Organic Ices in Titan’s Stratosphere

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Abstract Titan’s stratospheric ice clouds are by far the most complex of any observed in the solar system, with over a dozen organic vapors condensing out to form a suite of pure and co-condensed ices, typically observed at high winter polar latitudes. Once these stratospheric ices are formed, they will diffuse throughout Titan’s lower atmosphere and most will eventually precipitate to the surface, where they are expected to contribute to Titan’s regolith.

Early and important contributions were first made by the InfraRed Interferometer Spectrometer (IRIS) on Voyager 1, followed by notable contributions from IRIS’ successor, the Cassini Composite InfraRed Spectrometer (CIRS), and to a lesser extent, from Cassini’s Visible and Infrared Mapping Spectrometer (VIMS) and the Imaging Science Subsystem (ISS) instruments. All three remote sensing instruments made new ice cloud discoveries, combined with monitoring the seasonal behaviors and time evolution throughout Cassini’s 13-year mission tenure.

A significant advance by CIRS was the realization that co-condensing chemical compounds can account for many of the CIRS-observed stratospheric ice cloud spectral features, especially for some that were previously puzzling, even though some of the observed spectral features are still not well understood. Relevant laboratory transmission spectroscopy efforts began just after the Voyager encounters, and have accelerated in the last few years due
to new experimental efforts aimed at simulating co-condensed ices in Titan’s stratosphere. This review details the current state of knowledge regarding the organic ice clouds in Titan’s stratosphere, with perspectives from both observational and experimental standpoints.

Keywords Ices · Clouds · Titan · Infrared spectroscopy · Infrared observations · Thermal infrared · Stratospheric composition

1 Setting the Stage

It comes as no surprise that ices thrive on the surfaces of many airless bodies in the outer solar system, e.g., comets, gas and ice giant planet satellites, Kuiper Belt objects, etc. Ices are, however, also observed on solar system objects that, besides having discernible surfaces, also contain moderate atmospheres. We define this collection of objects to be the terrestrial group, with four primary members: Venus, Earth, Mars, and Saturn’s moon Titan. Titan is naturally grouped with the other three terrestrial planets since it is more similar to Earth than dissimilar, despite Titan’s distance of ∼10 AU from the sun. Titan has an atmosphere that is about ten times more massive than Earth’s, albeit Titan is less than half the size of Earth. Titan also has weather patterns with rain, clouds, and wind (see for example Porco et al. 2005; Roe et al. 2005; Rodriguez et al. 2011; Turtle et al. 2011a,b, and references therein), and even rivers, lakes, and seas (see for example Stofan et al. 2007; Hayes et al. 2008, 2011; Sotin et al. 2012; Hayes 2016, and references therein).

The four members of the terrestrial group have both similarities and differences. For example, the atmospheres of Venus and Mars are dominated by carbon dioxide (CO2), and those of Earth and Titan are predominantly molecular nitrogen (N2). Earth and Mars, however, are both fast rotators, resulting in latitudinal transport of heat and momentum preferentially through cyclonic-anticyclonic motion, whereas Venus and Titan are slow rotators, resulting in latitudinal transport of heat and momentum through cross-equatorial meridional circulation (Leovy and Pollack 1973; Flasar et al. 1981). On Titan, it is this slow atmospheric meridional circulation pattern that transports trace organic vapors from southern to northern latitudes in Titan’s mesosphere and stratosphere, with a return flow of the atmospheric bulk in the troposphere from north to south (around northern winter solstice). The meridional flow pattern is expected to reverse near southern winter solstice, although observations of Titan’s atmospheric vapors and temperatures have shown this reversal to occur a few years post equinox (see for example Teanby et al. 2012, 2017). For more information on Titan’s general circulation patterns, see for example Hourdin et al. (2004), Lebonnois et al. (2012), Lora et al. (2015), Newman et al. (2016), and references therein.

Titan is much colder than Earth, with significantly more atmospheric CH4. A warmer Titan during its formation would have driven off volatiles like CH4, so Titan’s lower temperature combined with its larger CH4 abundance, helps to explain the important role of Titan’s organic vapor inventory. The source of Titan’s CH4 vapor is the surface or subsurface, with a mole fraction of ∼1.5% transported through Titan’s cold trap, located at altitudes just below the tropopause, and on into the lower stratosphere. Assuming the CH4 mole fraction remains constant above the cold trap, this mole fraction of 1.5% would also be present in the mesosphere and thermosphere. At these upper atmospheric altitudes, both CH4 and N2 are dissociated from solar UV photons and energetic particles originating from Saturn’s magnetosphere, leading to complex organic chemistry (see Yung et al. 1984; Coates et al. 2009; Horst 2017, and references therein). The resulting photofragments, atoms, and ions recombine into more exotic trace organic vapors, some of which are transported down to
the stratosphere to condense and form ice clouds (Maguire et al. 1981; Sagan and Thompson 1984; Frere et al. 1990; Samuelson et al. 1997; Samuelson et al. 2007; Coustenis et al. 1999; Raulin and Owen 2002; Anderson and Samuelson 2011; Anderson et al. 2014, 2016), while others in turn combine, polymerize, and form Titan’s refractory aerosol particles (see Waite et al. 2004, 2007; Coates et al. 2009; Sciamma-O’Brien et al. 2010; Hörst 2017). Here we define aerosol as the refractory component of Titan’s atmospheric particulate material (i.e., haze), whereas the stratospheric ices are the volatile component of Titan’s haze. These aerosol particles are very small, leading to relatively large absorption cross-sections at solar wavelengths, and small absorption cross-sections at thermal IR wavelengths (Tomasko 1980; Tomasko and Smith 1982; Rages and Pollack 1980; Rages et al. 1983; Tomasko et al. 2009; Doose et al. 2016). Thus, the aerosol particles are easy to heat and difficult to radiatively cool. Temperatures build up, and heat is transferred by conduction to adjacent vapor molecules. The aerosol particles exponentially attenuate the solar radiation, giving rise to an extensive stratosphere defined by the largest positive temperature gradient of any object in the solar system (Hanel et al. 2003), with a tropopause temperature on average of \( \sim 70 \) K (at \( \sim 42 \) km) and a stratopause temperature on average of \( \sim 190 \) K (at \( \sim 300 \) km) (Hanel et al. 1981; Flasar et al. 2005; Achterberg et al. 2008, 2011). Methane vapor also plays a non-negligible role in radiatively heating Titan’s stratosphere.

From their formation altitudes, Titan’s aerosol particles eventually drift downward into the stratosphere, providing heterogeneous nucleation sites for vapor condensation. It is condensation in Titan’s stratosphere that is the ultimate sink mechanism for most of Titan’s trace organic vapors. First, the Voyager 1 InfraRed Interferometer Spectrometer (IRIS; see Sect. 2), and later from its successor, the Cassini Composite InfraRed Spectrometer (CIRS; see Sect. 3), observed trace quantities of the hydrocarbon vapors ethane \( (\text{C}_2\text{H}_6) \), acetylene \( (\text{C}_2\text{H}_2) \), propane \( (\text{C}_3\text{H}_8) \), methylacetylene \( (\text{C}_3\text{H}_4) \), diacetylene \( (\text{C}_4\text{H}_2) \), and of the nitrile vapors hydrogen cyanide \( (\text{HCN}) \), cyanoacetylene \( (\text{HC}_3\text{N}) \), and cyanogen \( (\text{C}_2\text{N}_2) \) (see for example Hanel et al. 1981; Coustenis and Bezard 1995; Samuelson et al. 1997; Coustenis et al. 1999, 2007, 2016, 2018; Teanby et al. 2007; Vinatier et al. 2010, 2015; Sylvestre et al. 2018), all capable of condensing to form ice clouds in Titan’s cold mid to low stratosphere. The other three terrestrial atmospheres are incapable of generating such a diversity of trace organic vapors. Figure 1 gives an example showing the very rough approximate condensation altitudes for eight of Titan’s stratospheric organic vapors during late southern fall at 79° S. The vapor abundances and temperature structure are determined from CIRS analyses during the Cassini flyby in July 2015. The saturation vapor pressure expressions for \( \text{HC}_3\text{N}, \text{C}_2\text{N}_2, \text{HCN}, \text{C}_2\text{H}_2, \text{C}_2\text{H}_6, \text{and } \text{C}_6\text{H}_6 \) are taken from Fray and Schmitt (2009), while those for \( \text{C}_3\text{H}_4 \) and \( \text{C}_4\text{H}_2 \) come from Lara et al. (1996). For an overview on the evolution of Titan’s stratospheric ice particles, see Sagan and Thompson (1984), Frere et al. (1990), Raulin and Owen (2002), Anderson et al. (2016).

For most of Titan’s stratospheric organic trace vapors, CIRS-measured abundances and temperatures (see for example Coustenis et al. 2007, 2016, 2018; Teanby et al. 2007; Achterberg et al. 2008, 2011; Vinatier et al. 2010, 2015; Sylvestre et al. 2018) require condensation directly into the solid phase; the vapors will never condense directly to the liquid phase. Thus ice clouds, and never rain clouds, are always formed in Titan’s stratosphere. As a generalization, the specific vapor compound and altitude location estimate of condensation depends on the vapor abundances and the temperatures at which saturation for each vapor occurs during its downward journey—cooling as it descends—through Titan’s stratosphere. Once formed, these ice cloud particles will precipitate through the mid and lower regions of Titan’s stratosphere, continually adding layers as they pass through altitude regions where the individual organic vapors successively become saturated. As the temperature continues to decrease...
Fig. 1 Titan’s pressure-temperature-altitude profile (black curve) at 79°S during late southern fall (July 2015). Superimposed are the vertical distributions for eight of Titan’s stratospheric organic vapors (color-coded curves), showing the approximate altitude location for each individual vapor where condensation is possible—the intersections between the saturation vapor pressure curves and the temperature profile give a rough estimate of cloud top altitude. The vapor abundances and temperature structure are determined from CIRS analyses during the July 2015 Cassini flyby, and the saturation vapor pressures for HC₃N, C₂N₂, HCN, C₂H₂, C₂H₆, and C₆H₆ are taken from Fray and Schmitt (2009), while those of C₃H₄ and C₄H₂ come from Lara et al. (1996). The gray color-coded rectangles represent the approximate altitude regions for three of Titan’s observed stratospheric ice clouds: the upper rectangle indicates Titan’s ISS-discovered south polar cloud (West et al. 2016), the middle rectangle depicts the CIRS-discovered High-Altitude South Polar (HASP) cloud, and the lower rectangle shows the CIRS Haystack.

With decreasing altitude, overlapping altitude regions will contain more than one condensing vapor at a time, leading to the onset of co-condensation (see for example Anderson and Samuelson 2011; Anderson et al. 2016, 2018). Co-condensation affects the developing physical structure of the ice particles being produced, and as laboratory experiments indicate (Anderson and Samuelson 2011; Anderson et al. 2016, 2018; Nna-Mvondo et al. 2018), considerably modifies the spectral characteristics from those of layered ices, most notably in the low energy part of the far-IR (see Anderson et al. 2018). Detailed examples for Titan ice analogs containing C₆H₆, HCN, and/or C₂N₂ are given in Anderson et al. (2018), with an emphasis on the extensive modifications seen in the resulting co-condensed ice refractive indices and particle extinction cross sections for wavenumbers lower than ~1000 cm⁻¹ (wavelengths longer than ~10 µm).

Moreover, the resulting saturation vapor pressure for the combination of two or more vapors that undergo simultaneous condensation (i.e., co-condensation) will be different than those of their individual compounds when considered separately. The condensation efficiency of Titan’s stratospheric ices is much more complicated than typically considered,
and is a function of many parameters such as the ice and aerosol sticking coefficients, the binding energies of the different molecules, etc. (see for example Barth 2017, and references therein). Thus, the expected altitude of the co-condensed ice will be different than those of the individual compounds when considered separately. Phase diagrams representing the saturation levels of mixed vapors, rather than solely from their pure counterparts, remain to be determined.

As Titan’s stratospheric ice particles continue to subside, they will eventually cross the tropopause and enter into the troposphere below, where some will disperse to lower latitudes due to atmospheric circulation and overall diffusion, and most will eventually sediment on to Titan’s surface. The exceptions are ethylene (C₂H₄), which suffers from several strong photochemical sinks in Titan’s lower stratosphere, thus reducing its mole fraction enough to inhibit vapor condensation, as well as the compounds CH₄, C₂H₆, and C₃H₈, which will convert from the solid to liquid phase (melt) near and on Titan’s surface. The majority of Titan’s stratospheric organic ices will remain on the surface in the solid phase, and contribute—along with the aerosol—to the surface particulate inventory. Quantitative determinations of the relative abundance of these two organic particulates remain to be determined.

The remainder of this review is devoted to the observed vertical and latitudinal structure of Titan’s stratospheric ice clouds as functions of season, their impact on Titan’s atmosphere, and to their chemical identifications with the aid of laboratory thin ice film transmission spectroscopy. Some consideration of solid-state photochemistry as a stratospheric ice cloud formation mechanism will also be discussed.

2 The Pre-Cassini Era

In 1973, ground-based positive linear polarization white light measurements provided the first observational evidence that Titan had a scattering atmosphere due to clouds and/or aerosol particles (Veverka 1973). Six years later, the Pioneer 11 spacecraft flew by the Saturn system, returning the first high spatial resolution data of Titan’s atmosphere. Although there were no cameras or spectrometers onboard, Pioneer 11 did contain a scanning photopolarimeter with red and blue broad band filters. Strong polarization near 90° phase angle—always unavailable from Earth—suggested that small particulates were suspended in Titan’s atmosphere (Tomasko 1980; Tomasko and Smith 1982). One year later in 1980, Voyager 1 flew by the Saturn system, and using the onboard thermal IR spectrometer (IRIS), the organic molecules CH₄, deuterated methane (CH₃D), C₂H₂, C₂H₄, C₂H₆, C₃H₄, C₃H₈, C₄H₂, C₂N₂, HCN, HC₃N, as well as carbon dioxide (CO₂) and molecular hydrogen (H₂), were all detected in Titan’s atmosphere (Hanel et al. 1981; Coustenis and Bezard 1995; Samuelson et al. 1997; Coustenis et al. 1999). Excluding H₂, which was seen in absorption (residing in the troposphere), all the remaining vapors were observed in emission (residing in the stratosphere), and analyses of these data indicated that most of the vapors were abundant enough to condense and form a profusion of organic ice clouds in Titan’s cold mid to low stratosphere.

At this time, the possibility of organic ice clouds forming in Titan’s stratosphere was unforeseen. The season on Titan was early northern spring, as Titan’s north polar region had just emerged from winter, and it was fortuitous that IRIS was scheduled to perform a mosaic of Titan’s north polar region—a spatially crude limb scan of the stratosphere—at high northern polar latitudes (see Fig. 2). The circles in Fig. 2 represent the 36-different tangent height positions of the IRIS fields-of-view (FOVs), which represents the FOVs footprint size on Titan’s north polar limb. The FOV diameters range from roughly 220 km at the beginning

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Fig. 2  Adapted from Samuelson et al. (1997). Schematic illustrating the Voyager 1 IRIS effective fields-of-view (FOVs) positioning on Titan’s limb during the north polar mosaic observation recorded in November 1980. The black circles represent the 36-different limb positions of the effective IRIS FOV sizes. The 30-minute mosaic began high in Titan’s stratosphere (top most circles) near 75°N when the spacecraft was furthest out from Titan, and then the FOV size steadily diminished, i.e., better spatial resolution on the limb, as the mosaic continued to sample deeper into Titan’s stratosphere, and eventually onto the disk at the end of the mosaic (bottom most circles) near 55°N when the spacecraft was closest to Titan. Background Titan image credit: NASA/JPL-Caltech/Space Science Institute.

of the observation (top most circles) to ~80 km near the end of the 30-minute observation (bottom most circles) (see Samuelson and Mayo 1991; Samuelson et al. 1997; Mayo and Samuelson 2005). The gradation in FOV size represents Voyager 1’s relative distance from Titan—the spacecraft was further out at the beginning of the mosaic and closer to Titan at the end of the mosaic. Figure 3 shows some resulting limb spectra from this north polar limb mosaic at averaged tangent heights of 297 km, 186 km, and 55 km (each spectrum contains three averaged spectra), in which the original tangent height positions have been vertically-corrected by 65 km. Easily discernible are stratospheric emission signatures from the ν6 band of HC3N ice near 506 cm$^{-1}$, the ν8 band of dicyanoacetylene (C4N2) ice at 478 cm$^{-1}$, and a spectrally broad and intense unidentified emission feature—presumed to be an ice cloud—that spectrally peaks at 221 cm$^{-1}$ (see Samuelson et al. 1997; Coustenis et al. 1999; Mayo and Samuelson 2005). Over the last few decades, this latter spectral feature has been called by many names, such as the far-infrared cloud (Jennings et al. 2015), a polar condensate cloud (Jennings et al. 2015), the far-infrared 220 cm$^{-1}$ cloud (Jennings et al. 2012a), the far-infrared haze (Jennings et al. 2012b), Haze B (de Kok et al. 2007), the 200–240 cm$^{-1}$ feature (Samuelson et al. 1997), unidentified feature 2 (Coustenis et al. 1999), condensate X (Flasar and Achterberg 2009), and the Haystack (Anderson et al. 2014, 2018). Hereafter, we refer to Titan’s unidentified 221 cm$^{-1}$ stratospheric particulate feature as the Haystack. Had the IRIS mosaic instead occurred near Titan’s south pole, the stratospheric ice detections in Titan’s north polar region would have gone undiscovered, and the upcoming planning for the Cassini mission may have gone very differently for Titan.

Even though the ν6 band of HC3N ice near 506 cm$^{-1}$ and its respective vapor at 499 cm$^{-1}$ were observed as a single blended emission feature (see middle and lower panels in Fig. 3;
Fig. 3  Voyager 1 IRIS limb spectra recorded during the north polar limb mosaic in November 1980. Each of the three spectra, shown at tangent heights 297 km, 186 km, and 55 km, contain three averaged spectra. The tangents heights were vertically-corrected by 65 km from the original published values (Samuelson 1985, 1992; Coustenis et al. 1999; Mayo and Samuelson 2005) based on the observed continuum spectral dependences. Easily seen are stratospheric emission signatures from the vapors C₄H₂, C₂N₂, C₃H₄, HC₃N, HCN, C₂H₂, C₃H₈, C₂H₆, C₂H₄, CH₃D, and CH₄, as well as the ν₆ band of HC₃N ice near 506 cm⁻¹, the ν₈ band of C₄N₂ ice at 478 cm⁻¹, and a spectrally broad and intense unidentified emission feature that spectrally peaks at 221 cm⁻¹ (see Samuelson et al. 1997; Coustenis et al. 1999; Mayo and Samuelson 2005). The latter spectral feature is called the Haystack (Anderson et al. 2014, 2018).

Samuelson 1985, 1992; Coustenis et al. 1999), a consequence of IRIS’ fixed 4.3 cm⁻¹ spectral resolution, the ice abundance relative to the gas was still observed to be more abundant in Titan’s north polar lower stratosphere than at higher stratospheric altitudes (Samuelson 1985, 1992). This observed behavior in ice abundance is consistent with vapor condensation formation processes, in which HC₃N vapor forms higher up in Titan’s atmosphere, cools as it diffuses/subsides downwards, and then condenses into ice cloud particles in Titan’s cold, lower stratosphere.

IRIS also discovered the Haystack (Fig. 3), and observed it to behave similarly to that of HC₃N ice, appearing more abundant in Titan’s lower stratosphere compared to those at higher stratospheric altitudes. The Haystack was presumed to be a stratospheric ice cloud due to 1) its spectrally broad and intense emission feature spanning wavenumbers 200 to 250 cm⁻¹, with a 221 cm⁻¹ spectral peak, 2) its limited vertical dependence in Titan’s north polar lower stratosphere, and 3) its restriction to latitudes poleward of ~50°N (see Samuelson 1985; Coustenis et al. 1999). Initial analyses of these data surmised the chemical composition of the Haystack was due to the 45-µm transverse optic vibration mode (Erickson et al. 1981) of water ice (Samuelson 1985; Coustenis et al. 1999). However, the water ice identification was eventually rejected given the amount of stratospheric water vapor calculated was three orders of magnitude below that of the IRIS-determined water ice upper limit (Yung et al. 1984; Samuelson 1985), as well as the obvious spectral mismatch between water ice and the Haystack (Samuelson 1985; Coustenis et al. 1999; Samuelson et al. 2007, see also Fig. 16). Subsequent laboratory thin ice film transmittance spectra were used to tentatively identify the chemical composition of the Haystack as crystalline propionitrile (C₂H₅CN).
(Khanna 2005). However, as discussed in Sect. 3.2.3 with the aid of Cassini CIRS spectra, pure C$_2$H$_5$CN ice was easily discounted as a likely Haystack candidate (see Fig. 16 and also Samuelson et al. 2007; Nna-Mvondo et al. 2018).

The $\nu_8$ band of C$_4$N$_2$ ice at 478 cm$^{-1}$ was also observed by Voyager 1 IRIS in Titan’s north polar lower stratosphere (Khanna et al. 1987; Samuelson 1992; Samuelson et al. 1997; Coustenis et al. 1999); see middle and lower panels in Fig. 3. However, unlike with the observed behavior of HC$_3$N ice and its corresponding vapor, there was no observational evidence of C$_4$N$_2$ vapor in Titan’s stratosphere. This was puzzling since the C$_4$N$_2$ vapor emission feature at 471 cm$^{-1}$ would be easily detectable if C$_4$N$_2$ ice was condensing from the vapor to form the IRIS-observed ice clouds. The absent C$_4$N$_2$ vapor emission feature at 471 cm$^{-1}$ led to a vapor mixing ratio upper limit calculation of $4 \times 10^{-10}$, and the C$_4$N$_2$ ice abundance was determined to be $\sim 100$ times greater than if the ice had condensed out from its uniformly-mixed vapor (Samuelson et al. 1997). C$_4$N$_2$ vapor is therefore not in equilibrium with the ice under the assumption that C$_4$N$_2$ ice clouds form as a result of vapor condensation, since both vapor and ice must be in equilibrium to maintain a steady-state. Here we define steady-state to only include the contribution from condensation in the absence of kinetics, since the latter is most likely a minor contributor to the steady-state of the stratospheric ice clouds in Titan’s dense atmosphere. Photolytic decomposition of C$_4$N$_2$ vapor was later proposed to satisfy vapor-ice equilibrium (Samuelson et al. 1997). However, as discussed in Sect. 3.2.1, an alternate formation mechanism for C$_4$N$_2$ ice clouds was invoked 30-years later from analyses of CIRS far-IR spectra to explain this observed vapor-ice conundrum, while at the same time satisfying vapor-ice equilibrium (Anderson et al. 2016).

Continued analyses of Voyager 1 IRIS north polar limb far-IR spectra revealed the presence of a rather tenuous and uniformly-distributed ice cloud residing in Titan’s lower stratosphere at altitudes between 58 and 90 km (Mayo and Samuelson 2005). These ice particles contributed to Titan’s scattering continuum between 250–560 cm$^{-1}$, with calculated effective particle radii between 1 and 5 µm. Based on Mie calculation fits, the most likely candidates proposed for the chemical compositions were HCN ice and C$_2$H$_6$ ice, near the upper and lower boundaries of the cloud, respectively (Mayo and Samuelson 2005). However, from the spectral continuum analyses, no direct identifications of the chemical compositions were possible.

Lastly, an unidentified emission feature at 700 cm$^{-1}$ was also observed—presumed to be an ice cloud—in Titan’s northern polar stratosphere (see upper and middle panels in Fig. 3). C$_3$H$_4$ ice has been the only candidate proposed (Samuelson 1992; Coustenis et al. 1999). This feature was also observed 25 years later by CIRS in Titan’s northern winter polar stratosphere (see Fig. 9); however, a definitive identification of this stratospheric ice feature still remains to be determined.

### 3 The Cassini Era

Nearly 25 years after the Voyager 1 flyby of Titan, the Cassini spacecraft entered into Saturn’s orbit and began its remarkable 13-year journey through the Saturn system. During this time, Titan transitioned from northern to southern winter, and numerous stratospheric ice clouds comprising different chemical compositions were observed by three of Cassini’s remote sensing instruments: the Visible and Infrared Mapping Spectrometer (VIMS), the Imaging Science Subsystem (ISS), and CIRS.
3.1 VIMS and ISS Observed Stratospheric Ice Clouds

Early on in the mission, VIMS detected diffuse and homogeneously-distributed ice clouds in Titan’s north polar winter stratosphere, residing mostly at low stratospheric altitudes, spanning latitudes 51°N to 68°N (Griffith et al. 2006). Initially, between latitudes 48°N and 55°N, radiative transfer analyses were applied and clouds were inferred to reside between 30 and 60 km, with effective particle radii no larger than ~3 µm (Griffith et al. 2006). The low altitudes ruled out clouds comprising chemical compositions other than C2H6 or CH4 ice. Methane clouds, however, were ruled out due to the small particle size, the long-lasting and homogeneous cloud characteristics, along with a lack of opacity at 5 µm (see Fig. 4).

Subsequent VIMS observations of Titan’s northern winter polar cloud revealed the emergence of a second cloud morphology with a heterogeneous horizontal structure poleward of ~64°N; this was in addition to the homogeneously-distributed C2H6 ice clouds that appeared to diffusely extend to higher northern winter polar latitudes (see Fig. 4; Rannou et al. 2012; Le Mouélic et al. 2012). Concurrent CIRS observations, combined with Radio Science Subsystem (RSS) measurements, revealed Titan’s tropopause temperatures at these high northern winter polar latitudes had cooled by several degrees below 70 K, suggesting CH4 ice as the chemical composition of this second type of cloud morphology (Anderson et al. 2014). This second type of cloud was reported by Anderson et al. (2014) to form as a result of strong subsidence, and not convection, and were therefore named Titan’s Subsidence-Induced Methane Clouds (SIMCs) (Anderson et al. 2014). Titan’s SIMCs condense directly from the vapor to solid phase via subsidence, and at high northern winter polar latitudes in the very low stratosphere, these clouds were expected to form near 50 km at 85°N (in March 2005) and ~48 km at 74°N (in May 2007).

A more recent VIMS study detailed the time evolution of this northern winter polar cloud, and reported it to spatially cover the north polar region beginning at the onset of the mission in 2004 through spring equinox in 2009 (Le Mouélic et al. 2012, 2018). The investigators created VIMS mosaics orthographically re-projected over Titan’s north pole, using an orange-reddish color to enhance the 4.78 µm spectral feature—this is most likely the ν1 band of HCN ice (see Fig. 4 in Anderson et al. 2018). They also report the spectral feature at
3.21 µm, which is most likely the $\nu_3$ band of HCN ice (see Anderson et al. 2018). However, Le Mouélic et al. (2018) do not distinguish between the north polar cloud’s differing cloud morphologies or their chemical compositions as discussed above; rather, focus remains on the 3.21 and 4.78 µm spectral features. The core of the cloudy structure is reported to begin thinning (becomes less opaque) between December 2007 and November 2008, with significant cloud dissipation most notable in March 2009. Finally, in June 2010, a small cloud residual was observed above Titan’s fully illuminated north pole. This study did not provide any altitude information on this cloud system spanning the 2004 to 2009 time period since there was no radiative transfer modeling performed in this study.

Later on in Titan’s mid southern fall season, the ISS instrument observed the rapid onset of a chemically-unknown particulate feature appearing at altitudes slightly below Titan’s south polar stratopause (altitude location shown in Fig. 1; image shown in Fig. 5). This newly formed particulate feature was nominally observed by ISS at an altitude of 300 ± 10 km in May 2012, with its circulation pole located at 85.5°S, 263.7°W (West et al. 2016). Over the years, this upper stratospheric ice cloud feature has been referred to by many different names, such as the high-altitude southern polar cloud (de Kok et al. 2014), the south polar cloud (West et al. 2016), the winter polar vortex (Teanby et al. 2017), the vortex cloud (Jennings et al. 2017), the HCN ice cloud at Titan’s south pole (Hörst 2017), and the southern polar vortex (Le Mouélic et al. 2018). In an effort to distinguish this distinct cloud from the many other observed stratospheric ice clouds, most notably those observed and discovered by CIRS, we hereafter call this cloud the ISS-discovered south polar cloud.

Despite the unusually high formation altitude of the ISS-discovered south polar cloud—slightly below Titan’s stratopause—that was observed at altitudes typically too warm for ice cloud formation via vapor condensation (i.e., vapors are sub-saturated at these high altitudes), the study by West et al. (2016) suggested that the unusual morphology and texture of this cloud seen in the ISS images was indeed characteristic of vapor condensation formation processes. ISS time monitoring of this cloud revealed its formation to have occurred as
early as late 2011 (roughly two years post northern spring equinox), and visible in the ISS images up until late 2014 (West et al. 2016), depicting a rather long-lived and dynamically stable cloud—this implies relatively small ice particle radii. A recent VIMS study, however, monitored the time evolution of the ISS-discovered south polar cloud, revealing a significant increase in its spatial extent between 2012 and end of mission in 2017 (Le Mouélic et al. 2018). Specifically, the investigators report a spatial extension from 85° S down to 75° S in 2013, then down to 68° S in 2014, then further extending down to 60° S in mid-2016, and finally expanding down to ~58° S in April 2017. The complete south polar spatial extent still remains unknown since the south polar region became more engulfed in darkness as end of mission approached. However, the reported spatial expansion of the ISS-discovered south polar cloud is consistent with the horizontal growth of Titan’s south polar vortex, reported to steadily increase in spatial extent from 90° S to 75° S in 2012 then down to 60° S in 2016 (Teanby et al. 2017).

The first identification of at least one of the chemical compounds contained within the ISS-discovered south polar cloud was reported by de Kok et al. (2014). These investigators analyzed a VIMS image cube acquired in June 2012 and identified the spectral feature at 3.21 µm to be that of HCN ice, with a derived ice cloud altitude of 300 ± 70 km, somewhat similar to the ISS-determined altitude location from May 2012 (West et al. 2016). Continued VIMS analyses between 2012 and 2017 identified both the HCN ice spectral signatures at 3.21 µm and 4.78 µm, and noted that these HCN ice spectral features were observed in most of the VIMS spectra spanning the 2012–2017 time period. This study, however, provided no information on the time-varying altitude locations of the ISS-discovered south polar cloud since the investigators did not perform any radiative transfer analyses. Thus, detailed analyses on the vertical extent of the time-varying ISS-discovered south polar cloud, its full chemical inventory, as well as its formation mechanism still remain to be determined.

### 3.2 CIRS-Observed Stratospheric Ice Clouds

Over the course of the Cassini mission, CIRS observed numerous ice clouds in Titan’s stratosphere with varying chemical compositions such as HC₃N ice (Sect. 3.2.1, Anderson et al. 2010), C₄N₂ ice (Sect. 3.2.1, Anderson et al. 2016), co-condensed HC₃N:HCN ice (Sect. 3.2.2, Anderson and Samuelson 2011), C₆H₆ ice (Sect. 3.2.5, Vinatier et al. 2018), co-condensed C₆H₆:HCN ice (Sect. 3.2.4, Anderson et al. 2017), and the chemically-unidentified Haystack (Sect. 3.2.3, Samuelson et al. 2007, de Kok et al. 2007, Jennings et al. 2012a,b, 2015, Anderson et al. 2014, 2018). Most of these detections resulted from CIRS’ expanded spectral range and higher spectral resolution over that of its predecessor IRIS, especially in the low energy part of the far-IR spectral region. IRIS had one interferometer, with the spectral range covering 200 to 1500 cm⁻¹ (50 to 6.67 µm), and a fixed spectral resolution (Δν) of 4.3 cm⁻¹. CIRS, on the other hand, was comprised of two interferometers, had broader spectral coverage (10–1500 cm⁻¹; 1000–6.67 µm), with variable spectral resolutions, although in practice, Titan spectra were recorded at Δν = 14.1 cm⁻¹, 2.56 cm⁻¹, and 0.48 cm⁻¹ (all apodised FWHM values). One interferometer contained the mid-IR focal planes FP3 and FP4, spectrally covering 600–1100 cm⁻¹ (16.67–9.09 µm) and 1100–1500 cm⁻¹ (9.09–6.67 µm), respectively, while the second interferometer consisted of a single-element far-IR focal plane sensitive to the spectral range 10 to 600 cm⁻¹ (1000 to 16.67 µm). The FP1 far-IR FOV was circular, with decreasing sensitivity from the center (at 0 mrad) to the edge, 3.2 mrad from the center, where the sensitivity dropped to zero (see Flasar et al. 2005; Anderson and Samuelson 2011). It is necessary to take the spatial FOV characteristics
Fig. 6 Conceptual CIRS far-IR limb scan of Titan’s north polar region. The black circles represent the effective fields-of-view (FOVs) sizes of the CIRS far-IR detector (FP1). Altitude separation between contiguous FOVs is not drawn to scale. Typical CIRS limb scan durations were $\sim 30$ minutes, recording $\sim 300$ individual spectra at varying tangent heights, beginning near 350 km (top most circles) and then steadily sampling deeper into Titan’s stratosphere, ending at tangent heights near $-150$ km (below the surface). There is a 98%–to–99% overlap of contiguous FOVs, enabling sub-FOV-diameter spatial resolution. Background Titan image credit: NASA/JPL-Caltech/Space Science Institute

into account wherever data from the limb of Titan are being analyzed since spatial deconvolution in any radiative transfer model is essential. For a review of the CIRS instrument, see Flasar et al. (2005) and Jennings et al. (2017).

The vertical distributions and spectral dependences of the majority of the CIRS-observed stratospheric ice clouds were determined from analyses of CIRS far IR-targeted low spectral resolution limb scans. The black circles in Fig. 6 represent the placement of the CIRS far-IR FOV on Titan’s limb during such a limb scan, and although not drawn to scale, are essentially contiguous footprints on Titan’s limb that overlap the FOV by 98% to 99% at each tangent height. Each CIRS far-IR limb scan targeted $\sim 300$ individual tangent heights, beginning near Titan’s lower mesosphere ($\sim 350$ km) and then steadily sampling deeper into Titan’s stratosphere, ending at tangent heights near $-150$ km, which is below Titan’s solid surface horizon. The FOV diameter ranged from roughly 220 km at the beginning of the observations to $\sim 100$ km near the end of the 30-minute limb scan (see Anderson and Samuelson 2011). The significant overlap of contiguous FOVs enables a much narrower vertical resolution ($\sim 30$ km) than FOV size would otherwise imply, and is optimized in the 100 to 250 cm$^{-1}$ spectral region where the CIRS far-IR interferometer has the highest signal-to-noise ratio. Figure 7 gives a limb scan example at 15°S at 160 cm$^{-1}$ acquired during northern winter in September 2006 and Fig. 8 gives a limb scan example at 75°S at 215 cm$^{-1}$ acquired nine years later in late southern fall in July 2015.

The detailed procedure for determining sub-FOV-diameter spatial resolution is described in Anderson and Samuelson (2011). The vertical distributions for Titan’s aerosol over three-to-four contiguous altitude regions (between 0 and 500 km), are each represented by ana-
lytic solutions. Eventually the stratospheric ices are also included as an opacity source. The number of free parameters included in the separate analytic expressions for the different opacities are kept to a minimum. To determine the numerical values for the free parameters, the Levenberg-Marquardt nonlinear least squares fitting procedure is utilized. Anderson and Samuelson (2011) have empirically determined that the wavenumbers spanning $\sim$100 to 250 cm$^{-1}$ have sufficient signal-to-noise ratios to determine accurate solutions. One reason the deconvolution of the intensity function (the average intensity over the FOV) works so effectively is because the 98–99% overlap in the CIRS FP1 FOVs during the far-IR limb scans essentially yields the derivative of the intensity function as well as the intensity function itself. This is true provided there is an adequate signal-to-noise ratio, which has been found in practice to occur at wavenumbers between $\sim$100 and 250 cm$^{-1}$. For a detailed account of how the aerosol and stratospheric ice cloud vertical distributions are determined from CIRS far-IR limb scans, see Anderson and Samuelson (2011).

The extension of the CIRS far-IR spectral range down to 10 cm$^{-1}$ (1000 µm; IRIS cut-off at 200 cm$^{-1}$ [50 µm]) opened up a previously unexplored far-IR spectral region containing a wealth of nitrile ice signatures, many of which are expected to comprise the chemical compositions of Titan’s stratospheric ice clouds. The top panel in Fig. 9 shows a CIRS composite limb spectrum of Titan, with the FP1 FOV centered at a tangent height of $\sim$125 km (mid-IR spectra are averaged over 100 to 150 km), recorded near 65°N during northern winter in 2006. In the bottom panel, the wavenumber dependence of the Planck intensity curve computed at 130 K is shown, which is comparable to the condensation temperatures in Titan’s mid-to-low stratosphere. The Planck intensity is maximized in the low energy part of the far-IR at wavenumbers surrounding 200 cm$^{-1}$, coinciding with the spectral location for many nitrile ice cloud signatures (some hydrocarbons too) that reside at altitudes in Titan’s mid to low stratosphere, and therefore contribute to the local environment there. On the other hand, the Planck intensities for wavenumbers higher than $\sim$700 cm$^{-1}$ are much reduced at 130 K, indicating that these ice signatures will be much more difficult to detect in Titan’s mid to low stratosphere. The strong vapor emission features seen in the upper panel for
Fig. 8 CIRS far-IR limb-scan tangent height intensity profile at 215 cm\(^{-1}\) during late southern fall in July 2015. Filled black hexagons represent the CIRS limb scan at 215 cm\(^{-1}\), with each symbol representing a tangent height. Indicated in the figure are the altitude locations and intensity contributions from the HASP cloud and the Haystack. The gray curve shows a radiative transfer model fit that does not restrict the Haystack with latitude, resulting in a very poor fit. On the other hand, the orange curve represents a radiative transfer model that restricts the Haystack to latitudes poleward of \(\sim 70^\circ\)S, resulting in a very satisfactory fit. The Haystack, therefore, appears to be restricted to Titan’s south polar high latitudes prior to the onset of Titan’s southern winter season.

Wavenumbers higher than \(\sim 700 \text{ cm}^{-1}\) simply means that these vapors are originating and contributing to Titan’s atmosphere at altitudes higher in the stratosphere than those of the ices. Thus, the low energy part of the far-IR is an ideal spectral location to detect Titan’s mid to lower stratospheric ice clouds due to the combination of the Planck intensity maximizing, the spectral location of many low energy nitrile ice signatures, the CIRS high signal-to-noise ratio in this part of the spectrum, and the increase in stratospheric ice signal due to the observing geometry of the CIRS far-IR limb scans, \textit{i.e.}, integrations along long path lengths of atmosphere. In order to determine the various ice cloud chemical compositions in Titan’s stratosphere, as well as their formation mechanisms, the vertical distributions and spectral dependences of Titan’s retrieved stratospheric particulate opacities must be combined with subsequent laboratory thin ice film transmission spectroscopy experiments. The remainder of this section details the various CIRS-observed stratospheric ice clouds.

3.2.1 \textit{HC}_3\textit{N} and \textit{C}_4\textit{N}_2 Ice Clouds

CIRS observed the \(v_6\) band of \textit{HC}_3\textit{N} ice at 506 cm\(^{-1}\) in Titan’s lower stratosphere during three different seasons: northern winter, early northern spring, and late southern fall (Anderson et al. 2010, 2016). The \(v_6\) band of \textit{HC}_3\textit{N} ice at 506 cm\(^{-1}\) was easily separated from its corresponding gas at 499 cm\(^{-1}\) (unlike with IRIS) due to CIRS’ higher spectral resolution. The upper left quadrant in Fig. 10 shows the first detection of the \(v_6\) band of \textit{HC}_3\textit{N} ice at 62°N during northern winter in July 2006, with the FP1 FOV centered on a tangent height of 125 km (Anderson et al. 2010). The investigators (Anderson et al. 2010) reported a mean particle radius of 3.0 \(\pm\) 0.4 \(\mu\)m, an ice column abundance of \(3.1 \times 10^{14}\) molecules cm\(^{-2}\)
Fig. 9 Upper curve: CIRS far- and mid-IR limb spectra plotted as functions of wavenumber, recorded near 65°N during mid northern winter (July 2006) and averaged over tangent heights covering 100 to 150 km. Stratospheric organic vapors are labeled, as is the ν6 band of HC3N ice at 506 cm\(^{-1}\), an unidentified ice feature at 700 cm\(^{-1}\), the unidentified broad ice emission feature centered at 221 cm\(^{-1}\) (the Haystack), and the broad ice emission feature that peaks at 160 cm\(^{-1}\), with its contribution to the continuum indicated by the light gray shaded region. Without the 160 cm\(^{-1}\) ice feature, the continuum would reduce to the dashed curve level. Lower curves: spectral dependences of laboratory-measured absorbances or imaginary part of the refractive index for various compounds that may contribute to ice clouds observed in Titan’s stratosphere. The ices C2N2, HCN, and co-condensed C6H6:HCN (39% HCN, 61% C6H6) are taken from (Anderson et al. 2018), C2H5CN from Nna-Mvondo et al. (2018), H2O ice is taken from Hudson and Moore (1995), HC3N ice comes from Moore et al. (2010), and C4N2 comes from Dello Russo and Khanna (1996). The Planck intensity curve computed at 130 K is superimposed (thin black curve), which is representative of the lower stratospheric temperatures at which ices are forming (abundance value revised from Anderson et al. 2010), and a cloud top near 150 km. The upper right quadrant in Fig. 10 shows the second observation of HC3N ice at 70°N in August 2007. This observation, however, recorded the ice emission features at two different tangent heights, with the FP1 FOV centered on 140 and 160 km, thus allowing for a vertical distribution determination of the HC3N ice cloud (Anderson et al. 2010). A mean particle radius of 2.23 ± 0.42 μm and a corresponding column abundance of 2.7 × 10\(^{14}\) molecules cm\(^{-2}\) (abundance value revised from Anderson et al. 2010) were determined, with the cloud top residing at ∼165 km. The best fit radiative transfer model found an HC3N ice cloud thickness between ∼10 and ∼20 km. The limiting factors on cloud thickness and ice particle size are vapor abundance and large particle precipitation near the cloud bottom. The cloud thickness is consistent with calculations showing that about half of the HC3N vapor is removed from the altitude where the cloud begins to condense (cloud top) at ∼4 km below the cloud top altitude. Moreover, the small HC3N ice particle sizes are indicative of long-lived and dynamically-stable stratospheric ice clouds; large ice particles will precipitate out in relatively short times compared to that of the stable HC3N ice clouds. The observed abundance...
Fig. 10  CIRS limb spectra showing stratospheric HC$_3$N ice clouds, along with other stratospheric particulates, at three different Titan seasons. The upper left quadrant shows an average of 24 spectra, recorded at $62^\circ$N during northern winter (2006), with the FP1 FOV centered at 125 km. Easily discernible are stratospheric particulate emission features from the 160 cm$^{-1}$ ice cloud (Sect. 3.2.2), the Haystack (Sect. 3.2.3), a tentative detection of C$_4$N$_2$ ice, and the first detection of HC$_3$N ice (Anderson et al. 2010, 2014; Anderson and Samuelson 2011). The upper right quadrant shows an average of 25 spectra, recorded at $70^\circ$N during northern winter (2007), with the FP1 FOV centered at $\sim 150$ km. The same stratospheric ice features are evident as those at $62^\circ$N (Anderson et al. 2010). The lower left quadrant shows an average of 254 spectra, acquired at $\Delta \nu = 2.56$ cm$^{-1}$, recorded at $70^\circ$N in early northern spring (2010), with averages over the FP1 FOVs centered on 100 km, 150 km, and 200 km. Evident are stratospheric ice emission features from the 160 cm$^{-1}$ ice cloud, the Haystack, HC$_3$N ice, albeit significantly reduced in abundance, and the first definitive detection of C$_4$N$_2$ ice (Anderson et al. 2016). The lower right quadrant shows an average of 35 spectra, recorded at $79^\circ$S during late southern fall (2015), with the FP1 FOV centered on 125 km. At this time, there is no evidence of the 160 cm$^{-1}$ ice cloud, and instead, the emergence of the HASP cloud is visible (Sect. 3.2.4), appearing at this tangent height as a broad shoulder to the left of the very strong Haystack. There is no trace of C$_4$N$_2$ ice, although the beginning of HC$_3$N ice cloud formation is observed (Sect. 3.2.1). The 2006, 2007, and 2015 spectra were recorded at $\Delta \nu = 0.48$ cm$^{-1}$, and shown here convolved with a gaussian kernel to $\Delta \nu = 2.56$ cm$^{-1}$.

of HC$_3$N vapor at the cloud top, compared to that of the HC$_3$N ice abundance, is consistent with vapor condensation as the formation mechanism.

Following the first two HC$_3$N ice observations, a customized CIRS limb integration was designed to target a more precise vertical distribution and abundance determination of Titan’s HC$_3$N ice clouds. In early northern spring (April 2010), CIRS recorded Titan limb spectra at $70^\circ$N at a spectral resolution of 2.56 cm$^{-1}$, with the FP1 FOVs centered on tangent heights of 100 km, 150 km, and 200 km (Anderson et al. 2016). Each limb integration contained $\sim 84$ spectra, shown as a weighted average in the lower left quadrant in Fig. 10. However, the strength of the HC$_3$N ice emission feature at 506 cm$^{-1}$ was significantly reduced compared to that observed during northern winter, and instead, the $\nu_8$ band of C$_4$N$_2$...
Fig. 11 Schematic illustrating an alternate C$_4$N$_2$ ice formation mechanism to that of vapor condensation. In this formation mechanism, photolysis within extant HCN and HC$_3$N ice particles has been proposed to form C$_4$N$_2$ ice—via solid-state chemistry (Anderson et al. 2016). In this case, only $\sim$1% of C$_4$N$_2$ ice molecules will reside on the particle surface (in other words, unobservable), instead of 100% required by vapor condensation, thus satisfying vapor-ice equilibrium. For more details, see Anderson et al. (2016). Image credit: Jay Friedlander

ice at 478 cm$^{-1}$ had emerged. Since both the C$_4$N$_2$ vapor emission features at 471 cm$^{-1}$ and 107 cm$^{-1}$ were absent, and with the rather large abundance of C$_4$N$_2$ ice observed, it was concluded that C$_4$N$_2$ vapor must be highly sub-saturated and therefore not in equilibrium with the pure ice—this unequivocally rules out vapor condensation as the formation mechanism. This observation was also the first indication that C$_4$N$_2$ ice formation may result from depletion of the HC$_3$N ice inventory, shedding some light on the formation mechanism of Titan’s C$_4$N$_2$ stratospheric ice.

With C$_4$N$_2$ vapor observed by CIRS to be absent in Titan’s stratosphere (this was also confirmed by Jolly et al. 2015), Anderson et al. (2016) postulated that solid-state photochemistry provides an alternate mechanism besides that of vapor condensation for producing Titan’s observed C$_4$N$_2$ stratospheric ice clouds. In this scenario, HC$_3$N vapor subsides to condense at the initial cloud top. HCN vapor, being more than two orders of magnitude more abundant than HC$_3$N vapor (Coustenis et al. 2007, 2016), is also subsiding and soon afterwards, both HC$_3$N and HCN vapors enter into altitude regions where they undergo simultaneous saturation, i.e., co-condensation. Once the two ices form a co-condensate, the individual molecules are very nearly in contact, so photochemistry is readily induced between the two molecules (see schematic in Fig. 11). On the other hand, if you only have a layered ice, then the HCN and HC$_3$N particles are close together only at their common boundary, so photochemistry is unable to promote interactions between the two ice layers very efficiently.

As detailed in Anderson et al. (2016), the UV irradiation initiates the solid-state photochemical reaction HCN + HC$_3$N $\rightarrow$ C$_4$N$_2$ + H$_2$ within extant HC$_3$N:HCN co-condensed ice particles (see Fig. 11 and also Anderson et al. 2016, 2018). In order to inhibit diffusion of C$_4$N$_2$ molecules through the ice particle, the formation of an HCN ice barrier between the surface and interior of the particle is needed to keep the C$_4$N$_2$ ice in the interior, separate from the C$_4$N$_2$ vapor outside the particle. This is needed to maintain a low C$_4$N$_2$ vapor pressure in the presence of a relatively high C$_4$N$_2$ ice abundance buried deeper toward the center of the particle. Once all HC$_3$N ice is photolytically destroyed, the photochemical reaction will terminate. UV-irradiation studies of extant HCN:HC$_3$N co-condensed ices are still needed to confirm this formation mechanism.

Such a solid-state photochemical reaction has similarities to the terrestrial reaction HCl + ClON$_2$ $\rightarrow$ HNO$_3$ + Cl$_2$, which is known to produce HNO$_3$ coatings on water ice particles in Earth’s stratosphere, a byproduct of the catalytic chlorine chemistry that produces
ozone holes in Earth’s polar stratosphere (see for example Molina et al. 1987; Solomon 1999). The cold temperatures in Titan’s lower stratosphere combined with the associated strong circumpolar winds that isolate polar air act in much the same way as on Earth, giving rise to compositional anomalies and stratospheric condensate clouds that are the sites of heterogeneous chemistry. While there are differences in detail—the terrestrial case entails oxygen and halogen compounds, whereas Titan’s chemistry involves organics—the overall picture is remarkably similar on the two worlds.

The final CIRS observation revealing the $\nu_6$ band of HC$_3$N ice in Titan’s stratosphere was discovered at 79$^\circ$S in late southern fall (July 2015), following the observed reversal of the meridional circulation pattern (see for example Teanby et al. 2012, 2017). The lower right quadrant in Fig. 10 shows the resulting limb integration spectrum, with the CIRS far-IR FOV centered on a tangent height of 125 km. This limb integration shows the emergence of stratospheric HC$_3$N ice as Titan’s south polar region was heading into winter; particle size and abundance are currently being determined.

3.2.2 The 160 cm$^{-1}$ Ice Clouds

The 160 cm$^{-1}$ ice cloud is the name given to a spectrally broad ($\sim$100–300 cm$^{-1}$), quasi-continuum ice emission feature, spectrally peaking at 160 cm$^{-1}$, located in Titan’s northern winter lower stratosphere (Anderson and Samuelson 2011; Anderson et al. 2014, 2018, see Figs. 9, 10, 12). Its significant contribution to Titan’s continuum is shown by the gray shaded region in the upper panel in Fig. 9 and also evident in Fig. 10. This ice cloud was discovered using the CIRS low spectral resolution far-IR limb scans (Fig. 7; Anderson and Samuelson 2011), which enable separation of Titan’s vertically-restricted stratospheric ice clouds from the surrounding ubiquitous aerosol continuum. As discussed in Sect. 3.2, the significant overlap in FOVs during a CIRS far-IR limb scan allows for not only the average intensity over each FOV but also the derivative of these intensities (at wavenumbers where there is sufficiently high signal-to-noise ratios), enabling a sub-FOV-diameter vertical resolution. Figure 7 gives an example of such a limb scan at 160 cm$^{-1}$, recorded during the northern winter season in September 2006 at 15$^\circ$S. Specifically, the figure shows the tangent height-intensity profile containing $\sim$300 individual limb spectra (solid black hexagons), with the FOV center—for each tangent height observation—separated by a few km at most. It became clear very early in the analysis that the limb scans spanning wavenumbers $\sim$100–$\sim$300 cm$^{-1}$ required an additional, vertically-limited, opacity source to that of the ubiquitous aerosol in order to better fit the CIRS limb scan observations. The gray curve in Fig. 7 shows the best fit radiative transfer model at 160 cm$^{-1}$ consisting of two vertically diffuse aerosols with different chemical compositions. It is clear that the fit is rather poor, and instead, when one aerosol and one vertically restricted stratospheric ice cloud—the latter residing at altitudes between $\sim$60 and 90 km in the lower stratosphere—are included, the fit dramatically improves, as shown by the orange curve (Anderson and Samuelson 2011). The 160 cm$^{-1}$ ice cloud was found by Anderson and Samuelson (2011) to be a geometrically thin and effectively continuous sheet of ice cloud that tends to increase monotonically from roughly 60$^\circ$S to high northern polar latitudes during northern winter. The 160 cm$^{-1}$ ice cloud’s derived spectral dependence at 15$^\circ$S is shown by the black curve in Fig. 12. From the radiative transfer analysis along with predicted cloud top locations for the various organic vapors in Titan’s stratosphere, this cloud was expected to contain both HCN and HC$_3$N ice due to its altitude location in Titan’s lower stratosphere.

The 160 cm$^{-1}$ ice emission feature is located in a spectral region that contains overlapping, low-energy lattice vibration features (librations) of multiple nitrile ices (see bottom
Fig. 12 Modified from Anderson and Samuelson (2011) and Anderson et al. (2018). Black curve represents the CIRS-derived spectral dependence of Titan’s 160 cm\(^{-1}\) ice cloud discovered at lower stratospheric altitudes (~60 to 90 km) during northern winter at 15°S. The green curve shows the weighted sum of individual HCN and HC\(_3\)N ices from thin ice film transmission spectroscopy (vapors deposited at 30 K; see also Moore et al. 2010). As discussed in Anderson et al. (2018), in the low-energy part of the far-IR, the sum of the individual ices is nearly identical to layered ice experiments. The orange curve depicts a co-condensed thin ice film containing mixed HC\(_3\)N and HCN, demonstrating that co-deposition (i.e., co-condensation) experiments better reproduce the CIRS observations.

panel in Fig. 9 and also Fig. 16), and as laboratory experiments indicate (Anderson and Samuelson 2011; Anderson et al. 2018), is consistent with an ice mixture dominated by HCN and HC\(_3\)N ices formed via co-condensation processes in Titan’s stratosphere (Anderson and Samuelson 2011; Anderson et al. 2014, 2018). Thin ice film transmission spectroscopy reveals that layered HC\(_3\)N and HCN ices do not reproduce the 160 cm\(^{-1}\) ice cloud feature; rather, the ice cloud is consistent with ice particles formed via co-condensed HC\(_3\)N:HCN (see Fig. 12; Anderson and Samuelson 2011, Anderson et al. 2018), resulting from combinations of the \(\nu_7\) band and lattice modes of HC\(_3\)N and the libration mode of HCN (Anderson and Samuelson 2011; Anderson et al. 2018).

3.2.3 The Haystack

As discussed in Sect. 2, the Haystack is a name given to one of Titan’s long-lived stratospheric ice clouds with an unknown chemical composition (Anderson et al. 2014, 2018). The Haystack is easily identifiable due to its broad emission feature spectrally peaking at 221 cm\(^{-1}\), and spanning wavenumbers ~200–250 cm\(^{-1}\) (see Figs. 9, 13, 14, 15). The Haystack’s spectral signature is quite strong, suggesting that whatever it is made out of is very abundant. The large spectral extent and its spectral location in the far-IR further suggests that the Haystack contains more than one chemical compound. Some chemical compounds that have been proposed as the identity of the Haystack are water ice (Samuelson 1985; Coustenis et al. 1999), crystalline C\(_2\)H\(_5\)CN (Khanna 2005), predominantly pure HCN ice (de Kok et al. 2008), and complex nitriles that have –C–CN bending vibrations (Jennings et al. 2012a,b).
Fig. 13  CIRS far-IR limb integrations spanning 30°N to 85°N during northern winter, all acquired at tangent heights of ~125 km. The 85°N, 62°N, 55°N, and 30°N spectra were recorded in early 2005, mid 2006, early 2006, and late 2006, respectively. The spectra at 55°N, 62°N, and 85°N were vertically offset from the 30°N spectrum. With the exception at 30°N, the Haystack is evident at the remaining latitudes, with its strongest signature near the north pole.

As discussed above in Sect. 2, the Haystack was first observed by Voyager 1 IRIS in Titan’s early northern spring stratosphere (November 1980) at latitudes poleward of ~50°N (Samuelson 1985; Coustenis et al. 1999) (Fig. 3). Later, with Cassini CIRS, the Haystack was first observed at the beginning of the mission in 2004 (mid northern winter), and then continuously observed at high northern polar latitudes throughout northern winter (see Fig. 13), and well into northern spring (Samuelson et al. 2007; de Kok et al. 2007; Jennings et al. 2012b; Anderson et al. 2014). As with IRIS, the Haystack appeared to be restricted to latitudes poleward of ~50°N during northern winter. A few years later during mid southern fall, the Haystack made its debut at high southern polar latitudes (Figs. 10, 14, 15; Jennings et al. 2012a, 2015), and was observable in CIRS far-IR disk-viewing spectra up until end of mission (data were acquired through May 2017).

Figure 16 shows a CIRS far-IR limb integration averaged spectrum ($\Delta \nu = 0.48 \text{ cm}^{-1}$) recorded at 85°N at a tangent height of ~135 km during mid northern winter in March 2005. This figure shows the very strong emission feature of the Haystack (solid black curve), ob-
Fig. 14 CIRS far-IR limb integrations spanning 50°S to 85°S during late southern fall, all recorded at tangent heights of ~125 km. The spectra were recorded in late 2015 (85°S), mid 2015 (79°S), early 2016 (65°S and 57°S), and mid 2016 (54°S and 50°S). The spectra at 54°S, 57°S, 65°S, 79°S, and 85°S were vertically offset from the 50°S spectrum served when it was at its strongest during northern winter. The varying color-coded continuum intensity curves were generated from Mie scattering calculations (for a 2 µm particle radius), coupled with our radiative transfer model (model detailed in Anderson and Samuelson 2011), using the appropriate aerosol and collision induced absorption (CIA) opacities and the temperature structure at 85°N (see Anderson et al. 2014). The optical constants for H2O ice (aqua curve) were taken from Hudson and Moore (1995); vapor was deposited at 140 K, C2H5CN ice (orange curve) were taken from Nna-Mvondo et al. (2018); vapor was deposited at 135 K, HCN ice (green curve) were taken from Anderson et al. (2018); vapor was deposited at 110 K, co-condensed C6H6:HCN ice (purple curve) containing a 4:1 mixing ratio were taken from Anderson et al. (2017); vapor was deposited at 110 K, and co-condensed C6H6:C2H5CN:HCN ice (pink curve) containing a 3.6:1.4:1.0 mixture ratio were determined from Nna-Mvondo et al. (2018); vapor mixture was deposited at 110 K. The obvious spectral mismatch between the Haystack and all five ice examples is evident.
Specifically, for pure $\text{C}_2\text{H}_5\text{CN}$ ice, it was ruled out as a likely Haystack compound due to its very strong low-energy lattice mode (Samuelson et al. 2007). However, as seen in the figure, this lattice mode is easily damped out when $\text{C}_2\text{H}_5\text{CN}$ ice becomes part of a co-condensed ice. This is expanded on in Nna-Mvondo et al. (2018). $\text{C}_2\text{H}_5\text{CN}$ ice may still indeed contribute to the Haystack’s identity, as part of a co-condensate, although the formation mechanism will most likely be something other than vapor condensation. This quashing trend of low-energy ice modes is also evident with the strong libration mode of HCN ice. The libration mode of pure HCN ice spectrally peaks near 168 cm$^{-1}$, but when co-condensed with $\text{C}_6\text{H}_6$ ice (purple curve) or co-condensed with $\text{C}_6\text{H}_6$ and $\text{C}_2\text{H}_5\text{CN}$ ice (pink curve), the libration mode of HCN is easily modified. A detailed discussion on co-condensed ices, as they specifically pertain to Titan’s stratospheric ices, are found in Anderson et al. (2018).

The first detection of the Haystack in Titan’s south polar stratosphere occurred in the mid southern fall season (July 2012; Jennings et al. 2012a). The investigators used CIRS weighted-averaged disk-viewing spectra—targeting 75°S—recorded at Cassini-Titan distances $>50,000$ km. This larger distance from Titan results in a larger FP1 FOV diameter size ($\sim320$ km), in which the FOV subtended both Titan’s limb and disk. Such an observation was unavoidable, however, since Cassini’s orbit at this time was in an inclined orientation so the dedicated far-IR limb scans required to constrain the vertical and latitudinal distributions (discussed in Sect. 3.2) at high polar latitudes were not possible. Thus, the limitations imposed by this type of disk-viewing observation prevents detailed vertical and latitudinal information. However, radiative transfer analyses could still improve the current state of knowledge especially at this critical moment in time when CIRS serendipitously caught the onset of the Haystack’s formation.

By July 2015, Cassini had returned to a lower inclination, allowing for far-IR limb-viewing of Titan’s south polar stratospheric regions, recorded at distances between $\sim15,000$ km.
and 35,000 km off Titan’s limb. CIRS FP1 high spectral resolution limb integrations (typically centered on 125 km tangent height) at multiple latitudes during Titan’s late southern fall season are shown in Fig. 14. Although this figure depicts a strong latitudinal dependence of the Haystack, in limb integration mode (or sit-and-stare mode), the large CIRS FP1 FOV collects and integrates the signal from many altitudes simultaneously. This causes the Haystack’s spectral signature to appear in lower latitude spectra, where the Haystack actually does not reside. From analysis of far-IR limb scans acquired in July 2015 (example limb scan at 215 cm$^{-1}$ is shown in Fig. 8), our best fit radiative model (model details given in Anderson and Samuelson 2011) reveals the Haystack to reside at latitudes poleward of $\sim$70°S. This point is further illustrated at 215 cm$^{-1}$ in Fig. 8. When no attempt is made to restrict the Haystack with latitude, the resulting fit is very poor (gray curve). However, when the presence of the Haystack is restricted to latitudes poleward of $\sim$70°S, a very satisfactory fit (orange curve) results. From these analyses, the Haystack’s altitude location is found to reside fairly deep in Titan’s stratosphere, with a cloud top near $\sim$110 km. This is evident in the limb scan at 215 cm$^{-1}$ (Fig. 8) and also indicated in Fig. 1.

Observations of the Haystack have spanned many seasons, including its disappearance from Titan’s northern spring stratosphere, with its reemergence in Titan’s southern fall stratosphere (Jennings et al. 2012a,b, 2015). No matter the latitude or season, the Haystack’s spectral dependence remains the same. This lack of spectral variation, combined with no known corresponding vapor in the stratosphere, implies that the Haystack—similarly to C$_4$N$_2$ ice clouds—has a different formation mechanism from that of vapor condensation. Continued experimental efforts are needed to identify the chemical composition of Titan’s Haystack.

3.2.4 The HASP Ice Cloud

The High-Altitude South Polar (HASP) cloud is a name given to the CIRS-discovered massive cloud system that developed at altitudes throughout Titan’s mid stratosphere at high southern polar latitudes during late southern fall in 2015 (Anderson et al. 2017, 2018, see Figs. 1, 8, 10, 15). The CIRS HASP cloud was first observed in the far-IR near Titan’s south pole is March 2015 and the last observation occurred in February 2016, when Cassini began to ramp back up into a higher inclined orbital configuration, preventing further close-in far-IR limb observations of Titan’s south polar stratosphere. Between March 2015 and February 2016, the CIRS HASP cloud was always observed at high southern polar latitudes, but at distinctly lower altitudes than the ISS-discovered south polar cloud, which was observed near 300 km.

Figures 15 and 17 show far-IR limb integrations in late southern fall (2015) at 79°S (upper spectra in Figs. 15 and 17) compared to those recorded 10 years earlier in Titan’s mid northern winter (2005) at 85°N (lower spectra in Figs. 15 and 17), with the CIRS FP1 FOVs centered on $\sim$125 km in Fig. 15 and $\sim$225 km in Fig. 17. Noticeable immediately in Fig. 15 is the appearance of a spectral mismatch between the Haystack at 79°S compared to the Haystack at 85°N. The spectral dependence of the Haystack is actually unaltered—the broad shoulder appearing to the lefthand side of the Haystack at 79°S is due to the CIRS HASP cloud. Likewise, in Fig. 17, in which the FP1 FOV is centered at a tangent height 100 km higher in the stratosphere than in Fig. 15, trace amounts of the Haystack material in Titan’s stratosphere are observed in these limb integration spectra, both at 85°N and at 79°S, whereas the strong spectral signature from the CIRS HASP cloud is only evident at 79°S. All of the apparent spectral pollutions result from CIRS operating in the limb integration mode. In this scenario, the large CIRS FP1 FOV collects the signal from many altitudes simultaneously during a limb integration, picking up on the HASP cloud opacity from 100 km.
Fig. 16 CIRS far-IR limb integration spectrum at 85°N recorded during mid northern winter in March 2005 (black curve), showing the spectral dependence and intensity contribution to Titan’s stratosphere from the Haystack. The various color-coded curves are synthetic spectra generated from our continuum model at 85°N (Anderson and Samuelson 2011; Anderson et al. 2014, 2018) for five different ices (Mie scattering calculations were performed with a 2 µm particle radius): water ice (aqua curve; vapor was deposited at 140 K; from Hudson and Moore 1995), pure C$_2$H$_5$CN ice (orange curve; vapor was deposited at 135 K; from Nna-Mvondo et al. 2018), pure HCN ice (green curve; vapor was deposited at 110 K; from Anderson et al. 2018), co-condensed C$_6$H$_6$:HCN ice in a 4:1 mixing ratio (purple curve; mixed vapors were deposited at 110 K; from Anderson et al. 2017), and co-condensed C$_6$H$_6$:C$_2$H$_5$CN:HCN ice in a 3.8:1.4:1.0 mixing ratio (pink curve; mixed vapors were deposited at 110 K; from Nna-Mvondo et al. 2018). One of the main reasons that pure C$_2$H$_5$CN ice was ruled out as a Haystack candidate was due to its strong low-energy lattice mode, easily seen in the figure (orange curve), and never observed by CIRS (Samuelson et al. 2007). However, during co-condensation formation processes, the C$_2$H$_5$CN lattice mode is easily damped out, as is the HCN libration mode.

higher up in the stratosphere when the FOV is centered on 125 km and also picking up on the Haystack’s opacity 100 km deeper down in the stratosphere when the FOV is centered on 225 km. As radiative transfer analyses of far-IR limb scans show the CIRS HASP cloud and the Haystack are vertically separated by ~100 km (seen also in Fig. 8)—the CIRS HASP cloud vertically resides near 210 km and the Haystack is located near 110 km in July 2015. CIRS data analyses of the low spectral resolution far-IR limb scans must be utilized to unambiguously retrieve the vertical distributions and spectral dependences for all of Titan’s stratospheric particulates.

The spectral dependence of the CIRS HASP cloud is directly connected to its chemical composition (as is true for all other stratospheric particulates). The CIRS-derived spectral dependence of the HASP cloud is shown in Fig. 18, while Fig. 19 depicts the spectral dependences between the HASP cloud and the 160 cm$^{-1}$ ice cloud. The disparate spectral dependences amongst the two clouds—observed 10 years apart—demonstrates that the two ice clouds are comprised of different chemical compositions.

The vertical extent of the CIRS HASP cloud is observed at stratospheric altitudes where the pure compounds HCN, HC$_3$N, and C$_6$H$_6$ are all expected to condense and form stratospheric ice clouds (Fig. 1). As with the northern winter 160 cm$^{-1}$ ice cloud, whose formation mechanism is consistent with simultaneous saturation of HCN and HC$_3$N, i.e., co-condensation, a similar formation scenario is expected for the CIRS HASP cloud, poten-
Fig. 17  CIRS far-IR limb integrations recorded at 79°S in late southern fall (black curve; July 2015) and at 85°N in mid northern winter (orange curve; March 2005), with the FP1 FOV centered on ~225 km. Noticeable immediately is a spectral difference between the two latitudes, with only trace amounts of Haystack material contributing to these limb integration spectra. The strong spectral signature observed at 79°S is due to the HASP cloud, which resides ~100 km higher in the stratosphere than the Haystack. The observed spectral pollutions result from the large range in stratospheric altitudes averaged over the FP1 FOV during the limb integrations.

3.2.5 Other Potential Ice Clouds

More efforts have been put forth to describe additional opacity signatures from presumed stratospheric ice clouds. One example was found at 58°S during northern winter, and in addition to the 160 cm\(^{-1}\) ice cloud that was observed near ~90 km, an additional ice cloud was found about 30 km deeper in the stratosphere, peaking at 60 km (Anderson and Samuelson 2011). The altitude location combined with the far-IR spectral dependence peaking near 80 cm\(^{-1}\) indicated that this ice cloud was composed of condensed C\(_2\)H\(_6\) ice. However, far-IR targeted thin ice film transmission spectroscopy experiments are still needed to confirm this chemical identify.

The second example involves the \(v_6\) band of pure C\(_6\)H\(_6\) ice at 682 cm\(^{-1}\). This ice spectral feature was reported to explain some of the stratospheric ice emission features observed in CIRS mid-IR nadir (in May 2013) and limb (in March 2015) spectra (Vinatier et al. 2018). Most notably for the limb spectra spanning 168 km to 278 km, the observed width of the observed ice emission features were reported to be significantly broader than those shown for the \(v_6\) band of pure C\(_6\)H\(_6\) ice at 682 cm\(^{-1}\). Even so, an upper limit of 1.5 \(\mu\)m for the C\(_6\)H\(_6\) particle radius was reported. Three additional spectral peaks at ~687, 695, and ~702 cm\(^{-1}\)
Fig. 18  Black curve represents the CIRS-derived spectral dependence of Titan’s HASP cloud discovered at mid-stratospheric altitudes during late southern fall at $79^\circ$S (July 2015). The magenta curve depicts a co-condensed thin ice film (vapors deposited at 110 K) containing $\sim$20% HCN and $\sim$80% C$_6$H$_6$. This result demonstrates that the HASP cloud’s chemical identification is consistent with a mixed C$_6$H$_6$:HCN ice, formed via co-condensation processes.

Fig. 19  CIRS-derived spectral dependences of Titan’s late southern fall HASP cloud (black solid curve) and the northern winter 160 cm$^{-1}$ ice cloud (black dashed curve). The dissimilar spectral dependences demonstrate that the two ice clouds—observed 10 years apart in opposite hemispheres—are comprised of different chemical compositions were also reported, although the investigators could not reproduce these spectral features with the refractive indices found in the literature from the pure nitrile ices HCN, HC$_3$N, CH$_3$CN (acetonitrile), C$_2$H$_5$CN, or C$_2$N$_2$. The four observed spectral signatures may indeed result from co-condensed ices, although dedicated thin ice film transmission spectroscopy
for mixtures containing C₆H₆ are still needed—at various deposition temperatures—to spectrally match the opacity spectral signatures reported by Vinatier et al. (2018).

4 Titan Stratospheric Ice Experimental Efforts

The first experimental laboratory study to interpret ice emission features observed in Titan’s north polar stratosphere by IRIS was performed in the far-IR by Khanna et al. (1987). The investigators identified the IRIS-observed ice emission feature at 478 cm⁻¹ as the ν₈ band of crystalline C₄N₂ ice. The C₄N₂ vapor was deposited at 70 K, warmed to 90 K, and then annealed. Experimental studies continued in the mid-IR with C₂H₂ and C₄H₂ ice (Khanna et al. 1988). Even though these hydrocarbon ices were not observed by IRIS, Khanna et al. (1988) claimed that they may reside in Titan’s lower stratosphere given their very strong observed vapor emission features in the mid-IR. Subsequent experiments in the mid-IR were performed involving HCN, HC₃N, and C₄N₂ (Masterson and Khanna 1990). HCN vapor was deposited at 60 K, then the ice was warmed and annealed at 80 K for 30 minutes. Both HC₃N and C₄N₂ vapors were deposited at 70 K, then warmed and annealed at 90 K for 30 minutes. The fundamental vibrational modes and combination or overtone bands of the ice spectral peaks detected were assigned across the wavenumber range 450–5000 cm⁻¹, and also their complex refractive indices were calculated.

Next, an extensive laboratory effort was initiated involving HCN, HC₃N, CH₃CN, C₂H₅CN, C₂H₃CN, C₂N₂, and C₄N₂, spanning wavenumbers 80 to 5000 cm⁻¹ (Dello Russo and Khanna 1996). The vapors were deposited somewhere between 50 and 100 K, and annealed at various temperatures (see Table 1), then cooled to 95 K and 35 K, where their transmittance spectra were recorded. This was the first detailed laboratory effort demonstrating the strong low-energy vibrational modes of Titan-relevant nitrile ices in the far-IR, especially below 200 cm⁻¹. Following this experimental effort, a focused study on C₂H₅CN ice was put forth, in an effort to chemically identify Titan’s Haystack (Khanna 2005). Whereas earlier work regarding C₂H₅CN ice performed by Dello Russo and Khanna (1996) showed C₂H₅CN ice had a strong spectral feature at 225 and 226 cm⁻¹ (due to the ν₁₃ C–C≡N bending mode), the re-examination of C₂H₅CN ice by Khanna (2005) revealed that the ν₁₃ band had shifted to 221 cm⁻¹. The 4 to 5 cm⁻¹ wavenumber shift resulted from annealing C₂H₅CN ice at 120 K (compared to 140 K from the earlier experiment) and also the annealing time duration was much longer (4 to 5 hours). Additional discrepancies at higher energies were also found. More recently, a very detailed study of C₂H₅CN ice, with a focus on better understanding Titan’s CIRS-observed stratospheric ice clouds, especially in regards to the Haystack, was reported for vapor deposition temperatures ranging from 30 K to 150 K (Nna-Mvondo et al. 2018).

In the Cassini era, the first Titan-relevant experimental study was performed by Moore et al. (2010). Both the amorphous and crystalline phases of HCN, HC₃N, CH₃CN, C₂H₂CN, and C₂N₂, spanning wavenumbers 30 to 5000 cm⁻¹ were measured, as well as determinations of their complex refractive indices. As with all the previous Titan-related thin ice film transmission spectroscopy studies, all the vapors were deposited at cold temperatures (e.g., 30 K), ensuring the amorphous phase was achieved, and then annealed to warmer temperatures for their transition to the crystalline phase (see Tables 1 and 2 for details, as well as Anderson et al. 2018).

In order to interpret and chemically identify many of the CIRS-discovered stratospheric ice clouds, a new experimental approach, modified from all the previous Titan ice work, was needed. As discussed in Sects. 1 and 3, Titan’s stratospheric ice clouds commonly form as
Table 1  Summary of crystalline phase laboratory studies related to Titan’s stratospheric ice clouds

| Compound | Experimental conditions | Spectral range | Published work |
|----------|-------------------------|----------------|----------------|
| HCN      | deposited at 60 K       | 450–5000 cm\(^{-1}\) | Masterson and Khanna (1990) |
|          | annealed for 30 min at 80 K |                |                |
|          | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed for 30 to 180 min at 120 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
|          | deposited at 50 K       | 30–5000 cm\(^{-1}\) | Moore et al. (2010) |
|          | annealed at 120 K for 120 min in mid-IR |                |                |
|          | annealed at 120 K for 80 min in far-IR |                |                |
|          | cooled to 20 K, 35 K, 50 K, 75 K, 95 K, 110 K |                |                |
| HC\(_3\)N | deposited at 70 K | 450–4000 cm\(^{-1}\) | Masterson and Khanna (1990) |
|          | annealed 30 min at 90 K |                |                |
|          | cooled to 60 K          |                |                |
|          | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed 30 to 180 min at 110 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
|          | deposited at 50 K       | 30–5000 cm\(^{-1}\) | Moore et al. (2010) |
|          | annealed at 110 K for 120 min in mid-IR |                |                |
|          | annealed at 110 K for 65 min in far-IR |                |                |
|          | cooled to 20 K, 35 K, 50 K, 75 K, 95 K, 110 K |                |                |
| C\(_2\)N\(_2\) | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed 30 to 180 min at 90 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
|          | deposited at 20 K       | 30–5000 cm\(^{-1}\) | Moore et al. (2010) |
|          | annealed at 90 K for 60 min in mid-IR |                |                |
|          | annealed at 90 K for 90 min in far-IR |                |                |
|          | cooled to 20 K, 35 K, 50 K, 75 K |                |                |
| C\(_4\)N\(_2\) | deposited at 70 K | 400–4000 cm\(^{-1}\) | Khanna et al. (1987) |
|          | annealed 30 min at 90 K |                |                |
|          | also annealed at 155 K  |                |                |
|          | deposited at 70 K       | 450–4000 cm\(^{-1}\) | Masterson and Khanna (1990) |
|          | annealed 30 min at 90 K |                |                |
|          | cooled to 60 K          |                |                |
|          | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed 30 to 180 min at 120 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
| CH\(_3\)CN | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed 30 to 180 min at 130 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
|          | deposited at 50 K       | 30–5000 cm\(^{-1}\) | Moore et al. (2010) |
|          | annealed 66 min at 130 K in mid-IR |                |                |
|          | annealed 45 min at 130 K in far-IR |                |                |
|          | cooled to 20 K, 35 K, 50 K, 75 K, 95 K, 110 K |                |                |
|          | collisional cooling cell particles formed at 95 K, 110 K, 130 K | 50–5000 cm\(^{-1}\) | Ennis et al. (2017) |
| C\(_2\)H\(_3\)CN | deposited between 50 and 100 K | 80–5000 cm\(^{-1}\) | Dello Russo and Khanna (1996) |
|          | annealed 30 to 180 min at 130 K |                |                |
|          | cooled to 95 K, 35 K    |                |                |
Table 1 (Continued)

| Compound | Experimental conditions | Spectral range | Published work |
|----------|-------------------------|----------------|----------------|
| C$_2$H$_5$CN | deposited between 50 and 100 K annealed 30 to 180 min at 140 K cooled to 95 K, 35 K | 80–5000 cm$^{-1}$ | Dello Russo and Khanna (1996) |
| | deposited at 60 K annealed 4 to 5 hours at 120 K | 200–7800 cm$^{-1}$ | Khanna (2005) |
| | deposited at 50 K annealed at 140 K for 90 min in mid-IR annealed at 140 K for 92 min in far-IR cooled to 20 K, 35 K, 50 K, 75 K, 95 K, 110 K collisional cooling cell particles formed at 95 K, 110 K, 130 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |
| C$_3$H$_3$CN | deposited between 50 and 100 K annealed 30 to 180 min at 145 K cooled to 95 K, 35 K | 80–5000 cm$^{-1}$ | Dello Russo and Khanna (1996) |

Table 2 Summary of Amorphous phase laboratory studies related to Titan’s stratospheric ice clouds

| Compound | Experimental conditions | Spectral range | Published work |
|----------|-------------------------|----------------|----------------|
| HCN | deposited at 50 K warmed to 75 K, 95 K, 110 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |
| HC$_3$N | deposited at 50 K warmed to 50 K, 75 K, 95 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |
| C$_2$N$_2$ | deposited at 20 K warmed to 35 K, 50 K, 75 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |
| CH$_3$CN | deposited at 50 K warmed to 75 K, 95 K, 110 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |
| C$_2$H$_5$CN | deposited at 50 K warmed to 75 K, 95 K, 110 K | 30–5000 cm$^{-1}$ | Moore et al. (2010) |

As a result of the vapors diffusing/subsiding, cooling as they descend in Titan’s stratosphere, then condensing once saturation has been achieved. Many of these vapors will enter altitude regions where they will undergo simultaneous saturation, thus forming co-condensed ice clouds. Thin ice film transmission spectroscopy for the purpose of studying Titan’s stratospheric ice clouds must therefore be performed with vapor deposition at temperatures $>100$ K. This is necessary since the structure of the ice, for both pure and mixtures, is modified between annealing and warm-deposition experiments (this is expanded on in Anderson et al. 2018). The main reason is that during annealing experiments, a complete ordering of the ice structure is not achieved, so a portion of the amorphous structure is irretrievably retained in the final crystalline phase. These differences are more pronounced in the low-energy part of the far-IR. Using C$_2$H$_5$CN ice as an example, Khanna (2005) reported that the longer time required for annealing C$_2$H$_5$CN ice, compared to other nitriles, is necessary since there may be several possible orientations of the methyl (CH$_3$) and methylene groups (CH$_2$) in the solid phase when the samples are quenched at low temperatures, thus causing low barriers in the rotation of these groups around the single bonds. These effects
are most noticeable in the low energy lattice modes, as recently detailed in a comprehensive laboratory study on the ice phases of C2H5CN (Nna-Mvondo et al. 2018).

C2H5CN ice was also recently studied by Ennis et al. (2017), in which C2H5CN and CH3CN ice particles were generated at 95 K, 110 K, and 130 K under simulated Titan temperature and pressure conditions using collisional cooling cell experiments. The investigators measured transmittance spectra from 50 to 5000 cm\(^{-1}\), with a focus in the far-IR to study the morphology of the icy particulates, identify their crystalline phases, and also attempt to spectrally match the Haystack. While a metastable b-phase (monoclinic) ice was identified for CH3CN, there was no such phase observed for C2H5CN. Ennis et al. (2017) also reported the 200–250 cm\(^{-1}\) region was absent of an CH3CN spectral feature, but found the \(\nu_{13}\) band of crystalline C2H5CN ice at 110 K was located at 226 cm\(^{-1}\), which is in agreement with Dello Russo and Khanna (1996).

Reproducing Titan’s stratospheric ices in the laboratory requires forming the ice samples with both high purity and free of oxygen-bearing components. Whereas some of the nitriles and hydrocarbons necessary for Titan ice experiments are available commercially by chemical manufacturers (e.g., C2H5CN, CH3CN, C2H2), others like HCN, HC3N, C2N2, C4N2, and C4H2 must be synthesized in the laboratory. No matter the way in which the compounds are secured, all will likely contain some degree of contamination, such as the air contaminants H2O and CO2, byproducts from the syntheses, or additional organics produced from exposing nitriles to ambient air. The presence of any sample impurity must be purged since it can alter the spectral features of the Titan ice analogs, ultimately resulting in inaccurately-determined complex refractive indices. Purifying the sample becomes very crucial, and unfortunately, most of the published laboratory work focused on Titan’s stratospheric ices are impure. For example, some work efforts report CO2 impurity bands in the infrared spectra of their synthesized ices, e.g., in the HC3N spectra reported in Moore et al. (2010), in the C4N2 ice spectrum of Khanna et al. (1987), and in the CH3CN ice spectrum of Ennis et al. (2017). This latter work effort also noted the cryopump oil impurities in their samples. They further showed the presence of CO2 as an impurity in their CH3CN and C2H5CN ice particle spectra (both generated from the purified commercial nitriles). CO2 was introduced as an impurity in the dry N2 bath gas as well as an adsorbed formic acid remnant from previous experiments. Moreover, their C2H5CN ice particle spectra also contained traces of acetonitrile absorbed on the enclosed flow cooling cell or on the deposition line walls. Additionally, when purchased chemicals are used, and no purification is performed, the resulting absorbance spectra can contain air contaminants such as CO2, e.g., in the C2N2 ice spectra of Moore et al. (2010), or H2O, e.g., in the C2H5CN ice spectra of Khanna (2005). In recent work, however, Anderson et al. (2018) illustrates the difficulties in synthesizing the initial compounds needed for the Titan stratospheric ice experiments, the efforts and time dedication involved in purifying the sample, and the corresponding benefits that are reflected in the recorded transmittance spectra. These investigators also discuss their new Titan ice chamber, focus on simulating Titan’s icy stratospheric environment, with an emphasis on the formation of co-condensed ices.

5 Closing Remarks

Many of the extensive laboratory efforts focused on thin ice film transmission spectroscopy over the last three decades (see for example Allamandola et al. 1988; Sandford and Allamandola 1993a,b; Hudgins et al. 1993; Bohn et al. 1994; Bernstein et al. 1997; Allamandola et al. 1999; Moore et al. 2007; Hudson et al. 2009, 2014; Bouwman et al. 2010; Gerakines and
Hudson 2015) have been astrochemistry-driven and/or aimed at the interpretation of ground- and/or space-based observations of interstellar ices, ices in molecular clouds, as well as ices on airless bodies (or with tenuous atmospheres) in the solar system, e.g., comets, Triton, Pluto, asteroids, etc. Except for the experiments listed in Sect. 4, very few have focused on simulating and reproducing Titan’s stratospheric ice clouds, especially those observed by Cassini’s ISS, VIMS, or CIRS instruments. This is a direct result of limited observational data, with most laboratory efforts regarding Titan’s stratospheric ices, aimed at interpreting Voyager 1 IRIS-derived results obtained during the Titan north polar limb mosaic recorded in 1980. It was not until Cassini arrived in the Saturn system in 2004, combined with its 13-year mission tenure, that observations of ice clouds residing in Titan’s stratosphere over multiple seasons were acquired, resulting in the need for much more detailed experimental studies. Specifically, these experiments must be performed by depositing the vapor at temperatures that are representative of Titan’s stratosphere, ideally at temperatures between \( \sim 67 \) K and 150 K. Dedicated thin ice film transmission spectroscopy experimental efforts are critically needed to better understand the chemical compositions, optical properties, and formation mechanisms of Titan’s observed stratospheric ice clouds. This is especially true regarding co-condensed ices and for ices that do not form from vapor condensation processes, i.e., \( \text{C}_4\text{N}_2 \) ice, the Haystack, etc.

Titan’s stratospheric ice cloud structures are very complex, and these clouds play an especially important radiative role where they spatially exist in Titan’s stratosphere. They are significant sources of thermal infrared opacity, especially at winter polar latitudes, contributing significantly to Titan’s atmospheric radiative balance. For example, based on the sheer strength in intensity of the Haystack (it is \( \sim 50 \) times more opaque than the aerosol at 221 cm\(^{-1}\) during northern winter at 85\(^{\circ}\)N), it appears to provide an exceptionally strong radiative cooling mechanism where it resides spatially, with additional contribution from the other stratospheric ice clouds. Both the Haystack and Titan’s additional stratospheric clouds are all very efficient far-IR radiative coolers, which leads to significant and unexpected alterations to Titan’s vertical temperature structures, especially in the winter polar stratosphere (Schinder et al. 2011, 2012; Anderson et al. 2014). The vertical extent of the Haystack appears to correspond very closely to the location of Titan’s local stratospheric temperature minimum near 110 km at 85\(^{\circ}\)N during northern winter (Anderson et al. 2014). Other local stratospheric temperature minima occur as well in the late southern fall season. Continued radiative transfer analyses, radiative cooling calculations, along with ongoing experimental efforts, are critically needed to better understand the contributions from all of Titan’s stratospheric ice clouds.

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