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The effects of variations in nuclear interactions on nucleosynthesis in thermonuclear supernovae

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ABSTRACT

Context. Type Ia supernova explosions are violent stellar events important for their contribution to the cosmic abundance of iron peak elements and for their role as cosmological distance indicators.

Aims. The impact of nuclear physics uncertainties on nucleosynthesis in thermonuclear supernovae has not been fully explored using comprehensive and systematic studies with multiple models. To better constrain predictions of yields from these phenomena, we investigate thermonuclear reaction rates and weak interaction rates that significantly affect yields in our underlying models.

Methods. We have performed a sensitivity study by postprocessing thermodynamic histories from two different hydrodynamic, Chandrasekhar-mass explosion models. We have individually varied all input reaction and, for the first time, weak interaction rates by a factor of ten (up and down) and compared the yields in each case to yields using standard rates.

Results. Of the 2305 nuclear reactions in our network, we find that in either model the rates of only 53 reactions affect the yield of any species with an abundance of at least $10^{-8} M_{\odot}$ by at least a factor of two. The rates of the ${}^{12}C(\alpha, \gamma)$, ${}^{12}C+{}^{12}C$, ${}^{20}Ne(\alpha, p)$, ${}^{20}Ne(\alpha, \gamma)$, and ${}^{30}Si(p, \gamma)$ reactions are among those that modify the most yields when varied by a factor of ten. From the individual variation of 658 weak interaction rates in our network by a factor of ten, only the stellar ${}^{28}Si(\beta^+){}^{28}Al$, ${}^{32}S(\beta^+){}^{32}P$, and ${}^{36}Ar(\beta^+){}^{36}Cl$ rates significantly affect the yields of species in a model. Additional tests reveal that reaction rate changes over temperatures T > 1.5 GK have the greatest impact and that ratios of radionuclides that may be used as explosion diagnostics change by a factor of ≤ 2 from the variation of individual rates by a factor of ten.

Conclusions. Nucleosynthesis in the two adopted models is relatively robust to variations in individual nuclear reaction and weak interaction rates. Laboratory measurements of a limited number of reactions would, however, help to further constrain model predictions. In addition, we confirm the need for a detailed, consistent treatment of all relevant stellar weak interaction rates since simultaneous variation of these rates (as opposed to individual variation) has a significant effect on yields in our models.

Key words. supernovae: general - nuclear reactions, nucleosynthesis, abundances

1. Introduction

Type Ia supernovae (hereafter SNe Ia) have become valuable cosmological tools. Through calibrated light curve analysis, they have been used as probes to outline the geometrical structure of the Universe, unraveling its unexpected acceleration stage (Riess et al. 1998; Schmidt et al. 1998; Perlmutter et al. 1999; for constraints on dark energy and on the cosmic expansion history, see recent work by Astier et al. 2011; Blake et al. 2011; Suzuki et al. 2012, and references therein).

SNe Ia are spectroscopically classified by the absence of hydrogen (Balmer) emission lines and the presence of a prominent P-Cygni absorption feature near 6150 Å due to blueshifted Si II (Wheeler & Harkness 1990; Filippenko 1997)¹. Whereas the first observational constraint places limits on the maximum amount of hydrogen that can be present in the expanding atmosphere of the star (i.e., $M_{\rm H} \le 0.03-0.1~M_{\odot}$), the second feature suggests that the outermost ejected shells contain intermediatemass elements from nuclear processing.

The increasing number of supernovae discovered has revealed some diversity among SNe Ia, raising doubts on the historically postulated uniqueness of the progenitor system. Already two decades ago estimates had indicated that only about 85% of the observed SNe Ia belong to a homogeneous class of events (Branch et al. 1993), with a dispersion of only $\Delta M \leq$ 0.3 mag when normalized to peak luminosity (see Cadonau et al. 1985; Hamuy et al. 1996). These SNe Ia are known as "branch normals", with canonical examples such as SNe 1972E, 1981B, 1989B, or 1994D. A more recent classification of SNe Ia (Li et al. 2011, in a volume-limited sample) found that the number of SNe Ia deviating from this homogeneous class of objects is closer to ~30% (see also Li et al. 2000; Branch 2004; Kasen et al. 2009). Within the minority group, likely progenitor systems and explosion models have been identified recently for two subclasses. For SN 2002cx-like supernovae, observable features

¹ Some SNe Ia are, however, anomalous in this regard. SN 2002ic (Hamuy et al. 2003), for instance, while exhibiting other standard features common to all SNe Ia, unequivocally showed broad-line H_{α} emission. This has been interpreted as proof of a SN Ia interacting with H-rich circumstellar material. Other SN 2002ic-like events include SN 2005gj, PTF 11kx (Dilday et al. 2012), and SN 2008J (Taddia et al. 2012).

are well explained by weak deflagrations in near-Chandrasekharmass carbon–oxygen (CO) white dwarf stars (WDs) which leave bound remnants (Jordan et al. 2012; Kromer et al. 2013); for SN 1991bg-like supernovae, the peculiar spectra, colors, and low expansion velocities of this subluminous class are reproduced by mergers of two, relatively light CO WDs of nearly equal mass (Pakmor et al. 2010).

The most promising progenitor scenarios that have been proposed for spectroscopically "normal" SNe Ia (e.g., Livio 2000; Hillebrandt & Niemeyer 2000; Isern et al. 2011; Howell 2011; Hillebrandt et al. 2013) include: the single degenerate scenario, consisting of a nondegenerate companion star that transfers hydrogen-rich (or, possibly, helium-rich) matter onto a CO WD (Whelan & Iben 1973); the double degenerate scenario, consisting of two merging CO WDs, such that the total mass exceeds the Chandrasekhar limit (Iben & Tututkov 1984; Webbink 1984); and a scenario where a sub-Chandrasekhar-mass CO WD accretes stably from a companion star and explodes before reaching the Chandrasekhar limit (Taam 1980; Iben et al. 1987). Other scenarios may indeed be possible. In spite of the large number of published SN Ia explosion models for proposed scenarios, detailed predictions of the associated nucleosynthesis have been calculated and published only for a very limited subset of near-Chandrasekhar-mass WD models in the single degenerate scenario. To the best of our knowledge, 1D model yields exist in the literature only for variants of pure deflagrations (Thielemann et al. 1986; Nomoto et al. 1997; Woosley 1997; Iwamoto et al. 1999; Maeda et al. 2010); 2D model yields have been published for gravitationally confined detonation (Meakin et al. 2009) and delayed-detonation models (Maeda et al. 2010); and detailed tables of yields from 3D models are available only for pure deflagrations (Travaglio et al. 2004; Röpke et al. 2006) and a suite of 14 delayed-detonation models (Seitenzahl et al. 2013).

These models have revealed that the nucleosynthesis in SNe Ia depends critically on the peak temperature achieved and the density at which the thermonuclear runaway occurs. In addition, the specific composition of the WD (in particular, the amount and distribution of ¹²C and ²²Ne) plays a central role (see, e.g., Chamulak et al. 2007, 2008; Townsley et al. 2009, and references therein) as it influences properties such as the ignition density, the release of energy, the flame speed, and the specific density at which the initial deflagration may transform into a detonation. In general, the abundance pattern of the ejecta is the result of five burning regimes: "normal" and " α -rich" freeze-out from nuclear statistical equilibrium (NSE) in the inner regions, and incomplete Si-, O-, and C/Ne-burning in the outermost layers (Thielemann et al. 1986; Woosley 1986). SNe Ia are galactic factories of ⁵⁶Fe, producing about half of the iron content in the Milky Way (Acharova et al. 2012) and perhaps 65-70% of the iron content in the solar neighborhood (Mennekens et al. 2012). Hence, reproducing the chemical abundance pattern around the Fe-peak is a critical test in SN Ia modeling. For decades, models systematically overproduced neutron-rich species such as ⁵⁴Cr or ⁵⁰Ti with respect to solar system values (Woosley 1990; Thielemann et al. 1997; Nomoto et al. 1997). The agreement improved following the revision of key stellar weak interaction rates (Langanke & Martinez-Pinedo 2000) and through the use of more recent 3D explosion models (Seitenzahl et al. 2013). These results emphasize the critical roles played by both the nuclear physics input and the modeling techniques employed.

Nonetheless, in spite of their demonstrated importance, the impact of nuclear physics uncertainties on the nucleosynthesis

produced in SNe Ia has not been analyzed at the same level of detail as for other astrophysical scenarios, such as classical nova explosions or type I X-ray bursts. Most efforts have focused on determining the role of the ¹²C+¹²C reaction (e.g., Spillane et al. 2007; Bravo et al. 2011) since it triggers the explosion when the temperature exceeds \approx 700 MK. Progress has been reported in very recent work by Bravo & Martínez-Pinedo (2012), in which thermonuclear reaction rates of importance were investigated using a single 1D delayed-detonation (or, deflagration-to-detonation transition, DDT) model of a Chandrasekhar-mass WD.

The goal of this paper is to investigate the sensitivity of nucleosynthesis in thermonuclear supernovae to variations of both the nuclear reaction and weak interaction rates involved during the explosion. Since all currently available yield predictions are for near-Chandrasekhar-mass models, we have chosen for our study two representative cases in this class: the W7 model (Nomoto et al. 1984), which has been serving as the fiducial SN Ia explosion model in the community for almost three decades, and a standard 2D DDT model. Since our method involves recalculating the nucleosynthesis in each model every time a reaction or weak interaction rate is varied (resulting in over several million individual postprocessing calculations), performing our study for a large suite of explosion models is not yet feasible.

In the following, we begin by briefly describing the two underlying models employed for our studies. Next, for each of our two models, we present and compare results obtained from individually varying each of the rates in our nucleosynthesis network to assess its impact on SNe Ia yields. Finally, we discuss some additional tests we performed to motivate new experimental measurements and to link our results to SN Ia observables.

2. Explosion models

For our sensitivity studies we chose to postprocess the results from two SN Ia explosion models: a generic 2D Chandrasekharmass DDT model (similar to models from Kasen et al. 2009), and the fiducial W7 pure deflagration model (Nomoto et al. 1984). The ⁵⁶Ni mass ejected is 0.66 M_{\odot} in the W7 model and 0.68 M_{\odot} in the DDT model; these values are rather typical for normal SN Ia (see, e.g. Stritzinger et al. 2006). The details of the models are not the focus here; they merely serve to define a set of reasonable thermodynamic conditions that arise in thermonuclear supernovae for the purpose of examining the sensitivity of SN Ia yields to variations of the input reaction and weak interaction rates.

2.1. Delayed-detonation model

The DDT model employed was a 2D axisymmetric hydrodynamic explosion of a 1.40 M_{\odot} cold WD with a central density of 2.9 × 10⁹ g cm⁻³. The initial chemical composition was 47.5% ¹²C, 50% ¹⁶O, and 2.5% ²²Ne homogeneously distributed throughout the WD. The ²²Ne content parametrizes the neutron excess and corresponds to an electron fraction of $Y_e = 0.49886$. The simulation was performed with the LEAFS code, which integrates the discretized reactive Euler equations with a finite volume method. The hydrodynamics solver is essentially the PROMETHEUS implementation (Fryxell et al. 1989) of the "piecewise parabolic method" by Colella & Woodward (1984).

Subsonic deflagration flames and supersonic detonation fronts were modeled as level set discontinuities between nuclear



Fig. 1. Tracer particle positions for the adopted 2D DDT model at t = 100 s from first ignition of the deflagration. The tracer particles are colored (online) according to the mass fraction of ⁵⁶Ni present.

fuel and nuclear ash (Osher & Sethian 1988; Smiljanovski et al. 1997; Reinecke et al. 1999). Any material traversed by these fronts was burned to a composition that depends on fuel density and the mode of burning (deflagration or detonation). The corresponding energy was then immediately released behind the front. For detonations, we used the energy release data from the tables of Fink et al. (2010). The speed of the detonations was modeled as in Fink et al. (2010): at high densities ($\rho > 10^7 \text{ g cm}^{-3}$), speeds were taken from Gamezo et al. (1999); at low densities, Chapman-Jouguet-like speeds were calculated for the incomplete burning yields in our detonation tables. For deflagrations, we used the same energy release table that was employed by Seitenzahl et al. (2011). Since in two dimensions our usual subgrid-turbulence-based approach to model the deflagrationto-detonation transition probabilities is not applicable, we modeled the transition using the method of Kasen et al. (2009). For the critical Karlovitz number we chose $Ka_{crit} = 250$, and we limited the density at which a DDT may occur to the interval $0.6 < \rho/10^7 \text{ g cm}^{-3} < 1.2.$

The simulation method is similar to that employed in Kasen et al. (2009), except that here we have assumed reflectional symmetry across the equator as well. This is to facilitate analysis of the nucleosynthetic yields of the model using the tracer particle method (Travaglio et al. 2004), in which the temperaturedensity-time profile of each tracer particle is recorded and later used within postprocessing calculations. With the above assumption we attain twice the spatial sampling of the explosion ejecta with the same number of tracer particles. We used the variable mass tracer particle method of Seitenzahl et al. (2010), which allows us to better resolve the lower density regions of incomplete burning without greatly increasing the total number of tracer particles. The resolution was 512×512 computational cells, and 1010 tracer particles were distributed to sample the mass distribution of the star. This approach adequately samples the different nucleosynthesis regimes in a multipoint ignition DDT model and provides yields that are accurate to a few percent or better for the more abundant nuclides (Seitenzahl et al. 2010). To illustrate, Fig. 1 shows tracer particle positions for this model at t = 100 s from the first ignition of the deflagration.

2.2. W7 model

The W7 model of Nomoto et al. (1984) is commonly used as a reference when general features of SN Ia are discussed. W7 is a 1D, fast deflagration explosion model of a 1.38 M_{\odot} WD, consisting of 50% $^{\rm 16}{\rm O},$ 47.5% $^{\rm 12}{\rm C},$ and 2.5% $^{\rm 22}{\rm Ne}$ by mass, homogeneously distributed. To model the acceleration of the convectively driven deflagration wave in their hydrodynamics code, Nomoto et al. (1984) used the time-dependent mixing length theory of Unno (1967). They chose $\alpha = 0.7$ for the mixing length $l = \alpha H_p$, where H_p is the pressure scale height. In their hydrodynamical simulation, only an α -chain network was included to model the nuclear energy release. Later, Thielemann et al. (1986) calculated the detailed nucleosynthesis for the W7 model by postprocessing the thermodynamic histories of the 172 zones with a reaction network comprising 259 nuclear species. The strong overproduction of neutron-rich Fe-group nuclei initially noted from this model was later attributed to the relatively large Fe-group electron capture rates of Fuller et al. (1982) that had been used (see Iwamoto et al. 1999). Use of newer, reduced Fe-group electron capture rates (Langanke & Martínez-Pinedo 2000) improved the agreement between W7 model yields and solar system abundances for these nuclei (Brachwitz et al. 2000; Maeda et al. 2010; but see also Seitenzahl et al. 2013).

3. Sensitivity of nucleosynthesis to rate variations

3.1. Variation of all rates by a factor of ten

We postprocessed the thermodynamic histories from the DDT and W7 thermonuclear supernova models described above. This involved coupling an extended nuclear physics network to



Fig. 2. (Color online) Yields in M_{\odot} vs. mass number A, as calculated for the DDT (filled circle) and W7 (empty square) models. These yields were obtained using standard rates in our nuclear network.

the temperature-density-time profiles extracted from these two models. Final yields were determined by summing the (massweighted) contributions from either all zones (for the W7 model) or all tracer particles (for the DDT model). The nuclear physics network consisted of 443 species ranging from n to ⁸⁶Kr (see Travaglio et al. 2004; Seitenzahl et al. 2009a). Nuclear reactions such as (p, γ) , (α, γ) , (n, γ) , (p, n), (α, n) , (α, p) were included, as well as reactions such as ${}^{12}C+{}^{12}C$, ${}^{12}C+{}^{16}O$, and ${}^{16}O+{}^{16}O$ (along with all corresponding reverse processes). Sufficient experimental information is available to determine rates for only a limited number of these reactions; these rates have been adopted from Iliadis et al. (2010), REACLIB V1.0 (Cyburt et al. 2010), and some recent updates for selected reactions. Theoretical nuclear reaction rates, for the most part determined through Hauser-Feshbach models (e.g., Rauscher & Thielemann 2000; Arnould & Goriely 2003), were adopted when experimentally based rates were not available. For weak interactions, we used stellar (temperature and density dependent) rates from the largescale shell model calculations of Oda et al. (1994) and Langanke & Martínez-Pinedo (2000), supplemented with additional stellar rates from Pruet & Fuller (2003) and Fuller et al. (1982). Together, these sources provided stellar weak interaction rates for most species in our network with $A \ge 17$. NSE was assumed above T = 5 GK. The initial composition of the WD for both models was 47.5% ¹²C, 50% ¹⁶O, and 2.5% ²²Ne by mass. Yields from the DDT and W7 models using our standard nuclear physics network are plotted in Fig. 2. For each of the two models, we then varied each individual rate in our standard network (together with the reverse process for reaction rates) by a factor of ten (up and down), repeated the postprocessing calculations for all thermodynamic histories, and compared the resulting yields with the yields shown in Fig. 2.

For reaction rate variations, the results are summarized in Table 1. Stable and radioactive species are listed since, for each model, abundances were determined one hour after the beginning of the explosion. To emphasize the rate variations that most significantly affected the yields in each model, we restrict the discussion here to species that achieved an abundance of at least $10^{-8}~M_{\odot}$ and deviated from the abundances determined with standard rates (i.e., Fig. 2) by at least a factor of two (unless otherwise indicated). We realize that for some applications the variation of yields of particular species by less than a factor of two may be of interest. As such, full results on the specific effect of varying any rate in our network in either model are available upon request (but see also Sect. 3.3). Table 2 highlights important reactions from Table 1 by listing only those reaction rates that affected (i) the yields of at least three species (again, all with abundances greater than $10^{-8} M_{\odot}$) by at least a factor of two in either the DDT or W7 models; and/or (ii) affected the yield of at least one species in both models by at least a factor of two. To facilitate interpretation, we also present the results of Table 1 in Figs. 3-5. For both the DDT and W7 models, Fig. 3 shows the heavy product against the heavy reactant for all reactions in Table 1. Figures 4 and 5 show reaction rates from Table 1 whose variation affects the yields of at least three species in the DDT or W7 models, respectively. For each reaction, these plots show a ratio: the yield of each affected species when the rate was enhanced by a factor of ten divided by the yield of that species when the rate was reduced by a factor of ten.

From the individual variation of nuclear reaction rates by a factor of ten, only 53 of the 2305 reactions in our network affect yields of any species with an abundance of at least $10^{-8} M_{\odot}$ by at least a factor of two in either model. (We note that for reactions, forward and reverse processes are not counted separately here since they must always be varied together.) Of these, all but nine reactions involve exclusively species with A < 40. There are no reactions listed in Table 1 involving nuclei with Z > 24. As can be seen from Table 2 (case C) and Fig. 3, 24 of these 53 reaction rates affect yields in both models. The overall impact of these rate variations is limited, however. As seen in Table 2 (cases A and B), only 14 reactions affect the yields of three or more species in either model. Of these, the rates of the ${}^{12}C(\alpha,\gamma), {}^{12}C({}^{12}C,\alpha), {}^{12}C({}^{12}C,p), {}^{20}Ne(\alpha,p), {}^{20}Ne(\alpha,\gamma), and$ 30 Si (p, γ) reactions have the greatest impact, affecting the yields of at least five species when varied by a factor of ten; this is also illustrated in Figs. 4 and 5. Finally, only five reactions have an

Table 1. Nuclear reaction rates that significantly affect abundances in the DDT and W7 models when varied by a factor of ten (up or down).

Table 1. continued.

$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Reaction	Nuclide	W7 1	nodel	DDT	model	
$ {}^{12}C((\alpha, \gamma)^{16}O = {}^{16}Ar = 2.2 = 3.0$			×10	×0.1	×10	×0.1	
$ {}^{39} K = 2.2 & 4.6 \\ {}^{40} Ca = 2.2 & 4.6 \\ {}^{47} Ca = 2.4 & 2.4 \\ {}^{47} Ti = 2.5 & 7.1 \\ {}^{45} Sc = 4.3 & 2.5 \\ {}^{46} Ti = 2.5 & 0.17 \\ {}^{54} Mn = 0.21 & 0.17 \\ {}^{54} Mn = 0.21 & 0.21 \\ {}^{52} Cr = 0.17 & 2.6 & 2.3 \\ {}^{21} Ne = 11 & 3.7 & 2^2 N \\ {}^{21} Ne = 11 & 3.7 & 2^2 N \\ {}^{21} Ne = 11 & 3.7 & 2^2 N \\ {}^{21} Na = 2.5 & 3.3 & 2^2 Mg \\ {}^{24} Na = 2.5 & 3.3 & 2^2 Mg \\ {}^{24} Na = 2.5 & 3.3 & 2^2 Mg \\ {}^{24} Na = 2.5 & 3.3 & 2^2 Mg \\ {}^{24} Na = 2.5 & 3.3 & 2^2 Mg \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 N \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 N \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 N \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 N \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 M \\ {}^{24} Na = 2.6 & 0.41 & 0.45 & 2^3 M \\ {}^{26} AI = 2.1 & 2.5 & 2^4 M \\ {}^{26} AI = 2.1 & 2.5 & 2^4 M \\ {}^{26} AI = 2.1 & 2.5 & 2^4 M \\ {}^{26} AI = 2.1 & 2.5 & 2^4 M \\ {}^{26} AI = 2.1 & 2.5 & 2^4 M \\ {}^{26} AI = 2.1 & 2.5 & 2^5 M \\ {}^{16} O(a, \gamma)^{17} O & 1^7 O & 0.31 & 2^5 M \\ {}^{16} O(a, n)^{21} Ne & 2.8 & 2.0 & 2^5 M \\ {}^{16} O(a, n)^{21} Ne & 2.5 & 2^5 M \\ {}^{26} AI = 2.1 & 2.5 & 2^5 M \\ {}^{26} AI = 2.0 & 2.4 & 2^5 M \\ {}^{26} AI = 2.0 & 2^5 M \\ {}^{16} O(a, n)^{21} Ne & 2.5 & 2^5 M \\ {}^{26} AI = 0.35 & 2^5 M \\ {}^{26} AI = 0.16 & 0.32 & 2^5 M \\ {}^{26} AI = 0.16 & 0.32 & 2^5 M \\ {}^{26} AI = 0.16 & 0.32 & 2^5 M \\ {}^{26} AI = 0.16 & 0.32 & 2^5 M \\ {}^{26} AI = 0.16 & 0.32 & 2^5 M \\ {}^{26} AI = 0.16 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.16 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.22 & 0.24 & 4.6 & 0.16 & 2^7 A \\ {}^{26} AI = 0.29 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.29 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.29 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.29 & 0.36 & 0.42 & 2^5 M \\ {}^{26} AI = 0.22 &$	$^{12}\mathrm{C}(\alpha,\gamma)^{16}\mathrm{O}$	³⁶ Ar	2.2		3.0		
$ {}^{40}Ca = 2.2 & 4.6 \\ 4^{4}Ca & 2.4 & 2.4 \\ 4^{4}Ti & 2.4 & 4.5 \\ 4^{4}Ti & 2.5 & 4.3 & 4.5 \\ 4^{4}Ti & 2.5 & 4.3 & 4.5 &$		³⁹ K			2.0		
${}^{12}C({}^{12}C, \alpha)^{20}Ne = \begin{pmatrix} 4^{11}C_{4} & & & & & 2.4 \\ 4^{4T_{11}} & & & 2.2 & 7.1 \\ 4^{45}Se & & & 2.5 \\ 5^{2}Cr & & & 0.17 \\ 5^{4}Mn & & & 0.21 \\ 3^{5}Ne & 5.0 & 2.3 & & 2^{11}N \\ 2^{5}Ne & 11 & & & 3.7 \\ 2^{5}Na & & 2.5 & 3.3 \\ 2^{2}Mg & 2.1 & & & & 2^{2}Ne \\ 2^{3}Ne & 2.1 & & & & 2^{2}Ne \\ 2^{3}Ne & 2.1 & & & & & 2^{2}Ne \\ 2^{3}Ne & 2.1 & & & & & & & & & \\ 2^{3}Ne & 2.1 & & & & & & & & & \\ 2^{3}Ne & 2.6 & 0.41 & 0.45 & & & & & & & \\ 2^{3}Ne & 2.6 & 0.41 & 0.45 & & & & & & & \\ 2^{3}Na & 2.6 & 0.41 & 0.45 & & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & \\ 2^{4}Na & 2.6 & 0.41 & 0.45 & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & \\ 2^{4}Na & 2.6 & 0.41 & 0.45 & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & \\ 2^{4}Na & 2.7 & & 3.1 & & & & & & & \\ 2^{4}Na & 2.0 & & & & & & & & & \\ 2^{4}Na & 2.0 & & & & & & & & & \\ 2^{4}Na & 2.0 & & & & & & & & & & \\ 2^{4}Na & 2.0 & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & & \\ 1^{6}O(n, \gamma)^{17}O & {}^{17}O & 0.31 & & & & & & & & & & & \\ 1^{6}O(n, \gamma)^{20}Ne & {}^{21}Ne & 2.5 & & & & & & & & & & & & \\ 2^{4}Na & 4.0 & & & & & & & & & & & & & & \\ 1^{6}O(n, \gamma)^{21}Ne & {}^{18}O & 8.2 & 0.21 & & & & & & & & & & & & \\ 2^{4}Na & 0.47 & 2.2 & 0.48 & & & & & & & & & & & & & \\ 2^{9}Ne(\alpha, \gamma)^{21}Ne & {}^{21}Ne & 6.2 & 0.24 & 4.6 & 0.16 & & & & & & & & & & & & \\ 2^{9}Ne(\alpha, \gamma)^{21}Ne & {}^{21}Ne & 6.2 & 0.24 & 4.6 & 0.16 & & & & & & & & & & & & & & & & & & &$		⁴⁰ Ca	2.2		4.6		
${}^{12}C({}^{12}C, \alpha)^{20}Ne = \begin{pmatrix} 4^{47}Ti \\ 4^{5}Sc \\ 4^{67}Ti \\ 5^{32}Cr \\ 0.17 \\ 5^{32}Ne \\ 0.21 \\ 5^{25}Ne \\ 5.0 \\ 2.3 \\ 2^{3}Ne \\ 2^$		⁴¹ Ca			2.4		
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		⁴⁴ Ti			2.4		
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		⁴⁵ Ti	3.2		7.1		
${}^{12}C({}^{12}C, \alpha)^{20}Ne = \begin{pmatrix} 46T_1 \\ 5^{32}C_T \\ 0.17 \\ 5^{45}Mn \\ 0.21 \\ 0.42 \\ 0.23 \\ 0^{21}Ne \\ 11 \\ 0^{25}Ne \\ 11 \\ 0^{25}Ne \\ 11 \\ 0^{25}Ng \\ 2.1 \\ 0^{26}Ng \\ 5.2 \\ 0.2 \\ 0^{26}Ng \\ 5.2 \\ 0.2 \\ 0^{26}Ng \\ 4.0 \\ 0.2 \\ 0.4 \\ 0^{26}Al \\ 2.7 \\ 0.4 \\ 0.45 \\ $		⁴⁵ Sc			4.3		
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		⁴⁶ Ti			2.5		
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		⁵² Cr			0.17		
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$		⁵⁴ Mn			0.21		
$ {}^{12}C({}^{12}C, a)^{20}Ne = {}^{17}O = {}^{20}Ne = {}^{5.0} = {}^{2.6} = {}^{21}N = {}^{21}Ni = {}^{22}Ni = $		⁵⁶ Fe			0.42		
${}^{12}C({}^{12}C,p){}^{23}Na$ ${}^{24}Na$ ${}^{24}Na$ ${}^{25}Mg$ ${}^{21}Ne$ ${}^{24}Na$ ${}^{25}Mg$ ${}^{21}Na$ ${}^{24}Na$ ${}^{25}S2$ ${}^{25}Mg$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{20}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{22}Na$ ${}^{26}Mg$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{26}Mg$ ${}^{2.7}$ ${}^{25}Ma$ ${}^{26}Mg$ ${}^{21}Ne$ ${}^{26}Ng$ ${}^{2.7}$ ${}^{26}Mg$ ${}^{21}Ne$ ${}^{26}Ng$ ${}^{2.7}$ ${}^{26}Mg$ ${}^{21}Ne$ ${}^{26}Ng$ ${}^{2.7}$ ${}^{25}Mg$ ${}^{24}Na$ ${}^{40}O$ ${}^{26}Mg$ ${}^{2.7}$ ${}^{25}Mg$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{27}Na$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{27}Al$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{22}Ne(a,p){}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{25}S$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{26}Al$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{26}Al$ ${}^{21}Ne$ ${}^{22}Ne(a,p){}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{22}Ne(a,p){}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{21}Ne$ ${}^{22}Ne(a,p){}^{21}Ne$ ${}^{21}Ne$ ${}^{22}Ne$ ${}^{21}Ne$ ${}^{22}Ne$ ${}^{21}Ne$ ${}^{22}Ne$ ${}^{21}Ne$ ${}^{22}Ne$	${}^{12}C({}^{12}C,\alpha){}^{20}Ne$	¹⁷ O			2.6		
${}^{12}C({}^{12}C,p)^{23}Na = \begin{bmatrix} 2^{11}Ne & 11 & & & & & & & & & & & & & & & & &$		²⁰ Ne	5.0		2.3		^{21}N
${}^{12}C({}^{12}C,p)^{23}Na = \begin{array}{ccccccccccccccccccccccccccccccccccc$		²¹ Ne	11				21 No
${}^{12}C({}^{12}C,p){}^{23}Na = \begin{pmatrix} 2.4 \\ 2^{4}Mg \\ 2.1 \\ 2^{6}Mg \\ 5.2 \\ 1^{8}O \\ 2.1 \\ 2^{9}Ne \\ 3.7 \\ 2^{1}Ne \\ 2^{1}Ne \\ 2^{1}Na \\ 2.6 \\ 0.41 \\ 0.45 \\ 2^{3}Na \\ 2.6 \\ 0.41 \\ 0.45 \\ 2^{3}Na \\ 2.6 \\ 0.41 \\ 0.45 \\ 2^{3}Na \\ 2^{4}Na \\ 2.7 \\ 3.1 \\ 2^{4}Mg \\ 4.0 \\ 2^{6}Al \\ 2.8 \\ 2^{4}Na \\ 2.1 \\ 2^{5}Na \\ 2^{5}Na \\ 2^{5}Na \\ 2^{5}Na \\ 2^{5}Na \\ 2^{5}Na \\ 2.1 \\ 2.9 \\ 2^{5}Na \\ 2.1 \\ 2.9 \\ 2^{5}Na \\ 2.1 \\ 2.9 \\ 2.9 \\ 3^{5}S \\ 3.0 \\ 2^{5}Na \\ 2.9 \\ 3^{5}S \\ 3.0 \\ 3^{5}S$		²³ Na			3.7		^{22}Nc
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁴ Na	2.5		3.3		
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁵ Mg	2.1				
$ \begin{array}{c ccccc} {}^{12}{\rm C}({}^{12}{\rm C},p)^{23}{\rm Na} & \begin{array}{c} {}^{18}{\rm O} & \begin{array}{c} 2.1 \\ {}^{20}{\rm Ne} & \begin{array}{c} 3.7 \\ {}^{21}{\rm Ne} & \begin{array}{c} 6.7 \\ {}^{23}{\rm Na} & \begin{array}{c} 2.6 \\ {}^{26}{\rm Mg} & \begin{array}{c} 4.0 \\ {}^{26}{\rm Al} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 4.0 \\ {}^{26}{\rm Al} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 4.0 \\ {}^{26}{\rm Al} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.7 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.1 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.7 \\ {}^{28}{\rm Ma} & \begin{array}{c} 2.5 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.8 \\ {}^{21}{\rm Ne} & \begin{array}{c} 2.8 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.7 \\ {}^{26}{\rm Mg} & \begin{array}{c} 2.4 \\ {}^{20}{\rm Ne}(\alpha, \eta)^{21}{\rm Ne} & \begin{array}{c} 180 \\ {}^{27}{\rm A} & \begin{array}{c} 2.2 \\ {}^{26}{\rm Al} & \begin{array}{c} 2.1 \\ {}^{29}{\rm Ne} & \begin{array}{c} 2.9 \\ {}^{20}{\rm Ne}(\alpha, \eta)^{23}{\rm Na} & \begin{array}{c} 180 \\ {}^{31}{\rm Si} & \begin{array}{c} 0.36 \\ {}^{31}{\rm Si} & \begin{array}{c} 0.36 \\ {}^{30}{\rm Si} & \begin{array}{c} 0.36 \\ {}^{30}{\rm Si} & \begin{array}{c} 0.36 \\ {}^{30}{\rm Si} & \end{array} \end{array} \right) $		²⁶ Mg	5.2				^{22}Nc
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	${}^{12}C({}^{12}C,p){}^{23}Na$	¹⁸ O	2.1				
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁰ Ne	3.7				
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²¹ Ne	6.7				²³ Na
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²³ Na	2.6	0.41		0.45	²³ N
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁴ Na	2.7		3.1		
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Mg	4.0				
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Al	2.1		2.5		^{24}N
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁸ Mg			6.2		^{24}M
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $		³¹ Si			2.8		^{24}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{12}C({}^{12}C,n){}^{23}Mg$	²¹ Ne	2.9				
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Mg	2.0				^{25}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{16}{\rm O}(n,\gamma){}^{17}{\rm O}$	¹⁷ 0		0.31			
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²¹ Ne		2.8			
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁴ Na		4.0			
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $		²⁶ Mg		2.7			^{25}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{16}\mathrm{O}(\alpha,\gamma)^{20}\mathrm{Ne}$	²¹ Ne	2.5				^{25}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Al	6.4				
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		³⁵ S	3.9				^{26}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{16}\mathrm{O}({}^{16}\mathrm{O},p){}^{31}\mathrm{P}$	⁴⁵ Ti			2.4		
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{17}{ m O}(p,\gamma){}^{18}{ m F}$	¹⁸ O	8.2	0.21			²⁶ M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{17}{\rm O}(\alpha, n)^{20}{\rm Ne}$	¹⁷ O	0.35				^{26}M
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	${}^{18}{\rm O}(\alpha, n){}^{21}{\rm Ne}$	¹⁸ O	0.32				07
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	20 Ne $(n, \gamma)^{21}$ Ne	²¹ Ne	6.2	0.24	4.6	0.16	27 A
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	20 Ne $(\alpha, p)^{23}$ Na	¹⁸ O	0.44	2.4			27
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²³ Na	0.47	2.2	0.48		2 A
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Al		2.1			
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁸ Mg			5.4		28
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		³¹ Si			3.0		²⁸ Si
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	20 Ne $(\alpha, \gamma)^{24}$ Mg	²⁴ Mg	3.1	0.07	5.1	0.21	20
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁶ Mg			2.1		²⁹ Si
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$		²⁶ Al			2.9		⁵⁰ Si
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		²⁷ Al		0.12	3.4	0.21	
$\begin{array}{ c c c c c c c c c c c c c c c c c c c$		³⁰ Si		0.36		0.42	
$ \begin{array}{ c c c c c c c c } & & & & & & & & & & & & & & & & & & &$		^{32}P	2.2				
$\begin{array}{ c c c c c c c c c c c c c c c c c c c$		³³ P				0.49	20~
$\begin{array}{ c c c c c c c c } & & & & & & & \\ & & & 3^{5}\text{Cl} & & & & & & \\ & & & 3^{6}\text{Cl} & & & 0.38 & & & & \\ & & & & 3^{6}\text{Ar} & & 0.36 & & & 0.29 & & & & & & \\ & & & & & & 3^{0}\text{Cl} & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & & \\ & & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & $		³⁵ S		3.0			SoSi
$\begin{array}{ c c c c c c c c c c c c c c c c c c c$		³⁵ Cl		2.2			
^{30}Ar 0.36 0.29 ^{30}Si		³⁶ Cl	0.38				30
		³⁰ Ar	0.36		0.29		30 51

Notes. Yields are presented relative to those using standard rates (i.e., Fig. 2). Only species that achieve an abundance of at least $10^{-8} M_{\odot}$ and deviate from the standard abundances by at least a factor of two are presented here.

Reaction	Nuclide	W7 model		DDT model	
		×10	×0.1	×10	×0.1
	$^{37}\Delta r$	0.45		0.39	
	39 K	0.45		0.35	
	4012	0.52		0.25	2.1
	40 G			0.41	2.1
	4°Ca			0.41	2.1
	⁴¹ Ca	0.29		0.22	
	⁴² Ca			0.32	
	⁴³ Ca			0.44	
	⁴⁵ Sc				2.5
	⁴⁵ Ti			0.19	3.0
	⁴⁶ Ti	0.24		0.15	2.1
	⁴⁷ Ti			0.29	
	⁵⁶ Fe			0.43	
	⁶⁰ Ni			0.15	
21 Ne $(n,\alpha)^{22}$ Ne	21 No			0.24	
$^{21}Ne(n, \gamma)$ INC	21 No			0.44	2 1
$22N_{\rm e}(\alpha, n) = 101g$	180	0.42	2.2	0.17	5.1
$rac{p}{\gamma}$	23 1	0.42	2.3		
	²³ Na	2.3			
22	²⁴ Na	4.0	0.33	2.2	
22 Ne(α , n) 25 Mg	1/O		0.36		
	¹⁸ O	2.5	0.23		
	²⁴ Na				0.44
23 Na $(n, \gamma)^{24}$ Na	²⁴ Na	8.1	0.10	4.0	0.14
23 Na(α , p) 26 Mg	²³ Na			0.47	
	²⁴ Na			0.30	
	⁵³ Cr			2.1	
24 Na $(n n)^{24}$ Mg	^{24}Na	0.16		0.13	2.8
$^{24}Mg(n, y)^{25}Mg$	²⁶ Δ1	0.10		3.0	0.38
$^{24}Mg(\alpha, \gamma)^{28}Si$	$^{24}M\alpha$	0.31		0.3/	0.50
$\operatorname{Nig}(a, \gamma)$ SI	27 A 1	0.31		0.34	
2514-(21 N -	0.42		0.54	
$-sMg(n, \gamma)-sMg$	Ne			0.41	
	24Na			0.35	
	²³ Mg			0.30	
	²⁶ Mg	4.3	0.45		
25 Mg $(p, \gamma)^{26}$ Al	²⁶ Al	6.2		2.2	
25 Mg(α , n) 28 Si	²⁶ Al			0.34	2.1
	³¹ Si			2.2	
${}^{26}Mg(n,\gamma){}^{27}Mg$	^{28}Mg			5.4	
	³¹ Si			2.2	
${}^{26}Mg(n,n){}^{26}Al$	²⁶ A1	0.40		0.32	
$^{26}Mg(\alpha, n)^{29}Si$	²⁶ Mg			0.44	
$\operatorname{ing}(a,n)$ of	³¹ Si			3.0	
$^{27} A1(n x)^{28} Si$	$^{24}M\alpha$	0.28		0.34	
$AI(p, \gamma)$ SI	27 A 1	0.26		0.34	
27 • 1(24 M	0.20		0.20	
$AI(\alpha, p)$ SI	27 A 1	0.23	2.0	0.52	0.1
	36 C	0.10	2.0	0.14	2.1
28 0 : () 31 D	³⁰ S	2.1	0.1		
26 S1 $(\alpha, p)^{51}$ P	⁵¹ P		2.1		
20	³⁰ S		0.31		0.40
29 Si $(\alpha, n)^{52}$ S	²⁹ Si	2.1			
30 Si $(p, \gamma)^{31}$ P	³¹ P		0.49		
	^{32}P	2.0	0.25		
	³³ P		0.48		
	³⁵ S	2.4			
	³⁶ S		0.33		
30 Si $(\alpha, \gamma)^{34}$ S	³⁰ Si	0.48			
	³² P	0.49			
	³³ P	0.50		0.50	
30 Si $(n \ \gamma)^{31}$ Si	31 Si	0.50		51	
${}^{30}\text{Si}(\alpha, n){}^{33}\text{Si}$	33 p	0.48	31	5.1	3.0
SI(u,n) S	36 c	0.70	2.1		2.0
310:(31 0		2.0		2.0 2.0
$SI(p,n)^{32}P$	320	0.42		0.40	3.9
$^{22}P(p,n)^{32}S$	³² P	0.42		0.48	
$^{3}S(p,\gamma)^{3}Cl$	³³ S		2.1		

Table 1. continued.

Reaction	Nuclide	W7 model		DDT	model
		×10	×0.1	×10	×0.1
$^{34}S(\alpha, p)^{37}Cl$	³⁷ Cl	0.29		0.43	
34 S(α , n) ³⁷ Ar	³⁷ Cl	0.37			
$^{35}\mathrm{Cl}(\alpha, p)^{38}\mathrm{Ar}$	⁴⁴ Ca			2.6	
${}^{36}S(p,n){}^{36}Cl$	³⁶ S		0.35		0.44
${}^{37}\mathrm{Cl}(p,n){}^{37}\mathrm{Ar}$	³⁷ Cl		0.44		
38 Ar $(\alpha, \gamma)^{42}$ Ca	⁴⁴ Ca			2.2	
39 K(α , p) 42 Ca	⁴⁴ Ca			2.5	
42 Ca $(\alpha, \gamma)^{46}$ Ti	³⁸ Ar	0.50			
	⁴² Ca	0.42			
	⁴⁶ Ti			2.2	
	⁴⁷ Ti			2.1	
44 Ca $(p, \gamma)^{45}$ Sc	⁴⁴ Ca				2.7
${}^{45}\mathrm{Sc}(p,\gamma){}^{46}\mathrm{Ti}$	⁴⁴ Ca				3.7
	⁴⁵ Sc			0.41	4.4
	⁴⁵ Ti				2.4
${}^{45}\text{Sc}(p,n){}^{45}\text{Ti}$	⁴⁵ Ti		2.1	0.43	3.0
${}^{47}\mathrm{Ti}(n,\gamma){}^{48}\mathrm{Ti}$	⁴⁷ Ti			0.35	
${}^{48}{\rm Ti}(p,\gamma){}^{49}{\rm V}$	⁴⁸ Ti				2.4
49 V $(p, \gamma)^{50}$ Cr	⁴⁸ Ti				4.4

impact on the yields of the most abundant species produced (i.e., those with abundances greater than $10^{-2} M_{\odot}$ in Fig. 2): ${}^{12}C(\alpha, \gamma)$, ${}^{20}Ne(\alpha, \gamma)$, ${}^{24}Mg(\alpha, \gamma)$, ${}^{27}Al(p, \gamma)$, and ${}^{27}Al(\alpha, p)$. We note that each of these five reactions also satisfies conditions for "case C" in Table 2.

From the individual variation of weak interaction rates (A > A)17) by a factor of ten, we first note that we have not varied the electron/positron capture rate and β -decay rate for a particular nucleus independently; rather, we only varied the sum of these two contributions to gain a measure of sensitivity to these processes. Only three of the 658 weak interaction rates varied affect yields of species by at least 20% in either model: ${}^{28}\text{Si}(\beta^+){}^{28}\text{Al}$, ${}^{32}\text{S}(\beta^+){}^{32}\text{P}$, and ${}^{36}\text{Ar}(\beta^+){}^{36}\text{Cl}$. From the enhancement of the stellar ²⁸Si(β^+) rate, the largest effects were seen in the yields of 48 Ti, 50 V, 55 Mn, 57 Fe (which increased by $\approx 40-80\%$) and 51 V, ⁵³Cr, ⁵⁸Fe (which increased by a factor of \approx 2). Similarly, in the enhancement of the stellar ${}^{32}S(\beta^+)$ rate, the largest effects were seen in the yields of ${}^{44}Ca$, ${}^{50}V$, ${}^{52}Cr$, ${}^{54}Mn$, ${}^{56}Fe$, ${}^{57}Fe$, ${}^{59}Co$, ${}^{61}Ni$ (which increased by $\approx 40-80\%$) and ${}^{48}Ti$, ${}^{51}V$, ${}^{53}Cr$, ${}^{55}Mn$, ${}^{58}Fe$ (which increased by factors of $\approx 2-3$). Yield changes were more modest from the multiplication of the ${}^{36}\text{Ar}(\beta^+)$ rate by a factor of ten: the yields of ${}^{48}\text{Ti}$, ${}^{51}\text{V}$, ${}^{53}\text{Cr}$, ${}^{55}\text{Mn}$, ${}^{58}\text{Fe}$ all increased by $\approx 20\%$. No effects on yields were seen at levels greater than 20% when any of these three rates was reduced by a factor of ten. In addition, no significant effect on yields was seen from the individual variation of any β^- decay rate, in agreement with expectations (e.g., Brachwitz et al. 2000).

Finally, we note that since postprocessing calculations only track existing thermodynamic histories, results obtained from variations in rates that significantly affect the energy production should be interpreted carefully. Indeed, a hydrodynamic code capable of suitably adjusting both the temperature and the density of the environment in response to any changes in energy generation is required to reliably treat such cases. In the present calculations, only the ¹²C+¹²C rates were observed to modify the overall energy output by more than 5% at some point during the explosion in both models when varied by a factor of ten. In addition, variations of the ¹²C+¹⁶O and ¹⁶O+¹⁶O rates affected the calculated energy generation in the DDT model only. Of course, energy variation deduced from a postprocessing study

Table 2. Summary of important reactions from Table 1.

Reaction	Importance				
	Case A	Case B	Case C		
$^{12}C(\alpha,\gamma)^{16}O$	Х	Х	Х		
${}^{12}C({}^{12}C,\alpha){}^{20}Ne$	Х	Х	Х		
${}^{12}C({}^{12}C,p){}^{23}Na$	Х	Х	Х		
${}^{16}O(n,\gamma){}^{17}O$	Х				
${}^{16}\mathrm{O}(\alpha,\gamma)^{20}\mathrm{Ne}$	Х				
20 Ne $(n, \gamma)^{21}$ Ne			Х		
20 Ne $(\alpha, p)^{23}$ Na	Х	Х	Х		
20 Ne $(\alpha, \gamma)^{24}$ Mg	Х	Х	Х		
22 Ne $(p, \gamma)^{23}$ Na	Х		Х		
22 Ne(α , n) 25 Mg			Х		
23 Na $(n, \gamma)^{24}$ Na			Х		
23 Na(α , p) 26 Mg		Х			
24 Na $(p, n)^{24}$ Mg			Х		
24 Mg(α, γ) 28 Si			Х		
25 Mg $(n, \gamma)^{26}$ Mg		Х	Х		
25 Mg $(p, \gamma)^{26}$ Al			Х		
26 Mg $(p, n)^{26}$ Al			Х		
$^{27}\mathrm{Al}(p,\gamma)^{28}\mathrm{Si}$			Х		
27 Al $(\alpha, p)^{30}$ Si	Х		Х		
28 Si $(\alpha, p)^{31}$ P			Х		
${}^{30}\text{Si}(p,\gamma){}^{31}\text{P}$	Х				
30 Si $(\alpha, \gamma)^{34}$ S	Х		Х		
30 Si $(\alpha, n)^{33}$ S			Х		
${}^{32}\mathrm{P}(p,n){}^{32}\mathrm{S}$			Х		
${}^{34}S(\alpha, p){}^{37}Cl$			Х		
${}^{36}S(p,n){}^{36}Cl$			Х		
42 Ca $(\alpha, \gamma)^{46}$ Ti			Х		
${}^{45}\mathrm{Sc}(p,\gamma){}^{46}\mathrm{Ti}$		Х			
${}^{45}\text{Sc}(p,n){}^{45}\text{Ti}$			Х		

Notes. Variation of each of the listed rates by a factor of ten (up or down) modified yields of at least *three* species by at least a factor of two in the W7 model ("Case A") or the DDT model ("Case B"). If variation of the rate affected the yield of at least one species in *both* models, it is designated as "Case C".

does not necessarily imply substantial energy variation in a hydrodynamic study. Indeed, a recent hydrodynamic study (Bravo & Martínez-Pinedo 2012) found that variations of the ${}^{12}C+{}^{12}C$ and ${}^{16}O+{}^{16}O$ rates by a factor of ten had a negligible impact on the energy of the supernova in their DDT model.

3.2. Additional tests to motivate experiments

We have examined the effects of varying thermonuclear reaction rates by a uniform factor of ten over all stellar temperatures. To motivate and interpret experiments to better constrain the rates whose uncertainties have the largest impact on yields, we performed three additional sets of postprocessing calculations using the DDT and W7 models. For all rates in Table 2 that affect the yields of at least three species in a model (11 rates for W7 and eight rates for DDT), we investigated the nucleosynthesis from (i) varying the rate by a uniform factor of two over all temperatures and (ii) varying the rate by a factor of two over four temperature windows: 0.01-0.5 GK, 0.5-1.0 GK, 1.0-1.5 GK, and 1.5–2.0 GK. To be clear, when a reaction rate (along with, as always, the rate of the reverse process) was varied within one of the four temperature windows, we retained the standard rate for all temperatures outside the chosen window. These tests help to estimate particular temperature ranges over which the largest effects on yields may be expected from rate variations so



Fig. 3. Reactions whose rates affect the yield of at least one species in the DDT (filled circle) or W7 (empty square) models by at least a factor of two, when their rates are individually varied by a factor of ten. The heavy product is plotted against the heavy reactant for all reactions in Table 1. For example, the ²²Ne(α , n)²⁵Mg reaction (variation of which affects yields in both models) would correspond to both an empty square and a filled circle at (x, y) = (22, 25) here. (This figure is available is color in electronic form.)

Fig. 4. Reaction rates whose variation by a factor of ten affects the yields of at least three species in the DDT model (see Table 1). For each reaction, we plot the yield of each affected nuclide when the rate was enhanced by a factor of ten divided by the yield of that nuclide when the rate was reduced by a factor of ten.

as to encourage laboratory measurements of, e.g., reaction cross sections at the corresponding energies. Finally, (iii) we examined the effect of using experimentally based uncertainties (when available) for important reactions identified in Table 2.

When each member of this subset of reaction rates was individually varied by a factor of two over all temperatures, only the ${}^{12}C({}^{12}C, \alpha)$ and ${}^{12}C({}^{12}C, p)$ rates continued to significantly affect standard yields (by at least a factor of two for species with standard abundances of at least $10^{-8} M_{\odot}$). In the DDT model, the abundance of ²⁸Mg was enhanced by a factor of ≈ 2 when the ¹²C(¹²C, *p*) rate was increased by a factor of two; in the W7 model, the abundances of ²⁰Ne, ²¹Ne, and ²⁶Mg were enhanced by factors of 2–4 when the ¹²C(¹²C, α) rate was multiplied by a factor of two.

When each of these rates was individually varied by a factor of two within each of the four temperature windows mentioned above, only variations within the 1.5–2.0 GK window significantly affected yields for both models. In the DDT model,



Fig. 5. Same as Fig. 4, but for the W7 model.

variation of the rates of the ${}^{12}C({}^{12}C, \alpha)$, ${}^{12}C({}^{12}C, p)$, ${}^{20}Ne(\alpha, p)$, 23 Na(α , p), and 25 Mg(n, γ) reactions by a factor of two within the 1.5-2.0 GK window affected yields of species by at least a factor of two; in the W7 model, variation of the ${}^{12}C({}^{12}C, \alpha)$, ${}^{12}C({}^{12}C, p), {}^{16}O(n, \gamma), {}^{16}O(\alpha, \gamma), {}^{20}Ne(\alpha, p), and {}^{22}Ne(p, \gamma) rates$ within this window had an impact on yields. Interestingly, these variations affected more isotopes than observed when varying the rates by a *uniform* factor of two; moreover, for some species, the deviations from standard yields approached the level observed when varying rates by a uniform factor of ten. For example, in the DDT model, enhancement of the ${}^{25}Mg(n, \gamma)$ rate by a uniform factor of ten reduced the yields of ²¹Ne, ²⁴Na, and ²⁵Mg by factors of 0.3–0.4 (Table 1). As mentioned above, enhancement of this rate by a uniform factor of two did not change the yield of any species by at least a factor of two. However, enhancing this rate by a factor of two within the temperature window of 1.5–2.0 GK reduced the yields of ²¹Ne, ²⁴Na, and ²⁵Mg by factors of 0.4–0.5. These tests imply that one must clearly be cautious when using results from sensitivity studies to interpret experimental measurements where, e.g., a resonance may modify a given rate only over a limited range of temperatures. In addition, the lack of impact on yields from variations in the lower temperature windows suggests that useful measurements of these rates are probably best made at energies that correspond to temperatures above 1.5 GK.

A final set of tests focused on the impact of using experimentally based (temperature-dependent) uncertainties for important rates rather than variations by constant factors. For the ¹⁶O(α , γ), ²⁰Ne(α , γ), ²²Ne(p, γ), ³⁰Si(p, γ), and ²⁰Ne(α , p) rates (all of which are in Table 2), we calculated the nucleosynthesis in the DDT and W7 models assuming, for each rate, the "high rate" and "low rate" calculations of Iliadis et al. (2010). For the ²²Ne(α , n) rate, we calculated the nucleosynthesis in the two models using the "high" and "low" rate calculations from Longland et al. (2012). In general, between $\approx 1-5$ GK, the high and low rates for each reaction differ by only $\approx 10-50\%$, although we note respectively. At lower temperatures, differences can be as large as several orders of magnitude (for the ${}^{30}Si(p,\gamma)$ rate, from 0.01–0.04 GK) but are usually within factors of \approx 2–40 for temperatures between $\approx 0.01-1$ GK. We also examined the effect of using two different calculations of the ${}^{12}C(\alpha, \gamma)$ rate (Kunz et al. 2002, and a re-evaluation of the Buchmann et al. 1996 rate from the REACLIB database (Cyburt et al. 2010)). These two rates agree to within a factor of $\approx 2-3$ at relevant temperatures. We note that we did not test the effects of using upper and lower limits for the ¹²C+¹²C reactions because variation of these processes affects the nuclear energy generation rate in our models. As such, postprocessing studies cannot be used to examine such cases in detail (see Sect. 3.1). We found no significant effects (i.e., changes to any yields by a factor of at least two) when, in either model, yields determined using the high rate for a reaction were compared to yields determined using the low rate for the same reaction. In addition, no significant difference was observed when comparing yields using the two different ${}^{12}C(\alpha, \gamma)$ rates. This is consistent with the results from the tests using the temperature windows: although uncertainties at low temperatures may be comparable to or even larger than a factor of ten, these have little impact on the yields. For these cases, we encourage experimentalists to confirm the nuclear physics input presently used to determine these rates for $T \gtrsim 1.5$ GK or (for the ²²Ne(p, γ), ²²Ne(α , n), and ²⁰Ne(α , p) reactions) perform measurements to better determine these rates up to ≈ 5 GK (at which NSE was assumed in our calculations).

that the ${}^{22}Ne(p,\gamma)$, ${}^{22}Ne(\alpha,n)$, and ${}^{20}Ne(\alpha,p)$ rates are not well-

constrained experimentally above 4 GK, 1.25 GK, and 3.5 GK,

3.3. Impact on predicted isotopic ratios

In principle, abundance ratios of radioactive species produced in SN Ia may be inferred from late-time bolometric light curves. The longest lived members of the contributing nuclear decay chains (starting from ⁴⁴Ti, ⁵⁵Co, ⁵⁶Ni, and ⁵⁷Ni) have half-lives that are well separated. This leads to "ankles" in the leptondominated bolometric light curves when a longer lived chain becomes the dominant heat source (Seitenzahl et al. 2009b; Seitenzahl 2011). If the production factors are sufficiently different, this effect can in turn be used to distinguish between competing explosion models (Röpke et al. 2012). In the two models examined here, the net production of ⁴⁴Ti is dominated by the rates of the ⁴⁴Ti(α , p), ⁴⁰Ca(α , γ), ⁴⁴Ti(p, γ), ⁴⁴Ti(n, p), and ⁴⁴Ti(n, γ) reactions. The net production of ⁵⁵Co is dominated by the rates of the ⁵⁵Co(n, p), ⁵⁴Fe(p, γ), ⁵⁵Co(p, γ), ⁵⁶Ni(n, γ), and ⁵⁵Co(p, n) reactions. Finally, the ⁵⁶Ni(n, p), ⁵⁶Ni(n, γ), ⁵⁶Co(p, γ), and ⁵⁷Ni(p, γ) reactions produce ⁵⁷Ni.

It is of interest to test to what extent model predictions of these ratios are sensitive to the input reaction and weak interaction rates. While this could be inferred from the information in Table 1, the restrictions we have placed upon the contents of that table for the sake of brevity (i.e., changes of abundances with respect to standard yields by at least a factor of two) suggest that a separate analysis be used to account for changes in both members of a particular abundance ratio. We examined the effects of individual rate variations by a (uniform) factor of ten on ratios composed of the species ⁴⁴Ti, ⁵⁵Co, ⁵⁶Ni, and ⁵⁷Ni. In the DDT model, we find that variations in the ${}^{12}C(\alpha, \gamma)$ rate affect the ratios ${}^{44}\text{Ti}/{}^{55}\text{Co}$ and ${}^{44}\text{Ti}/{}^{56}\text{Ni}$ by a factor of ≈ 2 ; individual variation of the ²⁰Ne(α, γ) and ⁴⁴Ti(α, p) rates affects the same ratios by $\approx 30\%$. In addition, variation of the ⁵⁵Co(*p*, γ) rate modifies the ⁴⁴Ti/⁵⁵Co, ⁴⁴Ti/⁵⁶Ni, ⁵⁵Co/⁵⁶Ni, and ⁵⁵Co/⁵⁷Ni ratios by $\approx 30\%$. In the W7 model, variation of the ${}^{12}C(\alpha, \gamma)$ and ${}^{40}Ca(\alpha, \gamma)$ rates affects the ratios of ${}^{44}Ti/{}^{55}Co$ and ${}^{44}Ti/{}^{56}Ni$ by $\approx 30\%$. The ⁴⁴Ti(α , p) and ⁵⁵Co(p, γ) rates have somewhat larger influence when varied in W7. Both change the ratios ${}^{44}\text{Ti}/{}^{55}\text{Co}$ and ${}^{44}\text{Ti}/{}^{56}\text{Ni}$ by $\approx 40\%$, while the latter also modifies the ${}^{55}\text{Co}/{}^{56}\text{Ni}$ and ${}^{55}\text{Co}/{}^{57}\text{Ni}$ ratios at about the same level. No weak interaction rates have comparable effects on ratios of these radioactive species in either model when individually varied by a factor of ten.

The rates affecting the ratios of concern include those rates directly responsible for the net production of the relevant isotopes, as mentioned above (e.g., ${}^{44}\text{Ti}(\alpha, p)$, ${}^{55}\text{Co}(p, \gamma)$, and ${}^{40}Ca(\alpha, \gamma))$, as well as other rates such as ${}^{12}C(\alpha, \gamma)$ and ²⁰Ne(α, γ) which affect the abundances of numerous species when varied by a factor of ten (see Table 1). In addition, we find that most of the ratios affected by rate variations include ⁴⁴Ti. This is because the abundances of ⁵⁵Co, ⁵⁶Ni, and ⁵⁷Ni are not very sensitive to changes of rates by a factor of ten, even when one considers abundance changes below a factor of two. The abundances of ⁵⁶Ni and ⁵⁷Ni are robust at the 10% level to all rate changes; the abundance of ⁵⁵Co changes by $\approx 30\%$ due to variation of the ${}^{55}Co(p, \gamma)$ rate. We conclude by noting that for the two models tested here the predicted ratios of radionuclides are rather robust to rate uncertainties. Given that delayed-detonation and merger scenarios predict ratios of, e.g., ⁵⁵Co/⁵⁶Ni, that differ by a factor of ≈ 3 (Röpke et al. 2012), our results further support the idea of using observed ratios of radioactive species as a means of discriminating between explosion mechanisms.

4. Discussion

Of the 2305 nuclear reactions in our network, individual variation of only 29 rates by a factor of ten significantly affects yields of either at least three species in one model or at least one

species in both SN Ia models examined in this study (Table 2). Moreover, all but eight of these 29 reactions exclusively involve species with $20 \le A \le 36$ and $10 \le Z \le 17$ (that is, species outside the Fe group). No reaction involving any species with Z > 24 (Table 1) was found to affect yields in either model by at least a factor of two (for species with standard yields greater than $10^{-8} M_{\odot}$). Variations in only three reaction rates (those of ${}^{12}C(\alpha, \gamma), {}^{20}Ne(\alpha, \gamma), \text{ and } {}^{23}Na(\alpha, p))$ affect the yields of Fegroup species (Z > 23), see Table 1). Reactions that have the greatest influence on supernova nucleosynthesis in our models (when their rates are varied by a factor of ten) include ${}^{12}C(\alpha, \gamma)$, $^{12}C+^{12}C$, $^{20}Ne(\alpha, p)$, $^{20}Ne(\alpha, \gamma)$, and $^{30}Si(p, \gamma)$, all of which have a significant impact on the yields of at least five species in a model. Additional tests where the most influential rates were varied only within specific temperature windows indicate that rate variations for T > 1.5 GK have more impact on yields than variations at lower temperatures. These results are broadly consistent with the work of Bravo & Martínez-Pinedo (2012), in which thermodynamic histories from a single 1D DDT model were postprocessed to determine nuclear reactions whose rates influenced SN Ia yields when varied by a factor of ten. (The impact of variations in weak interaction rates was not examined in that study.) They found that nucleosynthesis in their model was most sensitive to the ${}^{20}Ne(\alpha, \gamma)$, ${}^{24}Mg(\alpha, p)$, and ${}^{30}Si(p, \gamma)$ rates and that reactions involving species with Z > 22 did not have considerable impact on yields when their rates were varied by a factor of ten. In addition, they allude to the relative robustness of the yields of Fe peak nuclei and claim that rate modifications have largest impact between about 2 < T < 4 GK. This general agreement between the results from independent studies involving three different underlying models is encouraging and suggests that SN Ia yields in single degenerate scenarios with Chandrasekhar-mass WDs are not very sensitive to uncertainties in nuclear reaction rates. This is further supported by how most of the influential reactions mentioned above (e.g., $^{12}C(\alpha, \gamma), ^{20}Ne(\alpha, p), ^{20}Ne(\alpha, \gamma), ^{30}Si(p, \gamma))$ have thermonuclear rates that are at least partially based upon experimental measurements and have uncertainties that do not significantly affect our calculated yields (see Sect. 3.2). Given this, as well as the increasing reliability of statistical model methods with increasing stellar temperature, it is likely that the impact of even the limited set of reactions in Table 2 is overestimated when a variation factor of ten is employed. Nonetheless, as stated in Sect. 3.2, we found that even rate variations by a factor of two within the 1.5 < T < 2.0 GK temperature window affected yields in our models.

Of the 658 weak interaction rates varied within our network, only three stellar rates (where contributions from electron/ positron capture and β -decay were summed) have even a modest impact on yields when individually varied by a factor of ten, and then only when enhanced. This is interesting in light of the study by Brachwitz et al. (2000), who examined the role of different libraries of electron capture rates on SN Ia yields. In general, they found that the use of libraries with larger electron capture rates resulted in larger yields (by factors of $\approx 2-10$ or more) of neutron-rich species in the Fe group (e.g., ⁵⁰Ti, ⁵⁴Cr, ⁵⁸Fe, ⁶⁴Ni). We note that the different libraries adopted in that study differed by more than simple uniform scalings of all rates and that the largest electron capture rate differences between libraries (up to several orders of magnitude) occurred for odd-A parent nuclei. In the present study, we did find some effect on neutron-rich nuclei (e.g., ^{57,58}Fe, ⁵³Cr, see Sect. 3.1) from the enhancement of individual rates. To investigate this issue further, we examined the effect on yields from multiplying all weak interaction rates

in our network (simultaneously) by a uniform factor of ten in the W7 model. This is obviously not equivalent to the study of Brachwitz et al. (2000) since we used different underlying models and different rate libraries. Moreover, we enhanced the total weak interaction rates for all species (β -decay and electron capture) as opposed to only the electron capture contributions. Nonetheless, we can test the general trend observed. Indeed, we found significant overproduction of neutron-rich species between $A \approx 49-65$ when all weak rates were enhanced. For example, ⁵⁰Ti, ⁵⁴Cr, ⁵⁸Fe, and ⁶⁴Ni (highlighted in Brachwitz et al. 2000) were overproduced by factors of ≈ 30 , 12, 4, and 8, respectively. In addition, ⁴⁹Sc, ⁴⁹Ti, ^{51,52}V, ⁵⁶Mn, ⁵⁹Fe, ^{60,61}Co, and ⁶³Ni (all neutron rich) exhibited large overproduction factors of \approx 5–10 relative to yields with the standard rates. At least with regard to weak interaction rates involved in SN Ia nucleosynthesis, these results both expose limitations of the individual variation method for sensitivity studies and confirm the need for a detailed, consistent treatment of all relevant rates.

5. Conclusions

We have investigated the sensitivity of yields from two different thermonuclear supernova models to variations in nuclear reaction and weak interaction rates. Thermodynamic histories from a DDT model and the canonical W7 pure deflagration model were postprocessed in conjunction with the individual variation of each rate in our network by a uniform factor of ten. The rates of only 14 reactions significantly affect the yields of at least three species in a model by a factor of two for species with standard yields greater than $10^{-8} M_{\odot}$. Of these, the rates of the ${}^{12}C(\alpha, \gamma)$, ${}^{12}C+{}^{12}C$, ${}^{20}Ne(\alpha, p)$, ${}^{20}Ne(\alpha, \gamma)$, and ${}^{30}Si(p, \gamma)$ reactions had the greatest impact on nucleosynthesis. Weak interaction rates had a relatively more modest impact on yields when individually varied. Enhancement of the stellar ${}^{28}Si(\beta^+){}^{28}Al$, ${}^{32}S(\beta^+){}^{32}P$, and ${}^{36}\text{Ar}(\beta^+){}^{36}\text{Cl}$ rates affected some yields by a factor of ≈ 2 ; on the other hand, no significant effect on yields was noted in either model when any weak interaction rate was reduced by a factor of ten. In general, rates that had an impact on the calculated nucleosynthesis involved nuclei with $Z \le 24$ and $A \le 40$; variation of these rates mostly affected nuclei with $20 \leq A \leq 45$, with the abundances of Fe-group nuclei being rather robust to rate variations. This is likely due to how, in models involving Chandrasekhar-mass WDs, most species in the Fe-group are synthesized in NSE (which is insensitive to rate variations). The abundances of Fe-group nuclei may indeed be less robust to rate variations in a merger scenario, where a larger fraction of the material in the Fe-group is synthesized in incomplete Si-burning. Additional tests involved the variation of important nuclear reaction rates by a factor of two (over all temperatures), by a factor of two only within specific temperature windows, and by experimentally based uncertainties. In the first of these additional tests, only the ${}^{12}C+{}^{12}C$ rates continued to affect yields. In the second, it was found that variations for T < 1.5 GK did not have a large impact on yields in our models, while in the third test, it was observed that for the reactions examined, experimentally based uncertainties did not result in significant changes to calculated yields in either model.

Overall, given the size of the reaction network employed (443 species from H to ⁸⁶Kr), nucleosynthesis in our two adopted models involving single degenerate scenarios with Chandrasekhar-mass WDs is rather robust to individual variations of the input rates. Laboratory measurements of the few important nuclear reaction rates (see Table 2), especially at energies corresponding to temperatures above 1.5 GK, would be welcome to further constrain the model predictions. Nonetheless, our results support the idea of using isotopic ratios of radioactive species as a means of discerning between single and double degenerate scenarios (which may differ by factors of up to ≈ 3 in their predictions of these ratios).

Finally, we stress the need for some caution in interpreting our results. We have examined the role of rate variations in DDT and pure deflagration models with Chandrasekhar-mass WDs. Given the scale of the required calculations, we limited the scope of this work to investigating to what degree nucleosynthesis predictions from these different, representative, explosion simulations are affected by varying the input nuclear reaction and weak interaction rates. We did not examine the role of rate variations in other proposed SNe Ia scenarios, such as those involving mergers or sub-Chandrasekhar-mass WDs, nor did we test the role of rate variations in determining, e.g., the early evolution of the progenitor in our models. In addition, we did not explore the impact of the initial composition of the WD. Such studies would be valuable and are encouraged. Moreover, we individually varied all rates but did not examine in detail the effects of simultaneous variations of rates. In other astrophysical scenarios, sensitivity studies using individual variation and simultaneous variation methods gave similar results (e.g., for Type I X-ray bursts, see Parikh et al. 2008). This should be confirmed for thermonuclear supernovae, especially given our observation of large abundance changes when all weak interaction rates in our network were simultaneously enhanced by a factor of ten. As such, it is clear that a consistent set of stellar weak interaction rates for all nuclei involved in models of Type Ia supernovae is urgently needed.

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