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1 Thermodynamics

1.1 Definitions

Definition 1.1 (*Moles*): A mole is the amount of mass of a substance equal to its molecular weight, i.e., if you have O_2 , than 32 kg of O_2 constitutes one kg-mole and 32 grams would constitute 32 gm-mole.

Definition 1.2 (*Avagadro's constant*): Avogadro's constant is the number of particles in one mole of that substance. It is denoted as ' N_A ' and is equal to $6.02 \cdot 10^{26}$ particles per kg-mole.

Definition 1.3 (*Universal gas constant*): The universal gas constant is denoted by \hat{R} and is equal to 8.314 J/mol-K. Its units are 'energy per amount of substance per temperature increment' as it is a proportionality constant that relates energy to temperature and amount of substance.

Definition 1.4 (*Boltzmann constant*): The Boltzmann constant is denoted by k_B and is equal to $1.381 \cdot 10^{-23}$ J/k. It is the equivalent of the universal gas constant but on a *per particle* basis.

Definition 1.5 (*Dalton's law of partial pressures*): The pressure of a system can be found my summing up the partial pressures of each individual species:

$$p = \sum_j p_j \quad (1.1)$$

Definition 1.6 (*Mass fractions*): The mass fraction of species j (Y_j) is defined as the ratio of the density of species j to the density of the mixture:

$$Y_j = \frac{\rho_j}{\rho} \quad (1.2)$$

Definition 1.7 (*Molar fractions*): The molar fraction of species j (X_j) can be taken as the ratio of the partial pressure of species j to the pressure of the mixture, or the ratio of the number of moles of species j to the total number of moles in the system:

$$\frac{p_j}{p} = \frac{N_j}{N} = X_j \quad (1.3)$$

Definition 1.8 (*Mixture molecular weight*): The mixture molecular

weight, \mathcal{M} can be defined as:

$$\mathcal{M} = \frac{1}{\sum_j Y_j / \mathcal{M}_j} = \sum_j X_j \mathcal{M}_j \quad (1.4)$$

Where \mathcal{M}_j is the molecular weight of species j .

Definition 1.9 (*Molar fractions to mass fractions*): The mass fraction of species j can be obtained from the molar fraction by using:

$$Y_j = X_j \frac{\mathcal{M}_j}{\mathcal{M}} \quad (1.5)$$

The sum of the mass fractions

Definition 1.10 (*Mixture gas constant*): The gas constant of the mixture can be found with:

$$R = \sum_j Y_j R_j \quad (1.6)$$

where R is the mixture specific gas constant (\hat{R}/\mathcal{M}) and R_j is the specific gas constant of species j .

Definition 1.11 (*Calorically perfect gas*): A calorically perfect gas is a gas where the specific heats c_p and c_v are taken to be constant. The enthalpy is then $h = c_p T$, and the internal energy is $e = c_v T$.

Definition 1.12: Thermally perfect gas A thermally perfect gas is one there c_p and c_v are variables are functions of T only. Differential changes in enthalpy and internal energy is then $dh = c_p dT$, and $de = c_v dT$

1.2 Equations

1.2.1 Equation of state

$$PV = mRT \quad (1.7)$$

$$pv = RT \quad (1.8)$$

$$pV = \mathcal{N}\hat{R}T \quad (1.9)$$

$$p = \rho RT \quad (1.10)$$

$$pV = \mathcal{N}\hat{R}T \quad (1.11)$$

$$p\mathcal{V} = \hat{R}T \quad (1.12)$$

$$p = C\hat{R}T \quad (1.13)$$

$$pv = \eta\mathcal{R}T \quad (1.14)$$

$$pV = NkT \quad (1.15)$$

$$p = nkT \quad (1.16)$$

Where

- P is the pressure
- V is the volume
- v is the specific gas constant
- m is the mass
- T is the temperature
- ρ is the density
- \mathcal{V} is the molar volume (volume per mole)
- C is the concentration of moles per unit volume.
- η is the mole-mass ratio
- n is the number density, or number of particles per unit volume.

2 Statistical thermodynamics

Definition 2.1 (Bose-Einstein (BE) statistics): For atoms and molecules made up of an *even* number of elementary particles such as electrons, protons, and neutrons, there is no limit to the number particles that can be in any energy state at a given instant.

Definition 2.2 (Fermi-Dirac (FD) statistics): For atoms and molecules made up of an *odd* number of elementary particles, no energy state can be occupied by more than one particle.

Definition 2.3 ($W(N_j)$): For these statistics, the number of microstates in a given macrostate is given by:

$$W(N_j) = \prod_j \frac{(N_j + C_j - 1)!}{(C_j - 1)! N_j!} \quad (2.1a)$$

$$W(N_j) = \prod_j \frac{C_j!}{(C_j - N_j)! N_j!} \quad (2.1b)$$

Here, Eq. [2.1a] is used for BE statistics while Eg. [2.1b] is used for FD statistics. C_j is the number of energy states, and N_j is the number of particles in a group.

Definition 2.4 (W_{max}): W_{max} is defined as the largest of the individual terms in Equations [2.1] and is sometimes referred to as the *most probable microstate*. By solving for this value by setting $\delta(\ln W) = 0$, we can find the particular values of N_j that maximize W , giving:

$$N_j^* = \frac{C_j}{e^{\alpha+\beta\epsilon_j} \pm 1} \quad (2.2)$$

Where the minus sign is for BE, and the plus for FD. α and β are Lagrange multipliers used to enforce the constraints:

$$\sum_j \delta N_j = 0, \quad \sum_j \epsilon_j = \delta N_j \quad (2.3)$$

Definition 2.5 (Boltzmann limit): The Boltzmann limit is used for conditions where the number of energy states far out numbers the number of group of particles. Therefore, $C_j \gg N_j$, and Eqn. [2.2] reduced to:

$$N_j^* = C_j e^{-\alpha - \beta \epsilon_j} \quad (2.4)$$

Definition 2.6 (Boltzmann distribution): The Boltzmann distribution, after solving for α and β in the above, gives:

$$N_j^* = N \frac{C_j e^{-\epsilon_j/kT}}{\sum_j C_j e^{-\epsilon_j/kT}} \quad (2.5)$$

Definition 2.7 (Molecular partition function): The molecular partition function, Q , is defined as the denominator in Eq. [2.5]:

$$Q = \sum_j C_j e^{-\epsilon_j/kT} \quad (2.6)$$

However, this is in terms of groups of energy states having the same energy level ϵ_j . If we transition to the use of individual energy states, ϵ_j becomes ϵ_i , and the C_j term is neglected. The equation becomes:

$$Q = \sum_i e^{-\epsilon_i/kT} \quad (2.7)$$

Definition 2.8 (Population of energy states): The population of energy state ϵ_i then becomes:

$$N_i^{*'} = N \frac{e^{-\epsilon_i/kT}}{\sum_i e^{-\epsilon_i/kT}} \quad (2.8)$$

Definition 2.9 (Energy levels and degeneracy): Energy levels are sometimes more convenient to use. An energy level can have multiple energy states in it. If a level contains more than one state, it is said to be *degenerate*, and the number of states inside an energy level ϵ_l is given as g_l . Therefore, the population of particles in energy level ϵ_l , and the partition function is given as:

$$Q = \sum_l g_l e^{-\epsilon_l/kT} \quad (2.9)$$

$$N_l^{*'} = N \frac{g_l e^{-\epsilon_l/kT}}{\sum_i g_i e^{-\epsilon_i/kT}} \quad (2.10)$$

Definition 2.10 (Boltzmann's relation): Boltzmann's relation finds the relationship between the number of microstates Ω and the entropy of a system

as:

$$S = k \ln \Omega \quad (2.11)$$

Definition 2.11 (Entropy): Entropy can then be shown to be of the form:

$$S = Nk \left(\ln \frac{Q}{N} + 1 \right) + \frac{E}{T} \quad (2.12)$$

Definition 2.12 (Helmholtz free energy): The Helmholtz free energy, F , is defined as:

$$F = E - TS \quad (2.13)$$

Here, E is the internal energy, T is the temperature, and S the entropy of the system. Differentiation of [2.13] yields:

$$dF = dE - TdS - SdT \quad (2.14)$$

Since $dE - TdS = \mu dN - pdV$, (μ is the chemical potential per molecule, N is the number of particles, and p , V is the pressure and volume of the system), this equation can be rewritten as:

$$dF = -SdT - pdV + \mu dN \quad (2.15)$$

Which yields the following results:

$$S = - \left(\frac{\partial F}{\partial T} \right)_{V,N} \quad p = - \left(\frac{\partial F}{\partial V} \right)_{T,N} \quad \mu = \left(\frac{\partial F}{\partial N} \right)_{T,V} \quad (2.16)$$

Definition 2.13 (Thermodynamic properties of a system): The thermodynamic properties of a system, assuming Q and ϵ_i 's are independant of N , are:

$$F = -NkT \left(\ln \frac{Q}{N} + 1 \right) \quad (2.17a)$$

$$S = Nk \left[\ln \frac{Q}{N} + 1 + T \frac{\partial(\ln Q)}{\partial T} \right] \quad (2.17b)$$

$$E = NkT^2 \frac{\partial(\ln Q)}{\partial T} \quad (2.17c)$$

$$p = NkT \frac{\partial(\ln Q)}{\partial V} \quad (2.17d)$$

$$\mu = -kT \ln \frac{Q}{T} \quad (2.17e)$$

Definition 2.14 (Total partition function and energy): As will be defined shortly, the total partition function is the product of all energy-type specific partition functions, i.e.:

$$Q = Q_{tr} Q_{rot} Q_{vib} Q_{el} = Q_{tr} \cdot \prod_{int} Q_{int} \quad (2.18)$$

The total internal energy of a molecule is:

$$\epsilon = \epsilon_{tr} + \epsilon_{rot} + \epsilon_{vib} + \epsilon_{el} \quad (2.19)$$

This also works for specific heat, c_v .

Definition 2.15 (Partition function for translation energy): The partition function for translation energy is:

$$Q_{tr} = V \left(\frac{2\pi mkT}{h^2} \right)^{\frac{3}{2}} \quad (2.20)$$

The translational internal energy can be found as:

$$E_{tr} = \frac{3}{2} NkT \quad e_{tr} = \frac{3}{2} RT \quad (2.21)$$

Where the first equation is per N molecules, and the second equation is on a per-unit mass basis. The specific heat is then:

$$c_{v_{tr}} = \left(\frac{\partial e_{tr}}{\partial T} \right)_v = \frac{3}{2} R \quad (2.22)$$

2.1 Electronic energies

Since monoatomic molecules do not have rotational or vibrational energies, the only internal energy is the electronic one. However, this still applies to diatomic molecules.

Definition 2.16 (Electronic internal energy): The electronic internal energy can largely be ignored at moderate temperatures for some species of gases. However, they can become important at higher temperatures. Because the spacing between the levels can be so large, usually only a few degenerate levels are accounted for. Electronic energy accounts for when electrons populate higher electronic energy levels.

Definition 2.17 (Electronic partition function): The electronic partition

function is:

$$Q_{el} = \sum_l g_l e^{-\epsilon_l/kT} = \sum_l g_l e^{\theta_l/T} \quad (2.23)$$

Where $\theta_l = \epsilon_l/k$ are the *characteristic temperatures for electronic excitation*. Energy of the ground level is taken to be 0, so the first term is just g_0 with no exponential. If we assume that just the first two terms are used, the internal energy becomes:

$$e_{el} = R\theta_1 \frac{(g_1/g_0)e^{-\theta_1/T}}{1 + (g_1/g_0)e^{-\theta_1/T}} \quad (2.24)$$

and the specific heat:

$$c_{v_{el}} = R \left(\frac{\theta_1}{T} \right)^2 \frac{(g_1/g_0)e^{-\theta_1/T}}{[1 + (g_1/g_0)e^{-\theta_1/T}]^2} \quad (2.25)$$

2.2 Diatomic internal energies

3 High Temperature Transport Models

References for this section

1. *A Review of Reaction Rates and Thermodynamic and Transport Properties for an 11-Species Air Model for Chemical and Thermal Nonequilibrium Calculations to 30,000 K*, Gupta, Yos, Thompson, Lee, 1990 NASA RP 1232.
2. *An assessment of Transport Property Methodologies for Hypersonic Flows*, Palmer, 1971, AIAA.

3.1 Collision integral definitions

Definition 3.1 (Collision cross sections): The collision cross section has infinitely many values. Four of them are used for fluid dynamics to determine transport properties. The collision cross sections are, in general, the weighted averages of the cross sections for collisions between species i and j (sometimes denoted s and r). They are defined as:

$$\pi\bar{\Omega}_{ij}^{(l,s)} = \frac{\int_0^\infty \int_0^\pi \exp(-\gamma^2) \gamma^{2s+3} (1 - \cos^l \chi) 4\pi \sigma_{ij} \sin \chi d\chi d\gamma}{\int_0^\infty \int_0^\pi \exp(-\gamma^2) \gamma^{2s+3} (1 - \cos^l \chi) \sin \chi d\chi d\gamma} \quad (3.1)$$

Variables are:

- $\sigma_{ij} = \sigma_{ij}(\chi, g)$ is the differential-scattering cross section for the pair (i, j) .
- χ is the scattering angle in the center of mass system.
- g is the relative velocity of the colliding particles.
- γ is the reduced velocity equal to $\left(\left[\frac{m_i m_j}{2(m_i + m_j) k T} \right]^{\frac{1}{2}} \right) g$

In the collision integral symbol, l controls how strong the contribution of the differential-scattering angle (χ) is. s is a weighting on the reduced velocity (γ) of the particles. Of course, the subscript (ii) refers to the collision integral for similar particles. Here is a list of where certain collision integrals show up:

- Viscosity (μ) is a function of $\pi\bar{\Omega}_{ij}^{(2,2)}$ only.
- Translational component of thermal conductivity (κ_{tr}) is a function of $\pi\bar{\Omega}_{ij}^{(2,2)}$ only.

- The internal component of thermal conductivity (κ_{int}) is a function of $\pi\bar{\Omega}_{ij}^{(1,1)}$ only or a function of $\pi\bar{\Omega}_{ij}^{(1,1)}$ and $\pi\bar{\Omega}_{ij}^{(2,2)}$, depending on the expression used.
- The diffusion coefficient D_{ij} is a function of $\pi\bar{\Omega}_{ij}^{(1,1)}$ only.

Definition 3.2 (Gupta's curve fits for collision integrals): Gupta utilizes curve fits for the collision integrals. They have the following form:

$$\pi\bar{\Omega}_{ij}^{(l,s)} = [\exp D] T^{[A(\ln T)^2 + B \ln T + C]} \quad (3.2)$$

$$B_{i,j}^{*(2,2)} = [\exp C] T^{[A(\ln T)^2 + B \ln T]} \quad (3.3)$$

Definition 3.3 (Chapman-Enskog procedure):

3.2 Transport properties of a single species

Definition 3.4 (Viscosity of a single species): The viscosity of a gas containing a single species is:

$$\mu_i = \frac{5}{16} \frac{\sqrt{\pi m_i k T}}{\pi\bar{\Omega}_{ii}^{(2,2)}} = 8.3861 \cdot 10^{-6} \frac{\sqrt{\mathcal{M}_i T}}{\pi\bar{\Omega}_{ii}^{(2,2)}} \quad \text{units} = \frac{\text{N} \cdot \text{s}}{\text{m}^2} \quad (3.4)$$

Where \mathcal{M}_i is the molecular weight of species i , and T is the temperature in Kelvin.

Definition 3.5 (Thermal conductivity of a single species): For the thermal conductivity of a single species, we have a number of contributions. The total κ of the species is the sum of its internal energy specific contributions, i.e.:

$$\kappa = \kappa_{\text{tr}} + \kappa_{\text{rot}} + \kappa_{\text{vib}} + \kappa_{\text{el}} = \kappa_{\text{tr}} + \kappa_{\text{int}} \quad (3.5)$$

Where the internal energies that are not translational have been grouped into κ_{int} .

$$\kappa_{\text{tr},i} = \frac{5}{2} \frac{\mu_i}{\mathcal{M}_i} (c_{v,i})_{\text{tr}} = \frac{15}{4} \frac{\hat{R}}{\mathcal{M}_i} \mu_i \quad \text{units} = \frac{\text{J}}{\text{m-s-K}} \quad (3.6)$$

$$\kappa_{\text{int},i} = \frac{6}{5} \frac{c_{p_{\text{int},i}}}{\hat{R}} \frac{\hat{R}}{\mathcal{M}_i} \mu_i \frac{\pi\bar{\Omega}_{ii}^{(2,2)}}{\pi\bar{\Omega}_{ii}^{(1,1)}} \quad (3.7)$$

Here, $c_{p_{int,i}}$ is the total specific heat with the translation specific heat subtracted.

3.3 Transport properties of a mixture

For species summation and Wilke's mixing rule, μ_j and κ_j are calculated using the above equations for single species viscosity and conductivity.

Definition 3.6 (Species summation): The concept of the species summation version of finding mixture transport properties is quite simple. It assumes no interaction between species and is expressed as:

$$\mu = \sum_{j=1}^{NS} X_j \mu_j \quad \kappa = \sum_{j=1}^{NS} X_j \kappa_j \quad (3.8)$$

Definition 3.7 (Wilke's mixing rule): Both viscosity and thermal conductivity can be approximated using Wilke's mixing rule. It uses a quantity, ϕ_i , which is expressed as:

$$\phi_i = \sum_{j=1}^{NS} \frac{X_j \left[1 + \sqrt{\frac{\mu_i}{\mu_j}} \left(\frac{\mathcal{M}_j}{\mathcal{M}_i} \right)^{\frac{1}{4}} \right]^2}{\sqrt{8 \left(1 + \frac{\mathcal{M}_i}{\mathcal{M}_j} \right)}} = \sum_{j=1}^{NS} \frac{X_j \left[1 + \sqrt{\frac{\pi \bar{\Omega}_{jj}^{(2,2)}}{\pi \bar{\Omega}_{ii}^{(2,2)}}} \right]^2}{\sqrt{8 \left(1 + \frac{\mathcal{M}_i}{\mathcal{M}_j} \right)}} \quad (3.9)$$

Then the viscosity and thermal conductivity is found as:

$$\mu = \sum_{j=1}^{NS} \frac{X_j \mu_j}{\phi_j} \quad \kappa = \sum_{j=1}^{NS} \frac{X_j \kappa_j}{\phi_j} \quad (3.10)$$

Definition 3.8 (Chapman-Enskog approximation method): For obtaining mixture transport quantities using Chapman-Enskog approximations, a number of new variables are used. This is the most rigorous method for finding the properties. The units of Δ is in centimeters-second, while α_{ij} is unitless.

$$\Delta_{ij}^{(1)} = \frac{8}{3} (1.5460 \cdot 10^{-20}) \left[\frac{2\mathcal{M}_i \mathcal{M}_j}{\pi N_A k T (\mathcal{M}_i + \mathcal{M}_j)} \right]^{\frac{1}{2}} \pi \bar{\Omega}_{ij}^{(1,1)} \quad (3.11)$$

$$\Delta_{ij}^{(2)} = \frac{16}{5} (1.5460 \cdot 10^{-20}) \left[\frac{2\mathcal{M}_i \mathcal{M}_j}{\pi N_A k T (\mathcal{M}_i + \mathcal{M}_j)} \right]^{\frac{1}{2}} \pi \bar{\Omega}_{ij}^{(2,2)} \quad (3.12)$$

For electron-electron collisions, we have:

$$\pi\bar{\Omega}_{ee}^{(2,2)} = 1.29Q_e \cdot 10^{16} \quad (3.13)$$

For ion-ion collisions, we have:

$$\pi\bar{\Omega}_{II}^{(2,2)} = 1.36Z^4Q_e \cdot 10^{16} \quad (3.14)$$

Then for both:

$$\pi\bar{\Omega}_{ee}^{(1,1)} = \pi\bar{\Omega}_{II}^{(1,1)} = 0.795Z^4Q_e \cdot 10^{16} \quad (3.15)$$

Here, $Z = 1$ for singly ionized species (for aerospace applications this is fine), and:

$$Q_e = \frac{e^4}{(kT)^2} \ln \Lambda \quad \text{cm}^2 \quad (3.16)$$

$$\Lambda = \left[\frac{9(kT)^3}{4\pi e^6 n_e} + \frac{16(kT)^2}{e^4 n_e^{2/3}} \right]^{1/2} \quad (3.17)$$

$$= \left[2.09 \cdot 10^{-2} \left(\frac{T^4}{10^{12} p_e} \right) + 1.52 \left(\frac{T^4}{10^{12} p_e} \right)^{2/3} \right]^{1/2} \quad (3.18)$$

Here, p_e is the electron pressure in atmospheres, n_e is the number density of electrons, and $e = 4.8 \cdot 10^{-10}$ esu. These equations are only useful under for $p_e <$ some limiting value, which should be true in aerospace applications:

$$p_{em} = 0.0975 \left(\frac{T}{10^3} \right)^4 \quad (3.19)$$

Depending on the data given, the Coulomb cross section may need to be altered. Gupta and Yos give collision cross sections at the above limiting electron pressure, so at any other pressure the cross section needs to be changed:

$$\frac{\pi \bar{\Omega}_{ij}^{(l,s)}(p_e)}{\pi \bar{\Omega}_{ij}^{(l,s)}(p_{em})} = \ln \Lambda(p_e) \quad (3.20)$$

$$= \frac{1}{2} \ln \left[2.09 \cdot 10^{-2} \left(\frac{T}{1000 p_e^{1/4}} \right)^4 + 1.52 \left(\frac{T}{1000 p_e^{1/4}} \right)^{8/3} \right] \quad (3.21)$$

Finally, α_{ij} is needed for the heavy-particle translational component of thermal conductivity:

$$\alpha_{ij} = 1 + \frac{[1 - (\mathcal{M}_i/\mathcal{M}_j)][0.45 - 2.54(\mathcal{M}_i/\mathcal{M}_j)]}{[1 + (\mathcal{M}_i/\mathcal{M}_j)]^2} \quad (3.22)$$

Assuming thermochemical nonequilibrium, the mixture viscosity *appears* to not be a function of vibrational or rotation energy, and is thus found with:

$$\mu = \sum_{i=1}^{NS-1} \left[\frac{\frac{\mathcal{M}_i}{N_A} X_i}{\sum_{j=1}^{NS-1} X_j \Delta_{ij}^{(2)}(T) + X_e \Delta_{ie}^{(2)}(T_e)} \right] + \frac{\frac{\mathcal{M}_e}{N_A} X_e}{\sum_{j=1}^{NS} X_j \Delta_{ej}^{(2)}(T_e)} \quad (3.23)$$

Where X_j is the molar fraction, T is the Temperature, and T_e is the electron temperature. In contrast to viscosity, the thermal conductivity is dependant on all energy modes, and therefore has contributions from each mode. They are:

Translational component without electron-heavy-particle and electron-electron collision:

$$K_{tr}^* = (2.3901 \cdot 10^{-8}) \frac{15}{4} k \sum_{i=1}^{NS-1} \left[\frac{X_i}{\sum_{j=1}^{NS-1} \alpha_{ij} \Delta_{ij}^{(2)}(T) + 3.54 X_e \Delta_{ie}^{(2)}(T_e)} \right] \quad (3.24)$$

Rotational component. "p.e." means "partial excitation" while "f.e." mean "full excitation". If it is in equilibrium with translational temperature, $T_{rot} = T$:

$$(K_{rot})_{p.e.} = (2.3901 \cdot 10^{-8}) k \sum_{i=mol.} \left[\frac{\left[\frac{c_{p,i}(T_{rot})}{\hat{R}} - \frac{5}{2} \right] X_i}{\sum_{j=1}^{NS-1} X_j \Delta_{ij}^{(1)}(T) + X_e \Delta_{ie}^{(1)}(T_e)} \right] \quad (3.25)$$

$$(K_{\text{rot}})_{\text{f.e.}} = (2.3901 \cdot 10^{-8})k \sum_{i=\text{mol.}} \left[\frac{X_i}{\sum_{j=1}^{\text{NS-1}} X_j \Delta_{ij}^{(1)}(T) + X_e \Delta_{ie}^{(1)}(T_e)} \right] \quad (3.26)$$

Vibrational component:

$$(K_{\text{vib}})_{\text{p.e.}} = (2.3901 \cdot 10^{-8})k \sum_{i=\text{mol.}} \left[\frac{\left[\frac{c_{p,i}(T_{\text{vib}})}{\hat{R}} - \frac{7}{2} \right] X_i}{\sum_{j=1}^{\text{NS-1}} X_j \Delta_{ij}^{(1)}(T) + X_e \Delta_{ie}^{(1)}(T_e)} \right] \quad (3.27)$$

$$(K_{\text{vib}})_{\text{f.e.}} = (2.3901 \cdot 10^{-8})k \sum_{i=\text{mol.}} \left[\frac{X_i}{\sum_{j=1}^{\text{NS-1}} X_j \Delta_{ij}^{(1)}(T) + X_e \Delta_{ie}^{(1)}(T_e)} \right] \quad (3.28)$$

Electronic component:

$$K_{\text{el}} = (2.3901 \cdot 10^{-8})k \sum_{i=1}^{\text{NS-1}} \left[\frac{\left[\frac{c_{p,i}(T_{\text{el}})}{\hat{R}} - \frac{9}{2} \right] X_i}{\sum_{j=1}^{\text{NS-1}} X_j \Delta_{ij}^{(1)}(T) + X_e \Delta_{ie}^{(1)}(T_e)} \right] \quad (3.29)$$

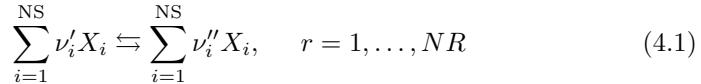
Free electron component:

$$K_{\text{e}} = (2.3901 \cdot 10^{-8})k \frac{X_e}{\sum_{j=1}^{\text{NS-1}} X_j \Delta_{ej}^{(2)}(T_e) + X_e \Delta_{ee}^{(2)}(T_e)} \quad (3.30)$$

4 Chemical Rate Kinetics

4.1 Rate equations

Definition 4.1 (Reactions): For multicomponent gas with NS reacting species and NR different reactions, the stoichiometric relations for the change in reactants and products are:



ν'_i and ν''_i are the stoichiometric coefficients for reactants and products respectively. $k_{f,r}$ governs the forward reaction (right arrow), and $k_{b,r}$ governs the backward reaction (left arrow).

Definition 4.2 (Mass production): The net rate of MASS production of a species per unit volume has a number of different expressions. Consider the reaction:



Here, M is an arbitrary collision partner, deemed a *third-body*. Concentration of species j is the moles of j per unit volume. $[O_2]$ is the concentration of O_2 . The time rate of change of the concentration of O is then:

$$\frac{d[O]}{dt} = 2k[O_2][M] \quad (4.3)$$

Where k is the reaction rate. This equation gives the rate at which Eq. [4.2] goes from left to right. Therefore, k is actually the *forward* reaction rate, k_f . The two previous equations should then be:



$$\frac{d[O]}{dt} = 2k_f[O_2][M] \quad (4.5)$$

Of course, the right to left reaction is the *backwards* reaction and is governed by the backwards rate coefficient:



$$\frac{d[O]}{dt} = -2k_b[O]^2[M] \quad (4.7)$$

You must combine these forward and backwards equations because both reactions happen. This gives:



$$\frac{d[O]}{dt} = 2k_f[O_2][M] - 2k_b[O]^2[M] \quad (4.9)$$

The net rate equation for an arbitrary reaction is then:

$$\frac{d[X_i]}{dt} = (\nu''_i - \nu'_i) \left\{ k_f \prod_i [X_i]^{\nu'_i} - k_b \prod_i [X_i]^{\nu''_i} \right\} \quad (4.10)$$

Where ν' is the stoichiometric for a reactant species, and ν'' is for product species.

Definition 4.3 (Reaction rate relations): The equilibrium constant is very useful to find one reaction rate from the other. There are two equilibrium constants: K_c is the equilibrium constant based on concentrations, and K_p is the equilibrium constant based on partial pressures. They are related through:

$$K_c = \left(\frac{1}{RT} \right)^{\sum_i \nu_i} K_p \quad (4.11)$$

Where it is raised to a sum due to the complicated nature of the equilibrium constant. Typically, the rate coefficient has variable units, since it depends on the number of molecules in the reaction, which gives rise to a key concept:

For most chemical kinetic models, the reaction's reactants always consist of 2 species, while the reaction's products may contain two or three species.

Therefore, the forwards rate coefficients k_f should always have the same units. Since the equilibrium constant is used to find k_b , no unit conversions are necessary besides perhaps from cgs to mks units.

Curve fits of the following form are used to find the equilibrium constant:

$$(K_{\text{eq,r}}) = \exp(\beta)$$

$$\beta = A_1 \frac{T}{10^4} + A_2 + A_3 \ln \frac{10^4}{T} + A_4 \frac{10^4}{T} + A_5 \left(\frac{10^4}{T} \right)^2$$

With the equilibrium constant of a reaction known, you can find the forward/backward reaction rates as stated above with:

$$\frac{k_{f,r}(T)}{k_{b,r}(T)} = K_{\text{eq,r}} \quad (4.12)$$

Definition 4.4 (Arrhenius equations): Both forward and backward rate coefficients can be found using the Arrhenius or modified Arrhenius equations, though usually the forward rate is computed and the backward rate is found with the equilibrium constant. The Arrhenius equation has the form:

$$k_{f,r} = C e^{-E_a/kT} \quad (4.13)$$

Where C is a constant, E_a is the activation energy, k is the Boltzmann constant, and T is the temperature. The modified Arrhenius form (more widely used) is:

$$k_{f,r} = CT^\eta e^{-E_0/kT} \quad (4.14)$$

Here, η is a reaction specific exponent, and E_0 depends on the reaction (could be a dissociation energy, could be 0 for recombination, etc.)

4.2 Reaction Types

Definition 4.5 (Dissociation Reaction): A dissociation reaction occurs when a molecule collides with some other species and one or more atoms "detatch" from the molecule. Examples are:



Eq. [4.18] is an example of a *bimolecular exchange* reaction. Eq. [4.21] is called a *dissociative-recombination* reaction since the recombination of the NO^+ ion with an electron produces a dissociated product.

To be continued...

Definition 4.6 (Efficiencies): For certain reactions like dissociation, the outcome of the reaction is the same regardless of the other reactant. These are called *third-body reactions*, since the secondary reactant does not change chemically. The third-body is usually represented with an M as we have already discussed:



This M can be any species, and the Arrhenius equation is nearly identical for any M , with only a slight change in the leading coefficient C . This coefficient change is called an *efficiency* (denoted here as ϵ), and the algebra will be shown as to how this is beneficial to use. A list of efficiencies in certain reactions are shown in Table 1.

Table 1: Change in modified Arrhenius constant C for some third-bodies in N_2 dissociation.

Third body	C	ϵ
$M = O_2$	7.0e21	1.0
$M = NO$	7.0e21	1.0
$M = O$	3.0e22	4.2857

Since $7.0e21$ is the smallest value, this is given an efficiency of 1.0, and then

any other reaction's efficiency is given as some multiplier of this base value. The physical meaning of efficiencies is how "efficient" that third body is at making that reaction happen. ϵ is larger for atoms, since they are more efficient in sending their energy into dissociation of a molecule. We can now do some algebra on the Arrhenius equation by letting $C = \epsilon C_c$, then $\gamma = C_c T^\eta e^{E_0/kT}$. The forward and backwards reaction constants are then:

$$k_f = \epsilon\gamma, \quad k_b = \frac{\epsilon\gamma}{K_{eq}} \quad (4.23)$$

For a third body reaction, the change in the concentration of N_2 due to a single third-body M is then:

$$\begin{aligned} \frac{d[N_2]}{dt} &= (0 - 1) \left\{ \epsilon\gamma[N_2][M] - \frac{\epsilon\gamma}{K_{eq}}[N]^2[M] \right\} \\ &= -1 \cdot \epsilon[M] \underbrace{\left\{ \gamma[N_2] - \frac{\gamma}{K_{eq}}[N]^2 \right\}}_R \end{aligned}$$

R is independant of the third-body, so it is a constant and only needs to be computed once. Summing over contributions from all independant reactions gives:

$$\frac{d[N_2]}{dt} = - \left[\sum_i \epsilon_i [M_i] \right] R$$

Where:

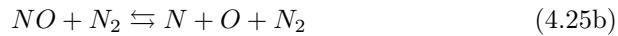
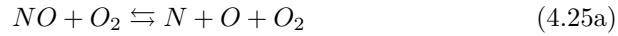
$$\sum_i \epsilon_i [M_i] = \epsilon_{N_2}[N_2] + \epsilon_{O_2}[O_2] + \epsilon_{NO}[NO] + \epsilon_N[N] + \epsilon_O[O] + \dots \quad (4.24)$$

R is retained for the production of N atoms as well, you only need to change the coefficient from the $\nu''_i - \nu'_i$ term:

$$\frac{d[N]}{dt} = \sum_i \epsilon_i [M_i] \underbrace{(2 - 0)}_{\nu''_N - \nu'_N} R$$

4.3 Example rate equation solution

Here, an example is done to find the rate of production of NO . Consider the following reactions that all contain NO :



The forward and backward rate coefficients for the reactions are $k_{f,a}$ and $k_{b,a}$ for the first reaction, all the way down with consistent lettering with the equation number. The entire equation is then:

$$\begin{aligned} \frac{d[NO]}{dt} = & - \left\{ \gamma[NO] - \frac{\gamma}{K_{eq}}[N][O] \right\} \sum_i \epsilon_i[M] \\ & + \{k_{f,f}[O_2][N] - k_{b,f}[NO][O]\} \\ & + \{k_{f,g}[N_2][O] - k_{b,g}[NO][N]\} \end{aligned}$$

5 Internal energy relaxation

5.1 Translational-Vibrational relaxation

The source term for vibrational energy due to transfer of energy between translational and vibrational energy modes from the Landau-Teller model is:

$$Q_{t-v}^{\text{LT}} = \sum_s Q_{t-v,s}^{\text{LT}} \quad (5.1)$$

Where

$$Q_{t-v,s}^{\text{LT}} = \rho_s \frac{e_{vs}^* - e_{vs}}{\langle \tau_s \rangle} \quad (5.2)$$

e_{vs}^* is the equilibrium vibrational energy, which is evaluated at the tran-rotational temperature, T_{tr} , while e_{vs} is the actual vibrational energy evaluated with T_v . The molar-averaged relaxation time, $\langle \tau_s \rangle$ is defined by:

$$\langle \tau_s \rangle = \frac{\sum_r X_r}{\sum_r X_r / \tau_{sr}} \quad (5.3)$$

This sum happens for all particles r , and molecules s . The term τ_{sr} is the relaxation time between species and has two parts - the Millikan-White part, as well as the high-temperature Park correction. The Millikan-White part uses the equations:

$$\tau_{sr}^{\text{MW}} = \frac{101325}{p} \exp [A_{sr}(T^{-1/3} - B_{sr}) - 18.42] \quad (5.4)$$

Where A_{sr} and B_{sr} are either set constants or computed from:

$$\mu_{sr} = \frac{\mathcal{M}_s \mathcal{M}_r}{\mathcal{M}_s + \mathcal{M}_r} \quad (5.5)$$

$$A_{sr} = 1.16 \times 10^{-3} \mu_{sr}^{1/2} \theta_{vs}^{4/3} \quad (5.6)$$

$$B_{sr} = 0.015 \mu_{sr}^{1/4} \quad (5.7)$$

The Park correction is:

$$\tau_{sr}^P = \frac{1}{\bar{c}\sigma n} \quad (5.8)$$

$$\bar{c} = \sqrt{\frac{8\hat{R}T}{\pi}} \quad (5.9)$$

$$\sigma = 3 \times 10^{-21} \left(\frac{50000}{T} \right)^2 \quad (5.10)$$

$$n = \frac{p}{k_b T} \quad (5.11)$$

5.2 Chemical-Vibrational relaxation

6 Thermochemical Nonequilibrium CFD

6.1 Species conservation equation

$$\underbrace{\frac{\partial}{\partial t} \rho_s}_{1} + \underbrace{\frac{\partial}{\partial x_j} \rho_s u_j}_{2} = \underbrace{\frac{\partial}{\partial x_j} (\rho D_s \frac{\partial}{\partial x_j} y_s)}_{3} + \underbrace{\dot{\omega}_s}_{4} \quad (6.1)$$

The terms are:

1. Rate of change of mass of species s .
2. Flux of mass due to convection.
3. Molecular diffusion flux of mass.
4. Chemical production rate.

The variables are:

- ρ_s is the density of species s
- u_j is the mass-averaged velocity in the j -direction
- D_s is the effective diffusion coefficient of species s
- y_s is the molar fraction of species s
- $\dot{\omega}_s$ is the chemical production rate of species s

6.2 Total continuity equation

For chemically reacting flows, the total continuity equation is unaffected. It is:

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{u}) = 0 \quad (6.2)$$

Since no mass can be created or destroyed, the condition of atomic conservation is applied, which yields:

$$\sum_s \dot{w}_s = 0 \quad (6.3)$$

6.3 Mixture momentum conservation

$$\underbrace{\frac{\partial}{\partial t} \rho u_i}_{1} + \underbrace{\frac{\partial}{\partial x_j} \rho u_i u_j}_{2} = - \underbrace{\frac{\partial p}{\partial x_i}}_{3} + \underbrace{\frac{\partial}{\partial x_j} \left[\mu \left(\frac{\partial u_i}{\partial x_j} + \frac{\partial u_j}{\partial x_i} \right) - \frac{2}{3} \mu \frac{\partial u_k}{\partial x_k} \delta_{ij} \right]}_{4} \quad (6.4)$$

The terms are:

1. Rate of change of i th component of momentum.
2. Flux of i th component of momentum due to convection.
3. Pressure forces in i th direction.
4. Viscous terms acting in i th direction.

Here, the mixture pressure is defined with Dalton's law of partial pressures, and the pressure of an individual heavy species is:

$$p_s = \frac{\rho_s \hat{R} T}{M_s} \quad (6.5)$$

For electron pressure, we get:

$$p_e = \frac{\rho_e \hat{R} T_e}{M_e} \quad (6.6)$$

Where the e subscript denotes an electron property.

6.4 Vibrational energy conservation

$$\begin{aligned} \underbrace{\frac{\partial}{\partial t} \rho e_v}_{1} + \underbrace{\frac{\partial}{\partial x_j} \rho e_v u_j}_{2} &= \underbrace{\frac{\partial}{\partial x_j} \left(\eta_v \frac{\partial T_v}{\partial x_j} \right)}_{3} + \underbrace{\frac{\partial}{\partial x_j} \left(\rho \sum_{s=1}^{NS} h_{v,s} D_s \frac{\partial y_s}{\partial x_j} \right)}_{4} \\ &+ \underbrace{\sum_{s=\text{mol.}} \rho_s \frac{(e_{v,s}^* - e_{v,s})}{<\tau_s>}}_{5} + \underbrace{\sum_{s=\text{mol.}} \rho_s \frac{(e_{v,s}^{**} - e_{v,s})}{<\tau_{es}>}}_{6} + \underbrace{\sum_{s=\text{mol.}} \dot{w}_s \hat{D}_s}_{7} \quad (6.7) \end{aligned}$$

The terms are:

1. Rate of change of vibrational energy per unit volume.
2. The flux of vibrational energy due to convection.

3. The conduction of vibrational energy due to vibrational temperature gradients.
4. The diffusion of vibrational energy due to molecular concentration gradients.
5. The energy exchange (relaxation) between vibrational and translational modes due to collisions within the cell.
6. The energy exchange between vibrational and electronic modes.
7. The vibrational lost/gained due to molecular depletion (dissociation) or production (recombination).

The variables are:

- e_v is the vibrational energy per unit mass, defined as $e_v = \sum_{s=1}^{NS} \frac{\rho_s e_{v,s}}{\rho}$. e_v is defined as a function of T_v , the vibrational temperature.
- η_v is the vibrational thermal conductivity.
- $h_{v,s}$ is the vibrational enthalpy. It is identical to $e_{v,s}$.
- $e_{v,s}^*$ is the vibrational energy of species s at the translational-rotational temperature.
- $e_{v,s}^{**}$ is the vibrational energy at the electron temperature.
- $\langle \tau_s \rangle$ is the characteristic relaxation times for translational-vibrational (T-V) energy exchange.
- $\langle \tau_{es} \rangle$ is the characteristic relation times for electron-vibrational (e-V) energy exchange.
- \hat{D}_s is the vibration energy level representative of those molecules of species s which are created or destroyed (recombined or dissociated) because of their high vibrational quantum numbers.

6.5 Electron and electronic excitation energy conservation

$$\begin{aligned}
& \underbrace{\frac{\partial}{\partial t} \rho e_e + \frac{\partial}{\partial x_j} [u_j (\rho e_e + p_e)]}_{1} = \underbrace{u_j \frac{\partial p_e}{\partial x_j}}_{2} + \underbrace{\frac{\partial}{\partial x_j} \left(\eta_e \frac{\partial T_e}{\partial x_j} \right)}_{4} + \underbrace{\frac{\partial}{\partial x_j} \left(\rho \sum_{s=1}^{NS} h_{e,s} D_s \frac{\partial y_s}{\partial x_j} \right)}_{5} \\
& + \underbrace{2\rho_e \frac{3}{2} \hat{R}(T - T_e) \sum_{s=1}^{NS-1} \frac{\nu_{es}}{\mathcal{M}_s}}_{6} - \underbrace{\sum_{s=\text{ion}} \dot{n}_{e,s} \hat{I}_s}_{7} - \underbrace{\sum_{s=\text{mol.}} \rho_s \frac{e_{v,s}^{**} - e_{v,s}}{\langle \tau_{es} \rangle}}_{8} - Q_{\text{rad}} \quad (6.8)
\end{aligned}$$

Terms are:

1. Rate of change of electronic energy per unit volume.
2. The flux of electronic enthalpy due to convection.
3. The work done on electrons by an electron field by an electron pressure gradient.
4. The conduction of electronic energy due to electron temperature gradient.
5. The diffuction of electronic energy due to concentration gradients.
6. The energy exchange due to elastic collisions between electrons and heavy particles.
7. Energy loss due to electron impact ionization (note the sum indices sum over ionized species only).
8. The relaxation due to inelastic collisions between electrons and molecules.
9. Rate of energy loss due to radiation caused by electronic transitions.

The variables are:

- e_e is the electronic energy per unit mass and contains contributions from NS electronic energy levels. It is defined as $e_e = \sum_{s=1}^{NS} \frac{\rho_s e_{e,s}}{\rho}$
- $e_{e,s}$ is the electronic energy per unit mass of species s and is defined as a function of T_e .

- T_e is the electron temperature.
 - η_e is the electronic thermal conductivity.
 - $h_{e,s}$ is the electronic enthalpy per unit mass and is identical to $e_{e,s}$ for all species except free electrons. For free electrons it is defined as:
- $$h_{e,e} = e_{e,e} + \frac{\hat{R}T_e}{\mathcal{M}_e}$$
- $\nu_{e,s}$ is the effective collision frequency of electrons with heavy particles.
 - $\dot{n}_{e,s}$ is the molar rate of production of species s by electron impact ionizations.
 - \hat{I}_s is the energy per unit mole lost by a free electron in producing species s through electron impact ionization.
 - Q_{rad} is the radiant energy transfer rate due to electron transitions.

6.6 Total energy conservation

$$\underbrace{\frac{\partial}{\partial t} \rho E}_{1} + \underbrace{\frac{\partial}{\partial x_j} \rho H u_j}_{2} = \underbrace{\frac{\partial}{\partial x_j} \left(\eta \frac{\partial T}{\partial x_j} + \eta_v \frac{\partial T_v}{\partial x_j} + \eta_e \frac{\partial T_e}{\partial x_j} \right)}_{3} + \underbrace{\frac{\partial}{\partial x_j} \left(\rho \sum_{s=1}^{\text{NS}} h_s D_s \frac{\partial y_s}{\partial x_j} \right)}_{4} + \underbrace{\frac{\partial}{\partial x_j} \left[u_i \mu \left(\frac{\partial u_i}{\partial x_j} + \frac{\partial u_j}{\partial x_i} \right) \right]}_{5} - \underbrace{Q_{\text{rad}}}_{6} \quad (6.9)$$

The terms are:

1. Rate of change of total energy per unit volume.
2. The flux of total enthalpy due to convection.
3. The conduction of energy due to temperature, vibrational temperature, and electron temperature gradients.
4. The diffusion of energy due to concentration gradients.
5. The work done by shear forces.
6. The rate of energy loss due to radiation caused by electronic transitions.

The variables are:

- E is the total energy per unit volume, defined as $E = \frac{u_i u_i}{2} + \sum_{s=1}^{NS} \frac{\rho_s e_s}{\rho}$
- H is the total enthalpy per unit mass, defined as $H = E + \frac{p}{\rho}$
- η is the frozen thermal conductivity of heavy particles. The part in which exchanges of translational and rotational energy occur.

6.7 Two temperature model

Everything before this assumes a three-temperature model, where there is a single temperature dedicated to translational-rotational energy, vibrational energy, and electronic-electron energies. The two temperature model assumes that one temperature can be used for describing the energy of translational-rotational energies, and then the rest (vib-electronic-electron) are described by the second temperature, T_V . Therefore:

$$T_v = T_e = T_V \quad (6.10)$$

By combining the vibrational and electronic energy equations, we get the vibrational-electronic energy conservation equation:

$$\begin{aligned} & \underbrace{\frac{\partial}{\partial t} \rho e_V}_{1} + \underbrace{\frac{\partial}{\partial x_j} \rho e_V u_j}_{2} = \underbrace{-p_e \frac{\partial u_j}{\partial x_j}}_{3} + \underbrace{\frac{\partial}{\partial x_j} \left[(\eta_v + \eta_e) \frac{\partial T_V}{\partial x_j} \right]}_{4} \\ & \quad + \underbrace{\frac{\partial}{\partial x_j} \left(\rho \sum_{s=1}^{NS} h_{V,s} D_s \frac{\partial y_s}{\partial x_j} \right)}_{5} + \underbrace{\sum_{s=mol.} \rho_s \frac{(e_{v,s}^* - e_{v,s})}{<\tau_s>}}_{6} \\ & \quad + \underbrace{2\rho_e \frac{3}{2} \hat{R}(T - T_V) \sum_{s=1}^{NS-1} \frac{\nu_{es}}{\mathcal{M}_s}}_{7} - \underbrace{\sum_{s=ion} \dot{n}_{e,s} \hat{I}_s}_{8} + \underbrace{\sum_{s=mol.} \dot{w}_s \hat{D}_s}_{9} - Q_{rad} \quad (6.11) \end{aligned}$$

Where:

$$e_V = e_v + e_e \quad (6.12)$$

$$h_{V,s} = h_{v,s} h_{e,s} \quad (6.13)$$

6.8 Derivatives

6.8.1 Partial derivatives of pressure (P)

$$\frac{\partial p}{\partial \rho_s} = \left(\frac{\hat{R}}{\mathcal{M}_s} - c_{vs} \frac{R}{c_v} \right) T + \frac{R}{c_v} \left[\frac{1}{2}(u^2 + v^2 + w^2) - h_s^\circ \right] \quad (6.14a)$$

$$\frac{\partial p}{\partial (\rho u)} = -u \frac{R}{c_v} \quad (6.14b)$$

$$\frac{\partial p}{\partial (\rho v)} = -v \frac{R}{c_v} \quad (6.14c)$$

$$\frac{\partial p}{\partial (\rho w)} = -w \frac{R}{c_v} \quad (6.14d)$$

$$\frac{\partial p}{\partial E_v} = -\frac{R}{c_v} \quad (6.14e)$$

$$\frac{\partial p}{\partial E} = \frac{R}{c_v} \quad (6.14f)$$

6.8.2 Partial derivatives of total energy (E)

$$\frac{\partial E}{\partial \rho_s} = \left(c_{vs} - \frac{c_v}{R} \frac{\hat{R}}{\mathcal{M}_s} \right) T + \frac{1}{2}(u^2 + v^2 + w^2) + e_v + h_s^\circ \quad (6.15a)$$

$$\frac{\partial E}{\partial u} = \rho u \quad (6.15b)$$

$$\frac{\partial E}{\partial v} = \rho v \quad (6.15c)$$

$$\frac{\partial E}{\partial w} = \rho w \quad (6.15d)$$

$$\frac{\partial E}{\partial e_v} = \sum_{s=1}^{NS} \rho_s \quad (6.15e)$$

$$\frac{\partial E}{\partial p} = \frac{c_v}{R} \quad (6.15f)$$

6.8.3 Partial derivatives of T

$$\frac{\partial T}{\partial \rho_s} = \frac{1}{\rho c_v} \left(-c_{vs}T + \frac{1}{2}(u^2 + v^2 + w^2) - h_s^\circ \right) \quad (6.16a)$$

$$\frac{\partial T}{\partial (\rho u)} = -\frac{u}{\rho c_v} \quad (6.16b)$$

$$\frac{\partial T}{\partial (\rho v)} = -\frac{v}{\rho c_v} \quad (6.16c)$$

$$\frac{\partial T}{\partial (\rho w)} = -\frac{w}{\rho c_v} \quad (6.16d)$$

$$\frac{\partial T}{\partial E_v} = -\frac{1}{\rho c_v} \quad (6.16e)$$

$$\frac{\partial T}{\partial E} = \frac{1}{\rho c_v} \quad (6.16f)$$

$$\frac{\partial c_r}{\partial \rho_s} = (\delta_{rs} - c_r) \frac{1}{\rho} \quad (6.16g)$$

6.8.4 Partial derivatives of vibrational temperature (T_v)

$$\frac{\partial T_v}{\partial \rho_i} = -e_{vi} / \sum_{s=1}^{NS} \left[\rho_s \left(\frac{\partial e_{vs}}{\partial T_v} \right) \right] \quad (6.17a)$$

$$\frac{\partial T_v}{\partial (\rho u)} = 0 \quad (6.17b)$$

$$\frac{\partial T_v}{\partial (\rho v)} = 0 \quad (6.17c)$$

$$\frac{\partial T_v}{\partial (\rho w)} = 0 \quad (6.17d)$$

$$\frac{\partial T_v}{\partial E_v} = 1 / \sum_{s=1}^{NS} \left[\rho_s \left(\frac{\partial e_{vs}}{\partial T_v} \right) \right] \quad (6.17e)$$

$$\frac{\partial T_v}{\partial E} = 0 \quad (6.17f)$$

6.8.5 Partial derivatives of vibrational energy (e_{vs})

$$\frac{\partial e_{vs}}{\partial \rho_i} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial \rho_i}, \quad \frac{\partial e_{vs}^*}{\partial \rho_i} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial \rho_i} \quad (6.18a)$$

$$\frac{\partial e_{vs}}{\partial (\rho u)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial (\rho u)}, \quad \frac{\partial e_{vs}^*}{\partial (\rho u)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial (\rho u)} \quad (6.18b)$$

$$\frac{\partial e_{vs}}{\partial (\rho v)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial (\rho v)}, \quad \frac{\partial e_{vs}^*}{\partial (\rho v)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial (\rho v)} \quad (6.18c)$$

$$\frac{\partial e_{vs}}{\partial (\rho w)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial (\rho w)}, \quad \frac{\partial e_{vs}^*}{\partial (\rho w)} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial (\rho w)} \quad (6.18d)$$

$$\frac{\partial e_{vs}}{\partial E_v} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial E_v}, \quad \frac{\partial e_{vs}^*}{\partial E_v} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial E_v} \quad (6.18e)$$

$$\frac{\partial e_{vs}}{\partial E} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v} \frac{\partial T_v}{\partial E}, \quad \frac{\partial e_{vs}^*}{\partial E} = \left(\frac{\partial e_{vs}}{\partial T_v} \right)_{T_v=T} \frac{\partial T_v}{\partial E} \quad (6.18f)$$

6.9 Jacobians

6.9.1 Flux Jacobians

6.9.2 Source term Jacobians - Third body reactions

As stated, for third body reactions, the Arrhenius rate coefficient k_f can be expressed as an efficiency times a base rate coefficient that is independant of the third body:

$$k_f(T) = \epsilon_i \gamma(T) \quad (6.19)$$

where

$$\gamma(T) = C_c T^\eta e^{E_a/T} \quad (6.20)$$

For third-body reactions, the rate of production of species s can be expressed as:

$$\frac{d[X_i]}{dt} = (\nu''_i - \nu'_i) \underbrace{\sum_{i=1}^{NS} \epsilon_i [M_i] \left\{ \gamma(T_f) \prod_i [X_i]^{\nu'_i} - \frac{\gamma(T_b)}{K_c(T_b)} \prod_i [X_i]^{\nu''_i} \right\}}_{E_{\text{eff}}} L_r \quad (6.21)$$

Here, T_f is the temperature used to compute the forward rate coefficient, and

T_b is the temperature used to compute the backward rate coefficient. For now, we are only interest in the two underbracketed sections, E_{eff} and L_r . The reaction rate (without the stoichiometric coefficients in the front) can then be described by:

$$\mathcal{R}_r = E_{\text{eff}} L_r \quad (6.22)$$

The partial derivative of this reaction rate with respect to some conserved variable U_j is then:

$$\frac{\partial \mathcal{R}_r}{\partial U_j} = \frac{\partial E_{\text{eff}}}{\partial U_j} L_r + E_{\text{eff}} \frac{\partial L_r}{\partial U_j} \quad (6.23)$$

The two derivatives on the right hand side are then:

$$\frac{\partial E_{\text{eff}}}{\partial U_j} = \begin{cases} E_{\text{eff}} + \frac{\epsilon_j}{M_j} (1 - \rho_j), & \text{if } U_j = \rho_j \\ 0, & \text{if } U_j = \rho u, \rho v, \text{ etc.} \end{cases} \quad (6.24)$$

$$\begin{aligned} \frac{\partial L_r}{\partial U_j} &= \left\{ \frac{\partial T}{\partial U_j} \frac{\partial}{\partial T} [\gamma(T_f)] \right\} \left[\frac{\rho_{s,f}}{M_{s,f}} \right]^{\nu'_s} + \gamma(T_f) \left\{ \frac{\partial}{\partial U_j} \left[\frac{\rho_{s,b}}{M_{s,b}} \right]^{\nu'_s} \right\} \\ &\quad - \left\{ \frac{\partial T}{\partial U_j} \frac{\partial}{\partial T} \left[\frac{\gamma(T_b)}{K_c(T_b)} \right] \right\} \left[\frac{\rho_{s,b}}{M_{s,b}} \right]^{\nu''_s} - \frac{\gamma(T_b)}{K_c(T_b)} \left\{ \frac{\partial}{\partial U_j} \left[\frac{\rho_{s,b}}{M_{s,b}} \right]^{\nu''_s} \right\} \end{aligned} \quad (6.25)$$

Of course, in the second equation, more needs to be expanded out. These expansions yield:

$$\frac{\partial}{\partial T} \gamma(T) = \gamma(T) \left[\frac{\eta q}{T_{tr}} - \frac{q E_a}{T_{tr}^{1+q} T_v^{1-q}} \right] \quad (6.26)$$

$$\frac{\partial}{\partial U_j} \left[\frac{\rho_s}{M_s} \right]^{\nu_s^*} = \nu_s^* \left[\frac{\rho_s}{M_s} \right]^{\nu_s^*-1} \cdot \frac{1}{M_s} \quad (6.27)$$

$$\frac{\partial}{\partial T} K_{c,m}(T) = \frac{K_{c,m}(T)}{T} \left[\sum_s (\nu_s'' - \nu_s') \frac{H_s^\circ}{\hat{R} T} + \nu_m \right] \quad (6.28)$$

Putting it together, the L_r derivative is:

$$\begin{aligned}
\frac{\partial L_r}{\partial U_j} = & \gamma(T_f) \left[\frac{\rho_{s,f}}{\mathcal{M}_{s,f}} \right]^{\nu'_s} \times \left[\frac{\partial T}{\partial U_j} \left(\frac{\eta q^f}{T_{tr}} - \frac{q^f E_a}{(T_{tr}^{1+q} T_v^{1-q})_f} \right) + \frac{\nu'_s}{\rho_{s,f}} \right] \\
& - \frac{\gamma(T_b)}{K_c(T_b)} \left[\frac{\rho_{s,b}}{\mathcal{M}_{s,b}} \right]^{\nu''_s} \times \dots \\
& \left\{ \frac{\partial T}{\partial U_j} \left[\frac{\eta q^b}{T_{tr}} - \frac{q^b E_a}{(T_{tr}^{1+q} T_v^{1-q})_b} - \frac{1}{T_b} \left(\sum_{i=1}^{\text{NS}} (\nu_i'' - \nu'_i) \frac{H_s^\circ}{\hat{R}T} + \nu_m \right) \right] + \frac{\nu''_s}{\rho_{s,b}} \right\} \\
& \quad \text{if } U_j = \rho_j
\end{aligned} \tag{6.29}$$

6.9.3 Source term Jacobians - Normal reactions

For normal reaction, we can revert to the usual notation of k_f and k_b for the forward and backward rate coefficients with no efficiencies. A reaction is then:

$$R_r = k_f(T_f) \prod_i \left[\frac{\rho_i}{\mathcal{M}_i} \right]^{\nu'_i} - \frac{k_f(T_b)}{K_c(T_b)} \prod_i \left[\frac{\rho_i}{\mathcal{M}_i} \right]^{\nu''_i} \tag{6.30}$$

The derivative of this reaction rate with respect to some conserved variable U_j is then:

$$\begin{aligned}
\frac{\partial R_r}{\partial U_j} = & k_f(T_f) \left[\frac{\rho_{s,f}}{\mathcal{M}_{s,f}} \right]^{\nu'_s} \times \left[\frac{\partial T}{\partial U_j} \left(\frac{\eta q^f}{T_{tr}} - \frac{q^f E_a}{(T_{tr}^{1+q} T_v^{1-q})_f} \right) + \frac{\nu'_s}{\rho_{s,f}} \right] \\
& - \frac{k_f(T_b)}{K_c(T_b)} \left[\frac{\rho_{s,b}}{\mathcal{M}_{s,b}} \right]^{\nu''_s} \times \dots \\
& \left\{ \frac{\partial T}{\partial U_j} \left[\frac{\eta q^b}{T_{tr}} - \frac{q^b E_a}{(T_{tr}^{1+q} T_v^{1-q})_b} - \frac{1}{T_b} \left(\sum_{i=1}^{\text{NS}} (\nu_i'' - \nu'_i) \frac{H_s^\circ}{\hat{R}T} - \nu_m \right) \right] + \frac{\nu''_s}{\rho_{s,b}} \right\} \\
& \quad \text{if } U_j = \rho_j
\end{aligned} \tag{6.31}$$

6.9.4 Source term Jacobians - Vibrational relaxation

Recall the vibrational energy source term:

$$Q_{t-v,s} = \rho_s \frac{e_{vs} - e_{vs}^*}{\langle \tau_s \rangle} \quad (6.32)$$

$$e_{vs} = \frac{\hat{R}}{\mathcal{M}_s} \frac{\theta_{vs}}{\exp(\theta_{vs}/T_v) - 1} \quad (6.33)$$

The derivative is then:

$$\begin{aligned} \frac{\partial Q_{t-v,s}}{\partial U_j} &= \left[\frac{\partial}{\partial U_j} \rho_s \right] \frac{e_{vs} - e_{vs}^*}{\langle \tau_s \rangle} \\ &+ \rho_s \left\{ \left[\frac{\partial}{\partial U_j} (e_{vs} - e_{vs}^*) \right] \frac{1}{\langle \tau_s \rangle} + (e_{vs} - e_{vs}^*) \left(-\tau_s^{-2} \left[\frac{\partial}{\partial U_j} \langle \tau_s \rangle \right] \right) \right\} \end{aligned} \quad (6.34)$$

Where:

$$\frac{\partial e_{vs}}{\partial T_v} = \left(\frac{\theta_{vs}}{T_v} \right)^2 \frac{\exp(\theta_{vs}/T_v)}{[\exp(\theta_{vs}/T_v) - 1]^2} \quad (6.35)$$

And

$$\frac{\partial e_{vs}^*}{\partial T_v} = \left. \frac{\partial e_{vs}}{\partial T_v} \right|_{T_v=T_{tr}} \quad (6.36)$$

We now aim to find the partial derivatives of the relaxation time:

$$\frac{\partial \langle \tau_s \rangle}{\partial U_j} = \frac{\partial}{\partial U_j} \frac{\sum_{r=1}^{NS} X_r}{\sum_{r=1}^{NS} X_r / \tau_{sr}} \quad (6.37)$$

This gives:

$$\frac{\partial \langle \tau_s \rangle}{\partial U_j} = \underbrace{\left[\frac{\partial}{\partial U_j} \sum_{r=1}^{NS} X_r \right]}_0 \frac{1}{\sum_{r=1}^{NS} X_r / \tau_{sr}} + \sum_{r=1}^{NS} X_r \left[\frac{\partial}{\partial U_j} \left(\sum_{r=1}^{NS} \frac{X_r}{\tau_{sr}} \right)^{-1} \right] \quad (6.38)$$

Since the first term is zero, this leaves us with:

$$\begin{aligned} \frac{\partial <\tau_s>}{\partial U_j} &= \sum_{r=1}^{\text{NS}} X_r \left[\frac{\partial}{\partial U_j} \left(\sum_{r=1}^{\text{NS}} \frac{X_r}{\tau_{sr}} \right)^{-1} \right] \\ &= - \left[\sum_{r=1}^{\text{NS}} X_r \right] \left[\sum_{r=1}^{\text{NS}} \frac{X_r}{\tau_{sr}} \right]^{-2} \left\{ \sum_{r=1}^{\text{NS}} \left[\left(\frac{\partial X_r}{\partial U_j} \right) \frac{1}{\tau_{sr}} + X_r \left(\frac{\partial}{\partial U_j} \frac{1}{\tau_{sr}} \right) \right] \right\} \quad (6.39) \end{aligned}$$

The first term inside the curly brace expands to:

$$X_r = \frac{\rho_r / \mathcal{M}_r}{\sum_{i=1}^{\text{NS}} \rho_i / \mathcal{M}_i} \quad (6.40)$$

$$\frac{\partial X_r}{\partial \rho_j} = \left[\frac{\partial}{\partial \rho_j} \frac{\rho_r}{\mathcal{M}_r} \right] \frac{1}{\sum_{i=1}^{\text{NS}} \rho_i / \mathcal{M}_i} + \frac{\rho_r}{\mathcal{M}_r} \left[\frac{\partial}{\partial \rho_j} \frac{1}{\sum_{i=1}^{\text{NS}} \rho_i / \mathcal{M}_i} \right] \quad (6.41)$$

Giving:

$$\boxed{\frac{\partial X_r}{\partial \rho_j} = \frac{1}{\sum_{i=1}^{\text{NS}} \rho_i / \mathcal{M}_i} \left[\frac{\delta_{jr}}{\mathcal{M}_r} - \frac{X_r}{\mathcal{M}_j} \right]} \quad (6.42)$$

And all other derivatives WRT conserved variables are 0. δ_{jr} is the Kronecker delta. Then $\rho_j = \rho_r$, it is 1, otherwise it is 0. The second term expands to the following without Park's correction:

$$X_r \left(\frac{\partial}{\partial U_j} \frac{1}{\tau_{sr}} \right) = -X_r \tau_{sr}^{-2} \frac{\partial \tau_{sr}}{\partial U_j} \quad (6.43)$$

$$= -X_r \tau_{sr}^{-2} \left[-\tau_{sr} \left(\frac{101325}{p} \frac{\partial p}{\partial U_j} + \frac{1}{3} \frac{\partial T}{\partial U_j} T^{-4/3} A_{sr} \right) \right] \quad (6.44)$$

$$= \frac{X_r}{\tau_{sr}} \left[\frac{101325}{p} \frac{\partial p}{\partial U_j} + \frac{1}{3} \frac{\partial T}{\partial U_j} T^{-4/3} A_{sr} \right] \quad (6.45)$$

With Park's correction we have:

$$\tau_{sr} = \tau_{sr}^{\text{MW}} + \tau_{sr}^{\text{P}} \quad (6.46)$$

So:

$$\frac{\partial \tau_{sr}}{\partial U_j} = \frac{\partial \tau_{sr}^{\text{MW}}}{\partial U_j} + \frac{\partial \tau_{sr}^{\text{P}}}{\partial U_j} \quad (6.47)$$

As we saw:

$$\frac{\partial \tau_{sr}^{\text{MW}}}{\partial U_j} = -\tau_{sr}^{\text{MW}} \left[\frac{101325}{p} \frac{\partial p}{\partial U_j} + \frac{1}{3} \frac{\partial T}{\partial U_j} T^{-4/3} A_{sr} \right] \quad (6.48)$$

The harder derivative is:

$$\frac{\partial \tau_{sr}^{\text{P}}}{\partial U_j} = \frac{\partial}{\partial U_j} \left[\frac{1}{\bar{c}\sigma n} \right] = -(\bar{c}\sigma n)^{-2} \left\{ \frac{\partial \bar{c}}{\partial U_j} \sigma n + \bar{c} \left[\frac{\partial \sigma}{\partial U_j} + \sigma \frac{\partial n}{\partial U_j} \right] \right\} \quad (6.49)$$

With:

$$\bar{c} = \sqrt{\frac{8\hat{R}T}{\pi \sum_{i=1}^{\text{NS}} X_i \mathcal{M}_i}}, \quad n = \frac{p}{kT}, \quad \sigma = \sigma'_v \left(\frac{50000}{T} \right)^2 \quad (6.50)$$

We begin with:

$$\frac{\partial \bar{c}}{\partial U_j} \quad (6.51)$$

So, we have:

$$\begin{aligned} \frac{\partial \langle \tau_s \rangle}{\partial U_j} &= - \overbrace{\left[\sum_r X_r \right] \left[\sum_r \frac{X_r}{\tau_{sr}} \right]^{-2}}^{-\langle \tau_s \rangle / [\sum_r X_r / \tau_{sr}]} \times \dots \\ &\left\{ \sum_r \left[\underbrace{\left(\frac{1}{\sum_i \rho_i \mathcal{M}_i} \left[\frac{\delta_{jr}}{\mathcal{M}_r} - \frac{X_r}{\mathcal{M}_j} \right] \frac{1}{\tau_{sr}} \right)}_{\text{only if } U_j = \rho_j} + \frac{X_r}{\tau_{sr}} \left[\frac{101325}{p} \frac{\partial p}{\partial U_j} + \frac{1}{3} \frac{\partial T}{\partial U_j} T^{-4/3} A_{sr} \right] \right] \right\} \end{aligned} \quad (6.52)$$

Taking this per conserved variable gives:

$$\frac{\partial \langle \tau_s \rangle}{\partial \rho_j} = -\langle \tau_s \rangle \frac{1}{\sum_r X_r / \tau_{sr}} \quad (6.53)$$

We can rearrange this to:

$$\frac{\partial \langle \tau_s \rangle}{\partial U_j} = - \frac{\langle \tau_s \rangle}{\sum_r X_r / \tau_{sr}} \left\{ \sum_r \left[\left(\frac{\partial}{\partial U_j} X_r \right) \frac{1}{\tau_{sr}} + X_r \left(\frac{\partial}{\partial U_j} \frac{1}{\tau_{sr}} \right) \right] \right\} \quad (6.54)$$