{1}{c_{\text{p}}}\frac{d}{d\hat{r}} = \left(\frac{n-\tilde{n}}{\tilde{n}+1}\right)\left(\frac{1}{\hat{r}}\right)\left[1 - \left(\frac{1-\beta}{\beta}\right)\left(\frac{\beta - e^{-N_\rho/n}}{1 - e^{-N_\rho/n}}\right)\hat{r}\right]^{-1}, \tag{3.54}$$

$$\frac{(\hat{r})}{c_{\text{p}}} = \left(\frac{n-\tilde{n}}{\tilde{n}+1}\right)\left\{\ln\left[\left(\frac{1-\beta}{\beta}\right)\hat{r}\right] - \ln\left[\left(\frac{e^{-N_\rho/n} - \beta}{\beta}\right)\hat{r} + \left(\frac{1 - e^{-N_\rho/n}}{1-\beta}\right)\right]\right\} \tag{3.55}$$

(non-dimensional polytrope).

The non-dimensional polytrope is purely characterized by the three numbers β, N_ρ, and n. (For completeness, we should also include γ, or $\tilde{n}$, but for most cases we are concerned with, $\gamma = 5/3$). Note that the dissipation number Di is *not* an independent parameter. For example, in the non-dimensionalization chosen here, we have

$$a = \frac{GM_*}{(n+1)\mathcal{R}T_{\mathrm{in}}r_{\mathrm{in}}}$$

$$= \left(\frac{\beta}{1-\beta}\right)\left(\frac{\gamma}{\gamma-1}\right)\left(\frac{1}{n+1}\right)\mathrm{Di},$$

$$= \left(\frac{\beta}{1-\beta}\right)\left(\frac{\tilde{n}+1}{n+1}\right)\mathrm{Di},$$

$$\text{where} \quad \mathrm{Di} := \frac{g_{\mathrm{in}}H}{c_{\mathrm{p}}T_{\mathrm{in}}}$$

This, combined with Equation (3.50), yields

$$\mathrm{Di} = \left(\frac{n+1}{\tilde{n}+1}\right)\left(\frac{1}{\beta}\right)(1 - e^{-N_\rho/n}). \tag{3.56}$$

If we instead using volume-averages for the representative values, then Equation (3.51) yields

$$\frac{\langle\rangle_{\mathrm{v}}}{{}_{\mathrm{in}}} = \left(\frac{3}{2}\right)\frac{\beta(1+\beta)[\beta(1-e^{-N_\rho/n})]}{1-\beta^3} - \frac{\beta - e^{-N_\rho/n}}{1-\beta}$$

and $g(r) \propto 1/r^2$ yields

$$\frac{\langle g\rangle_{\mathrm{v}}}{g_{\mathrm{in}}} = \frac{3(1-\beta)\beta^2}{1-\beta^3},$$

where $\langle\cdots\rangle_{\mathrm{v}}$ refers to a volume average over the full shell ($\hat{r}$ ranges from $\beta/(1-\beta)$ to $1/(1-\beta)$). Thus

$$\mathrm{Di}_{\mathrm{v}} := \frac{\langle g\rangle_{\mathrm{v}}/g_{\mathrm{in}}}{\langle\rangle_{\mathrm{v}}/{}_{\mathrm{in}}}\mathrm{Di}$$

$$= \left(\frac{n+1}{\tilde{n}+1}\right)\frac{3\beta(1-\beta)^2(1-e^{-N_\rho/n})}{(3\beta/2)(1-\beta^2)(1-e^{-N_\rho/n}) - (1-\beta^3)(\beta - e^{-N_\rho/n})}. \tag{3.57}$$

For $n = \tilde{n} = 1.5$, $\beta = 0.759$, and $N_\rho = 3$ (e.g., Hindman et al. 2020; Matilsky et al. 2022) Equation (3.57) yields their reported $\mathrm{Di}_{\mathrm{v}} = 1.72$. Note that Equation (3.57) is essentially the "gory" version of Hindman et al. 2020's Equation (A6). Either Equation shows that their are only three independent parameters characterizing the polytrope, so it may not make sense to give preferential reporting of Di_{v}, as was done Hindman et al. (2020); Matilsky et al. (2022).

Finally, note that Equation (3.55) gives the entropy difference across the layer ($\Delta := {}_{\mathrm{out}}$, since have chosen ${}_{\mathrm{in}} = 0$):

$$\boxed{\frac{\Delta}{c_{\mathrm{p}}} = \left(\frac{N_\rho}{\tilde{n}+1}\right)\left(\frac{n-\tilde{n}}{n}\right) \quad \text{(non-dimensional polytrope).}} \tag{3.58}$$

Note that for the two "extreme" atmospheres (isopycnic and isothermal), the entropy difference across the shell is

$$\frac{\Delta}{c_{\mathrm{p}}} = \begin{cases} -\infty & (n = 0; \text{ isopycnic atmosphere}) \\ +\frac{N_\rho}{\tilde{n}+1} & (n = \infty; \text{ isothermal atmosphere}). \end{cases} \tag{3.59}$$

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3.4 Non-Dimensional Polytrope (`Rayleigh reference_type = 3`)

For `Rayleigh`'s `reference_type = 3` (non-dimensional anelastic; e.g., [Heimpel et al. 2022](#)), we again consider a shell spanning $(r_{\mathrm{in}}, r_{\mathrm{out}})$ with aspect ratio $\beta = r_{\mathrm{in}}/r_{\mathrm{out}}$ and choose the shell depth $H = r_{\mathrm{out}} - r_{\mathrm{in}}$ to be the length-scale. We non-dimensionalize (r), (r), (r), and $g(r)$ with their values at the outer boundary, $_{\mathrm{out}}$, $_{\mathrm{out}}$, $_{\mathrm{out}}$, and g_{out} (respectively). Using hats for the non-dimensionalized background profiles (again defining $\hat{r} = r/H$), the hydrostatic Equation (1.1) becomes

$$\frac{_{\mathrm{out}}\mathcal{R}_{\mathrm{out}}}{H}\frac{d\hat{P}}{d\hat{r}} = -_{\mathrm{out}}g_{\mathrm{out}}\hat{\rho}(\hat{r})\hat{g}(\hat{r}), \tag{3.60a}$$

$$\text{or} \quad \frac{d\hat{T}}{d\hat{r}} = \mathrm{Di}_{\mathrm{alt}}\hat{g}(\hat{r}), \tag{3.60b}$$

$$\text{where} \quad \mathrm{Di} := \frac{g_{\mathrm{out}}H}{c_{\mathrm{pout}}}, \tag{3.60c}$$

$$\mathrm{Di}_{\mathrm{alt}} := \frac{g_{\mathrm{out}}H}{(n+1)\mathcal{R}_{\mathrm{out}}}$$

$$= \frac{\gamma}{(\gamma-1)(n+1)}\mathrm{Di}$$

$$= \left(\frac{\tilde{n}+1}{n+1}\right)\mathrm{Di}, \tag{3.60d}$$

$$\text{and} \quad \hat{g}(\hat{r}) = \frac{r_{\mathrm{out}}^2}{r^2} = \frac{1}{(1-\beta)^2}\frac{1}{\hat{r}^2}, \tag{3.60e}$$

and we have used the polytropic relations $\hat{\rho}(\hat{r}) = [\hat{T}(\hat{r})]^n$ and $\hat{P}(\hat{r}) = [\hat{T}(\hat{r})]^{n+1}$ in deriving Equation (3.60b). These Equations are easily integrated to yield

$$\hat{T}(\hat{r}) = \frac{\mathrm{Di}_{\mathrm{alt}}}{(1-\beta)^2}\frac{1}{\hat{r}} + 1 - \frac{\mathrm{Di}_{\mathrm{alt}}}{1-\beta}. \tag{3.61}$$

And since $\hat{\rho}_{\mathrm{in}} = [\hat{T}_{\mathrm{in}}]^n = e^{N_\rho}$, we also have

$$\mathrm{Di}_{\mathrm{alt}} = \beta(e^{N_\rho/n} - 1) \tag{3.62a}$$

$$\text{and} \quad \mathrm{Di} = \left(\frac{n+1}{\tilde{n}+1}\right)\beta(e^{N_\rho/n} - 1). \tag{3.62b}$$

For the derivatives, note that Equation (3.60b) yields

$$\frac{d\ln\hat{T}}{d\hat{r}} = -\mathrm{Di}_{\mathrm{alt}}\frac{\hat{g}(\hat{r})}{\hat{T}(\hat{r})} \tag{3.63}$$

$$\text{and} \quad \frac{d^2\ln\hat{T}}{d\hat{r}^2} = -\left(\frac{d\ln\hat{T}}{d\hat{r}}\right)^2 + \left[\frac{\mathrm{Di}_{\mathrm{alt}}}{\hat{T}(\hat{r})}\right]\left[\frac{2\hat{g}(\hat{r})}{\hat{r}}\right], \tag{3.64}$$

where we have noted that $d\hat{g}/d\hat{r} = -2\hat{g}(\hat{r})/\hat{r}$. The derivatives of density follow easily from $\hat{\rho}(\hat{r}) = [\hat{T}(\hat{r})]^n$.

Note that in the current formulation of `reference_type = 3` (and in Heimpel et al. 2022), Di_{alt} seems to be conflated with with Di, i.e., $n = \tilde{n}$. Since Heimpel et al. (2022) choose $n = 1$ and $n = 2$ for Jupiter, they effectively seem to be choosing a "strange" Jovian gas. In general,

$$\gamma - 1 = \frac{2}{f} \tag{3.65a}$$

$$\text{or} \quad \tilde{n} = \frac{f}{2}, \tag{3.65b}$$

where f is the (integer) number of degrees of freedom (d.o.f.) of the gas. It seems Jupiter should have $f = 5$ (diatomic cool gas; three translational d.o.f., two rotational d.o.f.), but Heimpel et al. (2022) have either $f = 2$ or $f = 4$.

3.5 Stable and unstable polytropes

Figure 1 (Figure 2) shows a sample of convectively stable (unstable) polytropes for various values of the polytropic index n in a solar-like convection zone. For very high values of n (Figure 1), the entropy-gradient profile asymptotes to its maximum value and the temperature profile becomes flat (since the atmosphere becomes isothermal; see Equation (3.21)). For very low values of n (highly unstable polytropes; Figure 2), the entropy-gradient profile asymptotes to its most-negative value and the density profile becomes flat (since the atmosphere becomes isopycnic; see Equation (3.26)). Note that the isopycnic atmosphere has limited validity, with $r_{\text{bound},\odot,n=0}/R_\odot = 0.809$.

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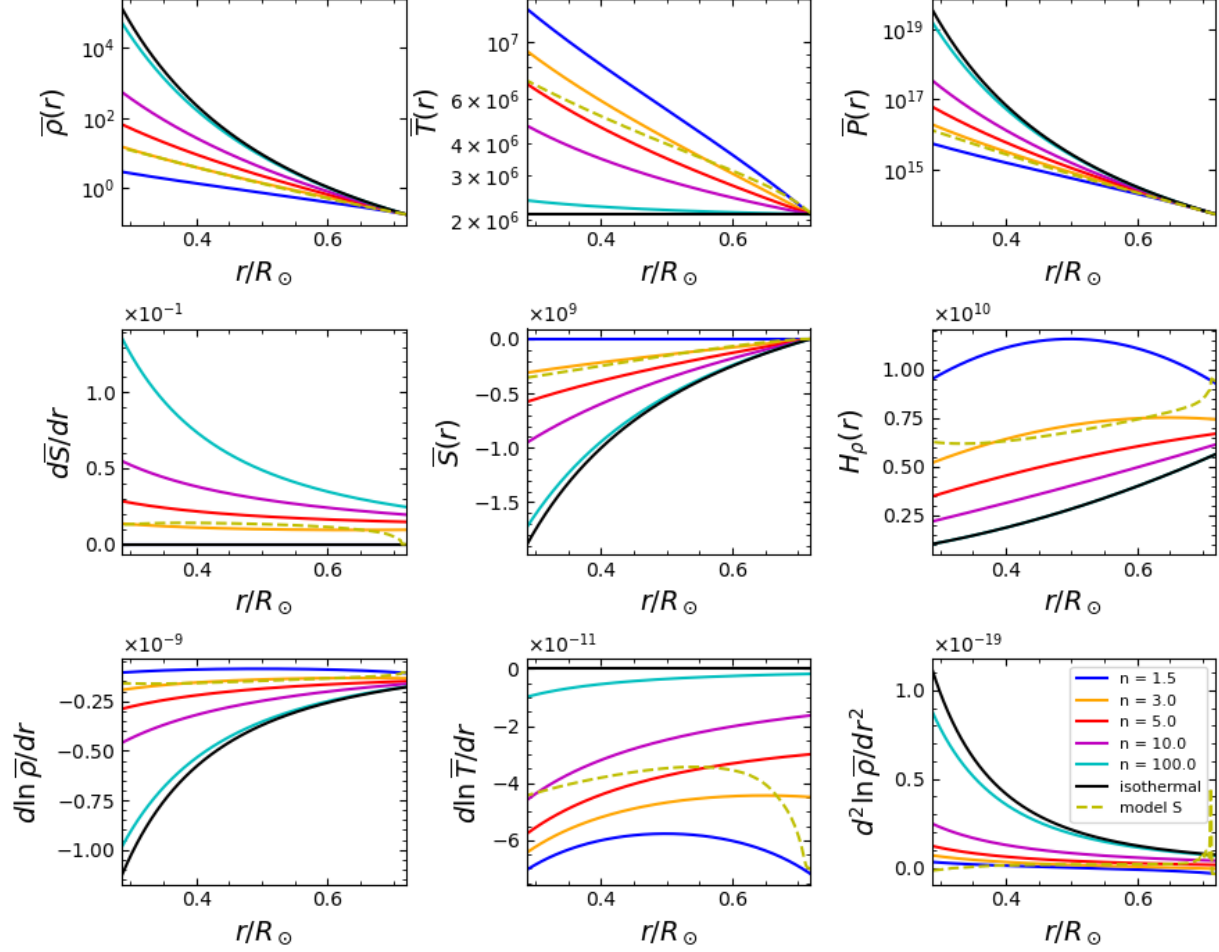
Figure 1: Sample of convectively stable polytropes for a solar-like radiative zone ($n \geq \tilde{n}$, with $\gamma = 5/3$ or $\tilde{n} = 1.5$). The dimensional parameters defining this polytrope are: $G = 6.67 \times 10^{-8}$ cm^3 g^{-1} s^{-2}, $M_* = M_\odot = 1.99 \times 10^{33}$ g, $r_0 = r_{\rm bcz} = 5.00 \times 10^{10}$ cm, $_0 =_{\rm bcz}= 2.11 \times 10^6$ K, $_0 =_{\rm bcz}= 0.181$ g cm^{-3}, and $c_{\rm p} = 3.50 \times 10^8$ erg g^{-1} K^{-1}. We set our bottom radius to $r_{\rm in} = 2 \times 10^{10}$ cm. For reference, we also plot the profiles from the isothermal atmosphere ($n = \infty$) and model S. Model S is remarkably similar (except for some of the radial-gradient profiles) to a polytrope with $n = 3$. All y-axes are measured in cgs units.
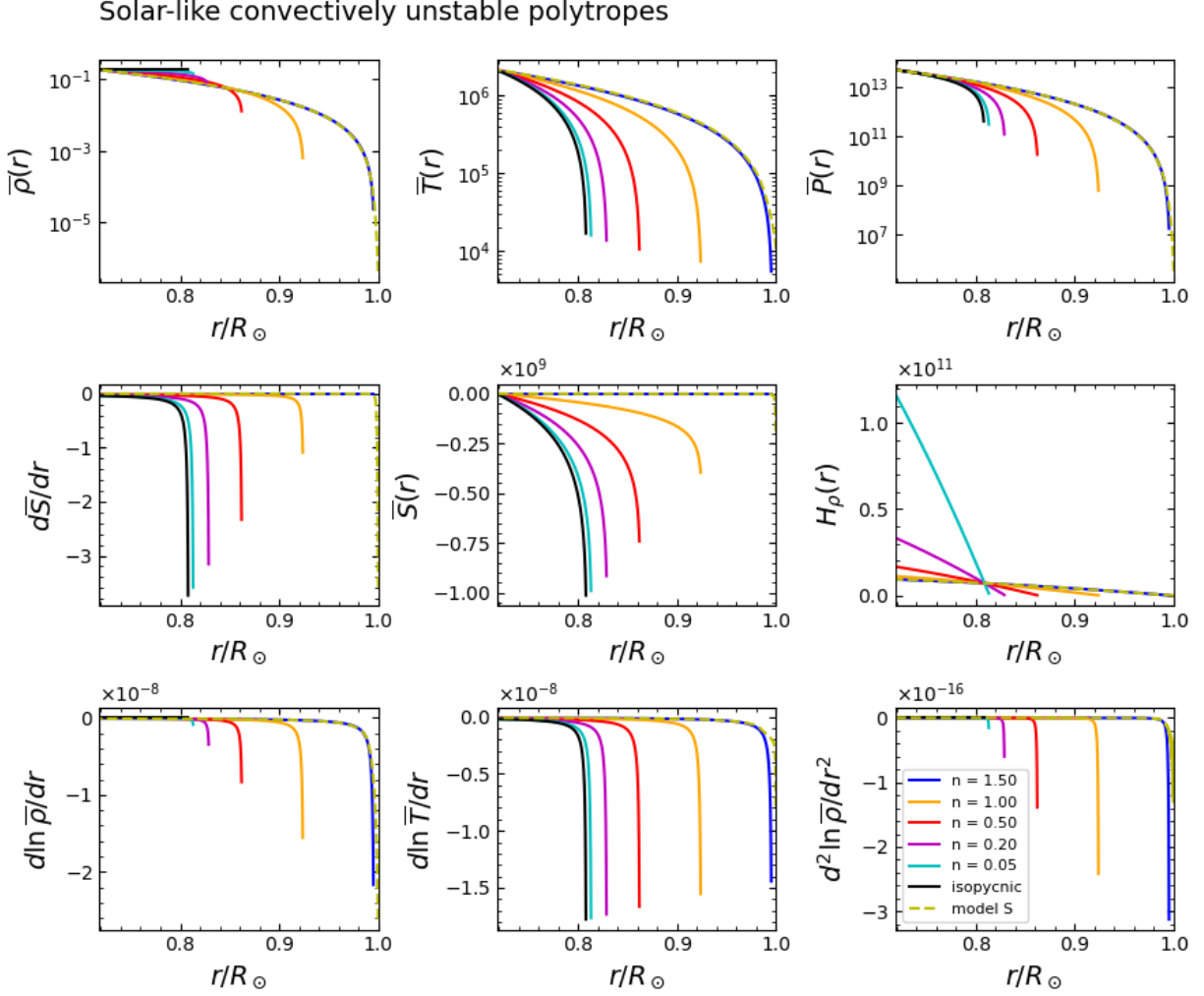
Figure 2: Sample of convectively unstable polytropes for a solar-like radiative zone ($n \leq 1.5$; same dimensional parameters as in Figure 1). For each polytrope, the outer radius is set to the limit r_{bound} defined in Equation (3.13) for $b = 8.98$ (see Equation (3.14) the calculation of b). For reference, we also plot the profiles from the maximally unstable isopycnic atmosphere ($n = 0$) and model S. Model S is similar (except near the photosphere) to an adiabatic polytrope with $n = 1.5$. All y-axes are measured in cgs units.