NEW INSIGHTS ON SATURN’S FORMATION FROM ITS NITROGEN ISOTOPIC COMPOSITION

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ABSTRACT

The recent derivation of a lower limit for the 14N/15N ratio in Saturn’s ammonia, which is found to be consistent with the Jovian value, prompted us to revise models of Saturn’s formation using as constraints the supersolar abundances of heavy elements measured in its atmosphere. Here we find that it is possible to account for both Saturn’s chemical and isotopic compositions if one assumes the formation of its building blocks at ~45 K in the protosolar nebula, provided that the O abundance was ~2.6 times protosolar in its feeding zone. To do so, we used a statistical thermodynamic model to investigate the composition of the clathrate phase that formed during the cooling of the protosolar nebula and from which the building blocks of Saturn were agglomerated. We find that Saturn’s O/H is at least ~34.9 times protosolar and that the corresponding mass of heavy elements (~43.1 M⊕) is within the range predicted by semi-convective interior models.

Key words: planets and satellites: atmospheres – planets and satellites: composition – planets and satellites: formation – planets and satellites: individual (Saturn) – protoplanetary disks – solid state: volatile

Online-only material: color figures

1. INTRODUCTION

The measurements of 14N/15N ratios throughout the solar system can be divided into three categories (see Mandt et al. 2014, and references therein): the solar wind and Jupiter have the lightest ratios, presumed to be representative of the protosolar ratio. Chondrites, grains coming from comet 81P/Wild 2, Earth’s mantle and atmosphere, Venus and Mars’ mantles, have moderately heavy ratios. Saturn’s moon Titan, Mars’ atmosphere, as well as NH3 and HCN in comets share the lowest 14N/15N values.

The recent derivation of a 1σ lower limit for the 14N/15N ratio in Saturn’s ammonia, which is found to be ~500 from TEXES/Infrared Telescope Facility ground-based mid-infrared spectroscopic observations (Fletcher et al. 2014), prompts us to revise models of Saturn’s formation that previously only used the supersolar abundances of heavy elements measured in the observable troposphere as constraints. This lower limit is formally consistent with the 14N/15N ratio (~435) measured by the Galileo probe at Jupiter (Fletcher et al. 2014) and implies that the two giant planets were essentially formed from the same nitrogen reservoir in the nebula, which is N2 (Owen et al. 2001; Fletcher et al. 2014). Any scenario depicting Saturn’s formation should match the 14N/15N ratio measured in its atmosphere and be consistent with disk temperatures greater than 30 K in the giant planet’s formation region. Lower temperatures have only been observed in regions located beyond ~30 AU in circumstellar disks (Qi et al. 2013).

Two scenarios of Saturn’s formation, aiming at matching the supersolar volatile abundances measured in its envelope, have been proposed. Both approaches determine the composition of the planet’s building blocks from a simple clathrate formation model and assume that all elements were in protosolar abundances in the disk’s gas phase. The first scenario, proposed by Hersant et al. (2008), assumes that Saturn formed at ~40–50 K in the protosolar nebula (hereafter PSN). In their model, NH3 was trapped in planetesimals, while the dominant N molecule in the PSN, N2, remained well mixed with H2 until the gas collapsed onto the core of the planet. This scenario has been ruled out because it suggests that Saturn’s supersolar N abundance essentially results from the delivery of NH3 trapped in solids, implying that its 14N/15N ratio should be substantially lower than the Jovian value (Hersant et al. 2008).

Alternatively, Moussis et al. (2009b) proposed that Saturn’s building blocks formed at a cooler temperature in the disk. In this scenario, planetesimals were agglomerated from a mixture of clathrates and pure ices condensed close to ~20 K, implying that both NH3 and N2 were trapped in solids. Their model is consistent with the measured 14N/15N ratio since N2 remains the main nitrogen reservoir delivered to Saturn. However, the formation of Saturn’s building blocks at such a low temperature in the PSN is questionable as the heating of the disk by proto-Sun’s UV radiation might prevent the temperature from decreasing down to ~40 K at 10 AU (D’Alessio et al. 1998).

Here we find that it is possible to account for both Saturn’s chemical and isotopic compositions if one assumes the formation of its building blocks at ~45 K in the PSN, provided that the O abundance was ~2.6 times protosolar in its feeding zone. To do so, we used a statistical thermodynamic model (Mousis et al. 2010, 2012) to investigate the composition of the clathrate phase that formed during the cooling of the PSN from the most abundant gaseous volatiles. These clathrates agglomerated with the other ices and rocks and formed the building blocks of Saturn. A fraction of these planetesimals accreted in the growing Saturn dissolved in its envelope and subsequently engendered the observed volatile enrichments.
Table 1

| Species | Measurements | Minimum and Maximum Fits of C |
|---------|--------------|-----------------------------|
| O       | . .          | 34.9–43.0                   |
| C       | 9.6 ± 1.0 b  | 8.6–10.6                    |
| N       | 2.8 ± 1.1 c  | 7.5–9.2                     |
| S       | 12.05 d      | 5.2–6.4                     |
| P       | 11.2 ± 1.3 e | 10.4–12.7                   |
| Ar      | . .          | 1.9–2.3                     |
| Kr      | . .          | 8.3–10.3                    |
| Xe      | . .          | 10.4–12.7                   |

Notes. Saturn’s formation temperature is considered at ~45 K. The observed values are derived from.

a Error is defined as \((\Delta E/E)^2 = (\Delta X_{\text{Saturn}}/X_{\text{Sun}})^2 + (\Delta X_{\text{C}}/X_{\text{C}})^2\).
b Fletcher et al. (2009b).
c Fletcher et al. (2011).
d Briggs & Sackett (1989).
e Fletcher et al. (2009a); using the protosolar abundances of Lodders et al. (2009).

2. USEFUL ELEMENTAL ABUNDANCES MEASURED IN SATURN

Table 1 summarizes the abundances of C, N, P, S, and O, normalized to their protosolar abundances, and measured in the forms of CH₄, NH₃, PH₃, H₂S (indirect determination), and H₂O in Saturn’s atmosphere. Note that the protosolar abundances correspond to the present day solar values corrected from elemental settling in the Sun over the past 4.56 Gyr (Lodders et al. 2009). The abundance of CH₄ has been determined from the analysis of high spectral resolution observations from Cassini/CIRS (Fletcher et al. 2009b). As methane does not condense at Saturn’s atmospheric temperatures, its atmospheric abundance can be considered as representative of the bulk interior. The NH₃ abundance is taken from the range of values derived at the equator by Fletcher et al. (2011) from Cassini/VIMS 4.6–5.1 μm thermal emission spectroscopy. The measured NH₃ abundance may be considered a lower limit since the condensation level of NH₃-bearing volatiles may be deeper than the sampled regions (Atreya et al. 1999, 2014), implying that there could be a large reservoir of ammonia hidden below the condensate cloud decks. PH₃ has been determined from Cassini/CIRS observations at 10 μm (Fletcher et al. 2009a). The H₂S abundance is quoted from the indirect determination of Briggs & Sackett (1989) from radio observations but remains highly uncertain.

3. MODEL DESCRIPTION

In our model, the volatile phase incorporated in planetesimals is composed of a mixture of pure ices, stoichiometric hydrates (such as NH₃–H₂O hydrate) and multiple guest (MG) clathrates that crystallized in the form of microscopic grains at various temperatures in the outer part of the disk. We assume that planetesimals have grown from collisional coagulation of the icy grains (Weidenschilling 1997). Here, the clathration process stops once crystalline water ice has been consumed by the trapping of volatile species in clathrate and hydrate phases. Only pure condensates can subsequently form if the disk cools down to very low temperatures. The process of volatile trapping in planetesimals formed in Saturn’s feeding zone follows the approach depicted in Mousis et al. (2012), who used a statistical thermodynamic model to compute the composition of MG clathrates formed in the PSN. We refer the reader to this Letter for further information on the employed model. We use typical PSN temperature and pressure profiles (Hueso & Guillot 2005) to compute the evolution of the thermodynamic conditions at the current location of Saturn. Here, our computations have been made in the case of the formation of structure I MG clathrates because CO, CO₂, and H₂S, namely, the most abundant volatiles in the PSN, also individually form structure I clathrates.

Our computations are based on a predefined initial gaseous composition in which all elemental abundances, except that of oxygen in some circumstances (see Section 5), are protosolar (Lodders et al. 2009). We assume that O, C, and N exist only in the form of H₂O, CO, CO₂, CH₃OH, CH₄, N₂, and NH₃. Hence, once the gaseous abundances of elements are defined, the molecular abundances are determined from the adopted CO:CO₂:CH₃OH:CH₄ and N₂:NH₃ gas-phase molecular ratios. The remaining O gives the abundance of H₂O. We set CO:CO₂:CH₃OH:CH₄ = 10:4:1.67:1 in the gas phase of the disk, values consistent with interstellar medium measurements (Pontoppidan 2006; Öberg et al. 2011) and measurements of production rates of molecules in Comet C/1995 O1 Hale–Bopp (Bockelé-Morvan et al. 2004). In addition, S is assumed to exist in the form of H₂S, with H₂S:H₂ = 0.5 × (S:H₂)⊙, and other S-rich refractory components (Pasek et al. 2005). We finally consider N₂:NH₃ = 10:1 in the nebula gas phase, a value predicted by PSN chemical models (Lewis & Prinn 1980).

Figure 1 shows two cases for the compositions of planetesimals condensed in Saturn’s feeding zone and represented as...
a function of their formation temperature. In both cases, NH$_3$ forms NH$_3$–H$_2$O hydrate and CH$_3$OH is assumed to condense as pure ice in the PSN because of the lack of thermodynamic data concerning its associated clathrate. In the first case (full clathration), all volatiles (except NH$_3$ and CH$_3$OH) are trapped in the clathrate phase as a result of an initial supersolar oxygen abundance ($\sim 2.6 \times (O/H)_{\odot}$) corresponding to H$_2$O/H$_2$ = 2.47 \times 10^{-3}$ in Saturn’s feeding zone. In the second case (limited clathration), we used a protosolar abundance for oxygen, corresponding to H$_2$O/H$_2$ = 5.55 \times 10^{-4}$ in Saturn’s feeding zone, and implying that the budget of available crystalline water is not sufficient to trap all volatiles in clathrates. In this case, significant fractions of CO, N$_2$, and Ar form pure ices if the disk cools down to very low temperatures ($\sim 20$ K) instead of being trapped in clathrates as it is the case for full volatile clathration. For example, Ar, N$_2$, and CO become substantially trapped in the clathrate phase at $\sim 38, 45$, and 48 K in the PSN, respectively. In contrast, these species form pure ices in the 22–26 K range in the PSN.

Assuming that the composition of the icy phase of planetesimals computed with our model is representative of that of Saturn’s building blocks, the precise adjustment of their mass accreted by the forming Saturn and vaporized into its envelope allows us to reproduce the observed volatile enrichments. Here, because of the lack of reliable measurements, our fitting strategy is to match the minimum carbon enrichment measured in Saturn. By doing so, this allows us to maintain a mass of solids accreted into Saturn’s envelope that is as small as possible in order to be compared to the mass of heavy elements predicted by interior models.

4. RESULTS

Figure 2 shows the evolution of the N$_2$/NH$_3$ ratio in Saturn as a function of the formation temperature of its building blocks, assuming that it was equal to 10 in the PSN prior to planetesimals formation. Depending on the temperature considered for Saturn’s formation, contributions of both N$_2$ and NH$_3$ in solid and gaseous phases have been taken into account in our computation. When not trapped or condensed, the species collapse with the nebula gas onto the forming planet and form its gaseous envelope. NH$_3$ is always in solid form at $T < 80$ K. In the full clathration case, the maximum temperature of Saturn’s formation yielding N$_2 \gg$ NH$_3$ in the envelope is $\sim 45$ K. Above this temperature, N$_2$ remains essentially in gaseous form. In contrast, in the limited clathration case, N$_2$ dominates in Saturn only at formation temperatures lower than $\sim 22$ K, a value corresponding to its condensation temperature in the PSN. In both situations, the amount of N$_2$ supplied to Saturn in gaseous form is less than that of NH$_3$ in solid form. A comparison between the two cases shows that the full volatile clathration favors a higher N$_2$/NH$_3$ in Saturn at temperatures below $\sim 45$ K. Because (1) disk temperatures as low as $\sim 22$ K at the formation location of Saturn are unlikely and (2) the full clathration scenario is fully consistent with a high N$_2$/NH$_3$ in Saturn, only this latter case is considered in the following.

Figure 3 represents the volatile enrichments in Saturn calculated from the fit of the minimum C abundance observed in the atmosphere ($\sim 8.6$ times protosolar—see Table 1) as a function of the formation temperature of its building blocks and in the case of the full clathration scenario. In the PSN temperature range ($T \leq 45$ K) consistent with the $^{14}$N/$^{15}$N constraint, we find that N is 7.5 times more enriched than the protosolar value in Saturn’s atmosphere, a value higher than the maximum inferred one (3.9 times protosolar), but still lower than the measured C enrichment (9.6 ± 1 times protosolar). In this case, O is predicted to be at least $\sim 34.9$ times more enriched than the protosolar value in Saturn’s envelope. This strong O enrichment is due to the assumption of a supersolar abundance of oxygen in Saturn’s feeding zone, which is required for the full trapping of guest molecules in clathrates ($\sim 6$ water molecules are needed to stabilize one guest molecule). Interestingly, the calculated P enrichment ($\sim 10.4$ times protosolar) matches the measured value (11.2 ± 1.3 times protosolar). S is found to be 5.2 times more enriched than the protosolar value, but remains lower than the indirect determination ($\sim 12$ times protosolar). Table 1 summarizes the enrichments calculated from the minimum and maximum fits of the C abundance observed in Saturn and gives predictions for the volatile species that have not yet been detected (Ar, Kr, Xe) or those whose sampling still needs to be investigated.
(O, N, S, P). The observed P abundance is matched by our model when C ranges between 8.6 and 10.4 times protosolar.

Figure 4 shows that ∼27.9–32.2 M⊕ of ices, including ∼21.9–25.3 M⊕ of water, are needed in Saturn to match the measured C enrichment in the full clathration scenario at ∼45 K in the PSN. Our calculated mass range must be seen as a minimum because planetesimals may harbor a significant fraction of the refractory phase. We find that 15.2–17.6 M⊕ of rocks are needed to match the observed C enrichment, assuming an ice-to-rock ratio of ∼1 for a protosolar composition of gas (Johnson et al. 2012). This implies that ∼43.1–49.8 M⊕ of heavy elements have been delivered to Saturn to match the observed C enrichment. The mass of heavy elements needed by the full clathration scenario then exceeds the maximum mass of heavy elements predicted in Saturn by homogeneous interior models (≤30 M⊕; Nettelmann et al. 2013). On the other hand, our calculations are consistent with the mass range (26–50 M⊕) of heavy elements predicted by semi-convective models (Leconte & Chabrier 2012).

5. DISCUSSION

The lower limit for the 14N/15N ratio found by Fletcher et al. (2014) implies that Saturn’s nitrogen was essentially accreted in N2 form at its formation time. However, this condition is not sufficient to match the measured 14N/15N ratio: our calculations suggest that N2 must have been accreted in solid form in Saturn, in order to match the observed C enrichment, otherwise NH3 would still remain the main N-bearing reservoir in the envelope.

We have explored two hypotheses to simultaneously account for the 14N/15N measurement and the volatile enrichments in Saturn by varying the O/H ratio in the giant planet’s feeding zone. Both possibilities were investigated by using a statistical thermodynamic approach, allowing us to compute the composition of clathrates formed in the PSN. In the first case (full clathration scenario), we assumed that oxygen was sufficiently abundant (∼2.6 × (O/H)⊙) to trap all volatiles as clathrates in Saturn’s feeding zone (except NH3 which forms a stoichiometric hydrate and CH3OH due to the lack of thermodynamic data concerning its associated clathrate), leading to N2 trapping in planetesimals at ∼45 K in the PSN. In the second case (limited clathration scenario), we assumed that the O abundance was protosolar, implying that planetesimals were agglomerated from a mixture of clathrates and pure condensates. The PSN had to cool down to ∼22 K in order to allow the trapping of solid N2 in planetesimals. The full clathration hypothesis is the only scenario allowing the formation of Saturn’s building blocks at temperatures consistent with our knowledge of the thermal structure of the PSN. The presence of a supersolar oxygen abundance in the giant planet’s feeding zone may be explained via its formation in the neighborhood of Jupiter, close to the water ice line location at earlier epochs of the PSN. At this location, the abundance of crystalline water ice may have been enhanced by diffusive redistribution and condensation of water vapor (Stevenson & Lunine 1988; Ali-Dib et al. 2014a, 2014b) thus easing the formation of clathrates. Recent volatile distribution models, elaborated by Ali-Dib et al. (2014a, 2014b) and taking into account the major dynamic and thermodynamic effects relevant to volatiles (turbulent gas drag, sublimation of solids, gas diffusion, and condensation), predict enhancements of the surface density of water up to several times the one derived from a protosolar O abundance at the location of the H2O ice line. If the region of the disk where the surface density of water is enhanced extends over several AU, then Jupiter and Saturn could form independently. This would require that the diffused H2O vapor condenses over large length scales, as shown by the simulations of Ros & Johansen (2013). Alternatively, recent disk models including midplane dead zone related effects (Martin & Livio 2012; Martin & Lubow 2013) show that the temperature profile might not be monotonic in the PSN. These models suggest the occurrence of several ice lines for the same species at some stages of the PSN evolution. In this context, Jupiter and Saturn may have formed at the locations of two distinct water ice lines, each of them providing enough crystalline water for enabling the full clathration of the other volatiles when the giant planets feeding zones reached lower temperatures. In any case, the building blocks accreted by Jupiter and Saturn during their formation should have close C/O and C/N ratios, given the similarity of their formation conditions and the bulk composition of the two planets should also reflect these ratios.

The full clathration hypothesis matches well the formation scenario of Jupiter proposed by Gautier et al. (2001), where the authors also proposed that the volatiles were fully trapped in clathrates and found O/H ∼ 2.5 × (O/H)⊙ in the giant planet’s feeding zone from a simple clathrate formation model. A higher abundance of water ice in Saturn’s feeding zone increases the ice-to-rock ratio in planetesimals and implies that the icy phase is dominant in the heavy elements accreted by Saturn’s envelope. Given that the full clathration scenario best fits the data, it favors the idea that Saturn’s interior is heterogeneous and may exhibit a continuous compositional gradient, as suggested by Leconte & Chabrier (2012). In order to match this model, one needs to argue that a fraction of the heavy elements sedimented onto Saturn’s core during its evolution (Fortney & Hubbard 2003, 2004). For example, if all rocks sedimented onto Saturn’s core (∼15.2–17.6 M⊕), then the mass of volatiles remaining in the envelope (∼27.9–32.2 M⊕) holds well within the mass range of heavy elements (10–36 M⊕) predicted by the semi-convective models of Leconte & Chabrier (2012). Given the high O enrichment predicted in Saturn (34.9 times the protosolar value), one should expect an increase relative to the D/H ratio in the envelope’s hydrogen, a prediction that is not supported by
existing observations (Lellouch et al. 2001; Bézard et al. 2003). However, if the interior of Saturn is semi-convective, the D/H ratio measured in its upper layers would not be representative of the planet’s global value.

Interestingly, our results are consistent with the fact that NH₃ must be the main primordial reservoir of nitrogen in Titan to explain its current ¹⁴N/¹⁵N ratio (Mandt et al. 2014). Indeed, formation scenarios predict that Titan’s building blocks must have experienced a partial devolatilization during their migration in Saturn’s subnebula, which would have induced the loss of the CO, N₂, and Ar captured from the nebula (Mousis et al. 2009a). Hence, Titan’s building blocks probably originate from Saturn’s feeding zone but they would have been subsequently altered by the subnebula. Finally, notwithstanding the conclusions of the present study, it should be kept in mind that only the in situ measurement of O below the condensation layer of water and the precise assessment of the C, N, P, and noble gas abundances will be able to shed light on the formation conditions of the ringed planet (Mousis et al. 2014).

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