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DETECTION OF HF TOWARD PKS 1830–211, SEARCH FOR INTERSTELLAR H₂F⁺, AND LABORATORY STUDY OF H₂F⁺ AND H₂Cl⁺ DISSOCIATIVE RECOMBINATION

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ABSTRACT

We report extragalactic observations of two fluorine-bearing species, hydrogen fluoride (HF) and fluoronium (H_2F^+) , in the z=0.89 absorber in front of the lensed blazar PKS 1830–211 with the Atacama Large Millimeter/ submillimeter Array. HF was detected toward both southwest and northeast images of the blazar, with column densities $> 3.4 \times 10^{14}$ cm⁻² and 0.18×10^{14} cm⁻², respectively. H₂F⁺ was not detected, down to an upper limit (3σ) of 8.8×10^{11} cm⁻² and an abundance ratio of $[H_2\hat{F}^+]/[H\hat{F}] \le 1/386$. We also searched for H_2F^+ toward the Galactic sources NGC 6334 I and W51C, and toward Galactic center clouds with the Herschel HIFI spectrometer.⁶ The upper limit on the column density was derived to be 2.5×10^{11} cm⁻² in NGC 6334 I, which is 1/68 of that for H₂Cl⁺. In constrast, the ortho transition of H₂Cl⁺ is detected toward PKS 1830–211. To understand the small abundance of interstellar H₂F⁺, we carried out laboratory experiments to determine the rate constants for the ionelectron recombination reaction by infrared time-resolved spectroscopy. The constants determined are $k_e(209 \text{ K}) =$ $(1.1 \pm 0.3) \times 10^{-7} \, \text{cm}^3 \, \text{s}^{-1}$ and $(0.46 \pm 0.05) \times 10^{-7} \, \text{cm}^3 \, \text{s}^{-1}$ for H_2F^+ and H_2Cl^+ , respectively. The difference in the dissociative recombination rates between H_2F^+ and H_2Cl^+ by a factor ~ 2 and the cosmic abundance ratio $[F]/[C1] \approx 1/6$ are not enough to explain the much smaller abundance of H_2F^+ . The difference in the formation mechanism of H₂F⁺ and H₂Cl⁺ in interstellar space would be a major factor in the small abundance of H₂F⁺.

Key words: galaxies: individual (PKS 1830–211) – ISM: individual objects (W31C, NGC 6334 I) – ISM: molecules – methods: laboratory: molecular

1. INTRODUCTION

The hydrogen fluoride (HF) molecule was first detected in space by Neufeld et al. (1997) toward Sgr B2, using the Infrared Space Observatory. Later, CF⁺ was detected toward the Orion bar region (Neufeld et al. 2006), where the detection was driven by a theoretical study (Neufeld et al. 2005) of fluorine-bearing molecules in the interstellar medium. HF and CF⁺ are significant fluorine-bearing molecules in interstellar space, while AlF was detected in a circumstellar envelope by Cernicharo & Guélin (1987). Recently, HF was detected in many interstellar clouds with the Heterodyne Instrument for the Far-Infrared (HIFI) on board the Herschel Space Observatory (Neufeld et al. 2010; Phillips et al. 2010; Sonnentrucker et al. 2010, 2015; Monje et al. 2011a). HF was also detected in four nearby galaxies: in Mrk 231 (van der Werf et al. 2010, in emission) and Arp 220 (Rangwala et al. 2011) with the Herschel SPIRE-FTS, and in NGC 253 and NGC 4945 using HIFI (Monje et al. 2014). The detection of HF toward a highredshift quasar at z = 2.56 was reported by Monje et al. (2011b) using the Caltech Submillimeter Observatory. The observed abundance of HF in diffuse clouds was found to be consistent with theoretical predictions of Neufeld et al. (2005), Neufeld & Wolfire (2009), and Sonnentrucker et al. (2015), where in the latest report they carried out some updates of reaction data and also included grain surface chemistry.

In the present paper, we report the detection of HF in the z =0.89 molecular absorber toward PKS 1830-211 and searches for H₂F⁺ toward PKS 1830–211 and other Galactic sources. H₂F⁺ is isovalent with H₂Cl⁺ and isoelectronic to H₂O. In 2010, Lis et al. reported the detection of H₂Cl⁺ toward NGC 6334 I and Sgr B2 with HIFI. Recently, H₂Cl⁺ was also detected toward PKS 1830-211 (Muller et al. 2014a) with the Atacama Large Millimeter/submillimeter Array (ALMA). The existence of the related H₂F⁺ ion in interstellar clouds would be interesting from the viewpoint of halogen chemistry. A purely rotational spectroscopic study of H₂F⁺ was reported by Fujimori et al. (2011), who measured five rotational lines of H₂F⁺ in the ground state by using a submillimeter-wave spectrometer based on a backward-wave oscillator (BWO) at the University of Waterloo. Since the BWO measurement was limited in the frequency region below 775 GHz, higher frequency transitions were covered by a tunable far-infrared spectrometer at the University of Toyama (Amano et al. 2012), and seven lines were detected in the region 1305-1851 GHz. These spectroscopic studies made it possible to search for H₂F⁺ in interstellar space. However, the rotational transition frequencies from the lowest state are not accessible to ground-based telescopes because of atmospheric absorption. Therefore, observations were carried out toward the z = 0.89molecular-rich absorber in front of PKS 1830-211 (see, e.g., Muller et al. 2011) using the ALMA and toward Galactic sources with the Herschel Space Observatory.

The Herschel Space Observatory detected many new interstellar molecules and molecular ions. In particular, H_2O^+ (Ossenkopf et al. 2010), HCl^+ (De Luca et al. 2012),

Herschel is an ESA space observatory with science instruments provided by European-led Principal Investigator consortia and with important participation from NASA.

 $\rm H_2Cl^+$ (Lis et al. 2010), and ArH $^+$ (Schilke et al. 2014) were first detected in relatively low-density clouds with $\rm H_2$ density of less than $10^3\,\rm cm^{-3}$. The proton affinity of HF (484 kJ mol $^{-1}$) is larger than that of $\rm H_2$ (422 kJ mol $^{-1}$), but smaller than those of CO (594 kJ mol $^{-1}$) and $\rm N_2$ (494 kJ mol $^{-1}$). Therefore, low-density clouds become potentially good candidates for searching for $\rm H_2F^+$. Since $\rm H_2F^+$ has been thought to be produced through the reaction of $\rm H_3^+$ and HF, $\rm H_2F^+$ may be a direct proxy for $\rm H_3^+$ in diffuse clouds.

In spite of sensitive searches for ${\rm H_2F^+}$ in the present study, the ion is still not detected in space. Its abundance depends on production and/or depletion mechanisms. In diffuse clouds, the depletion is dominated by dissociative recombination reactions with electrons. Neufeld et al. (2005) and Neufeld & Wolfire (2009) estimated the rate constant k_e for the following dessociative recombination reactions in their chemical study of fluorine-bearing molecules:

$$H_2F^+ + e \rightarrow HF + H$$

 $k_e = 3.5 \times 10^{-7} (T/300 \text{ K})^{-0.5} \text{ cm}^3 \text{ s}^{-1}$ (1)

$$H_2F^+ + e \rightarrow F + \text{products}$$

 $k_e = 3.5 \times 10^{-7} (T/300 \text{ K})^{-0.5} \text{ cm}^3 \text{ s}^{-1}.$ (2)

Various kinds of experimental methods have been reported for determining the recombination rate constants, as summarized by Plašil et al. (2002) and Geppert & Larsson (2008). Adam & Smith (1988) reported $k_e = 1 \times 10^{-7} \, \mathrm{cm}^3 \, \mathrm{s}^{-1}$ at 300 K for $\mathrm{H_2F^+} + \mathrm{e}$ by a flowing afterglow/Langmuir probe method. This value is seven times smaller than the value calculated from Equations (1) and (2) at 300 K. Nowadays storage ring experiments are well applied for measurements of dissociative recombination rate constants, such as those of $\mathrm{H_3^+}$ (McCall et al. 2004) and $\mathrm{HCl^+}$ (Novotny et al. 2013). The method has an advantage in measurement of the temperature dependence of the reaction rate. Neufeld et al. (2012) assumed the following k_e value of $\mathrm{H_2Cl^+}$, where storage ring data were cited as a private communication:

$$H_2Cl^+ + e \rightarrow Cl + products$$

 $k_e = 1.2 \times 10^{-7} (T/300 \text{ K})^{-0.85} \text{cm}^3 \text{ s}^{-1}.$ (3)

At 50 K (approximately the kinetic temperature of low-density clouds), Equations (1) and (2) give $k_e = 1.7 \times 10^{-6} \, \mathrm{cm^3 \, s^{-1}}$ for $\mathrm{H_2F^+}$, and Equation (3) gives $k_e = 5.5 \times 10^{-7} \, \mathrm{cm^3 \, s^{-1}}$ for $\mathrm{H_2Cl^+}$, which is about 1/3 of that of $\mathrm{H_2F^+}$. To check the difference in the recombination rate constants, the present paper reports measurements of the rate constants for the reactions of $\mathrm{H_2F^+}$ and $\mathrm{H_2Cl^+}$ ions with electrons by using an infrared OPO (optical parametric oscillator) laser coupled with a time-resolved method. Previously, the method of time-resolved infrared laser absorption has been applied to determine the rate constants of $\mathrm{H_3^+}$, $\mathrm{HCO^+}$, and $\mathrm{HN_2^+}$ by Amano (1988, 1990). Later it was found that the determined k_e value for $\mathrm{H_3^+}$ is close to the value obtained by storage ring experiments (McCall et al. 2004).

2. ASTRONOMICAL OBSERVATIONS

2.1. ALMA Observations

We obtained ALMA Cycle 2 observations of the ${\rm H_2F}^+$ ${\rm I}_{10^-}$ ${\rm I}_{01}$ transition (rest frequency of 760.929 GHz, redshifted to 403.5 GHz, i.e., in ALMA band 8) and HF J=1–0 transition (rest frequency of 1232.4762 GHz, redshifted to 653.5 GHz, i.e., in ALMA band 9) on 2015 May 14 and May 19, respectively. We adopt a redshift of $z_{\rm abs}=0.88582$, heliocentric frame, giving $v=0\,{\rm km\,s}^{-1}$ for the absorption toward the SW image of PKS 1830–211.

The ALMA correlator was configured with four different spectral windows of 1.875 GHz, each counting 960 channels separated by 1.938 MHz, providing velocity resolutions of $\sim\!1.4$ and 0.9 km s $^{-1}$ after Hanning smoothing, in bands 8 and 9, respectively. The spectral windows were centered at $\sim\!401.5$, 403.5, 413.8, and 415.5 GHz in band 8, and $\sim\!648.0$, 649.9, 651.7, and 653.6 GHz in band 9. In band 8, we also observed the ortho-H₂Cl⁺ 2₁₂–1₀₁ line (rest frequency of 781.6 GHz). No other lines were detected in the frequency coverage of band 9.

The data calibration was done within the CASA⁷ package, following a standard procedure. The bandpass response of the antennas was calibrated from observations of the bright quasar J1924–292. The complex gain solutions were determined by self-calibration of the visibilities of PKS 1830–211, using a simple model of two point sources at the positions of the two lensed images of the blazar. Using a Clean deconvolution (with Briggs weighting and robust parameter set to 0.5), we obtained synthesized beams of 0."39 × 0."29 (position angle PA = -84°) in band 8 and 0."21 × 0."19 (PA = 64°) in band 9. The two lensed images of PKS 1830–211, separated by ~1", were spatially well resolved. The spectra were extracted toward both images using the CASA-python task UVMULTIFIT (Martí-Vidal et al. 2014) to fit a model of two point sources to the visibilities.

2.2. Herschel Observations

On 2013 February 26–March 7, the H₂F⁺ ion was searched for toward four sources, W31C (star-forming region, α_{2000} : 18° 10'28.70, β_{2000} : $-19^{\circ}55'50.00$), NGC 6334 I (high-mass starforming region, α_{2000} : 17°20′53″32, β_{2000} : -35°46′58″5), GC IRS 21 (α_{2000} : 17°45′40″20, β_{2000} : -29°0′31″00), and 2Mass J17470898-2829561 (J1747) (α_{2000} : 17°47′8″980, β_{2000} : -28° 29'56."10), where the former two sources are known to have abundant HF and H₂Cl⁺. In the latter two sources near the Galactic center, the H₃⁺ ion has been detected strongly by infrared observations (Goto et al. 2008, 2011). We searched for two transitions, $J_{\text{KaKc}} = 2_{12} - 1_{01}$ (1850.082 GHz) and $1_{10} - 1_{01}$ (760.929 GHz), from the lowest energy state of ortho levels. We used the dual-beam switch mode and the wideband spectrometer of the Herschel/HIFI spectrograph with a spectral resolution of 1.1 MHz ($\Delta v = 0.43 \,\mathrm{km \, s^{-1}}$ at 760 GHz). The observed data of level 2 were provided as calibrated DSB spectra. The integration time was 8 minutes for each source in the 761 GHz region, and 34 minutes for W31C, 43 minutes for NGC 6334 I, 33 minutes for GC IRS 21, and 31 minutes for 2 Mass J1747 in the 1850 GHz region. The frequency region of the 1₁₁-0₀₀ line from the para ground state was not covered by Herschel/HIFI because of a lack of the detector system in the

⁷ http://casa.nrao.edu/

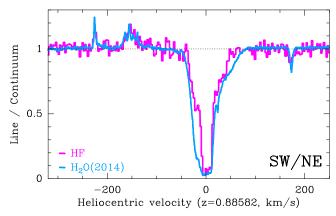


Figure 1. Spectrum of the HF J=1–0 line. Data were smoothed to a velocity resolution of 2.8 km s⁻¹. The normalized spectrum toward the SW image has been divided by that toward the NE image in order to remove small bandpass residuals. The SW and NE absorptions appear as features <1 and >1, respectively. The normalized spectrum of the ground-state transition of ortho- $\rm H_2O~1_{10}$ – $\rm 1_{01}$, observed in 2014 July, is shown for comparison. The velocity scale is referenced to the heliocentric frame taking z=0.88582.

region. The HF J = 1–0 transition at 1232.47 GHz was also observed toward NGC 6334 I and two sources near the Galactic center with an integration time of 7 minutes.

3. LABORATORY EXPERIMENTS FOR DISSOCIATIVE RECOMBINATIONS OF H₂F⁺ AND H₂Cl⁺

A continuous-wave OPO laser with a spectral width of about $0.8\,\mathrm{MHz}$ was used to cover the $3\,\mu\mathrm{m}$ infrared region in the present experiment (Okabayashi et al. 2015). It has a bow-tietype cavity, and a PPLN crystal is pumped by a fiber laser (iPG Photonics YAR-25 K-LP-SF), which is seeded by a DFB laser (Koheras Y10-PM). The output of the OPO laser in the 3 μ m region was 70 mW for a pumping power of the fiber laser of 6-8 W. The H₂F⁺ ion was produced by a pulsed hollowcathode discharge in a mixture of 5% F₂/He and H₂ gas with partial pressures of 70 mTorr (9.3 Pa) and 330 mTorr (44 Pa), respectively. A high-voltage transistor switch HTS 81 (Behlke Electronic GmbH) was used for the pulsed discharge with peak current of 200 mA. In the case of H₂Cl⁺, a mixture of HCl and H₂ was used with partial pressures of 50 mTorr (6.6 Pa) and 330 mTorr (44 Pa). The same absorption measurement system was used as in a previous Fourier-transform infrared (FTIR) study of H₂F⁺ (Fujimori et al. 2013) to attain an effective path length of 40 m with a multipath cell arrangement. The glass part of the cell had an inner diameter of 94 mm and a length of 1 m for the hollow-cathode discharge, and was connected to two glass tubes of inner diameter 144 mm and length of 250 mm on both sides, keeping a mirror separation of 150 cm. The cathode was made of a stainless plate of thickness 0.1 mm and mounted on the inner part of the glass cell. The discharge part was cooled by dry-ice. The transmitted laser beam was detected by an infrared mercury-cadmium-telluride detector (Vigo PVI-3TE-4, time constant of 20 ns) followed by a preamplifier (MIPDC-F-100, bandwidth of DC-100 MHz).

4. RESULTS

4.1. ALMA Observations of the z = 0.89 Absorber Toward PKS 1830-211

Figure 1 shows the observed absorption due to the HF J = 1-0 transition toward PKS 1830-211. Toward the

southwest (SW) image, the absorption spans a velocity range between about -50 and $+50 \, \mathrm{km \, s^{-1}}$ and is heavily saturated near $v = 0 \, \mathrm{km \, s^{-1}}$. Toward the northeast (NE) image, the absorption covers a velocity range between about -250 and $-100 \, \mathrm{km \, s^{-1}}$ and reaches at most $\sim 20\%$ of the continuum level. For comparison, we also show in Figure 1 the absorption spectrum of the ground-state transition of ortho-water, obtained in 2014 April with ALMA. The profiles are nearly identical along the NE line of sight, but the HF absorption has narrower wings than $\mathrm{H_2O}$ toward the SW image. Note, however, that time variations of the absorption profiles, with a timescale of months to years, have been observed in this absorber (Muller & Guélin 2008; Muller et al. 2014b) and somewhat limit the direct comparison of spectra taken at different epochs.

Assuming that all HF molecules are in the ground state of J=0, the column density is obtained by using the following formula:

$$N_{\rm HF} = \frac{8\pi^{3/2} \times \Delta v}{2\sqrt{\ln 2} \times \lambda^3 \times A} \frac{g_l}{g_u} \times \tau \tag{4}$$

where λ is the wavelength (243.2 μ m), A the Einstein A-coefficient (2.422 × 10⁻² s⁻¹), and g_l and g_u are the statistical weights of the lower and upper states. τ is the optical depth and is given by

$$\tau = -\ln\left(\frac{T_{\rm MB}}{T_{\rm back}}\right),\tag{5}$$

where $T_{\rm MB}$ and $T_{\rm back}$ are the measured brightness temperature and the background continuum temperature, respectively. In the SW direction, by using the linewidth of $20\,{\rm km\,s^{-1}}$, the lower limit of τ is estimated to be 4 from a simulation of the shape of the absorption line. On the other hand, the τ value of NE is obtained to be 0.21. The column densities are determined to be

$$N_{\rm HF}({\rm SW}) > 3.4 \times 10^{14} \, {\rm cm}^{-2} \quad {\rm [HF]/[H_2]} > 1.7 \times 10^{-8}$$

 $N_{\rm HF}({\rm NE}) = 0.18 \times 10^{14} \, {\rm cm}^{-2} \quad {\rm [HF]/[H_2]} = 1.8 \times 10^{-8}$

where the abundance relative to H2 is calculated by assuming an H₂ column density of $\sim 2 \times 10^{22}$ cm⁻² in SW and $\sim 1 \times$ 10²¹ cm⁻² in NE (Muller et al. 2014b). In the Milky Way, the relative abundance of HF is reported to be 2.5×10^{-8} in NGC 6334 I and 1.5 \times 10⁻⁸ in AFGL 2591 (Emprechtinger et al. 2012), 0.8×10^{-8} in IRC+10216 (Agúndez et al. 2011), $(1.1-1.6) \times 10^{-8}$ in W51 and W49N (Sonnentrucker et al. 2010), a conservative lower limit of 0.6×10^{-8} in W31C (Neufeld et al. 2010), and 1.6×10^{-10} in the Orion hot core (Phillips et al. 2010). Sonnentrucker et al. (2015) estimated the HF abundance in Galactic diffuse clouds by using a modified chemical reaction model and found $[HF]/[H_2] = (0.9-3.3) \times 10^{-8}$, which is consistent with the observed values including those of 12 newly observed sources. The currently obtained abundance in the z = 0.89 absorber is thus close to the values of low-density Galactic clouds.

The redshifted 403 GHz region for the H_2F^+ 1_{10} – 1_{01} transition was observed, as shown in Figure 2. The absorption due to H_2F^+ is not found, and the rms noise level (1σ) is estimated to be 0.5% with 4.2 km s⁻¹ channels. The absorption occurs from the lowest rotational level of the ortho H_2F^+ level, and the second lowest level 1_{10} is located at an energy 37 K

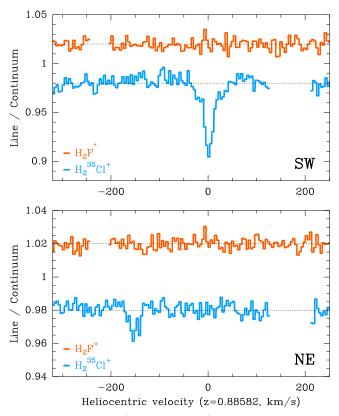


Figure 2. Spectra of the ${\rm H_2F^+}$ ${\rm I_{10}-I_{01}}$ and ${\rm H_2Cl^+}$ ${\rm 2_{12}-I_{01}}$ lines, toward the SW image (top) and NE image (bottom), observed simultaneously in the same tuning. Data were smoothed to a velocity resolution of 4.2 km s⁻¹. Regions with atmospheric lines were flagged. The velocity scale is referenced to the heliocentric frame taking z=0.88582.

higher than the 1_{01} level. Since the H_2 density of the absorbing gas is relatively low (a few $10^3 \, \mathrm{cm}^{-3}$), the excitation of polar species is coupled to the photons of the cosmic microwave background (CMB), with $T_{\rm CMB} = 5.14 \, \mathrm{K}$ at z = 0.89 (see Muller et al. 2013). We assume that all ortho state molecules are present in the lowest 1_{01} state. By using Equations (4) and (5), we derive the upper limit (3σ) for the ortho state of the ion to be

$$N_{H2F+}(SW) = 6.6 \times 10^{11} \, cm^{-2}$$

where the A-coefficient is $0.0200 \,\mathrm{s}^{-1}$ and λ is 394 $\mu \mathrm{m}$ for the 110-101 transition, and we assumed a velocity width of 20 km s⁻¹. Table 1 lists the upper limit for the total abundance, including the para state population by assuming an ortho-topara ratio of 3. It is known that some molecules with ortho-topara ratios smaller than 3 have been detected, and such molecules are thought to be produced on the surface of lowtemperature dust or grains. Some molecules produced by ionmolecule reactions in the gas phase have ratios smaller than 3 (Takakuwa et al. 2001; Morisawa et al. 2006; Faure et al. 2013), originating from abundant para-H₂ in cold clouds. However, in the case of H₂Cl⁺, ratios very different from 3 have not been reported (Neufeld et al. 2015), so the assumption of the high-temperature limit value may be valid for H₂Cl⁺ and also H₂F⁺. It is found that the abundance of H₂F⁺ is less than 1/386 of that of HF.

In the same line of sight, Muller et al. (2014a) reported a column density of H_2Cl^+ of $\sim\!1.4\times10^{13}\,\text{cm}^{-2}$ from the

Species	PKS 1830-211 ^a	W31C	NGC 6334 I
$\overline{\text{H}_2\text{F}^+}$	$\lesssim 8.8 \times 10^{11}$	$\lesssim 8.1 \times 10^{11}$	$\lesssim 2.5 \times 10^{11}$
H_2Cl^+	$\sim 1.4 \times 10^{13,b}$	$3.0 \times 10^{13,c}$	$1.7 \times 10^{13,d}$
HF	$> 3.4 \times 10^{14}$	$1.6 \times 10^{14,e}$	$(0.4-2.4) \times 10^{13,f}$
HCl	$3.4 \times 10^{13,g}$	$\sim 2 \times 10^{13,h}$	$1.9 \times 10^{14,i}$
H_2	$\sim 2 \times 10^{22,j}$	$1.2\times10^{22,k}$	$(0.6-1.8) \times 10^{21,f}$

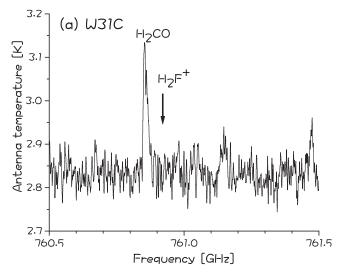
Notes.

- ^a SW line of sight.
- ^b Muller et al. (2014a).
- ^c Neufeld et al. (2012).
- ^d Lis et al. (2010).
- e Neufeld et al. (2010).
- ^f In the foreground components (Emprechtinger et al. 2012).
- g S. Muller et al. (2016, in preparation).
- ^h Monje et al. (2013).
- $T_{\rm ex} = 39 \, {\rm K}$ component (Zernickel et al. 2012).
- ^j Muller et al. (2011).
- ^k H₁ abundance (Godard et al. 2010).

observation of the para transition 1_{11} – 0_{00} and by assuming an ortho-to-para ratio of 3. In the present observation, the ortho transition 2_{12} – 1_{01} was detected, as shown in Figure 2. The integrated intensity (opacity) has been obtained as 2.03 km s⁻¹ and 0.71 km s⁻¹ toward SW and NE, respectively, and the column density of the ortho ions is estimated to be 9.1 × 10^{12} cm⁻² and 3.2×10^{12} cm⁻² by assuming that all ortho ions are in the lowest rotational 1_{01} level. The ortho-to-para ratio is 2.7 and 3 in SW and NE, respectively, which are consistent with the high-temperature limit value. Time variations of the absorption profiles (Muller & Guélin 2008) might affect this measurement of the ortho-to-para ratio for H_2CI^+ . Observations close in time would allow us to obtain a very robust ortho-to-para ratio in the two independent lines of sight. The H_2F^+ abundance is found to be less than 1/16 of that of H_2CI^+ .

4.2. Herschel Observations of Galactic Clouds: Search for Interstellar H_2F^+

The HF J = 1-0 spectral line was detected toward NGC 6334 I with many saturated absorption components, as reported by Emprechtinger et al. (2012). Spectra observed toward W31C and NGC 6334 I in the 760 GHz region are shown in Figure 3. In the two Galactic center sources GC IRS 21 and 2Mass J1747, no emission and absorption lines have been detected, where the background continuum level was 0.1 K and 0.03 K, respectively. The low continuum levels compared with 3σ noise levels of 0.1 K in both objects make it difficult to detect absorption lines. Many emission lines have been detected in W31C and NGC 6334 I, and assignments and analyses of these spectral lines will be reported in a separate paper. The H₂F⁺ line is expected to be observed as absorption, because the ion is thought to be present in low-density clouds as H₂Cl⁺, which is observed as absorption. Toward W31C and NGC 6334 I there was no feature of an absorption line in the 760 GHz region, as shown in Figure 3. In the 1850 GHz region, the noise level was 10-18 times larger than that in the 760 GHz region, so the sensitivity was not enough to detect the H₂F⁺ line. We estimated the upper limit for the column density to be 1.9 \times 10¹¹ cm⁻² in the ortho state toward NGC 6334 I, where we used three times (0.080 K) the rms noise and assumed a



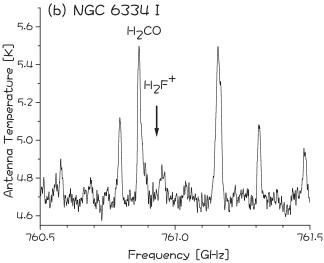


Figure 3. Search for the H_2F^+ 1_{10} – 1_{01} spectral line toward W31C and NGC 6334 I, where the frequency scale is given by using the systemic velocities of -3 km s^{-1} and -1.7 km s^{-1} , respectively. The H_2 CO line ($J_{KaKc} = 10_{19}$ – 9_{18}) is observed in the lower side band at 749.0721 GHz.

velocity width of 4.93 km s⁻¹. Similarly, the upper limit toward W31C was estimated by using $3\sigma = 0.090 \,\mathrm{K}$ and assuming a width of $9.5 \,\mathrm{km \, s^{-1}}$. The width corresponds to one velocity component among three or four components in the foreground gas of W31C. Table 1 lists total abundances, including the para state population of H₂F⁺ and related species. Since the abundance of H₂Cl⁺ has been reported to be 1.7 × $10^{13} \, \text{cm}^{-2}$ (Lis et al. 2010), which is 4.3% of that of HCl $(4 \times 10^{14} \text{ cm}^{-2})$ in NGC 6334 I, the abundance of H_2F^+ is found to be 1/68 of that of H₂Cl⁺. The solar abundance ratio of F and Cl is reported to be $804/5170 \sim 1/6$ (Lodders et al. 2009) in the solar photosphere, although an old value of 1/2 is quoted in our previous paper (Fujimori et al. 2011). The small abundance of H₂F⁺ compared with the cosmic atomic abundance may be explained by differences in production and/or depletion mechanisms.

4.3. Laboratory Time-resolved Infrared Laser Spectroscopy of H_2F^+ and H_2Cl^+

Figure 4 shows the observed time-resolved spectrum of the ${\rm H_2F}^+\nu_3$ band, $J_{\rm KaKc}=3_{13}\leftarrow4_{14}$ transition at 3251.99 cm⁻¹

(Fujimori et al. 2013). We also measured the time profile for the H₂Cl⁺ ν_3 , $J_{\text{KaKc}} = 6_{06} \leftarrow 5_{05}$ and $6_{16} \leftarrow 5_{15}$ transitions at 2691.24 cm⁻¹ (Lee et al. 1988), as shown in Figure 5. The time profile shows a rise in absorption due to the ion at the beginning of the pulsed discharge, and after turning off the discharge the absorption intensity of the ion signal decreases with a half-life of a few tens of microseconds. The rotational relaxation time of the ground state is expected to be less than $1 \mu s$, when we assume a pressure-broadening coefficient of 10–20 MHz Torr⁻¹. Therefore, the observed decay is thought to be due to the recombination reaction with an electron. There are many species in the discharge in addition to the reactant F_2 , He, and H₂ species. If a product species with a larger proton affinity than HF is present, the proton of H₂F⁺ moves to the species, and the decay of H₂F⁺ will be affected compared with the case of just the recombination reaction with an electron. However, the proton transfer occurs at the Langevin rate $(\sim 10^{-9} \text{ cm}^{-3} \text{ s}^{-1})$, which is two orders of magnitude smaller than the rate of the recombination reaction. Among reactants and products in the discharge, F, F2, He, H, and H2 have smaller proton affinities than HF. On the other hand, HF dimer and trimer are thought to have larger proton affinities than HF. However, the spectra of these species were not observed by FTIR (Fujimori et al. 2013) in the same discharge condition, indicating much smaller abundances than HF. So the effects of proton transfer reactions will be neglected in the analysis of the decay of H_2F^+ .

In a reaction $N^+ + e^- \rightarrow$ products, the time variation of the ion density is expressed as follows:

$$-\frac{d[N^{+}]}{dt} = k_{e}[N^{+}][e^{-}].$$
 (6)

When the density of positive ions is equal to that of electrons, the following relation is attained (the second-order reaction scheme):

$$\frac{1}{[N^{+}(t)]} = k_e \times t + \frac{1}{[N^{+}(0)]}.$$
 (7)

In the case of H_3^+ , Amano (1988) could explain the observed decay by Equation (7). On the other hand, the observed decay of H_2F^+ of Figure 4(a) was not fitted to the formula of Equation (7) as shown in Figure 4(b), and the decay was found to be fitted to an exponential function as shown in Figure 4(c). This indicates that the reaction proceeds in the pseudo-first-order scheme. That is, by assuming $[e^-] \gg [N^+(t)]$,

$$\ln[N^{+}(t)] = -k_{e}[e^{-}]t + \ln[N^{+}(0)]$$
 (8)

where $[N^+(0)]$ is the abundance at the starting time of decay t=0. For H_2Cl^+ , the same exponential fitting as H_2F^+ was found to be better. In the first-order reaction scheme, the electron density is higher than that of H_2F^+ or H_2Cl^+ . For neutrality of the discharge plasma producing H_2F^+ , other positive ions should carry the positive charge. Although positive ions such as F_2^+ , F^+ , H^+ , He^+ , and protonated $(HF)_2$ will be candidates, we did not monitor these species in the present experiment. In a pure hydrogen discharge, the H_3^+ ion was observed by FTIR (Fujimori et al. 2013), but the spectrum disappeared when the F_2/He gas mixture was added to the discharge. In the present discharge, the H_2F^+ ion is thought to be produced by reactions H_2^+ + HF and HF^+ + H_2 , where we monitored the HF absorption with FTIR. The HF^+ ion was observed by laser

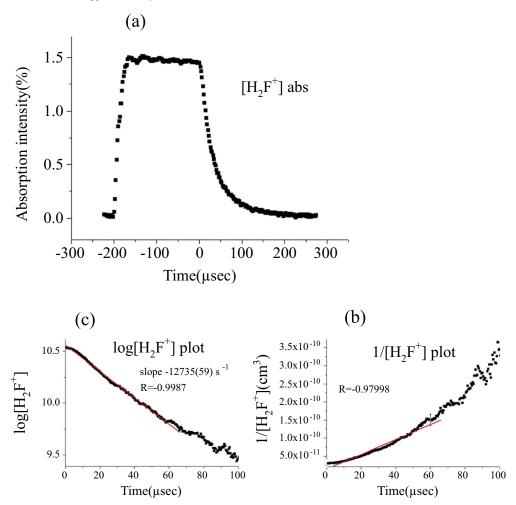


Figure 4. (a) Time-resolved spectrum of the $H_2F^+\nu_3$ band, $J_{KaKc}=3_{13}\leftarrow 4_{14}$ transition at 3251.99 cm⁻¹ obtained by a pulsed discharge with peak current of 200 mA, (b) 1/N fit, (c) exponential fit.

spectroscopy (Hovde et al. 1989) in a discharge of a mixture of a large amount of He and HF. In the present condition, the ion was not observed with FTIR in the 3.3 μ m region, because of rapid reaction with H₂.

By least-squares fittings of the observed decay curves, we determined the $k_e[e^-]$ values for H_2F^+ and H_2Cl^+ , as shown in Figure 6, where the error bars correspond to the range of several measurements at a constant value of current. As is known in a glow discharge (Teii 1986), Figure 6 shows that the electron density is proportional to the discharge current. To determine k_e , we have to obtain the electron density $[e^-]$, so we utilized a method with a single Langmuir probe (von Engel 1983; Teii 1986). When we set a discharge in a mixture of 9.3 Pa F₂/H₂ and 44 Pa H₂ with a DC current of 200 mA at the probe position between two electrodes, the density was obtained to be $(2.65-2.85) \times 10^{11}$ electrons cm⁻³. In the case of a pure hydrogen discharge (H₃⁺ production), a density of $(2.55-2.75) \times 10^{11}$ electrons cm⁻³ was obtained. Since the difference in the density between conditions producing H₃⁺ and H₂F⁺ was small, we assumed the same density for the production of H₂Cl⁺. By averaging three measurements, we obtained [e⁻] = $(2.78 \pm 0.56) \times 10^{11} \, \text{cm}^{-3}$, where the error corresponds to the range of the observed values.

We also derived the ion density at t = 0 for comparison with the electron density, although the value is not necessary for deriving the rate constant in the case of the first-order reaction scheme. The procedure is described by Bernath (2005), and we used the transition moment of 0.322 Debye for the $H_2F^+\nu_3$ band, reported by Bunker et al. (1990). The rotational temperature was fixed at the value of 209 K determined from Fourier-transform absorption spectroscopy of H_3^+ (Fujimori et al. 2013). The partition function was calculated numerically from the lowest energy level, until the new value does not contribute to 1% of the summed total value. The estimated abundance of H_2F^+ is $(3.3\pm0.7)\times10^{10}\,\mbox{cm}^{-3}$, where we confirmed that the ion density is an order of magnitude smaller than the electron density $[\mbox{e}^-]$ in the present experimental condition.

By using the $k_e[e^-]$ value of Figure 6 at 200 mA, we obtained the following dissociative recombination rate constants:

$$k_e(\text{H}_2\text{F}^+) = (1.08 \pm 0.29) \times 10^{-7} \,\text{cm}^3 \,\text{s}^{-1}$$

 $k_e(\text{H}_2\text{Cl}^+) = (0.46 \pm 0.05) \times 10^{-7} \,\text{cm}^3 \,\text{s}^{-1}$.

The present value for H_2F^+ is in agreement with that obtained by the flowing afterglow experiment, and eight times smaller than the value calculated from Equations (1) and (2) at 209 K. It is noted that in interstellar space the ion is distributed in only the lowest rotational levels of the ortho and para states. On the other

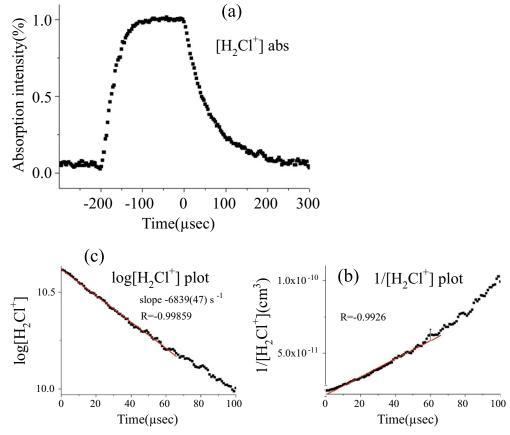


Figure 5. (a) Time-resolved spectrum of the $H_2Cl^+\nu_3$ band, $J_{KaKe}=6_{06}\leftarrow5_{05}$ and $6_{16}\leftarrow5_{15}$ transitions at 2691.24 cm⁻¹ obtained by a pulsed discharge with peak current of 300 mA, (b) 1/N fit, (c) exponential fit.

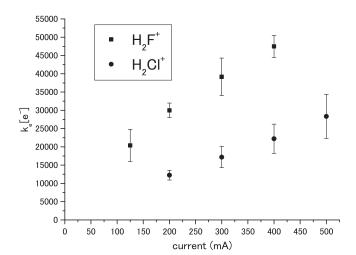


Figure 6. Product of the dissociative recombination constant (k_e) and electron density $[\mathrm{e}^-]$ obtained by the fittings of decay curves at various values of discharge current.

hand, the present laboratory measurement for H_2F^+ was carried out for the transition from the seventh lowest energy level of ortho, because of weak intensities of transitions from the lowest 1_{01} level. In the case of H_3^+ and HCO^+ , the k_e values for several rotaional states are in agreement within the measurement error limit (Amano 1990). So we assume the same rate for the laboratory and the astronomically observed levels. The k_e value of H_2Cl^+ is half of that given by Equation (3), and a factor 2 larger than that of HCl^+ at 209 K (Novotny et al. 2013).

5. DISCUSSION

The k_e value of H_2F^+ has been determined experimentally at 209 K. The present method is not applicable for lower temperatures, because of the adhesion of the reactants on the glass surface. In interstellar diffuse clouds, the translational temperature is expected to be around 50 K, so we estimated the k_e value by using temperature-dependent terms of Equations (1)–(3) and the present values for the magnitude terms, and found $k_e(50 \text{ K}) = 2.2(5) \times 10^{-7}$ and $1.6(4) \times 10^{-7}$ 10⁻⁷ cm³ s⁻¹ for H₂F⁺ and H₂Cl⁺, respectively. The difference in dissociative recombination rates between H₂F⁺ and H₂Cl⁺ is not large. Since the temperature-dependent terms of Equations (1)–(3) have not been confirmed experimentally, there may be ambiguity in the estimation. However, even with the same temperature dependence for both rates, the difference is only a factor of 2.3. Therefore, the observed small interstellar abundance of H₂F⁺ compared with that of H₂Cl⁺ is thought to be mainly due to the production mechanism.

The present value of $k_e(50 \text{ K})$ for H_2Cl^+ is 3.4 times smaller than that derived from Equation (3). This smaller rate increases the abundance of H_2Cl^+ and is in better agreement with astronomical observations, where the observed H_2Cl^+ ion is reported to be 5–10 times (Neufeld et al. 2012, 2015) more abundant than the estimate in previous chemical models of diffuse clouds.

The ionization potentials of Cl and HCl are smaller than that of hydrogen. On the other hand, F and HF have larger ionization energies than hydrogen. The $\rm H_2Cl^+$ ion is thought to

be produced via two routes in interstellar space:

$$Cl^+ + H_2 \rightarrow HCl^+ + H \tag{9}$$

$$HCl^{+} + H_{2} \rightarrow H_{2}Cl^{+} + H$$
 (10)

$$HCl + H_3^+ \rightarrow H_2Cl^+ + H_2.$$
 (11)

In diffuse clouds, the presence of HCl⁺ is important for production of H₂Cl⁺ by a reaction with H₂, and the ion has been detected by Herschel HIFI (De Luca et al. 2012). On the other hand, F and HF are not ionized by the interstellar radiation field, although the reaction of F⁺ with H₂ produces HF⁺. Therefore, a possible route for H₂F⁺ production is thought to be

$$HF + H_3^+ \to H_2F^+ + H_2$$
 (12)

with a formation rate $k_f(H_3^+) = 1.2 \times 10^{-8} (T/300 \text{ K})^{-0.15} \text{ cm}^3 \text{ s}^{-1}$ (Neufeld et al. 2005). Recently, the detection of CF⁺ toward PKS 1830-211 (SW) has been reported with a column density of $5.5 \times 10^{12} \, \mathrm{cm}^{-2}$ (Muller et al. 2016). If the reaction of HF and C+ is the main route for the production of CF⁺, the CF⁺ abundance is given in the steady-state approximation as follows:

$$[CF^{+}] = \frac{k_f(CF^{+})[HF][C^{+}]}{k_e(CF^{+})[e^{-}]}$$
(13)

where $k_f(CF^+)$ denotes the formation rate of CF^+ by the reaction $HF + C^+$, and $k_e(CF^+)$ the dissociative recombination rate for CF^+ + e. A similar formula is given for H_2F^+ :

$$[H_2F^+] = \frac{k_f(H_2F^+)[HF][H_3^+]}{k_e(H_2F^+)[e^-]}.$$
 (14)

From Equations (13) and (14) the abundance ratio of H_2F^+ and CF⁺ is given as follows:

$$\frac{[H_2F^+]}{[CF^+]} = \frac{k_f (H_2F^+)k_e (CF^+)[H_3^+]}{k_f (CF^+)k_e (H_2F^+)[C^+]}$$

$$= 1.7 \frac{k_e (CF^+)[H_3^+]}{k_e (H_2F^+)[C^+]} = 1.6 \frac{[H_3^+]}{[C^+]} \tag{15}$$

where the last part is obtained by using the present $k_e(H_2F^+)$. Gerin et al. (2015) reported an abundance ratio [C]/ [H] = $(1.5 \pm 0.4) \times 10^{-4}$ in diffuse clouds by Herschel observation of C^+ . The $[H_3^+]$ abundance is estimated to be ζ $[H_2]/k_e(H_3^+)[e^-]$ (McCall et al. 2002), where ζ is the cosmicray ionization rate and $k_e(H_3^+)$ the dissociative recombination rate constant. Considering the fraction of protons in H₂ (McCall et al. 2002), when we assume $\zeta = 2 \times 10^{-16} \text{ s}^{-1}$ (Indriolo et al. 2007), $k_e(H_3^+) = 2 \times 10^{-7} \text{ cm}^3 \text{ s}^{-1}$, and $[e^{-}] = [C]$, the H₂F⁺ abundance is expected to be 3.5% of that of CF^+ , that is, $[H_2F^+] = 1.9 \times 10^{11} \text{ cm}^{-2}$ toward PKS 1830-211, which is smaller than the present upper limit of $8.8 \times 10^{11} \, \mathrm{cm}^{-2}$. Non-detection of $\mathrm{H}_2\mathrm{F}^+$ toward W31C, NGC 6334 I, and PKS 1830-211 may be explained by the low abundance of H₃⁺. On the other hand, the Galactic center source GC IRS 21 has been reported with a large H₃⁺ column density of $61.3 \times 10^{14} \,\mathrm{cm}^{-2}$ (Goto et al. 2008). However, the continuum level in the 760 GHz region is too low to detect absorption lines of the H₂F⁺ ion. Only five sources have been observed in the present study, so further searches will be

necessary to detect the H₂F⁺ion in sources with high H₃⁺ abundance and a high continuum level.

6. SUMMARY

The HF molecule was detected in the z = 0.89 absorber in front of PKS 1830-211 with ALMA, and its abundance relative to H_2 (>1.7 × 10⁻⁸) is similar to that of other diffuse clouds in the Galaxy, indicating that HF is an excellent tracer of molecular hydrogen even in the interstellar medium of distant galaxies. In spite of the abundant HF, the protonated H_2F^+ ion is detected neither toward PKS 1830–211 nor toward a sample of four Galactic sources (W31C, NGC 6334 I, IRS 21, and 2Mass J1747). In contrast, the H₂Cl⁺ ion is detected with a column density more than an order of magnitude higher than that of H₂F⁺ in the observed sources. Laboratory experiments on dissociative recombination reactions provide a smaller rate for H₂F⁺ than the previous prediction.

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