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Water isotope and chemical records in a recent snow pit from Hercules Neve, northern Victoria Land, Antarctica

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Mineral dust effect

Climate

- Direct: the absorption of thermal radiation
- Indirect: the global carbon cycle and green gas content

Chemistry

- Chemical balance of the atmosphere through surface absorption and reaction

MINERAL DUST

Biogeochemical Cycle

- Providing nutrients on terrestrial and marine ecosystem

Health

- Mineral dust in the respirable size can cause health problem



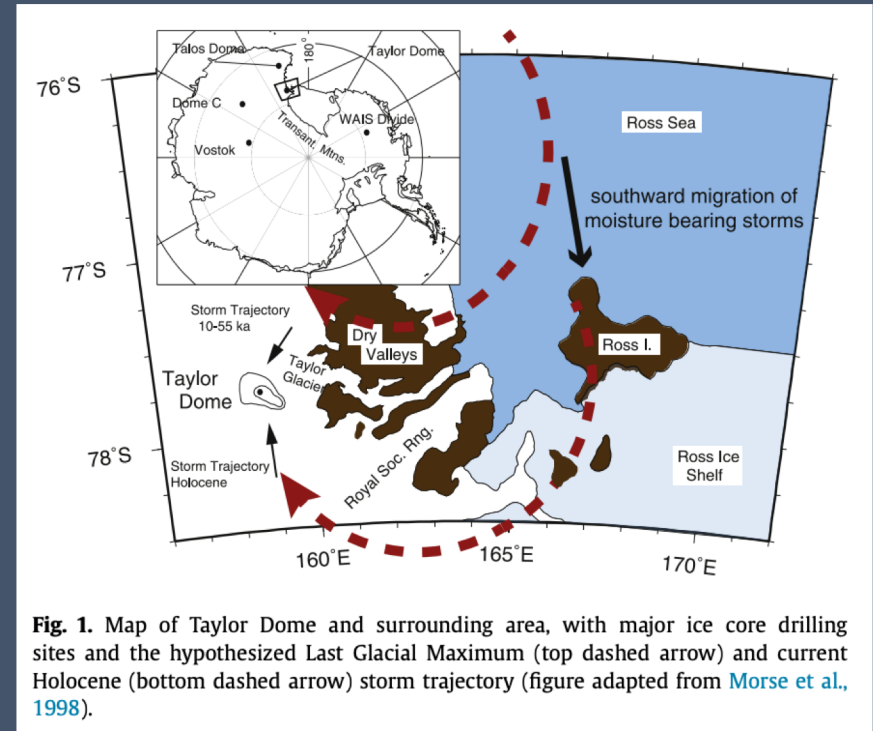
- Polar ice cores archive mineral aerosol exclusively of aeolian origin

- Since dust in snow does no weathering after deposition, it can give information about the past climate system.

The importance of local source

- During late Quaternary glacial periods on central and coastal east Antarctic ice cores identify southern South America (SSA) as the most likely source of mineral dust

Aarons et al., 2016



- During interglacial periods in coastal ice cores may potentially originate from **local sources** (Delmonte et al., 2007, 2010; Gabrielli et al., 2010; Aarons et al., 2016).

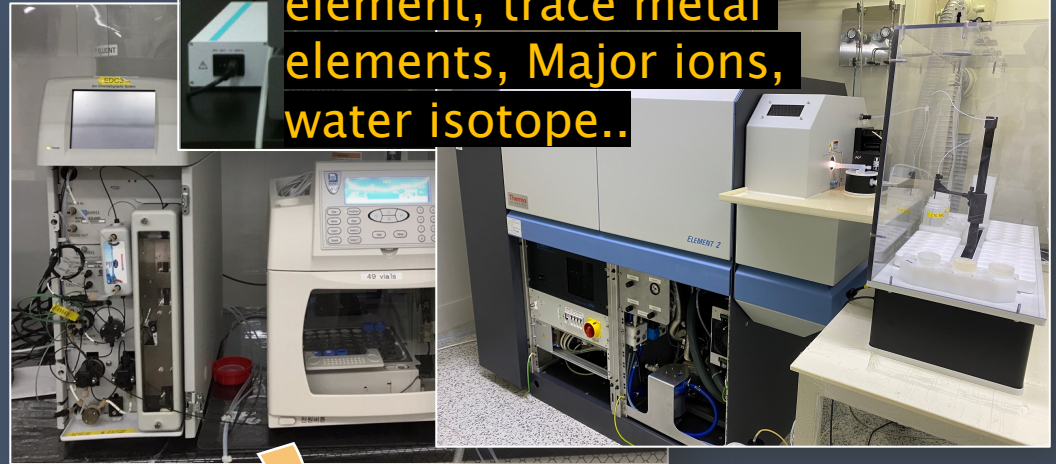
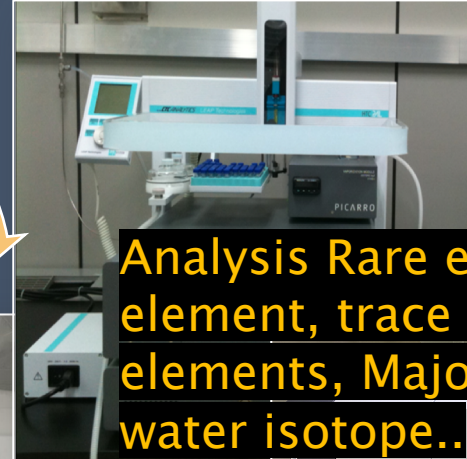
Hercules Neve

- ❖ Sampling date: Dec. 15, 2015
- ❖ Location: 73°03'S, 165°24'E
- ❖ Altitude: ~2900 m
- ❖ Distance from the sea: ~75 km
- ❖ Accumulation: 119~145 kg m⁻² a⁻¹ (Stenni et al., 1999; 2000)

- Hercules neve is surrounded by the mountains and close to the coastal.
- HN region in glacier ice also has the potential to provide information about the sources of aeolian dust reaching the Antarctic ice sheet at past climatic cycles



Research proposal



Identification of source

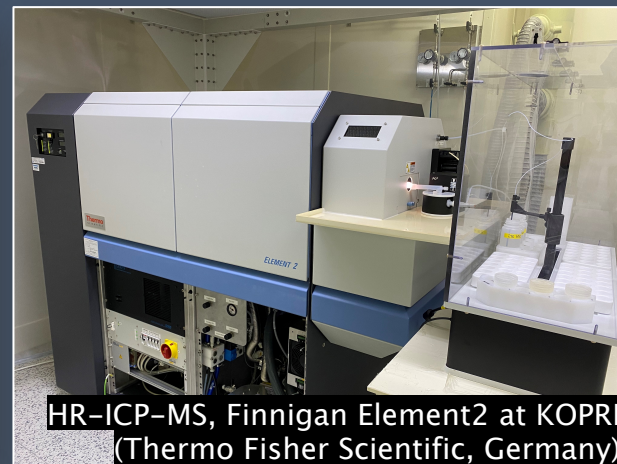
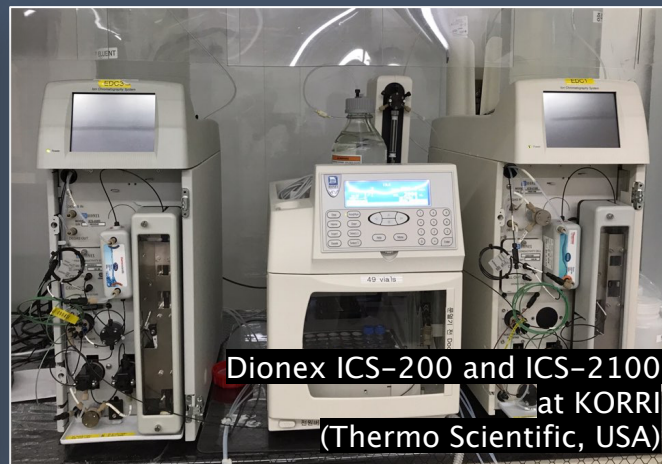
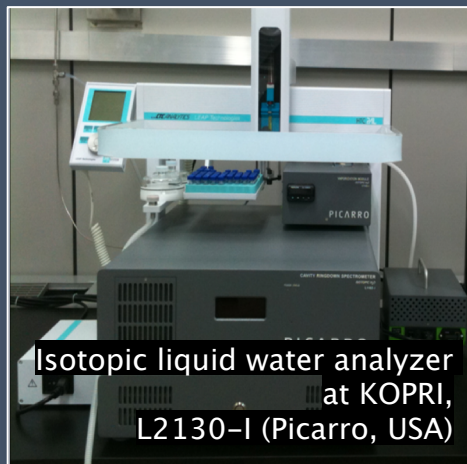


Source: 2014/15 Antarctic campaign by KOPRI

Snowpit sampling

- Depth: 2 m
- Sampling resolution: 5 cm
- Pre-cleaned equipment: polyethylene bottle, stain-less steel shovel, polyethylene gloves, Teflon cylinder, hammer, clean-suit, mask

Experimental method



	Precision	Number of sample	Standard
Water isotope ($\delta^{18}\text{O}$ and δD)	$\delta^{18}\text{O}$: 0.07 % δD : 0.6 %	40	VSMOW, GISP, SLAP and Lab-made standard
Major ion (Na^+ , K^+ , Mg^{2+} , Ca^{2+} , MSA , Cl^- , F^- , and SO_4^{2-})	<10 %	40	Cation; Product # 46070, Anion, Product # 57590
REE (La, Ce, Pr, Nd, Sm, Eu, Gd, Tb, Dy, Ho, Er, Yb, Lu), Trace metal (Y, Sb, Ba, Tl, Bi,Th)	<15 %	40	IV-ICPMS-71A, 71B, 71D

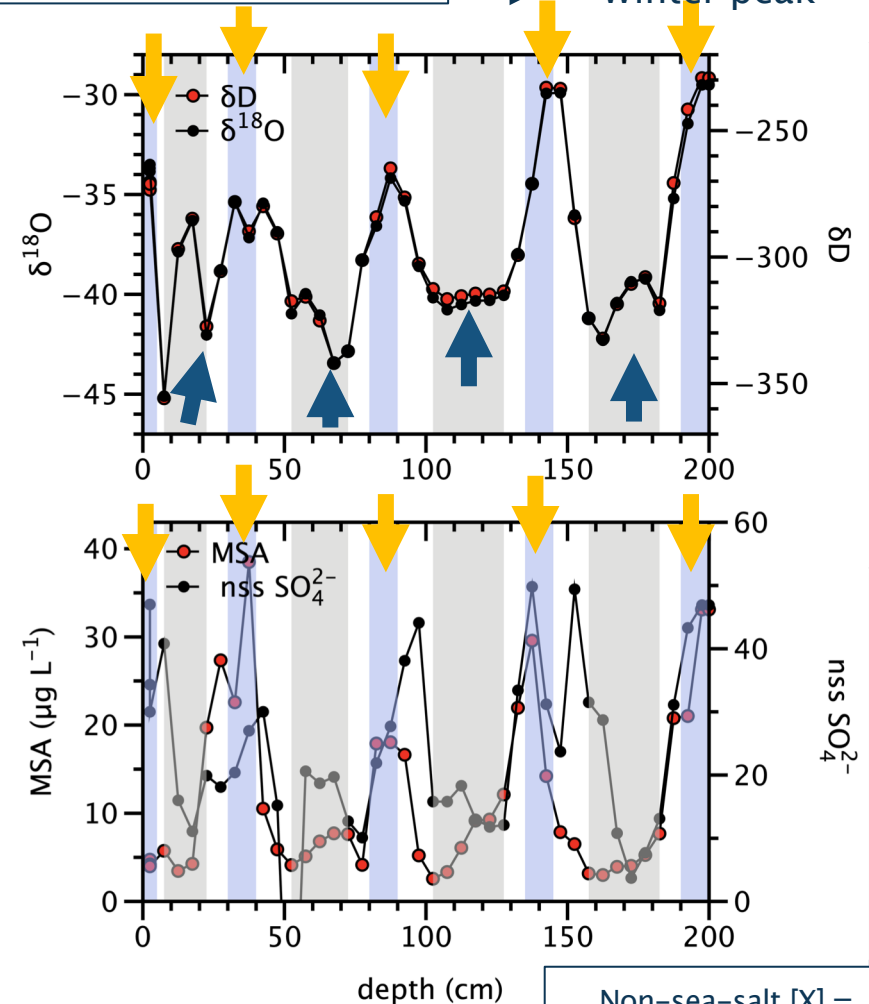
1.1 Temporal variation of water isotopes, MSA

- ❖ Water isotopes in snow → temperature proxy
 - Oxygen and hydrogen isotopes (water isotope ratios) are temperature related and can be used to confirm seasonality.
 - Summer enriched ratio, winter depleted ratio
- ❖ MSA and non-sea salt SO_4^{2-} → seasonality proxy
 - MSA (Methanesulfonic Acid), non-sea salt SO_4^{2-} is a by-product of marine biological activity and can be used as a seasonal proxy because it has high activity in summer and shows high concentration in snow.

$$\delta_{\text{Sample}} = (R_{\text{Sample}} - R_{\text{VSMOW}}) / R_{\text{VSMOW}} \times 1000 \text{ (‰)}$$

R is $\text{HDO}/\text{H}_2\text{O}$ or $\text{H}_2^{18}\text{O}/\text{H}_2^{16}\text{O}$

Summer peak
Winter peak



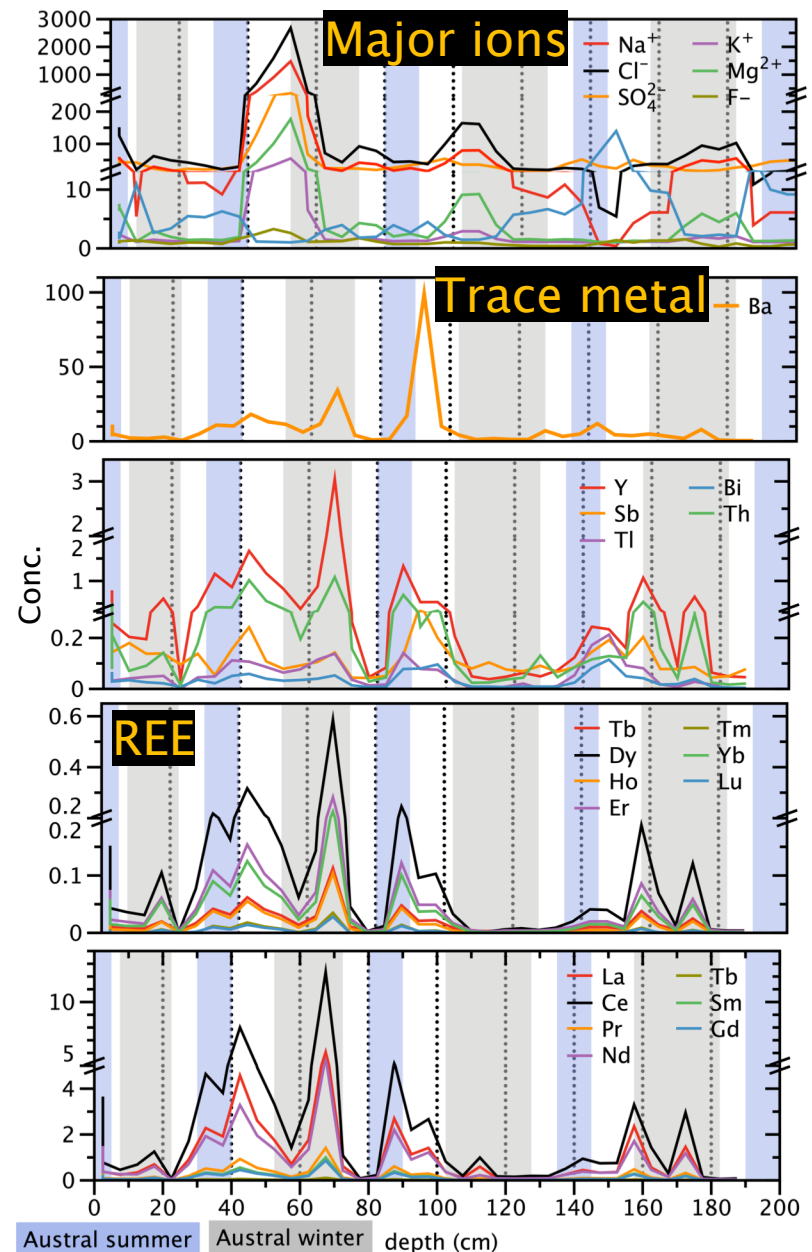
$$\text{Non-sea-salt [X]} = \text{tot[X]} - (\text{X/Na}^+)_{\text{sw}} \times \text{Na}^+$$

1.2 Temporal variation of major ions, REE and trace metal

The major ions, REE, and trace metal

- No consistent pattern indicate the various sources
 - Marine source (sea salt)
 - Terrestrial source (crust-derive dust)
 - Anthropogenic source
 - Volcanic source
- Sea salt ions (Na^+ , Cl^- , K^+ , Mg^{2+} , Ca^{2+} , SO_4^{2-}) show high correlations each others
- High correlation in REE and trace metal (except Ba, Bi, Sb)

Depth profile of the major ions, REE, and trace metal



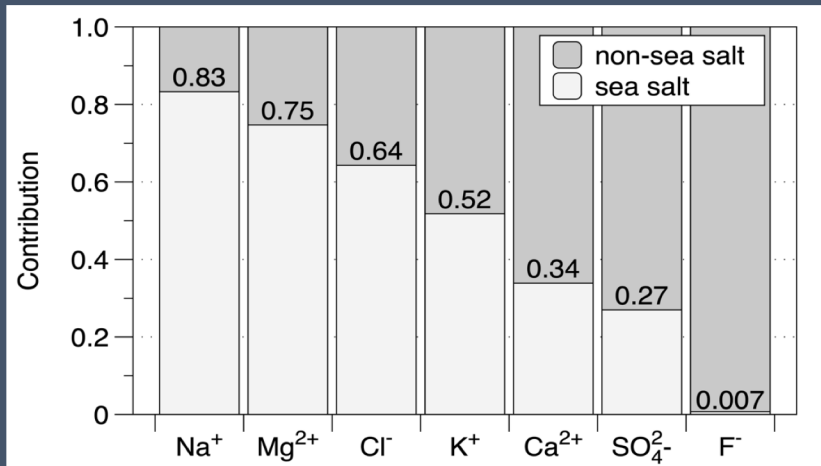
2.1 Effect of Marine source (sea salt)

Fraction of major ions, REE and trace metal

1) $\text{Non-sea-salt } [X] = \text{tot}[X] - (X/\text{Na}^+)_{\text{seawater}} \times \text{Na}^+$

2) $\text{Contribution of Non-sea salt } [X] = \text{nss}[X]/\text{tot}[X]$

Major ions

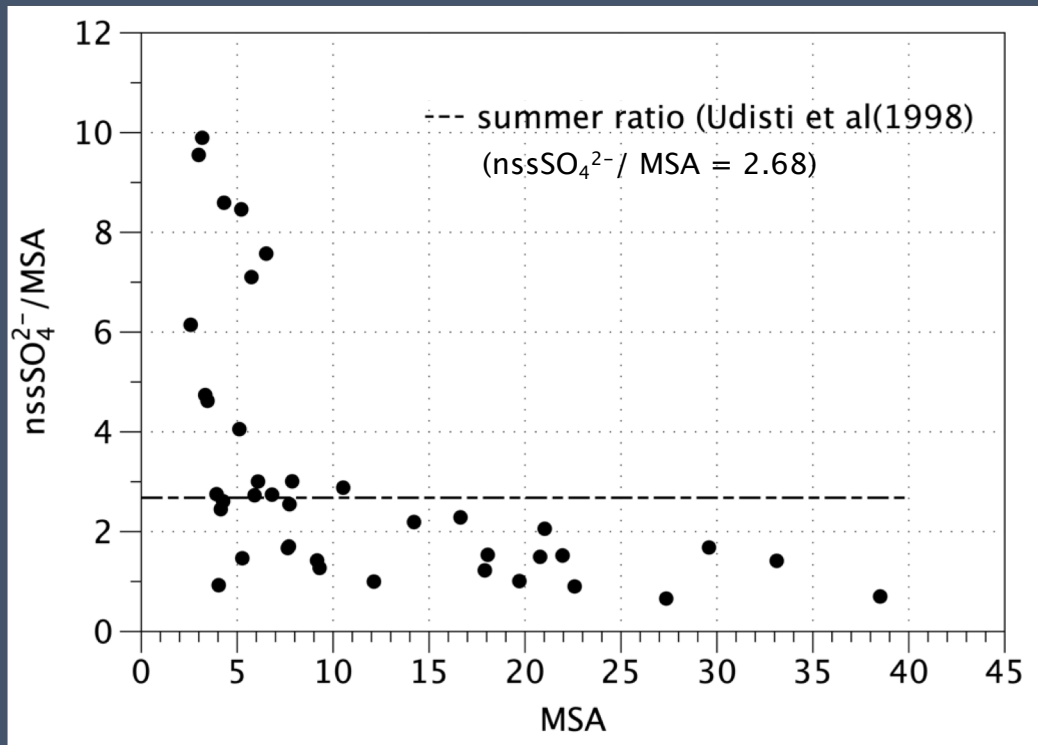


- Sea salt contribution was higher (>60 %) in Na⁺, Mg²⁺, Cl⁻
- Contributions of other sources (not only sea salt) shown in K⁺, Ca²⁺, and SO₄²⁻
- F⁻ no sea salt

REE & Trace metal

- Sea salt sourced REE is negligible, thus crustal and terrestrial sources were considered

2.2 Marine biogenic source (corr. nss SO_4^{2-} with MSA)



- Biogenic sulfur enhancement in spring and summer was detected in nssSO₄²⁻ and MSA ($r = 0.39$)
- Minor contribution of other sources (e.g. crust, volcanic) can be observed in nssSO₄²⁻ during winter (MSA conc. is low)

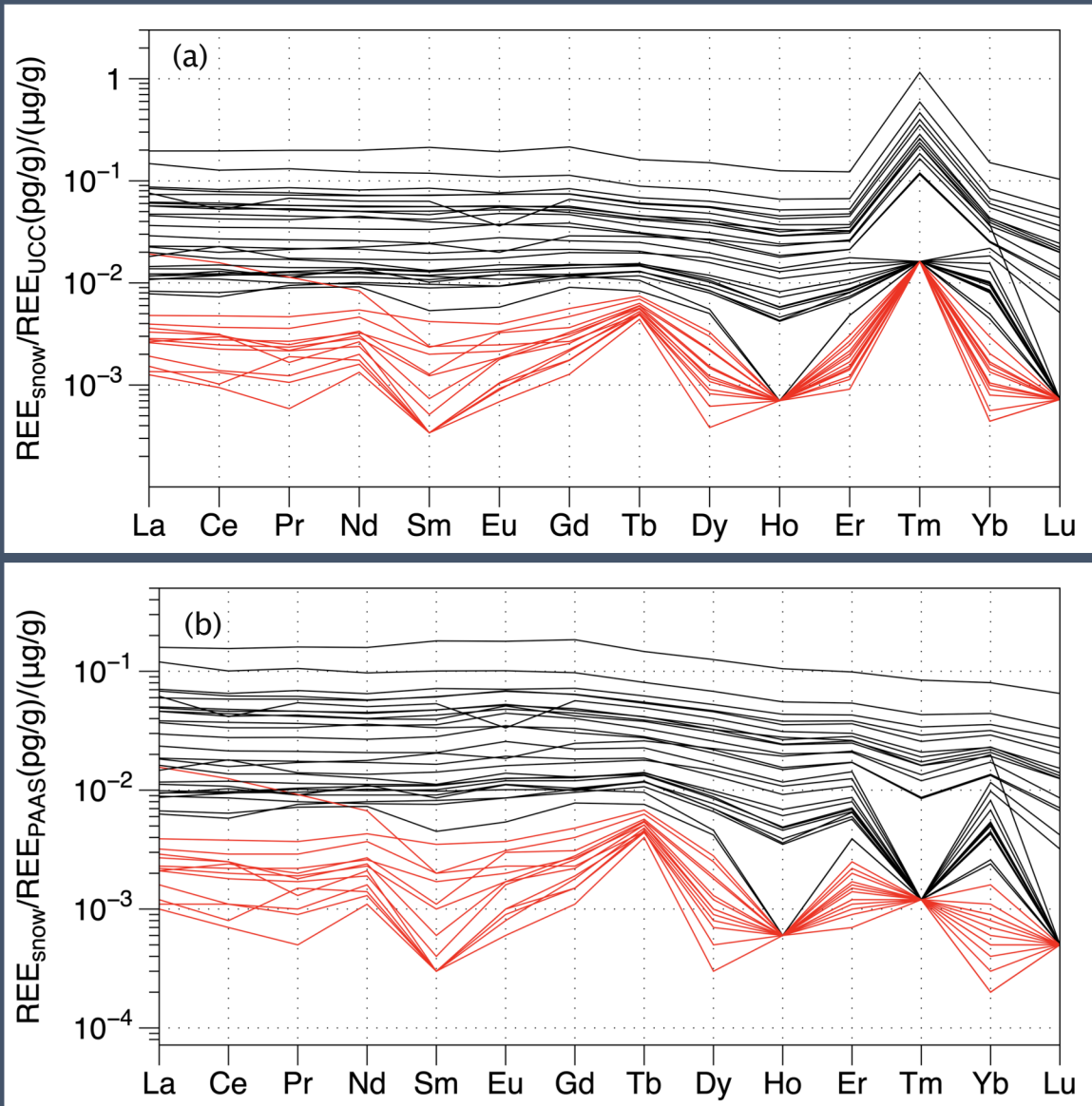
2.3 Potential Source – Trace metal

- ❖ Y and Th show high correlation ($r^2 = 0.99$) with the REE (La), while this correlation was not observed with the others including Sb, Ba, Bi, and Tl ($0.27 < R^2$).
- ❖ It can be considered the different sources for Sb, Ba, Bi, and Tl compared to potential sources of REE.
- ❖ Sb, Bi, Ba and Tl correlation (R^2)

	Sb	Bi	Tl	Ba
Sb	1			
Bi	0.36	1		
Tl	0.16	0.74	1	
Ba	0.22	0.10	0.10	1

- Bi show relation with Sb and Tl
- Sb, Bi and Tl were assumed to be sourced from other sources (e.g. anthropogenic)

2.4 Terrestrial Source – REE



❖ REE_{snowpit}/REE_{UCC} or PAAS

- The converging points of Ho, Tm, and Lu indicate the detection limit values (due to very low values).
- The anomalies of Sm and Tb were commonly seen in the low concentration section (red line) in (a) and (b), and the anomaly of the positive Tm in the high concentration part (black line) is observed in (a).
- It seems that there are at least two snow REE sources, continental sources, not marine or anthropogenic pollution.

Summary

- No consistent pattern (seasonal) was observed in major ions, REE, and trace metal
- The patterns of Sb, Tl and Bi are different from REE and are considered to be anthropogenic source as in previous studies.
- Considering concentration of REE, there are two crust-derived potential sources (maybe not marine and anthropogenic source)
- More records (long-term record (e.g. ice core)) in local scale are needed in the further research

Thank you.