

# Short-lived isotopes and $^{23}\text{Na}$ production in low mass AGB Stars

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**Abstract.** We discuss the synthesis of some short-lived isotopes and of  $^{23}\text{Na}$  in thermally pulsing AGB stars with initial mass of  $2 M_{\odot}$  and two different metallicities,  $Z=1.5\times 10^{-2}$ , corresponding to the metal amount in the present Sun, and  $Z=10^{-4}$ , representative of disk and halo stars, respectively.

The different nucleosynthesis channels are illustrated in some details. As previously found, the  $^{13}\text{C}$  formed after each third dredge up episode is usually completely consumed by  $\alpha$  captures before the onset of the subsequent thermal pulse, releasing neutrons. This is the most efficient neutron source in low mass AGB stars, and the resulting s-process nucleosynthesis is at the origin of the solar main component. However, in the solar metallicity model, we find that the temperature of the first formed  $^{13}\text{C}$  pocket remains too low during the interpulse and the  $^{13}\text{C}$  is not completely burnt, being partially engulfed in the convective zone generated by the following thermal pulse. Due to the rapid convective mixing in this zone, the  $^{13}\text{C}$  is exposed to a larger temperature and a nucleosynthesis characterized by a relatively high neutron density develops. The main effect is the strong enhancement of isotopes located beyond some critical branching in the neutron-capture path, like  $^{60}\text{Fe}$ , otherwise only marginally produced during a *standard* s-process nucleosynthesis.

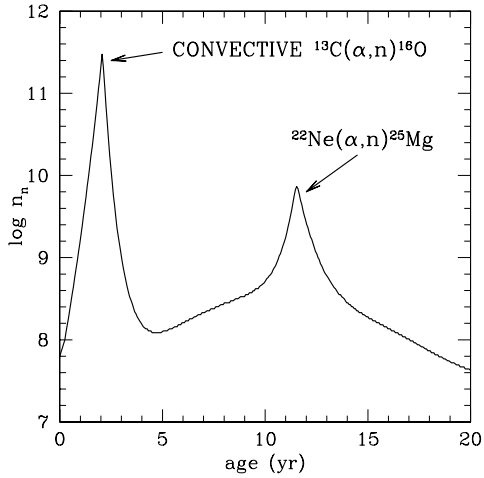
**Key words.** AGB stars – nucleosynthesis – s-process – short-lived isotopes

## 1. Introduction

Low mass AGB Stars ( $1 < M/M_{\odot} < 3$ ) are among the most important polluters of the Milky Way, because of the strong winds eroding their chemically enriched envelopes. In the AGB phase, the material processed during the alternating series of H and He burnings is recurrently mixed within the convective zones generated by thermal pulses (TPs) (Schwarzschild

et al. 1965) and then partially mixed with the surface by the so called third dredge up (TDU) episodes. As a matter of fact, MS, S, C(N) and some post-AGB stars are enriched in C and s-process elements and the spectroscopic detection of unstable Tc demonstrates that the s-process is at work in these stars (Merrill 1952).

Besides  $^{99}\text{Tc}$ , a number of short-lived isotopes are produced during the s-process, among which  $^{41}\text{Ca}$ ,  $^{60}\text{Fe}$  and  $^{107}\text{Pd}$ . Moreover,  $^{26}\text{Al}$



**Fig. 1.** Peak neutron density evolution after the ingestion of the  $^{13}\text{C}$ -pocket formed after the first TDU episode (solar metallicity model).

is an important product of the H-shell burning, by proton capture on  $^{25}\text{Mg}$ . In this work we want to study the impact of an improved version of the FRANEC code on the production of short-lived radioactivities, and discuss the challenging hypothesis that the protosolar nebula was polluted by the winds of a close-by AGB star (see Wasserburg et al. 2006 for a detailed review of this argument). A major improvement in the stellar evolution code was the introduction of a physical algorithm for the treatment of the convective/radiative interfaces (Cristallo et al. 2001) (see Section 2). In such a way we succeeded to overcome the discontinuity in the thermal gradients that otherwise would result at the border of the H-rich convective envelope and the inner radiative C and He-rich intershell region. Furthermore, the partial diffusion of protons in the top layers of the He shell gives naturally rise to the subsequent formation of a primary  $^{13}\text{C}$  pocket, partially overlapped with an outer  $^{14}\text{N}$  pocket, followed by a further minor  $^{23}\text{Na}$  pocket, both of primary origin. The new models presented here are for AGB stars of initial mass  $M = 2 M_{\odot}$  and two metallicities ( $Z=1.5 \times 10^{-2}$ ,  $Z=10^{-4}$ ), representative of disk and halo stars, respectively. In Section 2 we discuss some key characteris-

tics of the self-consistent formation of the  $^{13}\text{C}$  pocket and of the  $^{14}\text{N}$  pocket along the AGB phase. In Section 3 we examine in detail the impact of the new models on the production of short-lived radioactivities, while in Section 4 we reanalyze the production of  $^{23}\text{Na}$  in AGB stars.

## 2. The cool $^{13}\text{C}$ pocket

Two neutron sources have been identified as operating in thermally pulsing AGB stars. Concerning low mass AGBs, the main contribution to the s-process nucleosynthesis comes from the  $^{13}\text{C}(\alpha,n)^{16}\text{O}$  reaction, activated during the interpulse phase (Straniero et al. 1995), while a minor additional exposure is provided by the marginal activation of the  $^{22}\text{Ne}(\alpha,n)^{25}\text{Mg}$  reaction at the base of the convective zone generated by a TP. In our models, the introduction of an exponentially decaying profile of the average velocity of the turbulent eddies implies two important by-products. The first is a more efficient TDU, the total amount of dredged up material being about the double of the one obtained without imposing a smoothed velocity profile. The second by-product, very important for the nucleosynthesis, is the diffusion of a small amount of protons ( $\sim 5 \times 10^{-5} M_{\odot}$ ) in the underlying intershell region, mainly composed by  $^4\text{He}$  (80%) and  $^{12}\text{C}$  (20%). Later on, when the temperature in the contracting intershell region grows up, these protons are captured by the abundant  $^{12}\text{C}$ , leading to the formation of a tiny  $^{13}\text{C}$  pocket. Then, if the temperature attains  $T_8 \sim 0.9$  ( $T_8$  is the temperature in units of  $10^8$  K), the  $^{13}\text{C}$  burns, releasing neutrons and activating the s-process nucleosynthesis. Such a condition is normally reached during the interpulse phase and the  $^{13}\text{C}$  is fully consumed before the onset of the subsequent TP (Straniero et al. 1995). The resulting nucleosynthesis is characterized by a rather low neutron density, never exceeding  $10^7$  neutrons/cm $^3$  for the solar metallicity model, or  $10^8$  neutrons/cm $^3$  for the  $Z=10^{-4}$  model. In our solar metallicity model, a typical neutron exposure reached during a radiative  $^{13}\text{C}$  burning (weighted over the whole pocket) is  $\delta\tau \sim 0.25$  mbarn $^{-1}$ . The

s-rich pocket is then engulfed by the convective zone generated by the TP and, later on, dredged up by the next TDU. The partial activation of the  $^{22}\text{Ne}(\alpha, n)^{25}\text{Mg}$  reaction during the thermal pulse provides an additional minor neutron exposure. In this case a neutron burst with high peak neutron density ( $10^{10}$ - $10^{11}$  neutrons/cm<sup>3</sup>, depending on the maximum temperature reached at the base of the convective zone) is obtained.

As discussed in previous papers (see *e.g.* Cristallo et al. 2004), in the solar metallicity model, at the beginning of the AGB phase, the  $^{13}\text{C}$  pocket formed after the first TDU episode is not fully consumed and is partially engulfed by the convective shell generated by the next thermal pulse. The  $^{13}\text{C}$  burning takes place at the bottom of the convective shell ( $T_8 \sim 1.6$ ) producing a maximum neutron density  $n_n = 3.3 \times 10^{11}$  neutrons/cm<sup>3</sup>, about 30 times larger than the neutron density released by the subsequent  $^{22}\text{Ne}$  neutron burst (see Fig. 1). These numbers refer to the maximum neutron density attained at the bottom of the convective shell generated by the TP, while the average neutron density within the shell is one order of magnitude lower (Gallino et al. 1988). The contribution to the nucleosynthesis from this anomalous convective  $^{13}\text{C}$  burning reveals interesting peculiarities. In particular, some branchings, which remain closed during a standard radiative  $^{13}\text{C}$  burning and are marginally activated during the  $^{22}\text{Ne}(\alpha, n)^{25}\text{Mg}$  burning, are now open. In a very short temporal step ( $\Delta T < 3$  years, see Fig. 1) we obtain a consistent production of neutron-rich isotopes normally bypassed by the standard radiative  $^{13}\text{C}$  s-process, among which  $^{60}\text{Fe}$  (see Section 3),  $^{86}\text{Kr}$ ,  $^{87}\text{Rb}$  and  $^{96}\text{Zr}$ .

### 3. Radioactive Isotopes

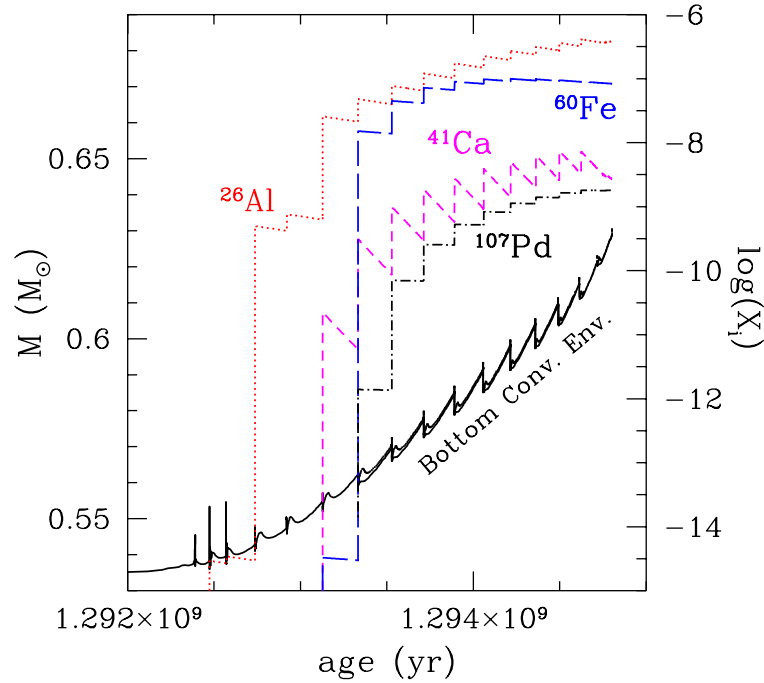
In this Section, we want to verify the hypothesis that a single low mass AGB star had contaminated the protosolar nebula right before its collapse (Wasserburg et al. 2006), by comparing the measured early solar system abundances of different short-lived isotopes to the predicted surface composition of our solar metallicity model. Before explaining

the procedure we follow, we describe the nucleosynthetic processes responsible of stellar surface enrichments of  $^{26}\text{Al}$ ,  $^{41}\text{Ca}$ ,  $^{60}\text{Fe}$  and  $^{107}\text{Pd}$ .

Ground state  $^{26}\text{Al}$  has a half-life of  $7.16 \times 10^5$  years and is efficiently produced in the H-burning shell by proton captures on  $^{25}\text{Mg}$ . The surface is firstly enriched of freshly synthesized  $^{26}\text{Al}$  when the convective envelope partially erodes the H-burning shell (first bump in the  $^{26}\text{Al}$  profile in Fig. 2) and, later on, as a consequence of TDU episodes. Note that a stellar layer exposed to a neutron flux is substantially depleted in  $^{26}\text{Al}$ , because of the large neutron-capture cross section of this isotope. This happens during both the radiative and convective  $^{13}\text{C}$  burnings as well as during the convective  $^{22}\text{Ne}$  burnings. When the temperature at the bottom of the convective shells generated by TPs exceeds  $T_8 \sim 2.7$ , the  $^{26}\text{Al}(n, p)^{26}\text{Mg}$  reaction efficiently destroys the  $^{26}\text{Al}$ . In our solar metallicity model this condition is attained only toward the end of the AGB phase, so that  $^{26}\text{Al}$  left in the H-burning shell ashes is largely preserved (with the exception of the thin layer where the radiative  $^{13}\text{C}$  burning takes place). For this reason we found a substantial surface enhancement of  $^{26}\text{Al}$ .

The appearance at the surface of  $^{41}\text{Ca}$  and  $^{60}\text{Fe}$  is simultaneous with the occurrence of the first TDU episode (see Fig. 2). As already recalled, the high  $^{60}\text{Fe}$  production we obtain is a direct consequence of the engulfment into the convective zone generated by the next TP of the  $^{13}\text{C}$  contained in the first  $^{13}\text{C}$ -pocket. The final surface isotopic ratio  $^{60}\text{Fe}/^{56}\text{Fe}$  we obtain in the solar metallicity model is equal to  $6.5 \times 10^{-5}$ , about 15 times higher than the value obtained in the case of the *standard* radiative  $^{13}\text{C}$ -burning (see for a comparison Table 4 of Wasserburg et al. 2006). On the contrary, in the  $Z=10^{-4}$  model, no cool  $^{13}\text{C}$  pockets are found.

Concerning  $^{41}\text{Ca}$ , the equilibrium value ( $^{41}\text{Ca}/^{40}\text{Ca}$ )  $\sim 10^{-2}$  is normally attained during an efficient s-process nucleosynthesis. Thus, an excess of  $^{41}\text{Ca}$  can only be obtained in the case of a very small neutron flux. This occurs when the  $^{13}\text{C}$  left in the He-rich intershell by



**Fig. 2.** TP-AGB surface evolution of  $^{26}\text{Al}$ ,  $^{41}\text{Ca}$ ,  $^{60}\text{Fe}$  and  $^{107}\text{Pd}$  (solar metallicity model).

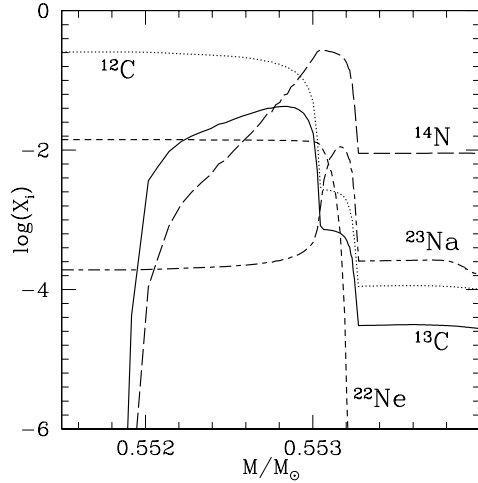
the H-burning during the Early-AGB phase is burnt by  $\alpha$  captures at the beginning of the thermally-pulsing phase, before the occurrence of the first TDU episode.

The  $^{107}\text{Pd}$  is mainly synthesized during the *standard* radiative  $^{13}\text{C}$  burning inside  $^{13}\text{C}$ -pockets, when isotopes are exposed to large neutron exposures. The more the number of radiative  $^{13}\text{C}$  burning episodes is, the larger the  $^{107}\text{Pd}$  abundance in the envelope is.

Under the hypothesis that the protosolar nebula was contaminated by a single AGB star, we follow the procedure described in Wasserburg et al. (2006): we adjust the isotopic ratios at the end of our model by assuming a dilution factor ( $f$ ) and a delay time ( $\Delta$ ) between the end of the AGB phase and the beginning of the pollution process (see Table 2). We use the  $\Delta$  assumed in Wasserburg et al. (2006) (see their Table 5) and we tune the dilution factor in order to match the Pd isotopic ratio. We find too low

Al, Ca and Fe isotopic ratios with respect ESS measurements. The situation is completely different if we analyse the "relative" isotopic enhancements just after the  $2^{\text{nd}}$  TDU: at that epoch the internal layers has suffered a unique low neutron exposure due to the convective  $^{13}\text{C}$  burning ( $\delta\tau \sim 0.05 \text{ mbarn}^{-1}$ ), avoiding in such a way the later contribution from radiative  $^{13}\text{C}$  burnings (which strongly increase the Pd isotopic ratio). We report in Table 2 (column 4) the isotopic ratios after the  $2^{\text{nd}}$  TDU, obtained by adjusting again the two free parameters. We found an agreement for Ca, Fe and Pd isotopic ratios, but a too low  $^{26}\text{Al}/^{27}\text{Al}$  turns out. We could however invoke the occurrence of Cool Bottom Process (Nollet et al. 2003), which produces  $^{26}\text{Al}$  at the base of the convective envelope enhancing its surface abundance.

Our guess is that the astrophysical source polluting the ESS was a lower mass AGB



**Fig. 3.** Abundances profiles in the  $^{13}\text{C}$  pocket formed after the first TDU episode in the solar metallicity model.

star ( $1.3 \div 1.5 M_{\odot}$ ): in that case the number and the strength of the TDU episodes would decrease and the final isotopic distribution could be similar to the one obtained after the 2<sup>nd</sup> TDU episode of our  $M=2 M_{\odot}$  (provided the occurrence of the first convective  $^{13}\text{C}$  burning, needed to increase the  $^{60}\text{Fe}$  abundance). However, this hypothesis has to be verified by a full stellar model.

#### 4. $^{23}\text{Na}$

In thermally pulsing AGB stars, sodium is synthesized by proton captures on  $^{22}\text{Ne}$ . We recall that the He-rich intershell is progressively enriched in  $^{14}\text{N}$ , left in the ashes of the H burning, and then fully converted to  $^{22}\text{Ne}$  in the convective zone powered by the He-shell flash, by the sequence  $^{14}\text{N}(\alpha, \gamma)^{18}\text{F}(\beta^-)^{18}\text{O}(\alpha, \gamma)^{22}\text{Ne}$ . In Fig. 3 we plotted, for the solar metallicity model, the chemical profiles of some trace isotopes in the region around the  $^{13}\text{C}$  pocket for a typical interpulse of the solar-metallicity model. The  $^{13}\text{C}$  pocket (solid line) partially overlaps with a more external  $^{14}\text{N}$  pocket (long-dashed line). The maximum of the  $^{14}\text{N}$  coincides with the region where the protons diffused from the convective envelope at the epoch of the TDU were abundant

enough to allow a full CN cycle. This is also the region where the  $^{23}\text{Na}$  (short-dashed-long-dashed line) has been efficiently produced by the  $^{22}\text{Ne}(p, \gamma)^{23}\text{Na}$  reaction. Indeed, at the end of the previous thermal pulse, the mass fraction of  $^{22}\text{Ne}$  in the top layer of the He-rich intershell is of the order of  $1.5 \cdot 10^{-2}$  and, around the  $^{14}\text{N}$  maximum, this  $^{22}\text{Ne}$  is almost completely converted into  $^{23}\text{Na}$ , leading to the formation of a tiny  $^{23}\text{Na}$  pocket. In the solar metallicity model, the  $^{23}\text{Na}$  production is dominated by this mechanism. In the  $Z=10^{-4}$  model, a second source of  $^{23}\text{Na}$  is represented by neutron capture on  $^{22}\text{Ne}$ , which occurs both during the radiative  $^{13}\text{C}$  burning and during the convective  $^{22}\text{Ne}$  burning. For this model, these additional sources contribute to 13% and 35% of the total sodium production, respectively.

While our solar metallicity model shows a negligible  $^{23}\text{Na}$  surface enrichment (+25%, equivalent to 0.1 dex) during the AGB phase, at the end of the  $Z=10^{-4}$  model we found a sodium surface enhancement of more than a factor 10 ( $[\text{Na}/\text{Fe}] \sim 1$ ). It is therefore worth to analyse the effects of varying some key reaction rates affecting the  $^{23}\text{Na}$  production in the low metallicity model. Recent measurements of  $^{22}\text{Ne}(p, \gamma)^{23}\text{Na}$  (Hale et al. 2002) and  $^{23}\text{Na}(p, \gamma)^{24}\text{Mg}$  reaction rates (Hale et al. 2004) significantly reduce the values quoted in the NACRE compilation (Angulo et al. 1999) around  $T_8=1$ . The reduction of the first reaction rate depresses the abundance peak of the  $^{23}\text{Na}$  pocket; this effect is marginally counterbalanced by the reduction of the second reaction rate. We then recalculate a series of TPs in the  $Z=10^{-4}$  model with rates from Hale et al. (2002, 2004), and we compare the resulting intershell sodium mass fraction with the one obtained with the NACRE rates (Angulo et al. 1999): a variation of about -30% is found. Moreover, we verified that the correction of a misprint in the  $^{23}\text{Na}(n, \gamma)^{24}\text{Na}$  reaction rate (5 keV data tabulated in Bao et al. (2000), see contribution from Heil et al. in these Proceedings) implies a further decrease by about 10%. This correction makes the contribution of the neutron capture on  $^{22}\text{Ne}$  during radiative  $^{13}\text{C}$  burning negligible, while it does not affect the  $\sim 23$  keV neutron capture

Isot. Ratio	ESS Inventory	END AGB ( $11^{\text{th}}$ TDU)		$2^{\text{nd}}$ TDU	
		$f=1.6\times 10^{-4}$	$\Delta=0.68$ Myr	$f=1.9\times 10^{-2}$	$\Delta=0.25$ Myr
$^{26}\text{Al}/^{27}\text{Al}$	$5\times 10^{-5}$		$3.3\times 10^{-7}$		$1.0\times 10^{-5}$
$^{41}\text{Ca}/^{40}\text{Ca}$	$1.5\times 10^{-8}$		$5.2\times 10^{-11}$		$1.5\times 10^{-8}$
$^{60}\text{Fe}/^{56}\text{Fe}$	$(2\div 20)\times 10^{-7}$		$4.4\times 10^{-9}$		$1.9\times 10^{-7}$
$^{107}\text{Pd}/^{108}\text{Pd}$	$2.0\times 10^{-5}$		$2.0\times 10^{-5}$		$2.0\times 10^{-5}$

**Table 1.** Measured and predicted surface isotopic ratios involving short-lived isotopes. (see text for details).

occurring in the convective  $^{22}\text{Ne}$  burning. As a whole, we estimate that with these changes of the relevant reaction rates the final overabundance of sodium will be  $[\text{Na}/\text{Fe}]\sim 0.7$ .

## 5. Conclusion

In this paper we study how short-lived isotopes are created in a low mass ( $M=2 M_{\odot}$ ) AGB star, showing the strong effects a single convective  $^{13}\text{C}$  burning episode implies. A possible explanation for the ESS short-lived isotopes has been proposed, this however implying strong *ad hoc* assumptions.

In addition, we present detailed nucleosynthetic results concerning the production of  $^{23}\text{Na}$  in low mass AGB stars at different metallicities ( $Z=1.5\times 10^{-2}$  and  $Z=10^{-4}$ ), representative of disk and halo stars respectively. Different processes responsible for the  $^{23}\text{Na}$  nucleosynthesis have been analyzed in detail,

stressing the importance of varying key nuclear reaction rates.

## References

- Angulo, C. et al. 1999, Nucl. Phys. A, 656, 3  
 Bao, Z.Y., & Käppeler, F. 2000, Atom. Data Nucl. Data Tables, 76, 70  
 Cristallo, S. et al. 2001, Nucl. Phys. A, 688, 217  
 Cristallo, S. et al. 2004, Mem. SAI, 75, 676  
 Gallino, R. et al. 1988, ApJ, 334, 45L  
 Hale, S.E. et al. 2002, Phys. Rev. C, 65, 015801  
 Hale, S.E. et al. 2004, Phys. Rev. C, 70, 045802  
 Merrill, P.W. 1952, ApJ, 116, 21  
 Nollett, K.M. et al. 2003, ApJ, 582, 1036  
 Schwarzschild, M. & Härm, R. 1965, ApJ, 142, 855  
 Straniero, O. et al. 1995, ApJ, 440L, 85S  
 Wasserburg, J.J. et al. 2006, Nucl. Phys. A, *in press*