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Nomenclature

\hat{R}	Universal gas constant $\approx 8.314 \text{ J/kg-K}$	F	Helmholtz energy
λ	Lagrange multiplier	G	Gibbs free-energy
\mathcal{L}	The Lagrangian	h'	Sum of contributions from each species towards enthalpy constraint, i.g. $\sum_{j=1}^{\text{NS}} \mathcal{N}_j H_j^\circ$
\mathcal{N}	Total number of moles in the system	h'_0	User-specified enthalpy
\mathcal{N}_j	Number of moles of species j in the system	H_j°	Standard state enthalpy of species j
μ_j	Chemical potential of species j	MW_j	Molecular weight of species j
μ_j°	Standard state chemical potential of species j	p_{ref}	Reference temperature used to compute chemical potential
ρ	Density (kg/m^3)	q_j	Charge of species j (0 for neutrals, 1 for ions, -1 for electrons)
a_{ij}	Stoichiometric coefficients, e.g. number of atoms of element i in species j	T	Temperature (K)
b_i	Sum of contributions from each species towards constraint for element i , i.g. $\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j$	u'	Sum of contributions from each species towards internal energy constraint, i.g. $\sum_{j=1}^{\text{NS}} \mathcal{N}_j U_j^\circ$
b_i°	Specified number of moles of element i	u'_0	User-specified internal energy
$c_{p,j}^\circ$	Standard state specific heat of species j	U_j°	Standard state internal energy of species j
$c_{v,j}^\circ$	Standard state specific heat of species j	V	Volume (m^3)

1 Minimization Procedures

1.1 Constraints for Both Procedures

There are two constraints that are used the most in both energy minimization procedures. The first, equation [1.1], is always used while the charge constraints in equation [1.2] is only used when ions are present:

$$\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j - b_i^\circ = 0 \quad (1.1)$$

$$\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j = 0 \quad (1.2)$$

Since these are used in abundance for both procedures, their constraints are placed in the Lagrangian term. Other constraints (such as specified volume, temperature, internal energy, and entropy) are given their rows in the Newton solve without addition into the Lagrangian.

Equation [1.1] says that the number of moles of element i must remain constant during the minimization process, as atoms can not be created or destroyed without nuclear physics being involved. b_i° is the number of moles of element i before minimizing, and a_{ij} is the stoichiometric coefficient, i.e. how many atoms of element i are in species j .

Equation [1.2] says that the charge of the system must remain neutral, i.e., if cations are formed, than there must be an equal number of free electrons in the gas to balance it out.

1.2 The Lagrangian

The Lagrangian is defined as;

$$\mathcal{L} = f + \sum_j \lambda_j g_j \quad (1.3)$$

where f is the function we want to minimize, λ_j are Lagrange multipliers, and g_j are constraint functions. Instead of minimizing just f , we aim to minimize the Lagrangian \mathcal{L} .

1.3 Newton-Raphson Iterative Method

A Newton-Raphson iterative procedure is used to minimize the energies. Since this is essentially a root finding system of equations, and we are seeking the solution update, the method looks like the following:

$$\sum_{i=1}^N \frac{\partial f}{\partial x_i} \Delta x_i = -f(x) \quad (1.4)$$

Here, $f(x)$ is the function that is being minimized. In the case of Gibbs and Helmholtz energy minimization, f takes the form of $\partial \mathcal{L} / \partial x$, with x being non-linear variables used to guide the convergence of the system. The non-linear correction variables are in the form $\Delta \ln \mathcal{N}_j$, $\Delta \ln \mathcal{N}$, $\Delta \ln T$, and $\pi_{i,q} = \Delta \pi_{i,q}$. For the last correction variable, it is argued that setting π to 0 at the beginning of each update does not influence the solution for the non-reduced equations, so the Δ is dropped for that term. This will influence the form of the update equations, which will be talked about in due time.

1.4 Gibbs Minimization

The Gibbs energy is a function of temperature, pressure, and composition of a gas:

$$G = G(T, p, \mathcal{N}_1, \mathcal{N}_2, \dots, \mathcal{N}_{NS}) = \sum_{j=1}^{NS} \mathcal{N}_j \mu_j \quad (1.5)$$

Where NS is the number of gas species, \mathcal{N}_j is the number of moles of species j , and μ_j is the chemical potential of species j defined by:

$$\mu_j = \frac{\partial G}{\partial \mathcal{N}_j} = \mu_j^\circ + \hat{R}T \left[\ln \left(\frac{p}{p_{ref}} \right) + \ln \left(\frac{\mathcal{N}_j}{\mathcal{N}} \right) \right] \quad (1.6)$$

Here, \hat{R} is the universal gas constant, T and p are the temperature and pressure of the system, p_{ref} is the reference pressure (usually takes as 101,325 Pa, or 1 bar depending on the literature), and \mathcal{N} is the total number of moles in the system. In order to find chemical equilibrium, the derivative of G wrt \mathcal{N}_j is set to 0 for all species. However, a number of constraints need to be added in.

1.4.1 Constraints

Firstly, the elemental constraint from equation [1.1] is added into the Lagrangian for our Gibbs function. If ions are present, the charge constraint is also put in. Therefore, our Lagrangian is:

$$\mathcal{L} = \sum_{j=1}^{NS} \mu_j \mathcal{N}_j + \sum_{i=1}^{NE} \lambda_i \left[\sum_{j=1}^{NS} a_{ij} \mathcal{N}_j - b_i^\circ \right] + \lambda_q \left[\sum_{j=0}^{NS} q_j \mathcal{N}_j \right] \quad (1.7)$$

The constraints available for the Gibbs energy are:

- Hold temperature (T) and pressure (P) constant.
- Hold enthalpy (H) and pressure (P) constant.
- Hold entropy (S) and pressure (P) constant.

For all Gibbs minimizations, we need the molar constraint:

$$\sum_{j=1}^{NS} \mathcal{N}_j - \mathcal{N} = 0 \quad (1.8)$$

For (TP) minimization, only the elemental and charge (if present) constraints are necessary. For (HP) minimization, the constraint is defined by:

$$h' - h'_0 = 0 \quad (1.9)$$

Where h'_0 is the specified enthalpy, and h' is solved for during the minimization process as:

$$h' = \sum_{j=1}^{NS} \mathcal{N}_j H_j^\circ \quad (1.10)$$

For (SP) minimization, the constraint is defined by:

$$s' - s'_0 = 0 \quad (1.11)$$

Where s'_0 is the specified entropy and s' is solved for in the minimization process as:

$$s' = \sum_{j=1}^{NS} \mathcal{N}_j S_j \quad (1.12)$$

where:

$$S_j = S_j^\circ - \hat{R} \ln \frac{\mathcal{N}_j}{\mathcal{N}} - \hat{R} \ln p \quad (1.13)$$

1.4.2 Derivatives

We now take the derivatives of \mathcal{L} wrt \mathcal{N}_j , and each λ and denote them as f :

$$f_1 = \frac{\partial \mathcal{L}}{\partial \mathcal{N}_j} = \mu_j + \sum_{i=1}^{NE} \lambda_i a_{ij} + \lambda_q q_j = 0 \quad (1.14)$$

$$f_2 = \frac{\partial \mathcal{L}}{\partial \lambda_i} = \sum_{i=1}^{NS} a_{ij} \mathcal{N}_j - b_i = 0 \quad (1.15)$$

$$f_3 = \frac{\partial \mathcal{L}}{\partial \lambda_q} = \sum_{j=0}^{NS} q_j \mathcal{N}_j = 0 \quad (1.16)$$

These three equations are our minimization functions. Through use of the aforementioned Newton-Raphson method, we can solve these equations. Constraint equations for enthalpy and entropy are:

$$f_4 = \sum_{j=1}^{NS} \mathcal{N}_j - \mathcal{N} = 0 \quad (1.17)$$

$$f_5 = \frac{h' - h'_0}{\hat{R}T} = \frac{\sum_{j=1}^{NS} \mathcal{N}_j H_j^\circ - h'_0}{\hat{R}T} \quad (1.18)$$

$$f_6 = \frac{s' - s'_0}{\hat{R}} = \frac{\sum_{j=1}^{NS} \mathcal{N}_j S_j - s'_0}{\hat{R}} \quad (1.19)$$

1.4.3 Derivatives of f_1

First, we must do some algebraic manipulation. We start by expanding equation 1.14 with equation 1.6:

$$\frac{\partial \mathcal{L}}{\partial \mathcal{N}_j} = \mu_j^\circ + \hat{R}T \left[\ln \left(\frac{p}{p_{ref}} \right) + \ln \left(\frac{\mathcal{N}_j}{\mathcal{N}} \right) \right] + \sum_{i=1}^{NE} \lambda_i a_{ij} + \lambda_q q_j = 0 \quad (1.20)$$

Dividing through by $\hat{R}T$ and setting $\pi = -\lambda/\hat{R}T$ gives:

$$f_1 = \frac{\mu_j^\circ}{\hat{R}T} + \ln \frac{p}{p_{ref}} + \ln \mathcal{N}_j - \ln \mathcal{N} - \sum_{i=1}^{NE} \pi_i a_{i,j} - \pi_q q_j = 0 \quad (1.21)$$

We now need to take the partial derivatives of our functions wrt our non-linear variables as well as each π , and then multiple it by the correction variables.

$$\frac{\partial f_1}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \Delta \ln \mathcal{N}_j \quad (1.22)$$

$$\frac{\partial f_1}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = -\Delta \ln \mathcal{N} \quad (1.23)$$

$$\frac{\partial f_1}{\partial [\ln T]} \Delta \ln T = -\frac{H_j^\circ}{\hat{R}T} \Delta \ln T \quad (1.24)$$

$$\frac{\partial f_1}{\partial [\pi_i]} \pi_i = -a_{ij} \pi_i \quad (1.25)$$

$$\frac{\partial f_1}{\partial [\pi_q]} \pi_q = -q_j \pi_q \quad (1.26)$$

Combining these into the form of equation 1.4 gives

$$\Delta \ln \mathcal{N}_j - \Delta \ln \mathcal{N} - \sum_{i=1}^{\text{NE}} a_{ij} \pi_i - q_j \pi_q - \frac{H_j^\circ}{\hat{R}T} \Delta \ln T = \mu_j + \sum_{i=1}^{\text{NE}} \pi_i a_{ij} + \pi_q q_j \quad (1.27)$$

Arguments are made that $\pi_i = 0$ at the start of every iteration, so this equation becomes:

$$\boxed{\Delta \ln \mathcal{N}_j - \Delta \ln \mathcal{N} - \sum_{i=1}^{\text{NE}} a_{ij} \pi_i - q_j \pi_q - \frac{H_j^\circ}{\hat{R}T} \Delta \ln T = -\frac{\mu_j}{\hat{R}T}}$$

There will be one of these equation for each species.

1.4.4 Derivatives of f_2

$$\frac{\partial f_2}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = a_{ij} \mathcal{N}_j \Delta \ln \mathcal{N}_j \quad (1.28)$$

$$\frac{\partial f_2}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = 0 \quad (1.29)$$

$$\frac{\partial f_2}{\partial [\ln T]} = 0 \quad (1.30)$$

$$\frac{\partial f_2}{\partial [\pi_i]} \pi_i = 0 \quad (1.31)$$

$$\frac{\partial f_2}{\partial [\pi_q]} \pi_q = 0 \quad (1.32)$$

Which gives:

$$\boxed{\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j \Delta \ln \mathcal{N}_j = b_i^\circ - \sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j}$$

1.4.5 Derivatives of f_3

$$\frac{\partial f_3}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = q_j \mathcal{N}_j \Delta \ln \mathcal{N}_j \quad (1.33)$$

$$\frac{\partial f_3}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = 0 \quad (1.34)$$

$$\frac{\partial f_3}{\partial [\ln T]} = 0 \quad (1.35)$$

$$\frac{\partial f_3}{\partial [\pi_i]} \pi_i = 0 \quad (1.36)$$

$$\frac{\partial f_3}{\partial [\pi_q]} \pi_q = 0 \quad (1.37)$$

Giving:

$$\boxed{\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \Delta \ln \mathcal{N}_j = -\sum_{j=1}^{\text{NS}} q_i \mathcal{N}_j}$$

1.4.6 Derivatives of f_4

$$\frac{\partial f_4}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j \Delta \ln \mathcal{N}_j \quad (1.38)$$

$$\frac{\partial f_4}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = -\mathcal{N} \Delta \ln \mathcal{N} \quad (1.39)$$

$$\frac{\partial f_4}{\partial [\ln T]} = 0 \quad (1.40)$$

$$\frac{\partial f_4}{\partial [\pi_i]} \pi_i = 0 \quad (1.41)$$

$$\frac{\partial f_4}{\partial [\pi_q]} \pi_q = 0 \quad (1.42)$$

$$\sum_{j=1}^{\text{NS}} \mathcal{N}_j \Delta \ln \mathcal{N}_j - \mathcal{N} \Delta \ln \mathcal{N} = \mathcal{N} - \sum_{j=1}^{\text{N}} \mathcal{N}_j$$

(1.43)

1.4.7 Derivatives of f_5

$$\frac{\partial f_5}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j \quad (1.44)$$

$$\frac{\partial f_5}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = 0 \quad (1.45)$$

$$\frac{\partial f_5}{\partial [\ln T]} = \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T \quad (1.46)$$

$$\frac{\partial f_5}{\partial [\pi_i]} \pi_i = 0 \quad (1.47)$$

$$\frac{\partial f_5}{\partial [\pi_q]} \pi_q = 0 \quad (1.48)$$

Giving:

$$\sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T = \frac{h' - h'_0}{\hat{R}T}$$

1.4.8 Derivatives of f_6

$$\frac{\partial f_6}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j \frac{S_j}{\hat{R}} \Delta \ln \mathcal{N}_j \quad (1.49)$$

$$\frac{\partial f_6}{\partial [\ln \mathcal{N}]} \Delta \ln \mathcal{N} = 0 \quad (1.50)$$

$$\frac{\partial f_6}{\partial [\ln T]} = \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T \quad (1.51)$$

$$\frac{\partial f_6}{\partial [\pi_i]} \pi_i = 0 \quad (1.52)$$

$$\frac{\partial f_6}{\partial [\pi_q]} \pi_q = 0 \quad (1.53)$$

Giving

$$\boxed{\sum_{j=1}^{NS} \mathcal{N}_j \frac{S_j}{\hat{R}} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T = \frac{s'_0 - s'}{\hat{R}} + \mathcal{N} - \sum_{j=1}^{NS} \mathcal{N}_j}$$

All five of the equations are now listed for convenience:

$$\Delta \ln \mathcal{N}_j - \Delta \ln \mathcal{N} - \sum_{i=1}^{NE} a_{ij} \pi_i - q_j \pi_q - \frac{H_j^\circ}{\hat{R}T} \Delta \ln T = -\frac{\mu_j}{\hat{R}T} \quad (1.54)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j \Delta \ln \mathcal{N}_j - \mathcal{N} \Delta \ln \mathcal{N} = \mathcal{N} - \sum_{j=1}^{NS} \mathcal{N}_j \quad (1.55)$$

$$\sum_{j=1}^{NS} a_{ij} \mathcal{N}_j \Delta \ln \mathcal{N}_j = b_i^\circ - \sum_{j=1}^{NS} a_{ij} \mathcal{N}_j \quad (1.56)$$

$$\sum_{j=1}^{NS} q_j \mathcal{N}_j \Delta \ln \mathcal{N}_j = - \sum_{j=1}^{NS} q_j \mathcal{N}_j \quad (1.57)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T = \frac{h'_0 - h'}{\hat{R}T} \quad (1.58)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j \frac{S_j}{\hat{R}} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \Delta \ln T = \frac{s'_0 - s'}{\hat{R}} + \mathcal{N} - \sum_{j=1}^{NS} \mathcal{N}_j \quad (1.59)$$

1.4.9 Reduced Gibbs Equations

A shortcut can be made to make this system of equations smaller. This is achieved by solving equation [1.54] for $\Delta \ln \mathcal{N}_j$ and pluggin it into equations [1.56] - [1.59]

$$\Delta \ln \mathcal{N}_j = \Delta \ln \mathcal{N} + \sum_{i=1}^{NE} a_{ij} \pi_i - \frac{\mu_j}{\hat{R}T} + q_j \pi_q + \frac{H_j^\circ}{\hat{R}T} \Delta \ln T \quad (1.60)$$

Substitution into equation [1.56] - [1.59] yields:

$$\begin{aligned}
& \sum_{i=1}^{\text{NE}} \left[\sum_{j=1}^{\text{NS}} a_{kj} a_{ij} \mathcal{N}_j \right] \pi_i + \left[\sum_{j=1}^{\text{NS}} a_{kj} \mathcal{N}_j \right] \Delta \ln \mathcal{N} + \left[\sum_{j=1}^{\text{NS}} a_{kj} q_j \mathcal{N}_j \right] \pi_q + \left[\sum_{j=1}^{\text{NS}} a_{kj} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \Delta \ln T \\
& = b_k^\circ - \sum_{j=1}^{\text{NS}} a_{kj} \mathcal{N}_j + \sum_{j=1}^{\text{NS}} a_{kj} \mathcal{N}_j \frac{\mu_j}{\hat{R}T} \quad (1.61)
\end{aligned}$$

$$\begin{aligned}
& \sum_{i=1}^{\text{NE}} \left[\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j \right] \pi_i + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j - \mathcal{N} \right] \Delta \ln \mathcal{N} + \left[\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \right] \pi_q + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \Delta \ln T \\
& = \mathcal{N} - \sum_{j=1}^{\text{NS}} \mathcal{N}_j + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{\mu_j}{\hat{R}T} \quad (1.62)
\end{aligned}$$

$$\begin{aligned}
& \sum_{i=1}^{\text{NE}} \left[\sum_{j=1}^{\text{NS}} a_{ij} q_j \mathcal{N}_j \right] \pi_i + \left[\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \right] \Delta \ln \mathcal{N} + \left[\sum_{j=1}^{\text{NS}} q_j^2 \mathcal{N}_j \right] \pi_q + \left[\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \Delta \ln T \\
& = \sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \frac{\mu_j}{\hat{R}T} - \sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \quad (1.63)
\end{aligned}$$

$$\begin{aligned}
& \sum_{i=1}^{\text{NE}} \left[\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \pi_i + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \Delta \ln \mathcal{N} + \left[\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \right] \pi_q \\
& + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j \left(\frac{H_j^\circ}{\hat{R}T} \right)^2 + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \right] \Delta \ln T = \frac{h'_0 - h'}{\hat{R}T} + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{H_j^\circ}{\hat{R}T} \frac{\mu_j}{\hat{R}T} \quad (1.64)
\end{aligned}$$

$$\begin{aligned}
& \sum_{i=1}^{\text{NE}} \left[\sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j \frac{S_j}{\hat{R}} \right] \pi_i + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{S_j}{\hat{R}} \right] \Delta \ln \mathcal{N} + \left[\sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j \frac{S_j}{\hat{R}} \right] \pi_q \\
& + \left[\sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{S_j}{\hat{R}} \frac{H_j^\circ}{\hat{R}T} + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{c_{p,j}^\circ}{\hat{R}} \right] \Delta \ln T = \frac{s'_0 - s'}{\hat{R}} + \mathcal{N} - \sum_{j=1}^{\text{NS}} \mathcal{N}_j + \sum_{j=1}^{\text{NS}} \mathcal{N}_j \frac{S_j}{\hat{R}} \frac{\mu_j}{\hat{R}T} \quad (1.65)
\end{aligned}$$

There are NE number of equation [1.62], and only one of all others. Once this system has been solved, you recover \mathcal{N}_j with equation [1.60].

The equations are colored according to when they are needed. Red coloring denotes terms that arise from the elemental constraint condition. These are included no matter what. Blue denotes terms that are added when you include the charge constraint. Violet terms are added for both HP and SP constraints. orange and teal are the rows added for enthalpy or entropy constraints, respectively.

1.5 Helmholtz Minimization

The Helmholtz energy is defined as:

$$F = G - pV \quad (1.66)$$

Where G is the Gibbs free energy, p is the pressure, and V is the volume. Substitution of G yields:

$$F = \sum_{j=1}^{NS} \mu_j \mathcal{N}_j - pV \quad (1.67)$$

Where the chemical potential is redefined as:

$$\mu_j = \mu_j^\circ + \hat{R}T \ln \left(\frac{\mathcal{N}_j R' T}{V} \right) \quad (1.68)$$

and $R' = \hat{R} \cdot 10^{-5}$ is the same as dividing \hat{R} by the reference pressure of 1 bar.

1.5.1 Constraints

The constraints available for the Helmholtz energy are:

- Hold temperature (T) and volume (V) constant.
- Hold internal energy (U) and volume (V) constant.
- Hold entropy (S) and volume (V) constant.

For (TV) minimization, only the elemental and charge (if present) constraints are necessary. For (UV) minimization, the constraint is defined by:

$$u' - u'_0 = 0 \quad (1.69)$$

Where u'_0 is the specified internal energy, and u' is solved for during the minimization process as:

$$u' = \sum_{j=1}^{NS} \mathcal{N}_j U_j^\circ = u'_0 \quad (1.70)$$

For (SV) minimization, the constraint is defined by:

$$s' - s'_0 = 0 \quad (1.71)$$

Where s'_0 is the specified entropy and s' is solved for during the minimization process as:

$$s' = \sum_{j=1}^{NS} \mathcal{N}_j S_j \quad (1.72)$$

where:

$$S_j = S_j^\circ - \hat{R} \ln \mathcal{N}_j - \hat{R} \ln \left(\frac{R' T}{V} \right) \quad (1.73)$$

While you may be able to include the constraints for internal energy and entropy inside the Lagrangian, we will exclude them from the Lagrangian and use our own residual equation for them. The Lagrangian for the Helmholtz energy minimization process is then defined as:

$$\mathcal{L} = \sum_{j=1}^{NS} \mu_j \mathcal{N}_j - pV + \sum_{i=1}^{NE} \lambda_i \left[\sum_{j=1}^{NS} a_{ij} \mathcal{N}_j - b_i^\circ \right] + \lambda_q \left[\sum_{j=1}^{NS} q_j \mathcal{N}_j \right] \quad (1.74)$$

1.5.2 Derivatives

We first find our function f that are equal to 0 by taking the derivatives wrt \mathcal{N}_j , λ_i , and λ_q :

$$f_1 = \frac{\partial \mathcal{L}}{\partial \mathcal{N}_j} = \mu_j + \sum_{i=1}^{NE} \lambda_i a_{ij} + \lambda_q q_j = 0 \quad (1.75)$$

$$f_2 = \frac{\partial \mathcal{L}}{\partial \lambda_i} = \sum_{j=1}^{NS} a_{ij} \mathcal{N}_j - b_i^\circ = 0 \quad (1.76)$$

$$f_3 = \frac{\partial \mathcal{L}}{\partial \lambda_q} = \sum_{j=1}^{NS} q_j \mathcal{N}_j - 0 = 0 \quad (1.77)$$

The negative of these functions ($-f$) are played on the RHS of the Newton solver. We now form the second derivatives that define our Jacobian in the Newton step. We expand the chemical potential term using equation 1.68 and non-dimensionalize the equation by $\hat{R}T$. We make sure to convert these equations into a form that can be differentiated by our non-linear variables (i.e. $N_j = \exp(\ln(\mathcal{N}_j))$). We also set $\pi = -\lambda/\hat{R}T$.

1.5.3 Derivatives for f_1

The derivatives for f_1 are then:

$$\frac{\partial f_1}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \Delta \ln \mathcal{N}_j \quad (1.78)$$

$$\frac{\partial f_1}{\partial [\ln T]} \Delta \ln T = -\frac{U_j^\circ}{\hat{R}T} \ln T \quad (1.79)$$

$$\frac{\partial f_1}{\partial \pi_i} \pi_i = -a_{ij} \pi_i \quad (1.80)$$

$$\frac{\partial f_1}{\partial \pi_q} \pi_q = -q_j \pi_q \quad (1.81)$$

The derivation for equation [1.79] is carried out in more depth in section [3.2]. Therefore, our final result is:

$$\Delta \ln \mathcal{N}_j - \sum_{i=1}^{NE} a_{ij} \pi_i - q_j \pi_q - \frac{U_j^\circ}{\hat{R}T} \Delta \ln T = -\frac{\mu_j}{\hat{R}T}$$

There will be NS number of these equations in our system.

1.5.4 Derivatives for f_2

Next, we take the derivatives of f_2 :

$$\frac{\partial f_2}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j a_{i,j} \Delta \ln \mathcal{N}_j \quad (1.82)$$

$$\frac{\partial f_2}{\partial [\ln T]} \Delta \ln T = 0 \quad (1.83)$$

$$\frac{\partial f_2}{\partial \pi_i} \pi_i = 0 \quad (1.84)$$

$$\frac{\partial f_2}{\partial \pi_q} \pi_q = 0 \quad (1.85)$$

Which gives:

$$\boxed{\sum_{j=1}^{\text{NS}} \mathcal{N}_j a_{ij} \Delta \ln \mathcal{N}_j = b_i^\circ - \sum_{j=1}^{\text{NS}} a_{ij} \mathcal{N}_j}$$

There will be NE of these equations.

1.5.5 Derivatives for f_3

For f_3 we get:

$$\frac{\partial f_3}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j q_j \Delta \ln \mathcal{N}_j \quad (1.86)$$

$$\frac{\partial f_3}{\partial [\ln T]} \Delta \ln T = 0 \quad (1.87)$$

$$\frac{\partial f_3}{\partial \pi_i} \pi_i = 0 \quad (1.88)$$

$$\frac{\partial f_3}{\partial \pi_q} \pi_q = 0 \quad (1.89)$$

Which gives only one equation:

$$\boxed{\sum_{j=1}^{\text{NS}} \mathcal{N}_j q_j \Delta \ln \mathcal{N}_j = - \sum_{j=1}^{\text{NS}} q_j \mathcal{N}_j}$$

1.5.6 Derivatives for f_4

For internal energy, we have:

$$f_4(\ln \mathcal{N}_j, \ln T) = \sum_{j=1}^{\text{NS}} \mathcal{N}_j U_j^\circ - u'_0 = u' - u'_0 \quad (1.90)$$

Using identity [3.1] helps with the temperature derivative. The derivatives are then:

$$\frac{\partial f_4}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \frac{\mathcal{N}_j U_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j \quad (1.91)$$

$$\frac{\partial f_4}{\partial [\ln T]} \Delta \ln T = \sum_{j=1}^{NS} [T \mathcal{N}_j c_{v,j}^\circ] \Delta \ln T \quad (1.92)$$

$$\frac{\partial f_4}{\partial \pi_i} \pi_i = 0 \quad (1.93)$$

$$\frac{\partial f_4}{\partial \pi_q} \pi_q = 0 \quad (1.94)$$

After non-dimensionalizing, we are left with:

$$\boxed{\sum_{j=1}^{NS} \frac{\mathcal{N}_j U_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{NS} \frac{\mathcal{N}_j c_{v,j}^\circ}{R} \Delta \ln T = \frac{u'_0 - u'}{\hat{R}T}}$$

1.5.7 Derivatives for f_5

For the entropy constraint we have:

$$f_5(\ln \mathcal{N}_j, \ln T) = \sum_{j=1}^{NS} \mathcal{N}_j S_j - s'_0 = s' - s'_0 \quad (1.95)$$

Again, identity [3.1] assists us. The derivates are:

$$\frac{\partial f_5}{\partial [\ln \mathcal{N}_j]} \Delta \ln \mathcal{N}_j = \mathcal{N}_j [S_j - \hat{R}] \Delta \ln \mathcal{N}_j \quad (1.96)$$

$$\frac{\partial f_5}{\partial [\ln T]} \Delta \ln T = \mathcal{N}_j c_{v,j}^\circ \Delta \ln T \quad (1.97)$$

$$\frac{\partial f_5}{\partial \pi_i} \pi_i = 0 \quad (1.98)$$

$$\frac{\partial f_5}{\partial \pi_q} \pi_q = 0 \quad (1.99)$$

Which, after non-dimensionalizing gives the singular equation:

$$\boxed{\sum_{j=1}^{NS} \mathcal{N}_j \left[\frac{S_j}{\hat{R}} - 1 \right] \Delta \ln \mathcal{N}_j + \mathcal{N}_j \frac{c_{v,j}^\circ}{\hat{R}} \Delta \ln T = \frac{s'_0 - s'}{\hat{R}}} \quad (1.100)$$

For convenience, they are all listed here together:

$$\Delta \ln \mathcal{N}_j - \sum_{i=1}^{NE} a_{ij} \pi_i - q_j \pi_q - \frac{U_j^\circ}{\hat{R}T} \Delta \ln T = -\frac{\mu_j}{\hat{R}T} \quad (1.101)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j a_{ij} \Delta \ln \mathcal{N}_j = b_i^\circ - \sum_{j=1}^{NS} a_{ij} \mathcal{N}_j \quad (1.102)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j q_j \Delta \ln \mathcal{N}_j = - \sum_{j=1}^{NS} q_j \mathcal{N}_j \quad (1.103)$$

$$\sum_{j=1}^{NS} \frac{\mathcal{N}_j U_j^\circ}{\hat{R}T} \Delta \ln \mathcal{N}_j + \sum_{j=1}^{NS} \frac{\mathcal{N}_j c_{v,j}^\circ}{R} \Delta \ln T = \frac{u'_0 - u'}{\hat{R}T} \quad (1.104)$$

$$\sum_{j=1}^{NS} \mathcal{N}_j \left[\frac{S_j}{\hat{R}} - 1 \right] \Delta \ln \mathcal{N}_j + \mathcal{N}_j \frac{c_{v,j}^\circ}{\hat{R}} \Delta \ln T = \frac{s'_0 - s'}{\hat{R}} \quad (1.105)$$

1.5.8 Reduced Helmholtz Equations

A "simplification" can be made to the above system of equations by solving equations [1.101] for $\Delta \ln \mathcal{N}_j$:

$$\Delta \ln \mathcal{N}_j = \sum_{i=1}^{NE} a_{i,j} \pi_i - \frac{\mu_j}{\hat{R}T} + q_j \pi_q + \frac{U_j^\circ}{\hat{R}T} \Delta \ln T \quad (1.106)$$

By substituting this into equations [1.102] - [1.104], we can shrink the system of equations dramatically. The resulting set of equations is:

$$\sum_{i=1}^{NE} \left[\sum_{j=1}^{NS} a_{kj} a_{ij} \mathcal{N}_j \right] \pi_i + \left[\sum_{j=1}^{NS} a_{kj} q_j \mathcal{N}_j \right] \pi_q + \left[\sum_{j=1}^{NS} a_{kj} \mathcal{N}_j \frac{U_j^\circ}{\hat{R}T} \right] \Delta \ln T = b_k^\circ - \sum_{j=1}^{NS} a_{kj} \mathcal{N}_j + \sum_{j=1}^{NS} a_{kj} \mathcal{N}_j \frac{\mu_j}{\hat{R}T} \quad (1.107)$$

$$\sum_{i=1}^{NE} \left[\sum_{j=1}^{NS} a_{ij} q_j \mathcal{N}_j \right] \pi_i + \left[\sum_{j=1}^{NS} q_j^2 \mathcal{N}_j \right] \pi_q + \left[\sum_{j=1}^{NS} q_j \mathcal{N}_j \frac{U_j^\circ}{\hat{R}T} \right] \Delta \ln T = \sum_{j=1}^{NS} q_j \mathcal{N}_j \frac{\mu_j}{\hat{R}T} - \sum_{j=1}^{NS} q_j \mathcal{N}_j \quad (1.108)$$

$$\begin{aligned} \sum_{i=1}^{NE} \left[\sum_{j=1}^{NS} a_{ij} \mathcal{N}_j \frac{U_j^\circ}{\hat{R}T} \right] \pi_i + \left[\sum_{j=1}^{NS} q_j \mathcal{N}_j \frac{U_j^\circ}{\hat{R}T} \right] \pi_q + & \left[\sum_{j=1}^{NS} \mathcal{N}_j \left(\frac{U_j^\circ}{\hat{R}T} \right)^2 + \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{v,j}^\circ}{\hat{R}} \right] \Delta \ln T \\ & = \frac{u'_0 - u'}{\hat{R}T} + \sum_{j=1}^{NS} \mathcal{N}_j \frac{U_j^\circ}{\hat{R}T} \frac{\mu_j}{\hat{R}T} \end{aligned} \quad (1.109)$$

$$\begin{aligned} \sum_{i=1}^{NE} \left[\sum_{j=1}^{NS} a_{ij} \mathcal{N}_j \left(\frac{S_j}{\hat{R}} - 1 \right) \right] \pi_i + \left[\sum_{j=1}^{NS} q_j \mathcal{N}_j \left(\frac{S_j}{\hat{R}} - 1 \right) \right] \pi_q + & \left[\sum_{j=1}^{NS} \mathcal{N}_j \left(\frac{S_j}{\hat{R}} - 1 \right) \frac{U_j^\circ}{\hat{R}T} + \sum_{j=1}^{NS} \mathcal{N}_j \frac{c_{v,j}^\circ}{\hat{R}} \right] \Delta \ln T = \frac{s'_0 - s'}{\hat{R}} + \sum_{j=1}^{NS} \mathcal{N}_j \frac{\mu_j}{\hat{R}T} \left(\frac{S_j}{\hat{R}} - 1 \right) \end{aligned} \quad (1.110)$$

There will be NE number of equations 1.107, and one of 1.108 if charge is added, and one of 1.109/1.110 depending on which one you made need.

The equations are colored according to when they are needed. **Red coloring** denotes terms that arise from the elemental constraint condition. These are included no matter what. **Blue** denotes terms that are added when you include the charge constraint. **Violet** terms are added for both UV and SV constraints. **orange** and **teal** are the rows added for either internal energy or entropy constraints.

1.6 Useful Relationships

$$\frac{c_v^\circ}{\hat{R}} = \frac{c_{p,j}^\circ}{\hat{R}} - 1 \quad (1.111)$$

$$\frac{\mu_j^\circ}{\hat{R}T} = \frac{H_j^\circ}{\hat{R}T} - \frac{S_j^\circ}{\hat{R}} \quad (1.112)$$

$$\frac{U_j^\circ}{\hat{R}T} = \frac{H_j^\circ}{\hat{R}T} - 1 \quad (1.113)$$

2 Univeristy of Minnesota Chemical Equilibrium Code UMCEC

The University of Minnesota code for computing chemical equilibrium of a gas mixture uses the well documented method of Gibbs and Helmholtz energy minimization with Lagrange multipliers as well as charge constrains for plasmas, and enthalpy, entropy, and internal energy constraints depending on the minimization method. It computes the composition and thermodynamic state of gas mixtures.

3 Standard State Chemical Potential Derivatives

Some important derivatives are derived here. First, we state a chain rule identity that is very useful:

$$\frac{\partial}{\partial T} = \frac{\partial[\ln(T)]}{\partial T} \frac{\partial}{\partial[\ln(T)]} \rightarrow \frac{\partial}{\partial[\ln(T)]} = T \frac{\partial}{\partial T} \quad (3.1)$$

Then, taking the derivative of μ_j^0 gives:

$$\frac{\partial}{\partial[\ln(T)]} \left(\frac{\mu_j^0}{RT} \right) = T \frac{\partial}{\partial T} \left(\frac{\mu_j^0}{RT} \right) \quad (3.2)$$

We expand μ_j^0/RT as:

$$\frac{\mu_j^0}{RT} = \frac{H_j^0}{RT} - \frac{S_j^0}{R} \quad (3.3)$$

$$\frac{\mu_j^0}{RT} = \frac{U_j^0}{RT} - \frac{S_j^0}{R} + 1 \quad (3.4)$$

3.1 Gibbs Derivative

First, we take the derivative of [3.3] for Gibbs minimization.

$$T \frac{\partial}{\partial T} \left(\frac{H_j^0}{RT} - \frac{S_j^0}{R} \right) = T \left(\frac{\partial_T(H_j^0) \cdot RT - H_j^0 \cdot R}{(RT)^2} - \frac{\partial_T(S_j^0)}{R} \right) \quad (3.5)$$

Here, ∂_T denotes $(\partial/\partial T)$. If:

$$c_{p,j}^0 = \frac{\partial H_j^0}{\partial T}, \text{ and } \frac{c_{p,j}^0}{T} = \frac{\partial S_j^0}{\partial T} \quad (3.6)$$

This reduces to:

$$T \left[\frac{c_{p,j}^0}{RT} - \frac{H_j^0}{RT^2} - \frac{c_{p,j}^0}{RT} \right] = -\frac{H_j^0}{RT} \quad (3.7)$$

Therefore:

$$\boxed{\frac{\partial}{\partial[\ln(T)]} \left(\frac{\mu_j^0}{RT} \right) = -\frac{H_j^0}{RT}} \quad (3.8)$$

3.2 Helmholtz Derivative

Using equation [3.4] gives us:

$$T \frac{\partial}{\partial T} \left(\frac{U_j^0}{RT} + 1 - \frac{S_j^0}{R} \right) = T \left(\frac{\partial_T(U_j^0) \cdot RT - U_j^0 \cdot R}{(RT)^2} - \frac{\partial_T(S_j^0)}{R} \right) \quad (3.9)$$

Recalling $\partial_T(S_j^0)$ from equation [3.6] and using:

$$c_{v,j}^0 = \frac{\partial U_j^0}{\partial T} \quad (3.10)$$

Equation [3.9] reduced to:

$$T \left[\frac{c_{v,j}^0}{RT} - \frac{U_j^0}{RT^2} - \frac{c_{p,j}^0}{RT} \right] \quad (3.11)$$

Since:

$$\frac{c_{v,j}^0}{R} - \frac{c_{p,j}^0}{R} = -1 \quad (3.12)$$

The final derivative is:

$$\boxed{\frac{\partial}{\partial[\ln(T)]} \left(\frac{\mu_j^0}{RT} \right) = -\frac{U_j^0}{RT} - 1} \quad (3.13)$$

4 Matrix Method for computation of Jacobian matrix for CFD

The reduced equations for Helmholtz energy minimization contains quite a large number of sums. These sums are necessary for every iteration since the sums contain \mathcal{N}_j , or some thermodynamic value like H_j° . However, this can be reduced even further with the introduction of the ξ matrix, and a separate matrix ω which does not reduce computation time.

Since each original Jacobian entry is essentially just the dot product of some vector with the \mathcal{N}_j vector, we can create these matrices such that, when multiplied by the column vector $\hat{\mathcal{N}}_j$, give a resultant column vector that contains terms for the Jacobian matrix and RHS vector of the Newton-Raphson iteration. This does not speed up the computation by a lot, but it certainly gets rid of some redundancy as well as providing some cache locality. The ξ matrix is defined as

$$\xi(a_{ij}, q_j) \begin{bmatrix} \mathcal{N}_1 & \mathcal{N}_2 & \dots & \mathcal{N}_{NS} \end{bmatrix}^T = \mathcal{J}(a_{ij}, q_j, \mathcal{N}_j) \quad (4.1)$$

Where the variables in parenthesis mean that that matrix or vector only contain sums with those variables. The ω matrix is defined as:

$$\omega(a_{ij}, q_j, H_j^\circ) \begin{bmatrix} \mathcal{N}_1 & \mathcal{N}_2 & \dots & \mathcal{N}_{NS} \end{bmatrix}^T = \mathcal{J}(a_{ij}, q_j, H_j^\circ, \mathcal{N}_j) \quad (4.2)$$

These matrices will be shown for each section of the Jacobian. The number of columns of the ξ and ω matrices is equal to the number of species of gas in the system.

4.1 The ξ matrix

4.1.1 Elemental Rows

The following matrix creates the k^{th} row of the Jacobian matrix, where k varies from 0 to $\text{NE} - 1$. The left hand matrix denotes which entry of the Jacobian matrix that row is associated with. There will be NE number of these stacked on top of each other:

$$\begin{vmatrix} J[k, 0] \\ \vdots \\ J[k, \text{NE} - 1] \\ J[k, \text{NE}] \end{vmatrix} = \begin{bmatrix} a_{k,1}a_{1,1} & \dots & a_{k,NS}a_{1,NS} \\ \vdots & \ddots & \vdots \\ a_{k,1}a_{\text{NE},1} & \dots & a_{k,NS}a_{\text{NE},NS} \\ a_{k,1}q_1 & \dots & a_{k,NS}q_{NS} \end{bmatrix} \begin{bmatrix} \mathcal{N}_1 \\ \vdots \\ \mathcal{N}_{NS} \end{bmatrix} \quad (4.3)$$

4.1.2 Charge rows

For the charge constraint, we have:

$$\begin{vmatrix} J[\text{NE}, 0] \\ \vdots \\ J[\text{NE}, \text{NE} - 1] \\ J[\text{NE}, \text{NE}] \end{vmatrix} = \begin{bmatrix} a_{1,1}q_1 & \dots & a_{1,NS}q_{NS} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1}q_1 & \dots & a_{\text{NE},NS}q_{NS} \\ q_1^2 & \dots & q_{NS}^2 \end{bmatrix} \begin{bmatrix} \mathcal{N}_1 \\ \vdots \\ \mathcal{N}_{NS} \end{bmatrix} \quad (4.4)$$

The elemental rows has already shown up in the previous section, so we can reuse its values and therefore the only additional row is only:

$$\left| J[\text{NE}, \text{NE}] \right| = \begin{bmatrix} q_1^2 & \dots & q_{\text{NS}}^2 \end{bmatrix} \begin{bmatrix} \mathcal{N}_1 \\ \vdots \\ \mathcal{N}_{\text{NS}} \end{bmatrix} \quad (4.5)$$

With:

$$\begin{vmatrix} J[\text{NE}, 0] \\ \vdots \\ J[\text{NE}, \text{NE} - 1] \end{vmatrix} = \begin{vmatrix} J[0, \text{NE}] \\ \vdots \\ J[\text{NE} - 1, \text{NE}] \end{vmatrix} \quad (4.6)$$

4.1.3 RHS

At the end of our ξ matrix, we place two terms that arrive on the RHS, namely the dot products of \mathcal{N}_j with elemental and charges. Therefore we get:

$$\begin{vmatrix} \text{RHS}[i] \\ \vdots \\ \text{RHS}[\text{NE} - 1] \\ \text{RHS}[\text{NE}] \end{vmatrix} = \begin{bmatrix} a_{11} & \dots & a_{1,\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1} & \dots & a_{\text{NE},\text{NS}} \\ q_1 & \dots & q_{\text{NS}} \end{bmatrix} \begin{bmatrix} \mathcal{N}_1 \\ \vdots \\ \mathcal{N}_{\text{NS}} \end{bmatrix} \quad (4.7)$$

4.1.4 Full ξ matrix

The full size of the ξ matrix is $[\text{NE}^2 + \text{NE} + \eta(\text{NE} + 2)] \cdot \text{NS}$, where η equals 0 for no ionization, and 1 for ionization. For 5-species air, this is a 30 entry matrix. For 11 species air with 2 elements, this is a 111 element matrix. Here, all charge entries have been brought to the bottom for easy indexing.

$$\xi \mathcal{N} = \begin{bmatrix} a_{k1}a_{11} & \dots & a_{k,\text{NS}}a_{1,\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{k,1}a_{\text{NE},1} & \dots & a_{k,\text{NS}}a_{\text{NE},\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1}a_{11} & \dots & a_{\text{NE},\text{NS}}a_{1,\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1}a_{\text{NE},1} & \dots & a_{\text{NE},\text{NS}}a_{\text{NE},\text{NS}} \\ a_{11} & \dots & a_{1,\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1} & \dots & a_{\text{NE},\text{NS}} \\ a_{k,1}q_1 & \dots & a_{k,\text{NS}}q_{\text{NS}} \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1}q_1 & \dots & a_{\text{NE},\text{NS}}q_{\text{NS}} \\ q_1 & \dots & q_{\text{NS}} \\ q_1^2 & \dots & q_{\text{NS}}^2 \end{bmatrix} \begin{bmatrix} \mathcal{N}_1 \\ \vdots \\ \mathcal{N}_i \\ \vdots \\ \mathcal{N}_{\text{NS}} \end{bmatrix} = \begin{bmatrix} \text{RHS}[k \cdot \text{NE}] \\ \vdots \\ \text{RHS}[k \cdot \text{NE} + (\text{NE} - 1)] \\ \vdots \\ \text{RHS}[(\text{NE} - 1)\text{NE}] \\ \vdots \\ \text{RHS}[(\text{NE} - 1)\text{NE} + \text{NE} - 1] \\ \text{RHS}[\text{NE} \cdot \text{NE}] \\ \vdots \\ \text{RHS}[\text{NE} \cdot \text{NE} + \text{NE} - 1] \\ \text{RHS}[\text{NE} \cdot (\text{NE} + 1)] \\ \vdots \\ \text{RHS}[\text{NE} \cdot (\text{NE} + 1) + \text{NE} - 1] \\ \text{RHS}[\text{NE} \cdot (\text{NE} + 2)] \\ \text{RHS}[\text{NE} \cdot (\text{NE} + 2) + 1] \end{bmatrix} \quad (4.8)$$

4.2 The ω matrix

The ω matrix is dedicated to terms that can't be created at the start of the loop and left as constants. These need to be updated since they are temperature dependant and temperature varies in our iterations.

For Helmholtz minimization we use $U_j^\circ/\hat{R}T$ which is calculated as $H_j^\circ/\hat{R}T - 1$. We can split many terms inside sums (such as the elemental one) as:

$$a_{kj} \left(\frac{H_j^\circ}{\hat{R}T} - 1 \right) \mathcal{N}_j = a_{kj} \frac{H_j^\circ}{\hat{R}T} \mathcal{N}_j - a_{kj} \mathcal{N}_j \quad (4.9)$$

Then, using the linearity of sums, the summation over species becomes:

$$\sum_j^{\text{NS}} \left[a_{kj} \frac{H_j^\circ}{\hat{R}T} \mathcal{N}_j - a_{kj} \mathcal{N}_j \right] = \boxed{\sum_j^{\text{NS}} a_{kj} \frac{H_j^\circ}{\hat{R}T} \mathcal{N}_j - \sum_{j=1}^{\text{NS}} a_{kj} \mathcal{N}_j} \quad (4.10)$$

This allows us so computational savings since we have already calculated the negative term in the ξ matrix.

4.2.1 Elemental Rows

The first NE rows of this matrix looks like:

$$\begin{bmatrix} a_{k1} H_1^\circ / \hat{R}T & \dots & a_{k,\text{NS}} H_{\text{NS}}^\circ \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1} H_1^\circ / \hat{R}T & \dots & a_{\text{NE},\text{NS}} H_{\text{NS}}^\circ \end{bmatrix} \quad (4.11)$$

4.2.2 Charge Rows

$$\begin{bmatrix} q_1 H_1^\circ / \hat{R}T & \dots & q_{\text{NS}} H_{\text{NS}}^\circ / \hat{R}T \end{bmatrix} \quad (4.12)$$

4.2.3 Internal Energy row

$$\begin{bmatrix} a_{i1} H_1^\circ / \hat{R}T & \dots & a_{i,\text{NS}} H_{\text{NS}}^\circ \\ \vdots & \ddots & \vdots \\ a_{\text{NE},1} H_1^\circ / \hat{R}T & \dots & a_{\text{NE},\text{NS}} H_{\text{NS}}^\circ \\ q_1 H_1^\circ / \hat{R}T & \dots & q_{\text{NS}} H_{\text{NS}}^\circ / \hat{R}T \\ H_1^\circ / \hat{R}T & \dots & H_{\text{NS}}^\circ / \hat{R}T \\ (H_1^\circ / \hat{R}T)^2 & \dots & (H_{\text{NS}}^\circ / \hat{R}T)^2 \\ c_{p,1}^\circ / \hat{R} - 1 & \dots & c_{p,\text{NS}}^\circ / \hat{R} - 1 \end{bmatrix} \quad (4.13)$$

Of course, we note again that the first $\text{NE} + 1$ rows have already been calcualted, so we can reuse their values. Therefore the system is only:

$$\begin{bmatrix} H_1^\circ / \hat{R}T & \dots & H_{\text{NS}}^\circ / \hat{R}T \\ (H_1^\circ / \hat{R}T)^2 & \dots & (H_{\text{NS}}^\circ / \hat{R}T)^2 \\ c_{p,1}^\circ / \hat{R} - 1 & \dots & c_{p,\text{NS}}^\circ / \hat{R} - 1 \end{bmatrix} \quad (4.14)$$

4.3 RHS

Many terms that show up on the RHS are calculated from temperature dependant variables.

5 Coding Tricks