

## Article

# The Effect of Elemental Abundances on Fitting Supernova Remnant Models to Data

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**Abstract:** Models for supernova remnant (SNR) evolution can be used to determine the energy of the explosion, the age of the SNR and the density of the surrounding medium when the model successfully reproduces the observed X-ray emission spectrum and other properties. Observed SNR properties include the shock radius, and the electron temperature ( $kT_e$ ) and emission measure ( $EM$ ) of the shocked-gas which are derived from the X-ray spectrum. The standard and XSPEC definitions  $kT_e$  and  $EM$  have an important difference, which is not well known. The XSPEC definition is superior for SNRs, which have components with low hydrogen abundance. SNR model calculations are based on hydrodynamic solutions for fluid variables of density, pressure and velocity. The relations between fluid variables and  $kT_e$  or  $EM$  depend on composition, ionization state and electron-ion temperature ratio ( $T_e/T_I$ ). Here the effects of composition, ionization and  $T_e/T_I$  on standard and on XSPEC versions of  $kT_e$  and  $EM$  are investigated.

**Keywords:** supernova remnants; supernova energetics; interstellar medium density

## 1. Introduction

Supernova remnants (SNRs) are a significant force in galaxies: they add energy to the interstellar medium (ISM) (e.g. [1]) and spread elements, which are created in the supernovae (SN), explosion around the ISM (e.g. [2]). From SNR studies we can learn about the end states of stellar evolution, the properties of ISM, and the impact of SN explosions on the Galaxy. SNR research has several purposes, including learning about SN explosions and the effects of SN mass and energy input to the ISM.

The observational data for a given SNRs depends on the brightness of emissions in different wavebands and by the instruments used for the observations. SNRs in our Galaxy have been mainly discovered by their radio emission [3]. To characterize a SNR, which has most of its energy contained in the X-ray emitting shocked gas, X-ray spectral observations are required.

Among the early important studies of X-ray spectra emitted by SNRs is that of [4]. The X-ray spectrum diagnoses the amount of shocked gas, via the emission measure ( $EM$ ), and the shocked gas electron temperature ( $kT_e$ ). X-ray observations of some historical SNR have been modelled with tailored hydrodynamic simulations (e.g. [5]). The majority of Galactic SNRs have less complete observations than the historical SNRs, including no observed ages. For these, usually a basic Sedov model with assumed energy and ISM density has been applied to obtain age estimates. However, SNRs have a wide range of energies and ISM densities ([6], [7], [8], [9]), which were obtained using more physically realistic models than the Sedov model. Some dispersion in energy is expected based on simulations of core-collapse SN (CCSN) ([10], [11] and references therein). As discussed by the comparison of simulations with observations of CCSN energies ([12]), both are heavily biased, implying our knowledge of SN energies is still quite incomplete.

To characterise SNRs, we have developed a set of models which are based on hydrodynamic calculations with scalings derived using the unified model of [13]. These

models assume spherical symmetry and are described in [14] and [15]. The observed quantities of electron temperatures and emission measures of the hot plasma, for forward-shocked and for reverse-shocked gas are calculated in the model. These models are important to enable the process of using X-ray observations, normally analysed using XSPEC [16] to determine  $EM$  and  $kT_e$ , to obtain the physical properties of a SNR, such as explosion energy, age and ISM density.

This work includes a detailed consideration of the effect of composition and partial ionization on the emission measures and shock temperatures that are calculated using hydrodynamic simulations. This is an important extension of the models presented in [14]. In Section 2.1 we present an overview of the SNR model. In Section 2.2 the standard and XSPEC definitions of emission measure ( $EM$ ) and  $EM$ -weighted gas temperature ( $kT_{EM}$ ) are compared and related to mean molecular weights. In Section 2.3.1 the relation of gas temperature to electron temperature ( $kT_{e,EM}$ ) is discussed. In Section 2.3 the dependence of temperature and emission measure of shocked gas on chemical composition and ionization state are determined. In Sections 3.2 and 3.3, the scaling relations for  $EM$  and  $kT_{e,EM}$  on mean molecular weights are given. The conclusions are summarized in Section 4.

## 2. Analysis

The assumption here is that a hydrodynamic model for SNR evolution is calculated, with fundamental variables of density, pressure and velocity of the gas. The aim of that model is to reproduce observed quantities of a given SNR, in particular  $EM$  and  $kT_{e,EM}$ , by relating the computed hydrodynamic variables to the observed variables. In order to make simplifying assumptions in the calculations of effects of chemical composition and ionization of the gas, a reference model is chosen. This is the spherically symmetric model of [14] (and references therein). However the results below apply to any SNR model subject to the particular assumptions made by that model.

### 2.1. The Model for SNR Evolution

A SNR is the interaction of the SN ejecta with the interstellar medium (ISM). The various stages of evolution of a SNR are labelled the ejecta-dominated stage (ED), the adiabatic or Sedov-Taylor stage (ST), the radiative pressure-driven snowplow (PDS) and the radiative momentum-conserving shell (MCS). These stages are reviewed in, e.g., [17], [18], [13], [3] and [15]. In addition, there are the transitions between stages, called ED to ST, ST to PDS, and PDS to MCS, respectively. The ED to ST stage is important because the SNR is still bright, and it is long-lived enough [14] that a significant fraction of SNRs are likely in this phase.

For simplicity our models assume that the SN ejecta and ISM are spherically symmetric. The ISM density profile is a power-law centred on the SN explosion, given by  $\rho_{ISM} = \rho_0 r^{-s}$ , with  $s=0$  (constant density medium) or  $s=2$  (stellar wind density profile). The unshocked ejecta has a power-law density  $\rho_{ej} \propto r^{-n}$  envelope with constant density core. With these profiles, the ED phase of the SNR evolution has a self-similar evolution ([19] and [20]) prior to the reverse shock hitting the core [13]. The evolution of SNR shock radius was extended for the ED to ST and ST phases by [13].

The model for SNR evolution that we use is partly based on the TM99 analytic solutions, with additional features. A detailed description of the model is given in [14]. Hydrodynamic variables for the interior structure of the SNR are calculated from hydrodynamic simulations which cover ED, ED to ST and ST phases. The scaling relations of the unified solution of [13] are used to keep the size of grid of hydro models feasible for calculation. Electron-ion temperature equilibration ( $T_e/T_I$ ) is included after the hydro calculations.  $T_e/T_I$  is calculated using the Coulomb collisional electron heating mechanism, consistent with the observational results of [21]. The emission measure ( $EM$ ) and emission measure-weighted electron temperature ( $T_{e,EM}$ ) are calculated from the hydrodynamic variables, gas composition and  $T_e/T_I$ . The inverse modelling problem was solved by [7].

This takes as input the SNR observed properties and determines the initial properties of the SNR.

The current version of the SNR modelling program calculates  $EM$  and  $kT_{e,EM}$  using the standard definition of these quantities. The current work presents the full dependence of  $EM$  and  $kT_{e,EM}$  on composition, ionization and  $T_e/T_I$ . It extends the calculations to include the definitions of  $EM$  and  $kT_{e,EM}$  used by the standard X-ray spectrum modelling program XSPEC, which differ in important ways from the standard definitions (see Section 2.2 below).

## 2.2. Emission measure (EM) and EM-weighted gas temperature ( $kT_{EM}$ )

Observed SNR quantities from the X-ray spectrum to be modelled are the emission measure,  $EM$ , and the  $EM$ -weighted electron temperature,  $kT_{e,EM}$ , for both forward-shocked ISM (FS) and reverse-shocked ejecta (RS). The relation between  $kT_{e,EM}$  and gas temperature  $kT_{EM}$  is discussed in Section 2.3.1 below. The standard definition of  $EM$  is:

$$EM = \int n_e(r)n_H(r)dV \quad (1)$$

with  $n_e$  and  $n_H$  the number densities of electrons and hydrogen nuclei, respectively, and the integration is over the volume of the emitting gas.  $kT_{EM}$  is given by

$$kT_{EM} = \int n_e(r)n_H(r)kT(r)dV/EM \quad (2)$$

However, the  $EM$  used by XSPEC and by observers, called  $EM_{XS}$  here, has a modified definition of  $n_H$  [23], called  $n_{H,XS}$  here, designed to include cases of very low hydrogen abundance without diverging. The total ion density  $n_I$  is converted to  $n_{H,XS}$  using cosmic abundances:  $n_{H,XS} = n_I \frac{x_{H,C}}{\sum_j x_{j,C}} = n_I x_{H,C}$ , where  $x_{j,C}$  are number fractions of all elements for cosmic abundances and the latter expression assumes they are normalized ( $\sum_j x_{j,C} = 1$ ). Thus one has

$$EM_{XS} = \int n_e(r)n_{H,XS}(r)dV = x_{H,C} \int n_e(r)n_I(r)dV \quad (3)$$

with  $x_{H,C} = 0.921^1$  the fractional number abundance of H.  $EM_{XS}$  defined in terms of the ion density whereas  $EM$  is defined in terms of hydrogen density.  $kT_{EM,XS}$  is found using  $EM_{XS}$ :

$$kT_{EM,XS} = \int n_e(r)n_{H,XS}(r)kT(r)dV/EM_{XS} \quad (4)$$

If  $n_I(r) \propto n_H(r)$  then one has  $kT_{EM,XS} = kT_{EM}$ . In general, spatial variations in composition violate a constant proportionality, so  $kT_{EM,XS}$  and  $kT_{EM}$  are different.

## 2.3. Dependence of $EM$ and $kT_{EM}$ on mean molecular weights and ionization

A hydrodynamic simulation of a SNR yields the variables of mass density, pressure and velocity. To convert to gas temperature  $T(r)$ , total number density  $n$ , hydrogen number density  $n_H(r)$ , ion number density  $n_I(r)$ , and electron number density  $n_e(r)$  needed for  $EM$  and  $kT_{EM}$ , one uses the definition of the molecular weights ( $\mu$ 's),

$$\rho = \mu m_H n = \mu_H m_H n_H = \mu_I m_H n_I = \mu_e m_H n_e \quad (5)$$

Temperature is determined using the ideal gas law,

$$T = \frac{\mu m_H}{k_B} \frac{P}{\rho} \quad (6)$$

The dependence on ionization state of the gas is included in the  $\mu$ 's.

<sup>1</sup> Calculated using the solar abundances from [22].

$EM$  and  $kT_{EM}$  for FS and RS gas in terms of mass density and pressure are:

$$\begin{aligned} EM_{FS} &= \int_{R_{CD}}^{R_{FS}} \frac{\rho(r)^2}{m_H^2 \mu_{e,FS}(r) \mu_{H,FS}(r)} dV \\ &= \frac{1}{\mu_{e,FS} \mu_{H,FS} m_H^2} \int_{R_{CD}}^{R_{FS}} \rho(r)^2 dV \\ EM_{RS} &= \int_{R_{RS}}^{R_{CD}} \frac{\rho(r)^2}{m_H^2 \mu_{e,RS}(r) \mu_{H,RS}(r)} dV \\ &= \frac{1}{\mu_{e,RS} \mu_{H,RS} m_H^2} \int_{R_{RS}}^{R_{CD}} \rho(r)^2 dV \end{aligned} \quad (7)$$

$$\begin{aligned} kT_{EM,FS} &= \int_{R_{CD}}^{R_{FS}} \frac{\rho(r)^2}{m_H^2 \mu_{e,FS}(r) \mu_{H,FS}(r)} \frac{P(r) \mu_{FS}(r) m_H}{\rho(r) k_B} dV / EM_{FS} \\ &= \frac{\mu_{FS} m_H}{k_B} \int_{R_{CD}}^{R_{FS}} \rho(r) P(r) dV / \int_{R_{CD}}^{R_{FS}} \rho(r)^2 dV \\ kT_{EM,RS} &= \int_{R_{RS}}^{R_{CD}} \frac{\rho(r)^2}{m_H^2 \mu_{e,RS}(r) \mu_{H,RS}(r)} \frac{P(r) \mu_{RS}(r) m_H}{\rho(r) k_B} dV / EM_{RS} \\ &= \frac{\mu_{RS} m_H}{k_B} \int_{R_{RS}}^{R_{CD}} \rho(r) P(r) dV / \int_{R_{RS}}^{R_{CD}} \rho(r)^2 dV \end{aligned} \quad (8)$$

where the  $\mu$ 's are assumed spatially uniform to simplify the integrals<sup>2</sup>. For  $EM_{RS}$  and  $kT_{EM,RS}$  the lower limit of integration is 0 after the RS reaches the center of the SNR.

For XSPEC defined quantities, one has:

$$\begin{aligned} EM_{XS,FS} &= x_{H,C} \int_{R_{CD}}^{R_{FS}} \rho(r)^2 / (m_H^2 \mu_{e,FS}(r) \mu_{I,FS}(r)) dV \\ &= \frac{x_{H,C}}{\mu_{e,FS} \mu_{I,FS} m_H^2} \int_{R_{CD}}^{R_{FS}} \rho(r)^2 dV \\ EM_{XS,RS} &= x_{H,C} \int_{R_{RS}}^{R_{CD}} \rho(r)^2 / (m_H^2 \mu_{e,RS}(r) \mu_{I,RS}(r)) dV \\ &= \frac{x_{H,C}}{\mu_{e,RS} \mu_{I,RS} m_H^2} \int_{R_{RS}}^{R_{CD}} \rho(r)^2 dV \end{aligned} \quad (9)$$

$$\begin{aligned} kT_{EM,XS,FS} &= \int_{R_{CD}}^{R_{FS}} \frac{\rho(r)^2 x_{H,C}}{m_H^2 \mu_{e,FS}(r) \mu_{I,FS}(r)} \frac{P(r) \mu_{FS}(r) m_H}{\rho k_B} dV / EM_{XS,FS} \\ &= \frac{\mu_{FS} m_H}{k_B} \int_{R_{CD}}^{R_{FS}} \rho(r) P(r) dV / \int_{R_{CD}}^{R_{FS}} \rho(r)^2 dV \\ kT_{EM,XS,RS} &= \int_{R_{RS}}^{R_{CD}} \frac{\rho(r)^2 x_{H,C}}{m_H^2 \mu_{e,RS}(r) \mu_{I,RS}(r)} \frac{P(r) \mu_{RS}(r) m_H}{\rho k_B} dV / EM_{XS,RS} \\ &= \frac{\mu_{RS} m_H}{k_B} \int_{R_{RS}}^{R_{CD}} \rho(r) P(r) dV / \int_{R_{RS}}^{R_{CD}} \rho(r)^2 dV \end{aligned} \quad (10)$$

where the  $\mu$ 's are assumed spatially uniform to simplify the integrals.

<sup>2</sup> There are other cases where the integrals simplify, which we do not discuss in detail. E.g. if  $\mu_{e,FS}(r) \mu_{H,FS}(r) = \text{constant}$  then  $EM_{FS}$  simplifies, and if  $\mu_{e,FS}(r) \mu_{H,FS}(r) / \mu_{FS}(r) = \text{constant}$  then  $kT_{EM,FS}$  simplifies. Similar special cases for other integrals are not discussed here.

### 2.3.1. Electron temperature and electron-ion equilibration

The *EM*-weighted electron temperature,  $T_e$ , is measured by the X-ray spectrum. The relation between gas  $T$ , electron  $T_e$  and ion  $T_I$ , from  $P=P_e+P_I$ , is:

$$T/\mu = T_e/\mu_e + T_I/\mu_I \quad (11)$$

In general, the spatial dependence of  $T(r)$ ,  $T_e(r)$  and  $T_I(r)$  is complex.

In the case that shocked SNR gas has  $T_e$  and  $T_I$  equilibrated by Coulomb collisions [24] and is not old enough to have radiative losses, the electron-to-ion temperature ratio  $g(t) = T_e(t)/T_I(t) = g(T_s(t), T_e(t), t)$  increases to unity with  $t$  the age of a given parcel of gas since it was shocked. The calculation of  $g$  is summarized in [15].

$T_{e,FS}(r)$  and  $T_{e,RS}(r)$  are found using Equations 6 & 11 which give  $T_e = \frac{1}{1/\mu_e+1/(g\mu_I)} \frac{T}{\mu}$ :

$$\begin{aligned} T_{e,FS}(r) &= f_{T,FS}(r) \frac{P(r)m_H}{\rho(r)k_B} \\ T_{e,RS}(r) &= f_{T,RS}(r) \frac{P(r)m_H}{\rho(r)k_B} \end{aligned} \quad (12)$$

with  $f_{T,FS}(r) = \frac{1}{1/\mu_{e,FS}(r)+g(r)/\mu_{I,FS}(r)}$  and  $f_{T,RS}(r) = \frac{1}{1/\mu_{e,RS}(r)+g(r)/\mu_{I,RS}(r)}$ .  $f_{T,FS}$  and  $f_{T,RS}$  are constants in the approximations of uniform  $\mu$ 's and uniform  $g_{FS}$  and  $g_{RS}$ .

The standard FS and RS *EM*-weighted electron temperatures are the same as those given by Equation 8 except for inclusion of extra factors  $f_{T,FS}(r)/\mu_{FS}(r)$  and  $f_{T,RS}(r)/\mu_{RS}(r)$  in the integrals for  $kT_{e,EM,FS}$  and  $kT_{e,EM,RS}$ . The XSPEC-defined FS and RS *EM*-weighted electron temperatures are:

$$\begin{aligned} kT_{e,EM,XS,FS} &= \int_{R_{CD}}^{R_{FS}} \frac{\rho(r)^2 x_{H,C}}{m_H^2 \mu_{e,FS}(r) \mu_{I,FS}(r)} \frac{f_{T,FS}(r) P(r) m_H}{\rho(r) k_B} dV / EM_{XS,FS} \\ &= \frac{f_{T,FS} m_H}{k_B} \int_{R_{CD}}^{R_{FS}} \rho(r) P(r) dV / \int_{R_{CD}}^{R_{FS}} \rho(r)^2 dV \\ kT_{e,EM,XS,RS} &= \int_{R_{RS}}^{R_{CD}} \frac{\rho(r)^2 x_{H,C}}{m_H^2 \mu_{e,RS}(r) \mu_{I,RS}(r)} \frac{f_{T,RS}(r) P(r) m_H}{\rho(r) k_B} dV / EM_{XS,RS} \\ &= \frac{f_{T,RS} m_H}{k_B} \int_{R_{RS}}^{R_{CD}} \rho(r) P(r) dV / \int_{R_{RS}}^{R_{CD}} \rho(r)^2 dV \end{aligned} \quad (13)$$

where the second expressions for  $kT_{e,EM,XS,FS}$  and  $kT_{e,EM,XS,RS}$  apply in the case of uniform  $\mu$ 's and  $g$ 's. In that case, the standard and the XSPEC versions are identical<sup>3</sup>:

$$\begin{aligned} kT_{e,EM,XS,FS} &= kT_{e,EM,FS} \\ kT_{e,EM,XS,RS} &= kT_{e,EM,RS} \end{aligned} \quad (14)$$

## 3. Results and Discussion

### 3.1. Definition of EM

The standard *EM* is defined in terms of hydrogen density, whereas  $EM_{XS}$  is defined in terms of the ion density then converted to an equivalent hydrogen normalization (Equation 9).  $kT_{e,EM}$  is weighted by  $n_e n_H$  whereas  $kT_{e,EM,XS}$  is weighted by  $n_e n_I$ . The measured  $kT$  is best estimated by weighting  $kT_e(r)$  by the X-ray emissivity integrated over the emission volume. In the case of high hydrogen abundance  $kT_{e,EM}$  and  $kT_{e,EM,XS}$  are nearly the same, but for low hydrogen abundance, X-ray emissivity is closer to proportional to  $n_e n_I$  than to  $n_e n_H$ . Thus  $kT_{e,EM,XS}$  is the better measure. Overall, for properties measured using

<sup>3</sup> We do not discuss other special cases which also give identical results, such as  $kT_{e,EM,XS,FS} = kT_{e,EM,FS}$  for  $(f_{T,FS}(r)\mu_{FS}(r)) / (\mu_{e,FS}(r)\mu_{I,FS}(r)) = \text{constant}$ .

the X-ray spectrum, sensitive to all electrons and ions, the XSPEC definitions of  $EM$  and  $kT_e$  are better, despite the artifact that it is labelled with  $n_H$  rather than  $n_I$ . 134  
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### 3.2. Scaling Relations for $EM$ and $kT_{EM}$ 136

For simplicity, spatially uniform  $\mu_{e,FS}$ ,  $\mu_{I,FS}$ ,  $\mu_{H,FS}$ ,  $\mu_{e,RS}$ ,  $\mu_{I,RS}$ ,  $\mu_{H,RS}$ ,  $g_{FS}$  and  $g_{RS}$  are assumed<sup>4</sup>.  $EM$  and  $kT_{e,EM}$  are compared between two sets of  $\mu$ 's and  $g$ 's, labelled set A and set B, using the equations above: 137  
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$$\begin{aligned} EM_{FS,B} \times \mu_{e,FS,B} \mu_{H,FS,B} &= EM_{FS,A} \times \mu_{e,FS,A} \mu_{H,FS,A} \\ EM_{RS,B} \times \mu_{e,RS,B} \mu_{H,RS,B} &= EM_{RS,A} \times \mu_{e,RS,A} \mu_{H,RS,A} \\ EM_{FS,XS,B} \times \mu_{e,FS,B} \mu_{I,FS,B} &= EM_{FS,XS,A} \times \mu_{e,FS,A} \mu_{I,FS,A} \\ EM_{RS,XS,B} \times \mu_{e,RS,B} \mu_{I,RS,B} &= EM_{RS,XS,A} \times \mu_{e,RS,A} \mu_{I,RS,A} \\ kT_{e,EM,FS,B} / f_{T,FS,B} &= kT_{e,EM,FS,A} / f_{T,FS,A} \\ kT_{e,EM,RS,B} / f_{T,RS,B} &= kT_{e,EM,RS,A} / f_{T,RS,A} \end{aligned} \quad (15)$$

The standard and XSPEC values of  $EM$  are related by: 140

$$\begin{aligned} EM_{FS,XS} \times \mu_{I,FS} / x_{H,C} &= EM_{FS} \times \mu_{H,FS} \\ EM_{RS,XS} \times \mu_{I,RS} / x_{H,C} &= EM_{RS} \times \mu_{H,RS} \end{aligned} \quad (16)$$

For the case that  $g_{RS,A}$  and  $g_{RS,B}$  are the same, 141

$$kT_{e,EM,RS,B} \left( \frac{1}{\mu_{e,RS,B}} + \frac{g_{RS}}{\mu_{I,RS,B}} \right) = kT_{e,EM,RS,A} \left( \frac{1}{\mu_{e,RS,A}} + \frac{g_{RS}}{\mu_{I,RS,A}} \right) \quad (17)$$

For older SNRs, one has  $g_{RS,A} = g_{RS,B} = 1$ , which gives: 142

$$kT_{e,EM,RS,B} / \mu_{RS,B} = kT_{e,EM,RS,A} / \mu_{RS,A} \quad (18)$$

For young SNRs (age less than a few hundred years)  $g_{2,A} \ll 1$  and  $g_{2,B} \ll 1$ , one has: 143

$$kT_{e,EM,RS,B} / \mu_{e,RS,B} = kT_{e,EM,RS,A} / \mu_{e,RS,A} \quad (19)$$

### 3.3. Chemical Composition and Partial Ionization Examples 144

Table 1 lists  $\mu$ 's for different ISM/FS and ejecta/RS compositions and different ionization levels. The  $\mu$ 's for FS (first two rows) have very little variation ( $\sim 1\%$ ) with composition. Thus  $kT_{e,EM,FS}$ ,  $EM_{FS}$  and  $EM_{FS,XS}$  show very little variation with composition, less than typical errors in measured  $EM$  and  $kT_e$ . For solar composition  $EM_{FS}$  and  $EM_{FS,XS}$  are the same:  $\mu_{I,FS} / x_{H,C} = \mu_{H,FS} = 1.356$ . 145  
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<sup>4</sup> Otherwise, the more complicated full integral expressions must be used

**Table 1.** Summary of Mean Molecular Weights.

Composition	$\mu_H$	$\mu_e^1$	$\mu_e^2$	$\mu_e^{c3}$	$\mu_I$	$\mu^{a1}$	$\mu^{b2}$	$\mu^{c3}$
Solar <sup>4</sup>	1.356	1.151	2.303	1.250	1.250	0.599	0.810	0.625
SMC <sup>5</sup>	1.340	1.145	2.290	1.236	1.236	0.594	0.803	0.618
CC-type ejecta <sup>5</sup>	1.810	1.289	2.578	1.542	1.542	0.702	0.965	0.771
Type Ia ejecta <sup>6</sup>	1327	2.093	4.187	35.57	35.57	1.977	3.746	17.78
Mixture 1 CC-Ia <sup>7</sup>	12.24	1.894	3.789	7.789	7.789	1.524	2.549	3.894
Mixture 2 CC-Ia <sup>8</sup>	3.615	1.596	3.191	2.956	2.956	1.036	1.535	1.478
Mixture ISM-CC <sup>9</sup>	1.551	1.216	2.433	1.381	1.381	0.647	0.881	0.690
Mixture ISM-Ia <sup>10</sup>	2.710	1.486	2.971	2.415	2.415	0.920	1.332	1.207
Pure oxygen	$\infty$	2.000	4.000	16.00	16.00	1.778	3.200	8.000
Pure iron	$\infty$	2.154	4.308	56.00	56.00	2.074	4.000	28.00

<sup>1</sup> Fully ionized plasma. <sup>2</sup> Each element 50% ionized. <sup>3</sup> Each element singly ionized. <sup>4</sup> Abundances from [22].

<sup>5</sup> Abundances from [25]. <sup>6</sup> Abundances from [14]. <sup>7</sup> Geometric mean of CC-type and Type Ia abundances. <sup>8</sup> Equal mass mixture of CC-type and Type Ia abundances. <sup>9</sup> Equal mass mixture of Solar and CC-type abundances. <sup>10</sup> Equal mass mixture of Solar and Type Ia abundances.

For RS/ejecta gas, the  $\mu$ 's can vary widely as shown in rows 3-10 of Table 1. The standard to XSPEC  $EM$  ratio  $\frac{EM_{RS}}{EM_{RS,XS}}$  varies from 0 (cases of no H in the ejecta), to 0.029 (for the adopted Type Ia abundances), to values near 1 for cases with significant H abundance (including the adopted CC-type and ISM-CC mixture). As noted earlier the XSPEC definition is superior. In particular, when the H abundance is 0 the standard  $EM$  is 0 and completely fails to represent the emission from the gas.

The effect of changing  $\mu$ 's on  $EM_{XS}$  of RS gas is shown by the ratios of  $\frac{EM_{RS,XS,B}}{EM_{RS,XS,A}}$ , with A= solar abundances. Table 2 gives this for two cases of ionization (last two columns): fully ionized plasma and singly ionized plasma. The composition for the fully ionized case results in a ratio ranging from 0.012 for pure Fe (smaller  $EM$  for pure Fe) to 1.017 (SMC abundances with more H than solar). For the singly ionized case, the ratio ranges from  $\sim 5 \times 10^{-4}$  to 1.023 (SMC abundances). To obtain  $\frac{EM_{RS,XS,B}}{EM_{RS,XS,A}}$  for A different from solar, one divides the ratio of B to solar to that for A to solar. E.g.,  $\frac{EM_{RS,XS,B=Ia}}{EM_{RS,XS,A=CC}}$ , fully ionized, is 0.0267 and  $\frac{EM_{RS,XS,B=PureFe}}{EM_{RS,XS,A=Ia}}$ , fully ionized, is 0.617.

**Table 2.** Emission Measure Ratios.

Composition <sup>1</sup>	$\frac{EM_{RS}}{EM_{RS,XS}}$	$\frac{EM_{RS,XS,B}}{EM_{RS,XS,\odot}}$ 2	$\frac{EM_{RS,XS,B}}{EM_{RS,XS,\odot}}$ 3
Solar	1	1	1
SMC	1.001	1.017	1.023
CC-type ejecta	0.925	0.724	0.657
Type Ia ejecta	0.0291	0.0193	$1.24 \times 10^{-3}$
Mixture 1 CC-Ia	0.691	0.0975	0.0258
Mixture 2 CC-Ia	0.888	0.305	0.179
Mixture ISM-CC	0.967	0.857	0.820
Mixture ISM-Ia	0.968	0.401	0.268
Pure oxygen	0	0.0450	$6.10 \times 10^{-3}$
Pure iron	0	0.0119	$4.98 \times 10^{-4}$

<sup>1</sup> See Table 1 for composition notes. <sup>2</sup> Case  $\odot$  is solar, Case B is from Composition row. Fully ionized and 50% ionized give the same ratio. <sup>3</sup> Case  $\odot$  is solar, Case B is from Composition row. Each element 50% ionized.

The H and heavy element abundances in Type Ia or CC-type could be different than assumed. As test cases,  $\mu$ 's are shown for two mixtures of CC-type and Ia-type ejecta: a geometric mean, and an equal mass mixture. The equal mass mixture has more H, resulting in significantly lower  $\mu$ 's.

The effect of partial ionization is illustrated in Table 1. Only  $\mu_e$  and  $\mu$  are affected. The ratio of  $\mu_e$  for singly ionized plasma to  $\mu_e$  for fully ionized ranges from 1.08 (SMC) to 26.0 (pure Fe). 50% ionized plasma has  $\mu_e$  larger than for fully ionized by a factor of 2, whereas  $\mu_e$  can be higher (cases with high H) or lower (cases with low H) than  $\mu_e$  for singly ionized.

The effect of ionization on  $\mu$  is smaller than for  $\mu_e$ , with ratio of  $\mu$ (singly ionized) to  $\mu$ (fully ionized) ranging from 1.04 (SMC) to 13.5 (pure Fe). 50% ionized plasma has  $\mu$  larger than for fully ionized by a factor of 1.35 (SMC) to 1.93 (pure Fe), whereas  $\mu$  can be higher (cases with high H) or lower (cases with low H) than  $\mu$  for singly ionized.

$kT_e$  depends on the  $\mu$ 's, as follows. For older SNRs ( $g = 1$ ) Equation 18 shows that RS temperatures for fully ionized gas can be larger than for solar composition up to a factor of 3.46 (pure Fe) or smaller by a factor of 0.99 (SMC abundances). In the extreme case of singly ionized gas,  $T_e$  can be larger than for solar composition up to a factor of 44.8 (pure Fe) or smaller by a factor of 0.99 (SMC abundances). For very young SNRs ( $g \ll 1$ ) Equation 19 applies, yielding smaller changes than for the fully ionized case:  $kT_{e,EM,RS,B}/kT_{e,EM,RS,\odot}$  varies from 0.995 (for SMC) to 1.87 (pure Fe). For singly ionized gas,  $kT_{e,EM,RS,B}/kT_{e,EM,RS,\odot}$  has the same range as for SNRs with  $g = 1$ : 0.995 to 44.8. Shock temperatures in SNRs are generally high enough that the gas is  $\sim 50\%$  or more ionized, so the ratios are less extreme than for the singly ionized case. In general, more heavy elements in the ejecta make the RS temperature higher.

### 3.4. Example application to a SNR with reverse shock measured

**Table 3.** SNR J0049-7314 CC and Ia reverse shock models<sup>1</sup>.

s,n	Composition <sup>2</sup>	$kT_{e,RS}(+,-)$ (keV)	$EM_{RS,XS}(+,-)$ ( $10^{56} \text{ cm}^{-3}$ )
Observed Models	n/a	0.91(+0.03,-0.03)	728(+125,-99)
0,7	CC	3.6(+1.1,-1.0)	0.14(+0.03,-0.03)
0,7	Ia	10.1(+3.1,-2.8)	0.0037(+0.0007,- 0.0007)
0,10	CC	2.2(+0.6,-0.6)	0.24(+0.04,-0.04)
0,10	Ia	6.2(+1.7,-1.7)	0.0065(+0.0010,- 0.0010)
2,7	CC	0.55(+0.16,-0.15)	130(+149,-68)
2,7	Ia	1.55(+0.45,-0.42)	3.5(+4.0,-1.8)
2,10	CC	0.11(+0.03,-0.03)	1660(+1900,-890)
2,10	Ia	0.31(+0.08,-0.08)	44(+51,-24)

<sup>1</sup> These are calculated by fitting the forward shock observed  $kT_{e,EM,FS}$  and  $EM_{FS}$ . <sup>2</sup> CC and Ia  $\mu$ 's are from Table 1.

As described above, the difference in standard and XSPEC values of  $kT_e$  and  $EM$  are largest for the RS and for cases with low H abundance in the ejecta. We illustrate the difference for an observed SNR using one with RS measured from the Small Magellanic Cloud: SNR J0049-7314 and for two cases of composition: CC and Ia. The measured  $kT_e$ 's and  $EM$ 's are from [26] and were compared to SNR models by [6]. We take the CC and Ia compositions adopted by [15].

The measured  $kT_{e,FS}$  and  $EM_{FS}$  for SNR J0049-7314 in [6] assuming CC composition and standard definitions of  $kT_e$  and  $EM$ . However the observed values were derived using XSPEC definitions. Our models assume uniform composition ISM and uniform composition ejecta, so that the standard and XSPEC values of  $kT_{e,FS}$  are the same (both use SMC composition). However  $kT_{e,RS}$  depends on composition as specified by Equation 17.

The age of SNR J0049-7314 is  $\sim$ 18 kyr if in a uniform ISM, or  $\sim$ 2-6 kyr if in a stellar wind. Thus it is old enough that we take the electron-ion temperature ratio close to 1. Then one obtains  $kT_{e,RS,Ia} = 2.82 \times kT_{e,RS,CC}$ . For  $EM_{RS}$ , we first obtain the relation between the normal and XSPEC RS values using Equation 16, which yields  $EM_{RS,XS,CC} = 1.27 \times EM_{RS,CC}$ . The relation between XSPEC values for CC and for Ia is given by the fourth line of Equation 15, which yields  $EM_{RS,XS,Ia} = 2.67 \times 10^{-2} EM_{RS,XS,CC}$ . The resulting model values of  $kT_{e,RS}$  and  $EM_{RS,XS}$  for the four cases  $s=0$  and 2 and  $n=7$  and 10 are given in Table 3.

Now we can assess the success of the various models.  $EM_{RS,XS}$  is most sensitive to  $s$ ,  $n$  and composition, so that is considered first. All Ia composition models too small  $EM_{RS,XS}$ , implying this is a CC SNR. All  $s=0$  models predict too small  $EM_{RS,XS}$ . The observed  $EM_{RS,XS}$  is between model values for CC composition for  $(s,n)=(2,7)$  and  $(2,10)$ . However the model  $kT_{e,RS}$  is below the observed one: by factor 0.60 for  $(s,n)=(2,7)$ . Because  $kT_{e,RS}$  and  $EM_{RS,XS}$  depend in different ways on the  $\mu$ 's, it is likely that adjustment of the composition could bring the model values into agreement with the observed ones within the uncertainties.

#### 4. Summary and Conclusion

The X-ray emission from a supernova remnant (SNR) is a powerful diagnostic of the state of the shocked plasma. Observed properties are shock radius, electron temperature ( $kT_e$ ) and emission measure ( $EM$ ) for forward-shocked ( $FS$ ) and in some cases for reverse-shocked ( $RS$ ) gas. Given a model, observations of  $FS$  gas can be used to determine the energy of the explosion, the age of the SNR, and the density of the surrounding medium. If  $RS$  gas emission is also detected, more properties of the SNR can be deduced [7] such as whether the SNR exploded in a uniform or stellar wind environment.

To calculate  $EM_{FS}$ ,  $EM_{RS}$ ,  $T_{e,FS}$  and  $T_{e,RS}$  from models (hydrodynamic solutions with variables  $\rho$ ,  $P$  and  $V$ ) to compare with observations, one requires the values of molecular weights ( $\mu_{FS}$ ,  $\mu_{e,FS}$ ,  $\mu_{I,FS}$ ,  $\mu_{H,FS}$ ) and electron-to-ion temperature ratio ( $g_{FS} = T_{e,FS} / T_{I,FS}$ ), with similar quantities for the  $RS$ . Here we investigated the effects of composition, ionization and  $g$  on model-derived  $kT_e$  and  $EM$  (XSPEC and standard definitions) and thus on derived SNR properties.

The  $\mu$ 's depend on the composition and ionization state of the shocked gas. For compositions with large H abundance, as expected for  $FS$  gas, the  $\mu$ 's don't depend strongly on composition and ionization. However, the expected H-poor compositions of  $RS$  gas give large variations in the  $\mu$ 's and the  $EM$  ratios.

The standard and XSPEC definitions of  $EM$  have an important difference. The XSPEC definition is superior for SNRs, which can have shocked gas with low hydrogen abundance. We presented formulae to calculate standard and XSPEC versions of  $EM$  and  $kT_e$ , including dependences on the  $\mu$ 's and  $g$ 's.

The formulae simplify in the case of spatially uniform  $\mu$ 's and  $g$ 's. In this case, the XSPEC and standard definitions for  $kT_e$  are the same and those for  $EM$  are different. These  $\mu$ 's and ratios of standard  $EM$  to XSPEC  $EM_{XS}$  are summarized in Table 1 for several different cases of composition and ionization. Generally for H-rich composition the ratio  $EM/EM_{XS} \simeq 1$ , and for H-poor compositions  $EM/EM_{XS} \ll 1$ .

The effects of electron-to-ion temperature ratio  $g$  on  $kT_e$  were shown to be composition dependent. For fully ionized gas and  $g = 1$ ,  $kT_{e,RS} / kT_{e,RS,\odot}$  varies from  $\simeq 1$  (for H-rich compositions) to 3.46 (pure Fe), and for the extreme case of singly ionized gas  $kT_{e,RS} / kT_{e,RS,\odot}$  varies from  $\simeq 1$  to 44.8. For young shocked gas with  $g = 1$ ,  $kT_{e,RS} / kT_{e,RS,\odot}$  varies much less than for  $g = 1$ , from  $\simeq 1$  to 1.87 for fully ionized gas, but the same range ( $\simeq 1$  to 44.8) as for  $g = 1$  for singly ionized gas.

Distances to Galactic SNRs have improved significantly over the past decade, allowing determination of radii (e.g. [25]). X-ray observations of SNRs have been carried out for a significant fraction of Galactic SNRs. We have improved the accuracy of spherically symmetric SNR evolution models by including results from a large grid of hydrodynamic simulations [14] to develop the the SNR modelling code SNRpy. Together, the new data

and the SNRpy code enable the application of SNR models to use radii, emission measures and temperatures to derive SNR properties (e.g. [7]). In previous work with SNRpy, we used the standard definition of  $EM$  and  $kT$ . Inclusion of XSPEC definitions will make the output of the code more realistic, particularly for gas with low H abundance, as described in this work. The process of including the XSPEC definitions in SNRpy is now underway, and a new version is planned for release in mid-2022 on GitHub and on the website <http://quarknova.ca>.

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