## Moisture sources and dynamics over southeastern Tibetan Plateau

# 2 reflected in dual water vapor isotopes

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#### Abstract

The southeastern Tibetan Plateau (SETP) has experienced a significant drying trend in recent decades, likely linked to changes in moisture sources. Water vapor isotopes are valuable tracers of the atmospheric water cycle, yet their interpretation is hindered by ambiguities in atmospheric controls. The Tibetan Plateau (TP) serves as a water tower for major rivers in Asia, and mountain valleys in southeastern TP are key channels for moisture entering the TP. Water resources on the TP are experiencing spatially opposite changes due to climate change, and understanding the sources and dynamics of atmospheric moisture is vital. To investigate the role of ocean surface evaporation, continental air mass intrusion, and rain-vapor interaction, we present a three-year daily time series of near-surface water-vapor—<u>isotope compositions</u> ( $\delta^{18}$ O and d-excess) from the <u>South East TPSETP</u> station. We find Our analysis reveals that apparent negative correlations between d-excess and relative humidity over the Indian Ocean <u>are mainly</u>

reflect primarily driven by their similar seasonalitysimilar seasonal patterns, which become . When analyzed for different seasons, the correlation is insignificant or only explains a marginal fraction of variancewhen examined seasonally. Therefore, This result underscores the need for caution is required when in interpreting the d-excess as a conservative tracer of ocean surface evaporation. Instead, we identify local and upstream specific humidity is as the main-primary factor determining determinant of non-monsoon season d-excess variability, due to influenced by the intrusion of cold and dry air from upper levels. During the summer monsoon season, both d-excess and  $\delta^{18}O$  mainly reflect the effect of raindrop evaporation—on humidity during transport, which decreases—lower vapor  $\delta^{18}O$  but increases d-excess values. These findings provide new insights into the significance of usinguse of water isotopes to track moisture sources and dynamics over the SETP<sub>2</sub> with-particularly under varying seasonally—seasonal alternating—circulation systems. Particularly, the findings for d-excess will improve contribute to our—the understanding of different moisture sources and guide-provide a framework for the interpreting ation of d-excess derived from other water bedies in various hydroclimatic applications, including and ice core studies.

## 1 Introduction

The Tibetan Plateau (TP) and its surrounding regions, also termedknown as the Third Pole and the Asian Water Tower, form the highest and largest plateau on Earth that influences climatic and hydrological systems at regional to global scales, such as the formation of the Asian Summer Monsoon (Wu et al., 2022; Yao et al., 2022). In addition, the TP stores the largest amount of frozen water outside of polar regions and sustainsplays a crucial role in supplying water freshwater supplies of to major river systems in Asian rivers Asia, including the Mekong, Salween, Ganges, Yarlung Zangbo, among others, sustaining ecosystems and populations across the continent (Immerzeel et al., 2020; Yao et al., 2022). HoweverIn recent decades, the water balance on the TP has experienced undergone significant

changes under the backdrop of global warming (Yao et al., 2022). Notably For instance, the southeastern TP (SETP) is experiencing a drying trend while wetting in the northern TP (Jiang et al., 2023; Zhang et al., 2023; Yao et al., 2022). Atmospheric water vapor is the primary input to the hydrological system, making it essential to understand its sources and dynamics to diagnose regional water imbalances. Atmospheric water vapor is the input of the water storage system and understanding its sources and dynamics is vital for understanding the imbalance of TP's hydrological system. Using a Lagrangian vapor tracking method, Zhang et al. (2023) suggested that the drying trend is associated with meteorological droughts propagating from moisture source regions. However, their conclusions and methodology are subjects of ongoing debate (Zhang et al., 2025; Zhao et al., 2025). — Water stable isotopes are natural tracers of the water cycle, offering valuable insights into moisture sources and dynamics (Bowen et al., 2019; Galewsky et al., 2016). and These isotopes have been intensively studied on the TP in precipitation, surface water, and ice cores (Yao et al., 2013; Thompson et al., 2024; Bershaw, 2018). However, the interpretation of these isotopic signals remains challenging due to complex fractionation processes and shifting circulation systems between summer monsoon and westerlies. In general, precipitation isotope ratios ( $\delta^{18}$ O and  $\delta^{2}$ H) over the southern TP have lower values during the summer monsoon season and higher values during the non-monsoon season under the influence of the westerlies (He et al., 2015; Tian et al., 2007; Guo et al., 2024; Yang et al., 2017). Recent studies have confirmed that monsoon convection at upstream along moisture transport pathways, rather than local precipitation amount, is the key process that controls summer monsoon season precipitation  $\delta^{18}$ O over southern TP (Cai et al., 2017; He et al., 2015). This is related to the "amount effect" (Dansgaard, 1964), Different processes have been proposed to elucidate the relationship between precipitation 8<sup>18</sup>O and convection where the amount effect (Dansgaard, 1964) is present (Bowen et al., 2019; Galewsky et al., 2016). A relatively classical interpretation is that where higher precipitation leads to lower  $\delta^{18}$ O values due to the continuous rainout associated with stronger convection could cause depleted

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precipitation, following the Rayleigh distillation (Cai and Tian, 2016; Scholl et al., 2009; Vuille et al., 2003). Additionally, Another interpretation emphasized the role of rain-vapor interaction interactions between rain and water vapor play a significant role that partial evaporation of raindrops formed at higher altitudes isotopicallyin depletes depleting the lower tropospheric water vapor and then affects subsequent precipitation isotope compositions isotopes (Risi et al., 2008a; Kurita et al., 2011; Cai et al., 2018; Lee and Fung, 2008). While the regional amount effect prevails during the monsoon season, this relationship weakens or reverses in the non-monsoon season when it is dominated by westerlies. This variability suggests additional controls such as moisture source variability, kinetic fractionation, or shifts in atmospheric circulation patterns (Breitenbach et al., 2010; Cai and Tian, 2020; Guo et al., 2024; Yao et al., 2013). Observations of vapor isotope-compositions could help disentangle the different processes involved in the amount effect, especially particularly through examining the secondary parameter deuterium excess (d-excess). The d-excess, is defined as  $\delta^2 H - 8\delta^{18}O$  by Dansgaard (1964) as  $\delta^2 H - 8\delta^{18}O$ , and mainly primarily reflects the effects of kinetic fractionation. The During rainout process, mostly involves equilibrium fractionation is the dominant mechanism, whereas while raindrop evaporation is associated with kinetic fractionation, and they can therefore have different d excess signatures in water vapor. Further, limited precipitation during non-monsoon seasons makes it challenging to study a full seasonal cycle of the atmospheric water cycle, which can be compensated by continuous monitoring of vapor isotopes. While a few stations on the TP have monitored Lisotopic compositions in the vapor phase have only been observed at a few stations on the TP and isotope ratios (δ values) have been the major focus of previous studies (Tian et al., 2020; Dai et al., 2021; Chen et al., 2024; Yu et al., 2016; Yu et al., 2015), less is knownthere is limited knowledge about vapor d-excess. It is less certain regarding what caused higher isotope ratios during the non-monsoon season. Following the regional amount effect (Galewsky et al., 2016; Bowen et al., 2019), high 8<sup>18</sup>O values could be explained by

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weakened convection during the non-monsoon season. However, precipitation δ<sup>18</sup>O is lower during late- to postmonsoon season in regions extending from the southeastern TP to the head of the Bay of Bengal (BOB), which is not consistent with the weakening of convection (Breitenbach et al., 2010; Cai and Tian, 2020). Shifts of moisture transport pathways between convection active and non-active regions have been invoked to explain this abnormal seasonal pattern (Cai and Tian, 2020; Lekshmy et al., 2022). On the other hand, higher precipitation of 180 accompanied by higher d excess during the non-monsoon season has been interpreted as more intense continental recycling or moisture from the Mediterranean delivered by the westerlies compared with moisture from the Indian Ocean during the summer monsoon (Tian et al., 2007; Yao et al., 2013; An et al., 2017; Breitenbach et al., 2010). In addition, understanding of the atmospheric water cycle for a full seasonal cycle is complicated by the lack of precipitation during the non-monsoon season which can be compensated by monitoring atmospheric water vapor isotopes as it is not limited by precipitation events. Both theoretical predictions and observations over ocean surfaces suggested indicate that d-excess reflects ocean surface evaporation conditions, such as sea surface temperature (SST) and relative humidity normalized to SST (RH<sub>SST</sub>) (Merlivat and Jouzel, 1979; Bonne et al., 2019; Liu et al., 2014; Craig and Gordon, 1965). These relationships are frequently invoked in interpreting Interpretations of d-excess over the TP-also frequently invoke these relationships with ocean evaporation conditions (Zhao et al., 2012; Shao et al., 2021; Chen et al., 2024; Liu et al., 2024). However, relationships with either RH<sub>SST</sub> or SST are much weaker than those observed over ocean surface. For instance, Shao et al. (2021) showed significant correlations between an ice core d-excess record derived from the central TP and RH<sub>SST</sub> over the northern Bay of Bengal (BOB) and Arabian Sea (AS). However, the correlation coefficient is was only -0.44 and the slope between d-excess and RH<sub>SST</sub> is as steep as with a steep slope of -0.99% % 1. This contrasts with oceanic regions where tThe slope over oceanic regions generally typically ranges from -0.3% %<sup>-1</sup> to -0.6% %<sup>-1</sup>-based on in-situ observations (Bonne et al., 2019; Liu et al., 2014; Benetti et al., 2014;

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Uemura et al., 2008), suggesting additional complexities over terrestrial areas. Moreover, mMany studies have suggested that d-excess at terrestrial sites is not a conservative tracer of evaporation conditions at from the oceanic source regions (Fiorella et al., 2018; Aemisegger et al., 2014; Welp et al., 2012; Wei and Lee, 2019; Samuels -Crow et al., 2014). Besides temporal variations In addition, ice core d-excess values at high altitudes are are generally higher than that those observed in precipitation at lower altitudes on the TP (Shao et al., 2021; Tian et al., 2001; Zhao et al., 2012; Joswiak et al., 2013; Zhao et al., 2017; Thompson et al., 2000). The reason for this discrepancy remains unclear, It is still unclear what caused the abnormally high d-excess in these high-altitude ice cores relative to precipitation and river isotope observations at lower altitudeshighlighting the need for further research to understand the mechanism driving these differences. Mountain valleys in the southeastern TPSETP are believed to beconsidered significant major moisture transport channelspathways for delivering water vapor toward the TPtransporting water vapor into the TP (Araguás-Araguás et al., 1998; Tian et al., 2007; Yao et al., 2013). Therefore To investigate these processes, we initiated started a water vapor sampling campaign at the South-East Tibetan Plateau Station for integrated observation and research of alpine environment (SETP station) in June 2015. We aim to study the moisture sources and dynamics and their influence influence on water vapor isotope compositions across different seasons. To achieve these goals, we explored the relationships between vapor isotopes and oceanic evaporation conditions, continental air mass intrusions, as well as rain-vapor interactions during different seasons. Finally, we discuss the implications of our findings for interpreting ice core records. Following the previous study (Yao et al., 2013), we define June-September (JJAS) as the summer monsoon season. In contrast, we define November-April (Nov-Apr) as the non-monsoon season and May and October as the transition between the two seasons. We will show distinct seasonal moisture sources and dynamics between the two seasons as reflected in our vapor isotope observations and Lagrangian moisture source diagnostic. Our results suggest that the apparent correlation between SETP vapor d-excess and oceanic surface evaporation

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conditions is mainly a result of their similar seasonality. Alternatively, we suggest the intrusion of dry and cold air by the westerlies from high altitudes contributes to high d-excess. In contrast, vapor  $\delta^{18}$ O and d-excess confirm the significant role of rain-vapor interaction in the amount effect during the summer monsoon season.

#### 2 Data and methods

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#### 2.1 Atmospheric water vapor sampling

Atmospheric water vVapor samples were collected at the SETP station (29°46'N, 94°44'E, 3326 m above sea level, and Fig. S1) using a cryogenic trapping method at the SETP station (29°46'N, 94°44'E, 3326 m above sea level, and Fig. S1). The sampling system includes an air pump pumping ambient air into the, cold trap, a linkedball-shaped glass cold trap, and an electric-powered system that creates and maintains a cold environment filled by 95% ethanol as cold as below -80 °C. Ambient air was pumped from an inlet at approximately positioned about 8 m above ground level through a Teflon tube to a glass trap maintained immersed in a cold environment with aat an operational temperature ofat -70 °C. The airflow rate was adjusted set to ~5 L/min, to allowing the collection of 10-20 ml of water samples throughout during each sampling session operation. Sampling durations were adjusted seasonally: During summers, the duration of each sampling operation is 24 hours in summer and extended to 48 hours in winter when necessary to ensure adequate sample volume. During dry winters, however, we increased the sampling duration to 48 hours if a 24-hour sampling period cannot guarantee enough sample amount. The sSamples were collected at 20:00 Beijing Standard Time (12:00 UTC). The efficiency of extracting water vapor from ambient airthe trapping method was tested-verified by connecting an additional cold trap to the outlet of the initial cold trapsystem, which showed and no visible condensation was noticed in the additional cold trap (Yu et al., 2015). Further comparison validation was achieved through comparisons with against direct measurements of vapor isotope composition by theusing a Picarro L2130-i Cavity Ring Down Spectroscopy (CRDS) at Lhasa, southern TP, also

confirmed confirming the reliability of this method in for sampling atmospheric water vapor sampling over the TP (Tian et al., 2020).

The sampling campaign was started on an from 25 June 2015 and ended on to 14 June 2018. In total, yielding a total of 742 samples were collected. These and all the collected samples were kept\_stored\_frozen until transportation to the laboratory for measurements analysis. Samples collected before 28 June 2016 were measured at the Key Laboratory of Tibetan Plateau Earth System, Environment and Resources, Institute of Tibetan Plateau Research, Chinese Academy of Sciences by a Picarro L2130-i analyzer. Samples collected after 28 June 2016 were measured at the Institute of International River and Eco-security, Yunnan University by a Picarro L2140-i analyzer. The isotopic values are ealibrated using three standard waters, with detailed calibration procedures described by Liu et al. (2024), and The measurements are expressed relative to Vienna Standard Mean Ocean Water 2 (VSMOW2). The precisions of measurements at both laboratories are, with precisions of 0.1% for δ<sup>18</sup>O, 0.4% for δ<sup>2</sup>H, and 1.2% for d-excess.

#### 2.2 Meteorological data

Daily local meteorological data before prior to 2018, including precipitation amount, air temperature, air pressure, and relative humidity, at the SETP station, were provided by the station at obtained from the National Tibetan Plateau-/-Third Pole Environment Data Center (Luo, 2018). Specific humidity (q) at the SETP station is was calculated from using air temperature, air pressure, and relative humidity at the stationdata following established equations outlined in (Huang, 2018). Consistent with Yao et al. (2013), we defined June-September (JJAS) as the summer monsoon season. In contrast, November-April (Nov-Apr) was designated as the non-monsoon season, with May and October considered transition periods between the two seasons.

To facilitate analyses on larger spatial seales, wWe further obtained meteorological variables (such as including

2-meter air temperature, 2-meter dew point temperature, and SST, etc.) and others at 0.25°×0.25° and hourly

resolution from the European Centre for Medium-Range Weather Forecasts fifth generation reanalysis (ERA5) (Hersbach et al., 2019). RH<sub>SST</sub> is estimated from using ERA5 2 meter meteorological data and SSTdata using:  $RH_{SST} = e_{air}/e_{sat}$ , where  $e_{air}$  is vapor pressure of air and  $e_{sat}$  is saturation vapor pressure with respect to SST. Additionally, We further obtained precipitation data at  $0.1^{\circ} \times 0.1^{\circ}$  and half-hourly resolution were obtained from the Integrated Multi-satellitE Retrievals for GPM (V07) dataset (Huffman et al., 2023). In additionMoreover, meteorological data at  $1^{\circ} \times 1^{\circ}$  and 3-hourly resolution from the Global Data Assimilation System (GDAS) are were used to calculate air massbackward trajectories (see sSection 2.4 for details).

Statistical analyses primarily involved linear correlations and regressions, with the coefficient of determination (R<sup>2</sup>) used to quantify the variance explained by each variable. In addition, we also used composite analysis to reveal relationships between variables. For example, to identify general patterns in backward trajectories associated with d-excess exceeding 30%, all the days with such high d-excess were compiled into a collection. A composite map of trajectories from this collection was then constructed to reveal typical pathways under these conditions.

## 2.3 Theoretical framework for the understanding of isotope compositions and humidity

Besides complex atmospheric circulation models, the evolution of vapor isotope compositions during different moistening and dehydration processes can be <u>understood predicted by through</u> a compilation of atmospheric processes, such as condensation, mixing, and raindrop evaporation, that lead to different pathways of isotopic evolution along atmospheric humidity (Noone, 2012; Worden et al., 2007; Galewsky et al., 2016). These process shape distinct pathways of isotopic evolution in relation to atmospheric humidity.

The Rayleigh distillation model describes  $\underline{T}_t$  the progressive condensation of water vapor and removal as rain droplets or ice is best described by the canonical Rayleigh distillation model (Dansgaard, 1964). In the Rayleigh distillation framework, condensate is removed from the air mass as soon as it forms, and  $\underline{t}_t$  is isotope ratio composition of remaining vapor, denoted as  $\delta$ , is described as  $\delta$  as  $\delta = (1 + \delta_0)(q/q_0)^{\alpha-1} - 1$ ,

where q is the specific humidity, and  $\alpha$  is the fractionation factor. A subscript of 0 refers to the initial condition of the air mass. The falling rRaindrop evaporation introduces further complexity may partially evaporate or exchange isotopes with ambient vapor. As raindrops are formed form at higher altitudes where water vapor is depleted in heavy isotopes, the their partial evaporation of raindrops would preferentially deplete its affects the surrounding water vapor, leading to isotope values lower than those predicted by Rayleigh models but increase the atmospheric humidity, which leads to values lower than that predicted by the Rayleigh distillation (Risi et al., 2008a; Worden et al., 2007). The evolution of along with under partial evaporation of raindrops can be described inis effect gives rise to a "super-Rayleigh" trajectory trajectories, characterized by an inflated ting the effective fractionation factor  $(\alpha_e)$ , defined as  $\alpha_e = (1 + \phi)\alpha$ , where  $\phi$  quantifies the degree to which deviates deviations from equilibrium. We note that Notably, Worden et al. (2007) and Noone (2012) have given different equations to quantify the deviations of from under different degrees of raindrop evaporation for such deviations, and the same deviation of from requires very different degrees of raindrop evaporation. In this study, we followthis study <u>aligns with</u> the formulations by Noone (2012). Finally, the Air mass mixing also influence of air mass mixing on humidity and isotopic compositions can be modeled through the mass balance perspective principles. When considering mixing a dry air mass mixes with a moist air massone, for instance, the specific humidity of the mixed air mass can be described as  $q = f_{dry}q_{dry} +$  $f_{moist}q_{moist}$ , where f is represents the fraction of each air mass, with  $f_{dry} + f_{moist} = 1$  the subscript denoting different air masses and. Isotopic compositions of the mixed air mass can be are similarly derived similarly by solving the mass balance equations for the light and heavy isotopes, respectively. The outcome of the mixing process leads to resulting in a hyperbolic relationship between  $\delta$  and q. In other words,  $\delta \times q$  and q should have a linear

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relationship in the mixing process (Fiorella et al., 2018). In a framework of the Keeling plots (Keeling, 1958), the

The intercept of the regression between  $\delta$  and 1/q or the slope between  $\delta \times q$  and q gives provides an

estimation estimate of the moist end member's isotope composition— (Keeling, 1958) of the moist end member.

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Assuming a surface temperature of 25 °C and relative humidity of 85%, following we utilize the evaporation model by Craig and Gordon (1965) we can derive thatto determine the isotopic composition of ocean evaporation. This results in  $\delta^{18}O$  of ocean surface evaporation is -11.5%,  $\delta^{2}H = -81.4\%$ , and d-excess = 10.6%. We use this isotopic signature of evaporated water vapor These values serve as a the wet end member to model the for modeling moistening process throughby mixing with ocean surface evaporation. For the dry end member, we consider a <u>dehydrated air mass A hypothetical dry end member</u> from the Rayleigh curve at q = 0.5 g/kg,  $\delta^{18}O = -60.3\%$ , and  $\delta^2$ H = -418.0\ldots is chosen to represent the dehydrated dry air(Fig. S2). The dehydration process by via condensation is modeled is initiated at by choosing a starting point at the mixing line with a relative humidity of 80% on the mixing line. Similarly, the "super-Rayleigh" distillation with involving partial rain evaporation is started from also begins from the samethis starting point. We explore two For the cases of "super-Rayleigh", we simulated the isotopic evolution under two\_scenarios: (Rain evap A assumes 2% rain evaporation, while and Rain evap B). \_assumes 5%, based on Following the equations from Noone (2012), Rain evap A represents that 2% of rain is evaporated and Rain evap B represents an evaporated fraction of 5%. Additionally, we consider the influence of Mixing with evapotranspiration over south Asia and the TP is another way that could modify theon atmospheric humidity and vapor isotope compositions over southeastern SETP. Accurate qQuantifyicationng of the isotopic compositions of land surface evapotranspiration is challenging. Given that the precipitation  $\delta^{18}$ O over south Asia is generally between ranges from -1.0% and to -5.0% (Bowen and Wilkinson, 2002; Terzer-Wassmuth et al., 2021) and transpiration may account constitutes two-thirds or more of evapotranspiration or more (Cao et al., 2022; Han et al., 2022; Good et al., 2015), we assume the a  $\delta^{18}$ O value of -5.0% as an upper bound for land surface evapotranspiration has a value of -5.0% as an upper bound. Similarly, we assume that thea d-excess of 15.0% for thise wet end member of land surface evapotranspiration is 15.0%.

#### 2.4 Air massBackward trajectory and moisture source diagnostic

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Air mass backward trajectories were calculated tTo investigate the air mass transport and diagnose moisture sources and transport pathways toward SETP, we calculated backward trajectories using the Hybrid Single-Particle Lagrangian Integrated Trajectory model (HYSPLIT) (Stein et al., 2015). Trajectory calculations is were driven by the meteorological data of from the GDAS. We released a Air parcels were released from 5 locations (: the studied study site and points displaced 0.2° in each cardinal direction) at. These releases occurred at 7 different vertical levels at: 10, 50, 100, 200, 300, 400, and 500 m above ground level. For each day during the sampling campaign, trajectories were initiated every 3 hours to calculate 10-day backward trajectories. In this setting, resulting in 280 trajectories were derived for each singleper day. Geographical and meteorological variables, including location, pressure, temperature, specific humidity, rainfall amount, boundary layer height, and the terrain height along the trajectories, were stored at hourly intervals. To quantify m<sub>M</sub>oisture contributions along air mass trajectories to the humidity at SETP's humidity, is quantified usingwe applied the Lagrangian moisture source diagnostic method of developed by Sodemann et al. (2008). The This method considers uses mass balance principles along the trajectories, and assigns interpreting increases in specific humidity (forward in time) as moisture uptake, and decreases in specific humidity as moisture lost loss due to precipitation. The method It also proportionally considers accounts for the decreased reduced contribution of early earlier moisture uptake due to the precipitation en route. We have previously adapted this method to quantify theidentify moisture sources forof precipitation in sub-regions of South Asia and East Asia (Cai et al., 2018; Cai and Tian, 2020). The diagnostic results suggest indicated that the approximately 5% of the moisture arriving at SETP remained unattributed fraction of moisture arriving at SETP is -5%, and therefore, confirming that 10-day trajectories are

eapable of diagnosing most of the sufficient to diagnose most moisture sources. Unlike previous applications focused

on Instead of focusing on quantifying identifying the evaporative moisture sources from the Earth's surface, in this study, we focus onthis study emphasizes the contribution of the air parcels itself themselves to the humidity at SETPSETP's humidity. This variable is readily available from the diagnostic method, and the where changes in of air parcel contributions within the boundary layer between time steps within the boundary layer is the represent moisture uptake from the Earth surface.

The moisture contribution by of an air parcel to <u>SETP's</u> humidity-at SETP gives is a measure of the importance of upstream air masses. We calculated weighted-mean values for key variables by using the moisture contribution of the air parcel along trajectories Using this variable as the weight, mean upstream geographical and meteorological variables are hence calculated as weighted means. We also applied <u>K-means clustering to group eluster analysis on</u> the trajectories, helping to visualize the identify major transport pathways using the K-means clustering method. When calculating the mean trajectory for each cluster and meteorological variables along each mean trajectory, the moisture contribution by of the air parcel is also considered as the weight to calculate weighted-weighted-means.

#### 3 Results

## 3.1 General characteristics of vapor $\delta^{18}$ O, d-excess, and local meteorological variables

In general, water vapor  $\delta^{18}$ O values are at a-lower levels during the summer monsoon season and a-higher levels during the non-monsoon season (Fig. 1a). Mean—vapor  $\delta^{18}$ O values are -18.4% for the non-monsoon season, -23.3% for the summer monsoon season, -16.9% for May, and -22.8% for October. During the onset of the summer monsoon, the vapor  $\delta^{18}$ O shows a dramatic decrease to lower values at the onset of the summer monsoon. Conversely, from the end of the summer monsoon season to spring and early summer, Without a sharp rebound to values before the summer monsoon, the  $\delta^{18}$ O value shows a gradual increase trend-from the end of the summer monsoon season toward the highest values during spring and early summer. Although the amount effect significantly influences this

region is significantly influenced by the amount effect, the scasonal variation of \_vapor δ<sup>18</sup>O does not strictly fellow the seasonal variation of align with local precipitation patterns. For instance, while local precipitation shows clear cessation ceases clearly after the summer monsoon (Fig. 1e)<sub>s</sub>—while δ<sup>18</sup>O does not rebound to the level before summer monsoon onset remains at relatively low levels. These seasonal characteristics of vapor δ<sup>18</sup>O This behavior is consistent with precipitation δ<sup>18</sup>O observed in southeastern SETP, northeast India, and Bangladesh (Yao et al., 2013; Cai and Tian, 2020; Yang et al., 2017).

In contrast to δ<sup>18</sup>OOverall, water vapor d-excess displays different seasonal dynamics, also has lower The d-excess values during are lower during the summer monsoon season and higher values during the non-monsoon season periods (Fig. 1b). Mean vapor d-excess values are 18.3% for the non-monsoon season, 11.9% for the summer monsoon season, 13.7% for May, and 14.9% for October. However, the timing of the seasonal transitions of in vapor d-excess is different differs from that of vapor \_ δ<sup>18</sup>O. The highest vapor d-excess values generally occur during winter months when the air temperature and relative humidity (RH) are at their lowest levels (Fig. 1c and 1d). Furthermore, \_-and the d-excess starts to decrease from in spring which is, earlier than the sharp drop of in vapor δ<sup>18</sup>O during at the onset of the summer monsoon.

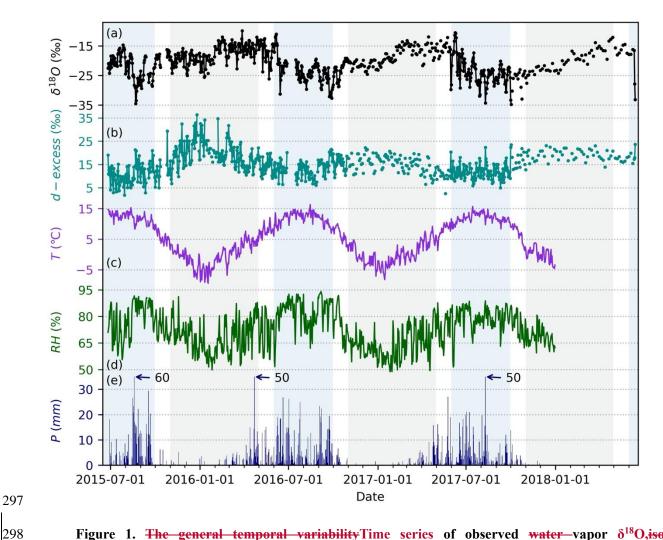
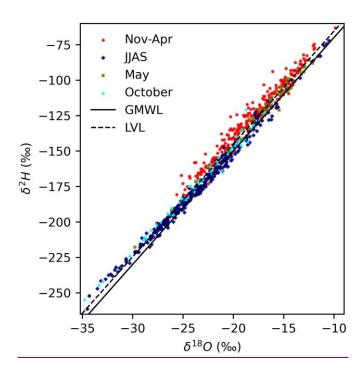


Figure 1. The general temporal variability Time series of observed water vapor  $\delta^{18}O_{*}$  isotope compositions <u>d</u>-excess, and daily local meteorological variables from 2015-2018: (a)  $\delta^{18}O_{*}$  (b) <u>d</u>-excess, (c) air temperature, (d) relative humidity (RH), and (e) precipitation amount. The <u>IL</u>ight blue shading highlights the summer monsoon season, and the <u>while</u> light steel blue shading highlights the non-monsoon season.

The linear relationship between paired  $\delta^{18}O$  and  $\delta^{2}H$  values, along with their data points and position their locations \_ relative to the global meteoric water line (GMWL,  $\delta^{2}H = 8\delta^{18}O + 10$ ) (Craig, 1961), generally provides additional insights into isotopic fractionation <u>processes</u> (Putman et al., 2019). The local <u>meteoric watervapor</u> line (LMWLLVL), estimated from all—vapor  $\delta^{2}H$  and  $\delta^{18}O$  data points, is  $\delta^{2}H = 7.96\delta^{18}O + 14.04$  (R<sup>2</sup> = 0.98). This LVL which plots above but approximately parallel with the GMWL. This relatively higher intercept of vapor LMWL LVL reflects the continental location of the site and further-additional kinetic fractionation after ocean surface

evaporation. The  $\delta^2 H$ - $\delta^{18} O$  relationship also varied seasonally. During the non-monsoon season, The vapor LMWL\_LVL for non-monsoon season is  $\delta^2 H = 7.58\delta^{18} O + 10.61$  ( $R^2 = 0.96$ ), while during the and for summer monsoon season, it shifts to is  $\delta^2 H = 7.53\delta^{18} O + 0.91$  ( $R^2 = 0.99$ ). Non-monsoon season vapor isotope compositions mainly data primarily plot above both the GMWL and even above the overall vapor LMWL\_LVL. Conversely, While the majority most of monsoon season isotope data plot fall below the overall vapor LVMWL, those data points that have though the lowest  $\delta$ - $\delta$ -values plot points during this period are positioned above the overall vapor LMWL\_LVL, indicating further suggesting additional kinetic fractionation such as rain evaporation (He et al., 2024). Vapor isotope compositions for May are more similar to resemble those during of the non-monsoon season but plot closer to align more closely with both the GMWL and LMWL\_LVL, while whereas data for October show a more exhibit similar behaviors similar to with the monsoon season observations.



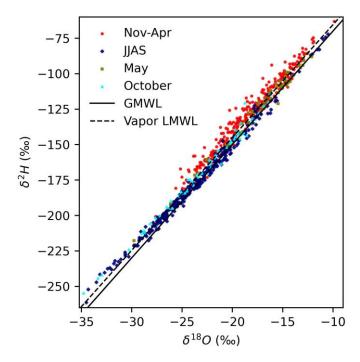


Figure 2. Relationship between vapor  $\delta^2 H$  and  $\delta^{18} O$ . The data is presented for different seasons: Data during the non-monsoon season (Nov-Apr) as red dots, the summer monsoon season (JJAS) as navy diamonds, May as olive squares, and October are shown as red dots, navy diamonds, olive squares, and cyan trangles, respectively. The solid line indicates the global meteoric water line (GMWL). The dashed line indicates the local meteoric watervapor line (LMWLLVL) estimated from all-vapor—  $\delta^2 H$  and  $\delta^{18} O$  data points.

The relationships between—wapor-δ<sup>18</sup>O and specific humidity (q) further indicate distinct seasonal patterns inseasonally contrasting moisture dynamics (Fig. 3a). Note that only data before 2018 are shown as local meteorological data are unavailable in 2018. In addition, the sampling frequency during 2018 was reduced (Fig. 1) due to logistic issues. Therefore, when analyzing relationships between vapor isotope compositions (δ<sup>18</sup>O and dexcess) and meteorological variables (both locally and regionally) we only focused on data before 2018 Due to unavilability of local meteorological data for 2018, our analyses focused on data collected before this year. For months dDuring the non-monsoon season, particularly in winter months, the majority of most data points fall-are positioned above the Rayleigh distillation line but below the mixing line that of represents an an upper bound of hypothetical evapotranspiration over South Asia, especially for the winter months. This suggests a mix between a

dry end member and a moist end member. In contrast, data forduring the summer monsoon season, months data predominately fall below the Rayleigh distillation \_\_line, influenced \_\_and are constrained by "super-Rayleigh" lines with different degrees of processes linked to rain evaporation.

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Futher insights come from examing  $\delta \times q$  The relationships between versus and q relationships, which highlight further indicate seasonal contrasts in moisture source signatures (Fig. S3). Distribution of non-monsoon season and suggests the mixing between a dry end member that has almost totally dehydrated through condensation and a moist end member of surface evaporation or moisture that has been partially dehydrated through Rayleigh distillation (Fig. S3). For the non-monsoon season, aA simple estimation through the linear regression between  $\delta \times q$  and q suggests a  $\delta^{18}$ O of the moist end member for the non-monsoon season is with an  $\delta^{18}$ O of - $13.9\% \pm 0.6\%$ . The amount weighted annual mean precipitation  $\delta^{18}$ O at this our site was about -14.5% (Yao et al., 2013). However, during the monsoon season, the overall estimation of  $\delta^{18}$ O for the moist end member through the linear regression between  $\delta \times q$  and q for the summer monsoon season suggests  $\delta^{18}O$  of the moist end member isis significantly lower at -30.9\sim \pm 1.8\significantly lower at -30.9\sim \pm 1.8\significantly lower than the estimation for the non-monsoon season. This exceptionally low value requires an additional moisture source of from rain evaporation that is much more depleted in heavy isotopes than surface evapotranspiration and is consistent with the distribution of  $\delta^{18}O$ -q below the Rayleigh distillation line (Fig. 3a). These results align with the distribution of  $\delta^{18}$ O-q data below the Rayleigh line during the summer monsoon season (Fig. 3a), underscoring the influence of different moisture sources and processes across seasons.

The relationships between vapor d-excess and q also suggest reflect seasonal contrasts in moisture dynamics (Fig. 3b). During non-monsoon season months, vapor d-excess shows a negative correlation is observed where lower with  $q_5$  corresponds to and the highest higher d-excess values (Figs. 1 and 3b). This relationship is particularly pronounced under dry and cold conditions are generally associated with the driest and coldest air (Figs. 1 and 3b).

In contrast, during the summer monsoon season, Howeverno clear relationship between, vapor d-excess does not show a clear relationship with and q is apparent, with d-excess and shows—showing a substantial considerable variability (-of approximately 20%—in range) at any given q-during the summer monsoon season. These relationships findings suggest that vapor d-excess is less predictable by using q compared to than  $\delta^{18}$ O, except for under low humidity levels.

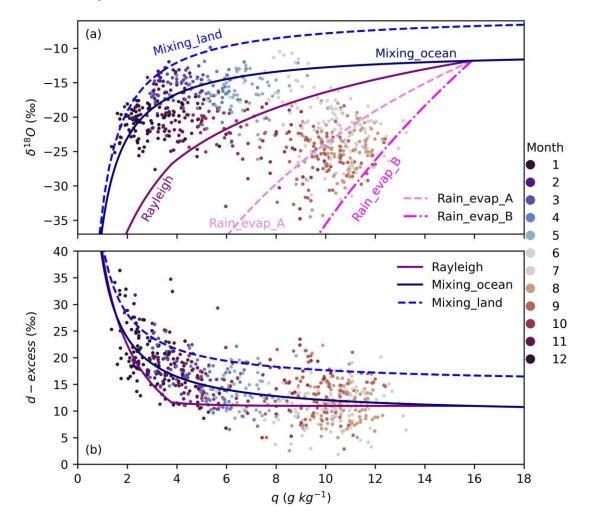


Figure 3. Relationships between vapor isotopes ( $\delta^{18}O$  and d-excess) and isotope compositions and specific humidity (q) from 2015-2017. (a) scatter plot of  $\delta^{18}O$  against specific humidity (q). (b) scatter plot of d-excess against q. Each data point is color-coded by month. The months for the data points are color-coded. The reference lines are the same ascorrespond to those in Fig. S2; and their interpretations of these reference lines are referred to is detailed in Fig. S2 and section 2.3. Note that only data before 2018 are shown (see

#### text for details).

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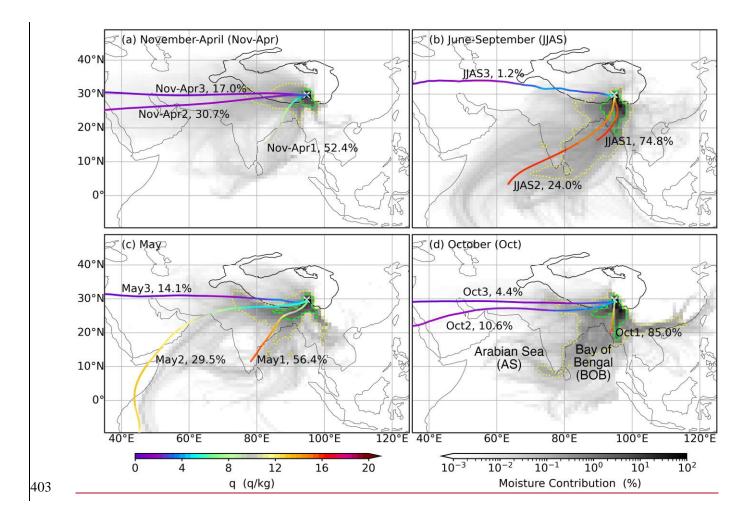
## 3.2 Seasonal variability in mMoisture sources and transport pathways for different seasons

To understand the drivers of behind the seasonal contrasting variations in moisture dynamics reflected in vapor isotope compositions, we first analyzed the moisture sources and transport pathways during different seasons (Fig. 4). Note again that we Our focus was on the contribution by of moisture from historical air masses (last 10 days) air mass to humidity at SETP-instead of moisture uptake from the Earth surface. During the non-monsoon season, moisture is mainly transported by via two pathways: branches with one originating from the west of SETP, carried by the westerlies (clusters Nov-Apr2 and Nov-Apr3), and the another from the south of SETP, such as the BOB (cluster Nov-Aprl). Quantitatively, the fraction of moisture from the south pathway and the sum of the two west branches contributions frm these pathways are is comparable, with (52.4%) from the southern pathway and vs. 47.7% from the western branches combined). Interestingly, We note that if moisture contributions by air parcels are not considered when considering only trajectories without accounting for moisture contributions, trajectories for all three clusters are from the west of SETP and they only reflect the transport of air masses appear to originate from the west of SETP (Fig. S4). These differences This discrepancy highlights the importance of distinguishing between pure air mass trajectories transport and considering moisture contribution by air masses call for cautionactual moisture sources when interpreting air mass trajectoriesy data. In contrast, dDuring the summer monsoon season, moisture transport is predominantly transported from the south of SETP, driven by the summer monsoon. The moisture sources and transport pathways for observed in May show someshare similarities with the results forthose of the non-monsoon season, but with notable differences. Compared with Specifically, the second transport-pathway during May (cluster May2) shifts southward toward the AS compared to its counterpart during the non-monsoon (cluster Nov-Apr2), the second transport pathway during

May (cluster May2) shows an overall southward shift toward the AS. Similarly, Although thewhile October's air

mass transport pattern direction during October is also similar to that mirrors that of during the non-monsoon, the moisture sources and transport pathways for October show similarities greater alignment with the results for those of the summer monsoon season, albeit with a slight eastward shift (Figs. 4d and S4d).

Another emerging feature of notable aspect of the moisture source distributions is that humidity at SETP is predominantly contributed by air masses over proximal terrestrial regions the dominant contribution from proximal terrestrial regions, especially particularly those regions in itsto the south of SETP (Fig. 4). In contrast, air masses over oceanic regions make a much smaller contribution to humidity at SETP. For instance example, the 1% contour of representing moisture contributions by from air parcels over each 1°×1° grid box does not reach extend into oceanic regions during all any of the four seasons. Therefore, moisture uptake of This indicates that surface evaporation from oceanic regions; such as from the BOB and AS; is also very limited contributes minimally. Instead, as most of the moisture in air masses originating over these oceanic regions is lost by through precipitation before reaching SETP, and what remains is replenished by evapotranspiration during the transport toward SETP over land. This result finding raises an important questions question: do whether the vapor isotopic compositions measured at SETP still preserve reflect the meteorological information conditions at their—ocean surface is sources?



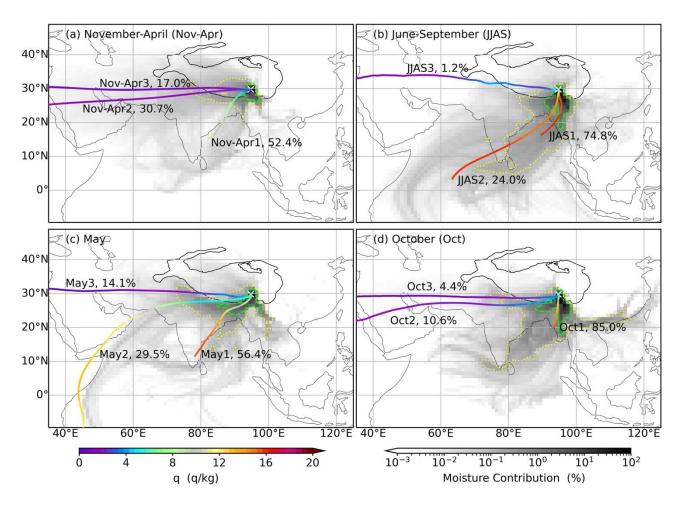


Figure 4. Moisture sources and transport pathways during different seasons from 2015-2017. (a) spatial distribution of relative contributions of moisture by from all air parcels overall each 1°×1° box (shading) to humidity at the SETP station, along with and specific humidity (q) along mean trajectories (weighted by the moisture contributions of air parcels) for the non-monsoon season of November-April (Nov-Apr). (b-d) are the same as (a), but for the monsoon season of June-September (JJAS, b), May (c), and October (d), respectively. The dotted yellow and dashed green contours indicate the moisture contribution at 0.1% and 1%, respectively. The yellow crosses indicate the location of the SETP station. The black solid lines denote the Tibetan Plateau with altitude contour at 3000 m.

#### 4 Discussion

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## 4.13.3 Role of ocean surface evaporation conditions at seasonal and intraseasonal time scales

Relationships between vapor d-excess and ocean surface evaporation conditions of, such as RH<sub>SST</sub> and SST, are first tested were examined using all the data from 2015-2017 (Fig. 5a and Fig. S5a). Results indeed show negative correlations between vapor d-excess and RH<sub>SST</sub> over northern Indian Ocean, especially particularly in the northern parts of AS and BOB (Fig. 5a). Quantitively Specifically, the regression slopes of the regression between d excess and RH<sub>SST</sub> for this relationship over across the northern Indian Ocean range vary from higher than -0.1% %<sup>-1</sup> to values below -0.6% %<sup>-1</sup>. Focusing on specific regions, The regression slopes over the northern BOB (10-22°N and 80-99°E) and the eastern AS (7-20°N and 65-78°E; Fig. 5a) fall exhibited regression slopes within the range (from -0.3% %<sup>-1</sup> to -0.6% %<sup>-1</sup>) previously reported in previous studies (Uemura et al., 2008; Benetti et al., 2014; Liu et al., 2014; Bonne et al., 2019). For instance, Vapor d-excess and the regional average RH<sub>SST</sub> in the eastern AS vields shows an overall <u>regression</u> slope of -0.49% %<sup>-1</sup> (r = -0.52 and p < 0.01) for the eastern AS (Fig. S66a) and, while the northern BOB has a slope of -0.52% %<sup>-1</sup> (r = -0.55 and p < 0.01) for the northern BOB (Fig.  $\frac{876b}{}$ ). However, upon closer inspection of the d-excess-RH<sub>SST</sub> plots (Fig. 6), it becomes evident that the distribution of data in the d-excess and RH<sub>SST</sub> space suggests a clustering of datadata points clustered according to that observations during summer months are mainly located in the lower right with high RH<sub>SST</sub> and low d-excess, but in the upper left part of the space for winter months different seasons. During each season, there is substantial variability in vapor d-excess for a given RH<sub>SST</sub>. These results suggest, implying that the apparent negative correlations between d excess and RH<sub>SST</sub> may mainlymight primarily arise stem from their oppositeng seasonalityseasonal trends. Similarly, apparent negative correlations between vapor d-excess and SST also emerge over the northern Indian Ocean (Fig. S5a). HoweverYet, both

theoretical prediction (Merlivat and Jouzel, 1979) and in-situ observations above the ocean surface (Bonne et al.,

2019; Liu et al., 2014) suggest a positive correlation between vapor d-excess and SST. Therefore, we argue These discrepancies lead us to speculate that the overall correlations between SETP vapor d-excess and surface evaporation conditions over the northern Indian Ocean are mainly a result of their seasonality likely driven by seasonal variability and do not hold realistic causal relationships. We further examined tThe relationship between vapor d-excess and RH<sub>SST</sub> was further analyzed by distinguishing betweenfor the summer monsoon and non-monsoon seasons, respectively. During the summer monsoon season, tThe negative correlation between vapor d-excess and RH<sub>SST</sub> almost totally diminishes significantly, especially during the summer monsoon season when absolute values of with correlation coefficients drop toping below 0.3 (Fig. 5b). In contrast, significant Stronger correlations present during the non-monsoon season (Fig. 5c), eould bepotentially due to the overall intraseasonal variations that where d-excess is the highest during peaks in winter and lower decreases at the beginning and ending stages of the non-monsoon season (Fig. 1b), which could be possibly accompanied with an by opposite opposing RH<sub>SST</sub> trends. Even so However, even during the non-monsoon season, the correlations during the non-monsoon still only explain a marginal fraction of the explained variance in d-excess remains low, at a maximum of (10%-16% at maximum over the northern BOB). Similarly, correlations with SST over the northern Indian Ocean also become trivial negligible when seasons are considered separately-considering the summer monsoon or non-monsoon season (Fig. S5). In summary, vapor d-excess at SETP is less likely a conservative tracer of surface evaporation conditions (neither RHSST nor SST) over the northern Indian Ocean. Therefore, it should be cautious when interpreting d-excess in meteoric water or paleo archives from the TP as a proxy of for Indian Ocean evaporation conditions over the Indian Oceanshould be approached with

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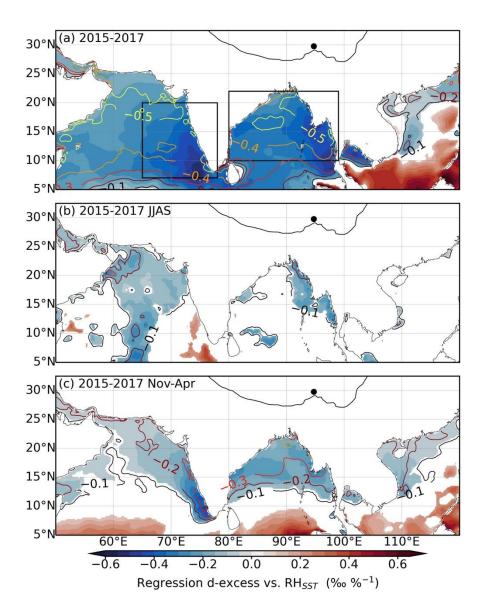


Figure 5. Relationships between water-vapor *d*-excess and relative humidity scaled to sea surface temperature (RH<sub>SST</sub>). (a) regression of vapor-*d*-excess against RH<sub>SST</sub> (shading and only values significant at the 95% significance level are shown) and correlation coefficients between them (contours at an interval of 0.1 and only negative correlations are shown) for all the data from 2015-2017. (b) and (c) are the same as (a) but only for the data within the summer monsoon season (JJAS) or the non-monsoon season (Nov-Apr), respectively. The black dots indicate the location of the SETP station. The black solid lines denote the Tibetan Plateau with altitude contour at 3000 m.

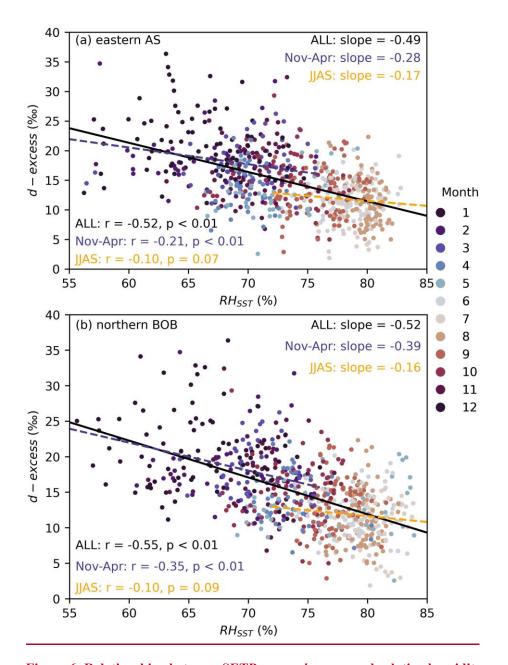


Figure 6. Relationships between SETP vapor *d*-excess and relative humidity normalized to sea surface temperature (RH<sub>SST</sub>) averaged over (a) eastern Arabian Sea (7-20°N and 65-78°E) and (b) Bay of Bengal (10-22°N and 80-99°E) from 2015-2017. Each data point is color-coded by month. Solid black lines indicate the linear regression between all data points. Dashed orange lines indicate linear regression for data during the non-monsoon season (Nov-Apr) and dashed dark blue lines for data during the summer monsoon (JJAS). The slope (% %-1), r, and p values for the three data groups are also shown.

## 4.23.4 Role of dry and cold air intrusion during the non-monsoon season

Both theoretical predictions by from the Rayleigh model and observations during the non-monsoon season suggest an increasing trend of that vapor d-excess increases as when q goes to extremely low values decreases when q reaches extremely low values (Fig. 2). In addition, results for both air mass transport and moisture transport show the dominant role of the westerlies (Figs. S4a and 4a). Therefore Based on these evidences, we hypothesize propose that during the non-monsoon season, vapor isotopes are influenced by the mixing of cold and dry air transported by the westerlies from higher altitudes with surface vapor controls vapor isotope compositions during the non-monsoon season. Furthermore, sSurface vapor influenced by recycled moisture from terrestrial evapotranspiration would further elevate vapor d-excess at a given q (Fig. 3b).

We first didperformed a composite analysis on moisture sources and transport pathways for the highest (higher than  $30\%_2$  and n = 10) and lowest d-excess observations (lower than  $10\%_2$  and n = 8) d-excess observations during the non-monsoon season (Fig. 67). For hHigh vapor d-excess values, moisture is are primarily associated predominantly with moisture transported by westerlies from the regions west of SETP, such as over the TP and northwestern India. In addition, air masses along backward trajectories for these cases are show air masses characterized by extremely low  $q_z$  i.e. reaching as low as below 2 g kg<sup>-1</sup> along the mean trajectories (weighted by the moisture contribution) over the TP (Fig. 6a7a). Conversely, fFor low d-excess cases, a substantial amountsignificant portion of moisture transport pathways (account for 39.2% by the L1 cluster, Fig. 6b) shifts toward more humid areas regions, including of northeast India, Bangladesh, and the BOB, with the L1 cluster accounting for 39.2% (Fig. 7b). This contrasting moisture transport pattern between high and low d-excess cases agrees aligns with our hypothesis that the high d-excess is associated with dry and cold air transported by the westerlies.

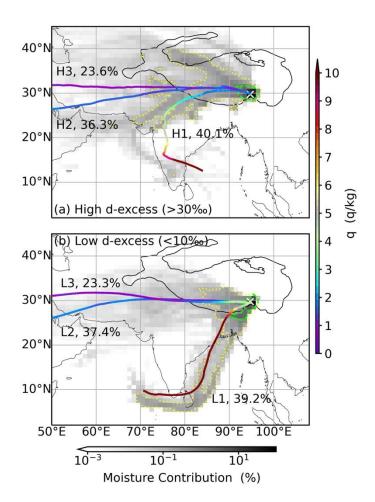


Figure 67. Composite of moisture sources and transport pathways for high and low d-excess days during the non-monsoon season of November-April. (a) spatial distribution of relative contribution of moisture by from all air parcels overall each  $1^{\circ}\times1^{\circ}$  box (shading) to humidity at the SETP station, and along with specific humidity (q) along mean trajectories (weighted by the moisture contribution—sof air parcels) for d-excess values higher than 30% during the non-monsoon season (n = 10). (b) is the same as (a) but for d-excess lower than 10% (n = 8). The yellow crosses indicate the location of the SETP station. The black solid lines denote the Tibetan Plateau with altitude contour at 3000 m.

The relationship between vapor d-excess and  $t\underline{T}$  he intrusion-influence of cold and dry air intrusions was is further investigated through an analysis of tested by relationships among involving vapor d-excess, local q, weighted-mean upstream air temperature, and weighted-mean upstream air altitude (Fig. 78).  $\underline{U}\underline{U}$  pstream variables are represent weighted averages mean values along the 10-day backward trajectory,

where weights correspond to weighted by the moisture contribution of the air parcel at each time step (sSection 2.4). The non-monsoon season vapor d-excess shows robust negative correlations with both local q (r = -0.65, p < 0.01) both at the local scale as well as at and upstream  $\underline{q}$  (r = -0.65 and -0.67, respectively, and p < 0.01 for both). At the same time Furthermore, low q is associated with air masses with characterized by low temperatures and from high altitudes (Figs.  $\frac{7e-8c}{}$  and  $\frac{7d8d}{}$ ). This effect, which could also have an impact on vapor  $\delta^{18}$ O. Indeed,  $\delta^{18}$ O during the high d-excess cases is lower than  $\delta^{18}$ O-during—the low d-excess cases (at a significance level of 95.3%), and the The overall correlation coefficient between  $\delta^{18}$ O and d-excess during the non-monsoon season is -0.29 (p < 0.01). Notably, c Correlations between  $\delta^{18}$ O and  $\underline{q}$  are weaker compared to those observed for  $\underline{d}$ -excess, with local  $\underline{q}$ (showing  $r = 0.42 \frac{\text{and}}{\text{(p}} < 0.01) \frac{\text{or and}}{\text{or and}} \text{ upstream } q \frac{\text{(showing } r = 0.38 \frac{\text{and}}{\text{(p}} < 0.01) \frac{\text{are weaker than the correlations}}{\text{or and } \text{(p)}}$ between d-excess and q. The relationship between non-monsoon season  $\delta^{18}$ O and humidity is mainly expressed as the relationship between  $\delta \times q$  and q (r = 0.82 for local q and r = 0.90 for upstream q). We further analyzed the sSpatial distribution of correlations between SETP vapor isotope compositions ( $\delta^{18}$ O and d-excess) and 2-meter air temperature as well as humidity measured by 2-meter dew point temperature during the non-monsoon seasonalso support these findings (Fig. 8S6). SResults show significant negative correlations between d-excess and dew point temperature at the regional scale exist over southeastern TP, northeast India, and northern Bangladesh (Fig. 8a). Correlations with air temperature are generally similar with correlations between dexcess and dew point temperature but the most significant correlations are in a smaller region (Fig. 8b). In contrast, correlations between  $\delta^{18}$ O and dew point temperature is not as strong as that for d-excess (Fig. 8c). Instead,  $\delta^{18}$ O shows stronger significant positive correlations with air temperature over the India subcontinent and the northwestern part of sSoutheast Asia (Fig. 8d). As shown in Fig. 3b, Eextremely high d-excess values are predicted at very low q levels are predicted in Fig.

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3b<sub>2</sub> and pPrevious study studies has have shown that as q approaches zero, vapor d-excess can approach 7000%

following the Rayleigh distillation trajectory (Bony et al., 2008), eaused by a behavior inherent to the definition of the *d*-excess (Dütsch et al., 2017). High vapor *d*-excess values have also been observed in low humidity conditions environments, such as in the polar regions (Bonne et al., 2014; Steen-Larsen et al., 2017) or at and high altitudes (Samuels - Crow et al., 2014; Webster and Heymsfield, 2003; Sayres et al., 2010; Sodemann et al., 2017). Therefore, we infer that the increasing trend of vapor *d*-excess along with decreasing local *q*, upstream *q*, and regional dew point temperature is a result of intensified due to enhanced mixing with dry and cold subsiding air transported by the westerlies from high altitudes. Relationships between upstream *q* and upstream air temperature as well as altitude further support this inference, indicating that low humidity conditions is are associated with the presence of subsiding dry and cold air from high altitudes (Figs. 7e-8c and 7d8d). Therefore, vapor *d*-excess during the nonmonsoon not only provides information on insights into the specific humidity levels but also indicates the source of humidity.

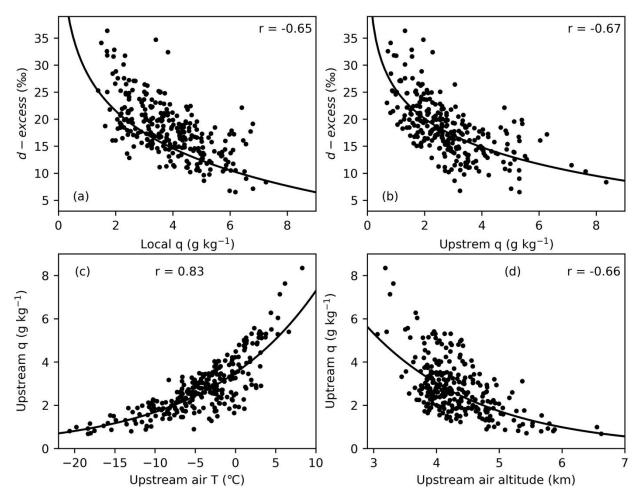


Figure 78. Relationships among water-vapor d-excess, local specific humidity (q), weighted-mean upstream q, weighted-mean upstream air temperature (T), and weighted-mean upstream air altitude during the non-monsoon season of November-April. (a) scatter plot of d-excess against local q. (b) scatter plot of dexcess against upstream weighted-mean q. (c) scatter plot of upstream q against upstream air T. (d) scatter plot of upstream q against upstream air altitude. All the upstream variables are mean values along backward trajectories weighted by the moisture contribution of air parcels. The solid curves indicate the log regression between the respective variables with the correlation coefficients indicated by the numbers.

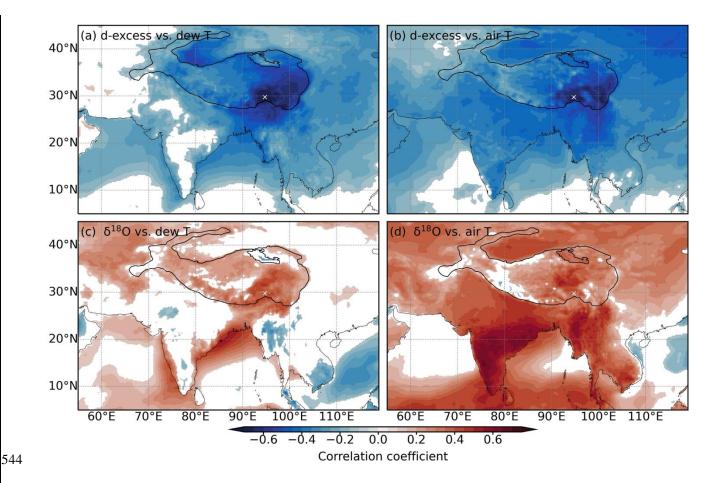


Figure 8. Spatial distribution of correlation coefficients among water vapor isotope compositions, dew point temperature, and air temperature during the non-monsoon season of November-April. (a) spatial distribution of correlation coefficients between SETP vapor d-excess and 2-meter dew point temperature. (b) the same as (a) but with 2-meter air temperature. (c) the same as (a) but between  $\delta^{18}$ O and 2-meter dew point temperature. (d) the same as (e) but with 2-meter air temperature. Only values significant at the 95% significance level are shown. The white crosses indicate the location of the SETP station. The black solid lines denote the Tibetan Plateau with altitude contour at 3000 m.

## 4.33.5 Role of rain-vapor interaction during the summer monsoon season

Different from In contrast to the significant dependence of vapor d-excess on qspecific humidity during the non-monsoon season, vapor d-excess is not correlated shows no correlation with qspecific humidity (r = 0.04 and p = 0.51) during the summer monsoon season. The behavior of  $\delta^{18}$ O is also distinct during differs between the two

seasons (Fig. 3). During the Distribution of summer monsoon season, observations in the  $\delta^{18}$ O-q plots space below the Rayleigh curve, suggests indicating that the vapor has undergone a certain degree of rain-vapor interaction by raindue to evaporation (Fig. 3a). On the other hand, pPartial rain evaporation in an unsaturated atmospheric environment is associated leads to with kinetic fractionation, which decreases the d-excess values of their raindrops but increases the while increasing d-excess in the of surrounding vapor (Risi et al., 2008b). This effect of rain-vapor interaction on vapor isotope compositions has been suggested as a major process responsible forprimary mechanism driving the amount effect in the tropicsal regions (Risi et al., 2008a; Kurita et al., 2011; Bowen et al., 2019; Galewsky et al., 2016). Therefore, we hypothesize that vapor isotope-compositions during the summer monsoon season at SETP are controlled influenced by the degree of rain-vapor interaction. The first evidence supporting this hypothesis is that the significant correlation between— vapor—δ<sup>18</sup>O is significantly correlated with and d-excess during the summer monsoon season (r = -0.55-and, p < 0.01-; Fig. 9a). In addition, there is a trend that where - vapor  $\delta^{18}$ O and d-excess are less correlated exhibit weaker correlations when  $\delta^{18}$ O is high and the opposite for low  $\delta^{18}$ O levels are high, and stronger correlations when  $\delta^{18}$ O levels are low (Figs. 9a and S8S7). If there is no rain, rain-vapor interaction is not possible. Therefore To explore this further, we flitted data during days when the categorized days with daily precipitation amount is not less than of at least 2 mm as "rainy days" and days those with precipitation less than 2 mm as no rain occurs locally ("non-rainy days"). This distinction is based on the premise that rain-vapor interaction cannot occur in the absence of rainfall. The analysis reveals that Vapor  $\delta^{18}$ O during rainy days is significantly higher-lower than that during non-rainy days, -and while d-excess show the opposite trend applies for d-excess (p < 0.01 for both  $\delta^{18}$ O and d-excess) (; Figs. 9b and 9c). Furthermore, <u>t</u>The correlation between—<u>vapor</u>  $\delta^{18}$ O and *d*-excess <u>becomes stronger during on</u> rainy days <del>becomes stronger</del> (r = -0.69-and, p < 0.01). However, vapor  $\delta^{18}$ O is still negatively correlated with d-excess during, though a weaker

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<u>negative correlation persists even on</u> non-rainy days (r = -0.40-and, p < 0.01). Even if when applying a stricter

threshold of 0 mm for daily precipitation amount is used for flittering-non-rainy days, there is still athe significant negative correlation between—vapor— $\delta^{18}$ O and d-excess remains significant (r = -0.37—and, p < 0.01). In additionMoreover, correlations with local precipitation amount are weak—both for both  $\delta^{18}$ O (r = -0.31—and, p < 0.01) and d-excess (r = 0.26—and, p < 0.01). Therefore, we further These findings lead us to infer that vapor isotopes during the summer monsoon season at SETP are influenced not only by local the effect of rain-vapor interactions is not only from the local scale—but also inherit by the history of rain-vapor interactions before vapor has been transported to SETP that occurred before the vapor reached the region.

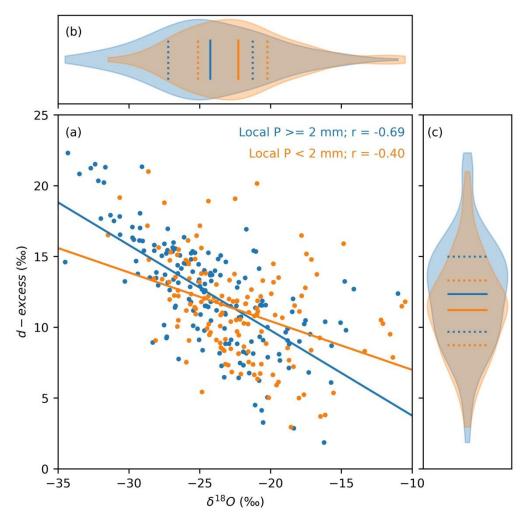


Figure 9. Relationships between SETP vapor d-excess and  $\delta^{18}O$  during the summer monsoon season. (a) scatter plot of d-excess against  $\delta^{18}O$  and linear regression lines between them. (b) distribution of  $\delta^{18}O$  values with the dashed lines indicate values at the lower and upper quartiles and the solid lines indicate the mean

values. (c) is the same as (b) but for *d*-excess. Orange colors indicate data observed during daily precipitation amount less than 2 mm and blue colors indicate data observed during days with precipitation amount not less than 2 mm. The r values for both lines are indicated in (a) and both of them are significant at the 0.01 level.

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If there is a larger amount of rainfall, the effect of rain-vapor interaction on atmospheric humidity would be stronger. Therefore To further investigate the role of rain-vapor interactions, we use total precipitation amount ( $P_{acc}$ ) as a measure of an indicator of rain-vapor interaction, considering the cumulative effect over several days preceding sampling. To account for the history during moisture transport, the total precipitation amount during several days before sampling is considered. We have tested Our analysis examined correlations the relationships between vapor isotope compositions ( $\delta^{18}$ O and *d*-excess) and  $P_{acc}$  over for periods ranging from 1-10 days prior to sampling (Figs. 89-88 and 81089). Vapor d-excess reaches an optimal correlation with  $P_{acc}$  when the total precipitation amount during considering 3 days before sampling ( $P_{acc\ 3d}$ ) is considered. Vapor  $\delta^{18}$ O shows a slightly longer memory and reaches an optimal correlation with Page when the total precipitation amount during around 5-6 days before sampling is considered. Fig. 10 shows the spatial distribution of these correlations, between vapor isotope compositions (δ<sup>18</sup>O and d excess) and  $P_{acc-3d}$ . Vaporwhere d-excess shows significant positive correlations positively correlates with Pacc 3d in the region surrounding SETP with a spatial scale of across a ~5°×5° region surrounding SETP and the positive correlation extendsing southwestward to the foothill of the Himalayas (Fig. 10a). In contrast, – vapor δ<sup>18</sup>O shows significant negative correlations with Pacc 3d in similar regions (Fig. 10b). Interestingly, even For on non-rainy days, vapor  $\delta^{18}$ O and d excess still show significant regional-scale correlations with  $P_{acc-3d}$  at regional scale persist, albeit with weaker correlation levels and with a smaller spatial extent (Fig. S11S10).

These significant correlations among vapor  $\delta^{18}$ O, d-excess, and  $P_{acc\_3d}$  findings provide further evidence for understanding processes that are responsible forthe mechanisms driving the amount effect. The negative correlation

between  $\delta^{18}$ O and  $P_{acc\_3d}$  has also been observed in precipitation and can be interpreted in terms of attributed to either continuous rainout (Scholl et al., 2009; Vuille et al., 2003; Ruan et al., 2019) or the effect of rain-vapor interactions (Lawrence et al., 2004; Risi et al., 2008a; Kurita et al., 2011; Worden et al., 2007). Although While continuous rainout, explained by the Rayleigh distillation model, with increased rainfall amount can explain accounts for the decreasing trend of  $\delta^{18}$ O with increased rainfall by the Rayleigh distillation model, d-excess stays-remains at a relatively stable level when unless specific humidity is not very low drops to very low levels (above  $\sim$ 4 g kg $^{-1}$  in Fig. 3b for example). Therefore, tThe positive correlation between vapor d-excess and  $P_{acc\_3d}$  provides an additional constraint, suggesting that the amount effect is not simply solely a result of rainout but rain vapor interaction plays an important role in altering lower tropospheric isotope compositions also involves rain-vapor interactions, which significantly influence vapor isotopes in the lower troposphere.

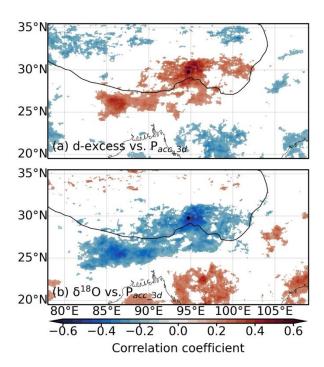


Figure 10. Relationships between vapor isotope compositions for rainy days (local daily precipitation amount not less than 2 mm) and total precipitation amount at the regional scale during the summer monsoon season. (a) spatial distribution of correlation coefficients between vapor d-excess and total precipitation amount during 3 days prior sampling ( $P_{acc\_3d}$ ). (b) is the same as (a) but for  $\delta^{18}$ O. Only values significant at

the 95% significance level are shown. The black dots indicate the location of the SETP station. The black solid lines denote the Tibetan Plateau with altitude contour at 3000 m.

Interpretationsing of d-excess in meteoric water and ice cores on the TP are is complicated by evaporation

## 4 Implications for interpreting TP ice core isotope data

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## 4.4 An alternative interpretation for the high d-excess in high-altitude TP ice cores

conditions over the northern Indian Ocean (RH<sub>SST</sub> and SST) and continental recycling (Shao et al., 2021; Zhao et al., 2012; Joswiak et al., 2013; Pang et al., 2012; An et al., 2017). Attempts have been made to establish a relationships between vapor d-excess and RH<sub>STT</sub> (Chen et al., 2024; Liu et al., 2024) as well as between ice core dexcess and RH<sub>STT</sub> (Shao et al., 2021) or SST (Zhao et al., 2012). Based on relationships between vapor d-excess and surface evaporation conditions discussed above results in Section 3.3, however, the apparent relationships are mainly <u>primarily</u> a result of similarities in the seasonality of these variables. Furthermore, the direct contribution of oceanic water vapor contained in air masses over oceanic regions to humidity at SETP is very limited (Fig. 4), which implies implying an even smaller contribution that the contribution to humidity over the TP since SETP is further decreased than at SETP as it is at the forefront of moisture transport toward TP (Fig. S1). The dominant terrestrial origin indicates that the moisture has undergone a certain degree of significant continental recycling. Terrestrial processes such as transpiration and evaporation introduce isotopically enriched moisture and high d-excess signatures, respectively. Mixing with terrestrial sources is also reflected in the relationship between <u>vapor</u> isotope <u>compositions</u> and q (Fig. 3). The degree of continental recycling also alters vapor isotope compositions that transpiration introduces isotopically enriched moisture and evaporation introduces moisture with high d excess. Seasonally changing isotope signatures in precipitation and ice cores as well as

variations at longer timescales Seasonal changes and long-term variations in precipitation and ice core isotopes have

been interpreted as <u>shifts in moisture</u> source <u>shift between recycled terrestrial moisture over terrestrial regions</u> and oceanic <u>moisture-sources</u> or their relative contributions (An et al., 2017; Yang and Yao, 2020). A <u>further inference</u> of this process is that the one moisture is <u>brought bytypically associated with</u> the summer monsoon, while the westerlies bring moisture from continental recycling or even the Mediterranean Sea. <u>Water isotope signatures on</u> the <u>TP were thought to reflect this and therefore water isotope signatures reflect the</u> interplay between the summer monsoon and westerlies (Joswiak et al., 2013; Pang et al., 2012; Tian et al., 2007). <u>Although our results also indicate Despite</u> seasonally shifting moisture sources, continental recycling prevails throughout the year (Fig. 4). <u>Besides focusing on moisture sources at the Earth surface, we provide a Ourn alternative perspective to explains the high d-excess induced by the westerlies as dry and cold air intrusions <u>rather than surface evaporation or evapotranspiration</u>. In this circumstance, the interpretation of <u>While</u> the interplay between the summer monsoon and westerlies is <u>still remains</u> valid, but we emphasize changes in air mass <u>property properties</u> driven by the different circulation systems.</u>

The proposed alternative interpretation could also help understandexplain the abnormally high d-excess in high-altitude ice cores mentioned in the Introduction, the increasing trend of precipitation and river water isotope observations toward ice cores at higher altitudes on the TP This is because specific humidity is very low at these ice core sites is extremely low, and prolonged interaction with cold and dry air may further modify snow isotope compositions (Ma et al., 2024; Wahl et al., 2022). In addition, intense rain-vapor interactions during the summer monsoon is represent another potential source of higher-elevated vapor d-excess (section Section 4.33.5). When this high Higher vapor d-excess vapor contributes to subsequent precipitation, its signal could can be inherited in subsequent the resulting precipitation—when it feeds the precipitation (Risi et al., 2008b). However, a clear relationship between TP precipitation d-excess and monsoon convection has not been yet to be established yet, partly due to less-limited attention—has been paid to d-excess in previous studies. On the other hand, local raindrop

evaporation may counteract this effect by reducing raindrop *d*-excess values. Nevertheless, summer monsoon rainfall *d*-excess observed on the TP is generally between 0-10% (Tian et al., 2001). Raindrop evaporation at upstream increases vapor *d*-excess and therefore could cause elevated *d*-excess in downstream rainfall. On the contrary, this effect can be compensated by on-site raindrop evaporation as it lowers raindrop *d*-excess values. The overall positive correlation between precipitation *d*-excess and altitude in across Asia has been sometimes been interpreted attributed to as stronger evaporation at lower altitudes (Bershaw, 2018). For snowfall on glaciers, however, evaporation is less likely for falling snowflakes is less likely due to cold temperatures and the short distance between the cloud base and the glacier surface. Therefore, elevated vapor *d*-excess signals caused by accumulated rain-vapor interactions at upstream associated with monsoon convection could be another source for a possible source of the high *d*-excess in ice cores.

#### **5 Conclusions**

We present a three-year-long daily near-surface water-vapor isotope compositions dataset observed at the South East TPSETP station, which is at the major channel for moisture entering the TP. Our vapor isotope compositions The paired measurements of vapor isotopes and with specific humidity reflect reveal distinct moisture sources and dynamics between the non-monsoon and summer monsoon seasons, consistent with findings from Lagrangian moisture diagnostic results. Despite significant negative correlations between vapor d-excess and relative humidity scaled to sea surface temperature existing over the northern Indian Ocean when data for all seasons are considered, such correlations with oceanic surface evaporation conditions largely disappear when separately considering each seasonthese correlations weaken or even disappear when analyzed within individual seasons. This result questions finding challenges the early earlier interpretation of TP d-excess as indicator of oceanic evaporation conditions and guarantees new interpretations in the future.

During the non-monsoon season, vapor d-excess is mainlyprimarily influenced by specific humidity both at both the local seale and upstream scales. Highly Air that has undergone significant dehydration, situateded air at the lower end of the Rayleigh distillation, is expected to have extremely high d-excess values. Air massBackward trajectory analyses and moisture source diagnostics suggest-reveal that the intrusion of cold and dry air intrusion driven by the westerlies during the non-monsoon season leads to the increasing trend of in vapor d-excess along with decreasing as specific humidity decreases. This process also contributes to a weak negative correlation between vapor d-excess and  $\delta^{18}O$ . Furthermore, vapor  $\delta^{18}O$  primarily reflects mixing processes with involving a relatively enriched moist end-member compared with to the summer monsoon season. These new insights into vapor d-excess on during the non-monsoon season vapor d-excess provides an alternative way framework for to interpreting the high d-excess in high-altitude TP ice cores.

During the summer monsoon season, rain evaporation is emerges as the dominant process determining shaping water vapor isotope compositions. First,—\_vapor \delta^{18}O systematically shifts below the Rayleigh distillation curve, aligning with predictions of falling in the region predicted by "super-Rayleigh" distillation driven caused by partial rain evaporation. Second,—\_vapor \delta^{18}O is anti-correlated with d-excess, pointing to an origin of depleted vapor by kinetic fractionation as a source of depleted vapor, which is not likely simply a result of cannot be attributed solely to rainout. Third, at the regional scale, vapor \delta^{18}O is shows significantly negatively correlated correlations with total precipitation amount—at the regional scale, but—while vapor d-excess positively correlates with total precipitation amount. These results—findings help usenhance our understanding of the dynamics of atmospheric humidity dynamics and also help disentangle the different effects of rainout and rain-vapor interactions in the context of the amount effect.

This study reveals distinct moisture sources and dynamics between non-monsoon and monsoon seasons over the southeastern Tibetan Plateau. These findings will aid in interpreting  $\delta^{18}$ O and d-excess records from Tibetan

Plateau glaciers, offering refined insights into past hydroclimatic conditions and challenging assumptions linking ice core isotopes to oceanic evaporation alone. Overall, the new findings from the study reveal different moisture sources and dynamics between the non-monsoon and monsoon seasons over the southeastern TP. The findings will also help the interpretation of ice core  $\delta^{18}$ O and d excess records derived from glaciers on the TP.

# **Competing interests**

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The authors declare that they have no conflict of interest.

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### Data availability

The NOAA ARL provided the HYSPLIT model and the **GDAS** data (https://www.ready.noaa.gov/HYSPLIT.php). The Copernicus Climate Change Service provided the ERA5 data (https://doi.org/10.24381/cds.adbb2d47 and https://doi.org/10.24381/cds.f17050d7). The GPM data are available through GES DISC (https://doi.org/10.5067/GPM/IMERG/3B-HH/07). Local meteorological data at the SETP station provided by National Tibetan Plateau Third Pole Environment Data Center are

- 732 (https://dx.doi.org/10.11888/AtmosphericPhysics.tpe.68.db). The observation data at the SETP station have been
- uploaded to Figshare and will be made publicly available after publication (10.6084/m9.figshare.27302871).

### **Author contributions**

- 735 **Zhongyin Cai**: Conceptualization, methodology, investigation, formal analysis, funding acquisition, writing-
- original draft, writing-review & editing; Rong Li: Investigation, data curation, writing-review & editing; Cheng
- 737 Wang: Validation; Qiukai Mao: Investigation, Lide Tian: Resources, project administration, funding acquisition.

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