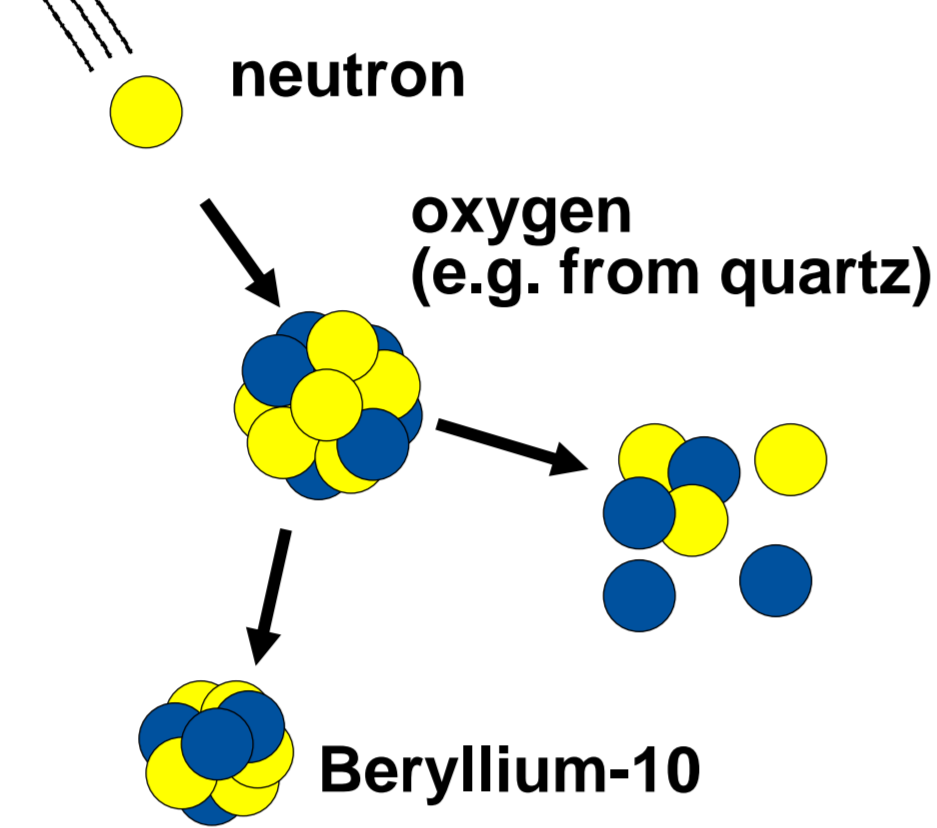
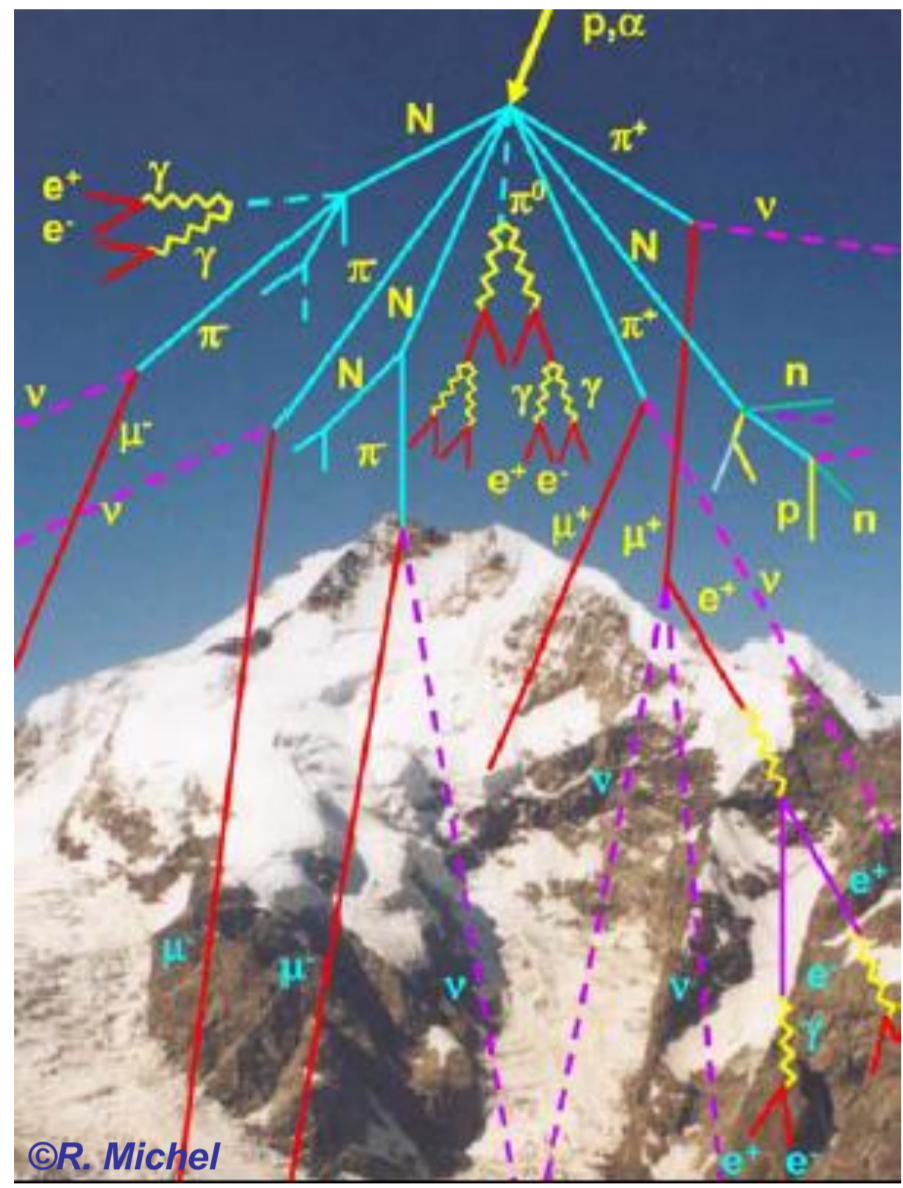


# SETTING-UP CHEMISTRY LABS FOR ACCELERATOR MASS SPECTROMETRY

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## Cosmic radiation on Earth

- MeV/GeV particles (p, α, heavy ions)
- shielding (flux & energy) and transformation e.g. into neutrons by the Earth's magnetic field & atmosphere
- elemental conversion by nuclear reactions in the atmosphere ( $^{14}\text{N}(n,p)^{14}\text{C}$ ) >>> radiocarbon-dating e.g. for archaeology or climate reconstruction from ice cores
- elemental conversion by nuclear reactions in terrestrial materials / rocks (so-called "in-situ"-production) >>> with time concentration of (radio-) nuclides increases
- irradiation of a "fresh" surface (e.g. after a volcanic eruption), reconstruction of "starting time" possible >>> "in-situ"-dating



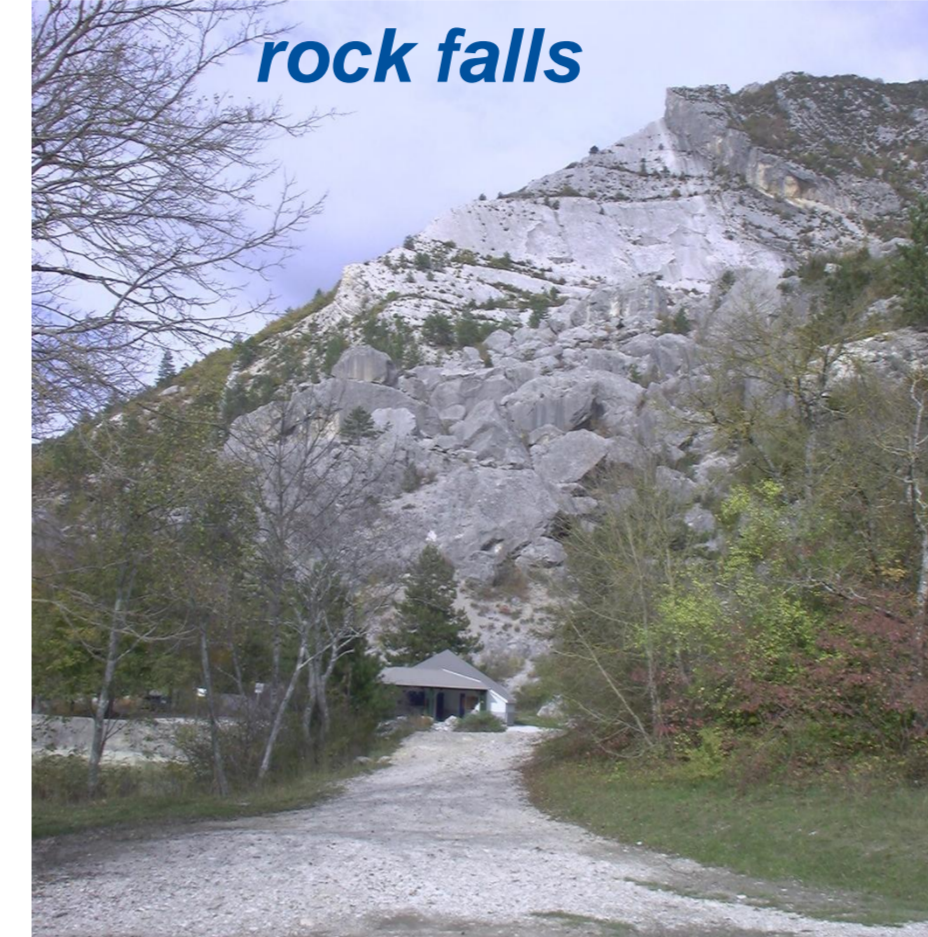
## "In-situ"-produced cosmogenic nuclides (CN)

- everything having a "fresh" surface can be dated

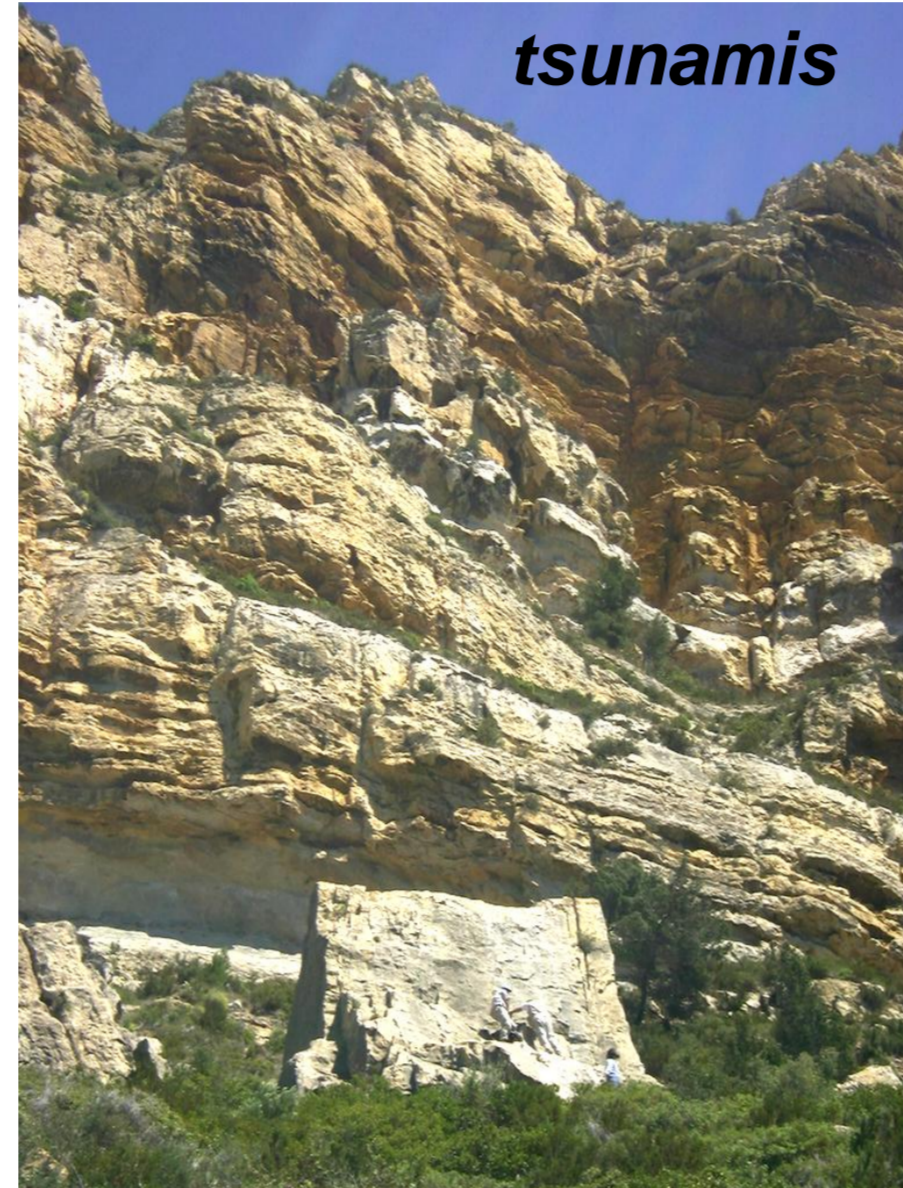
volcanic eruptions



rock falls



tsunamis

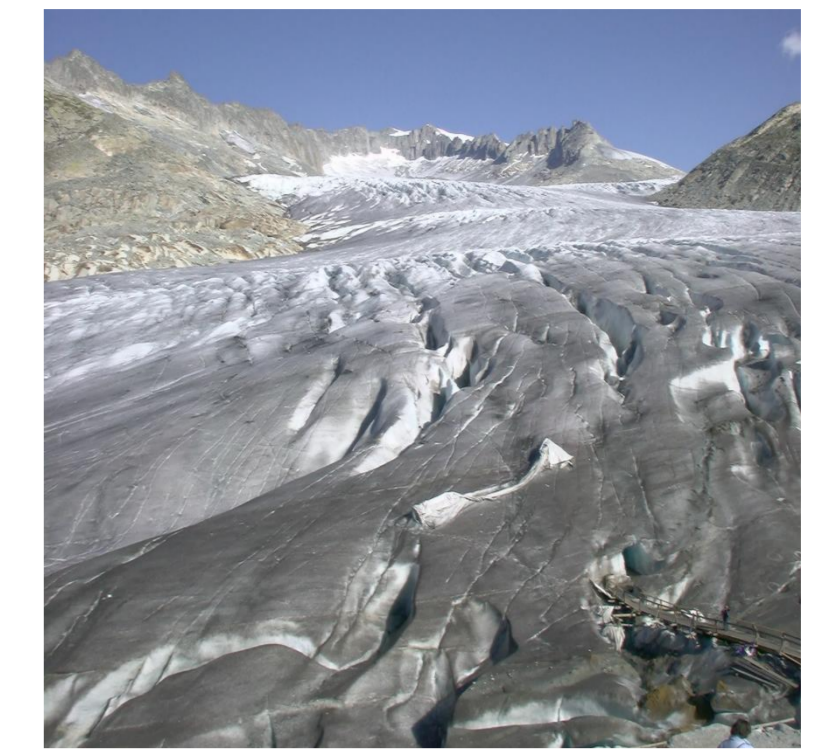


earthquakes



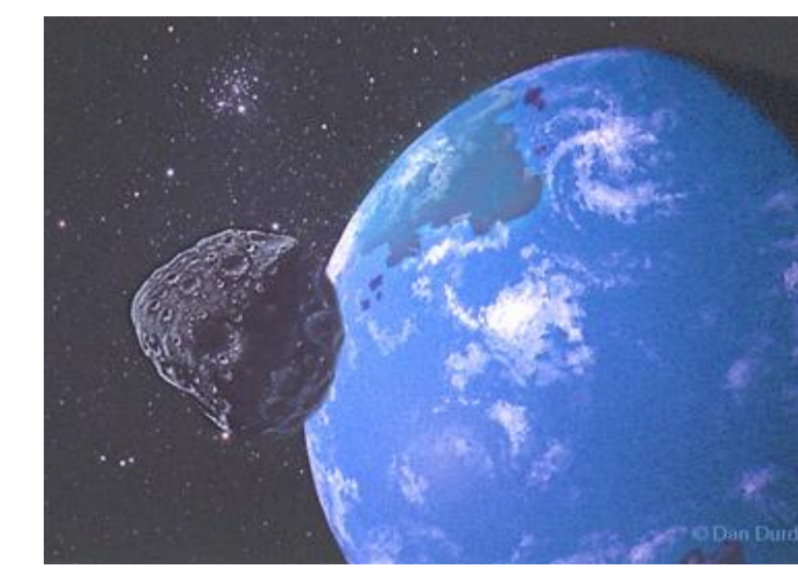
## Indirect dating by CN

- climate change, e.g. via glacier movements



## Special case (irradiation in space): Extraterrestrial material

- transfer times from the meteorite's parent body (asteroids, Moon, Mars) >>> irradiation age
- residence time at the place of discovery (e.g. hot desert, Antarctica) before somebody takes the meteorite home >>> terrestrial age



## Sample preparation for accelerator mass spectrometry (AMS)

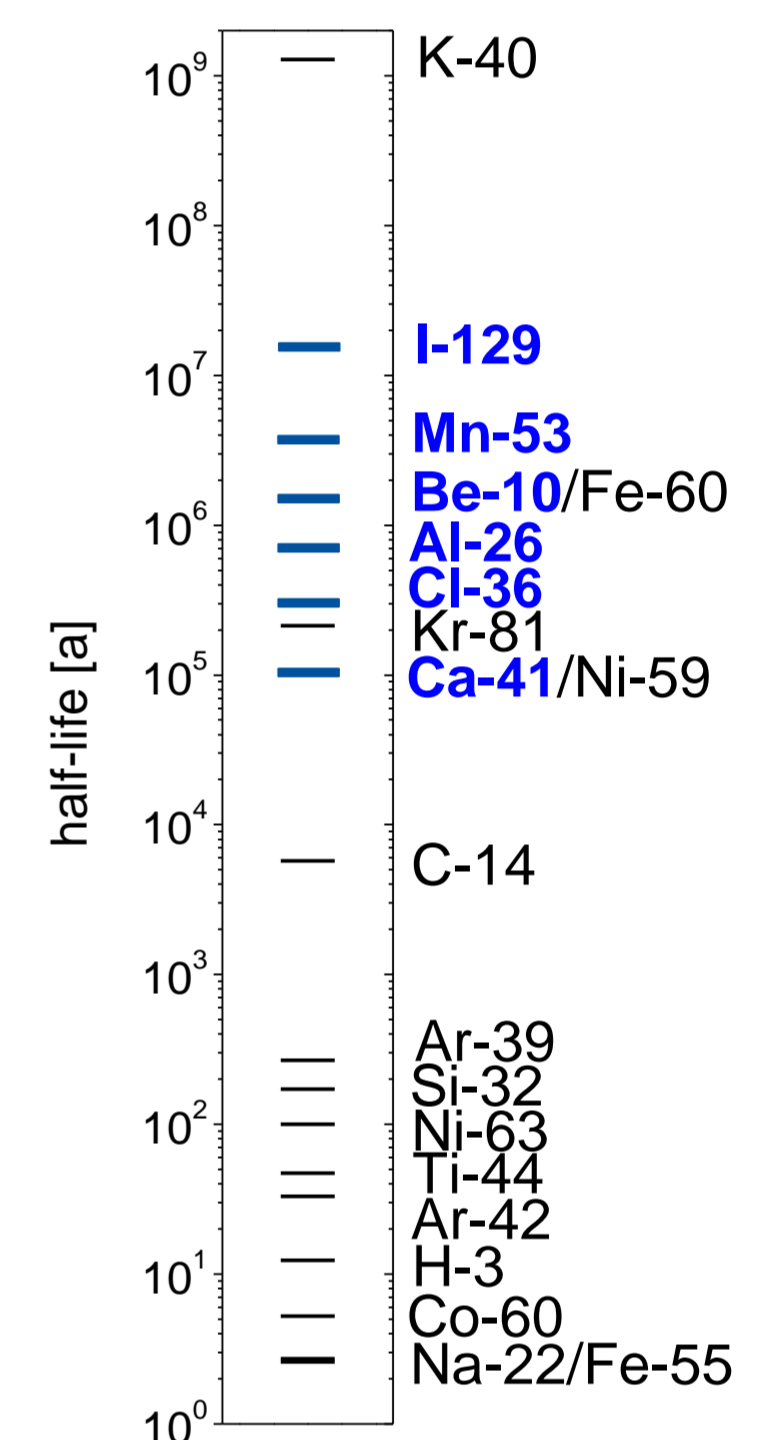
### Definition of "good" AMS-chemistry

- "good" standards, i.e.
  - traceable
  - different ratios for volatile elements to avoid cross-contamination ( $^{36}\text{Cl}$ ,  $^{129}\text{I}$ )
- low machine blanks ( $^{10}\text{Be}$ )
- low stable nuclide carrier for chemistry blanks ( $^{10}\text{Be}$ )
- real sample chemistry for different matrices & nuclides
  - fast and low-cost (chemicals, man power, lab space)
  - safe (also for non-chemists)
  - low risk of cross-contamination & contamination
  - producing "pure" targets, i.e. high stable nuclide current & low isobar concentration >>> good statistics, detection limits & high sample throughput

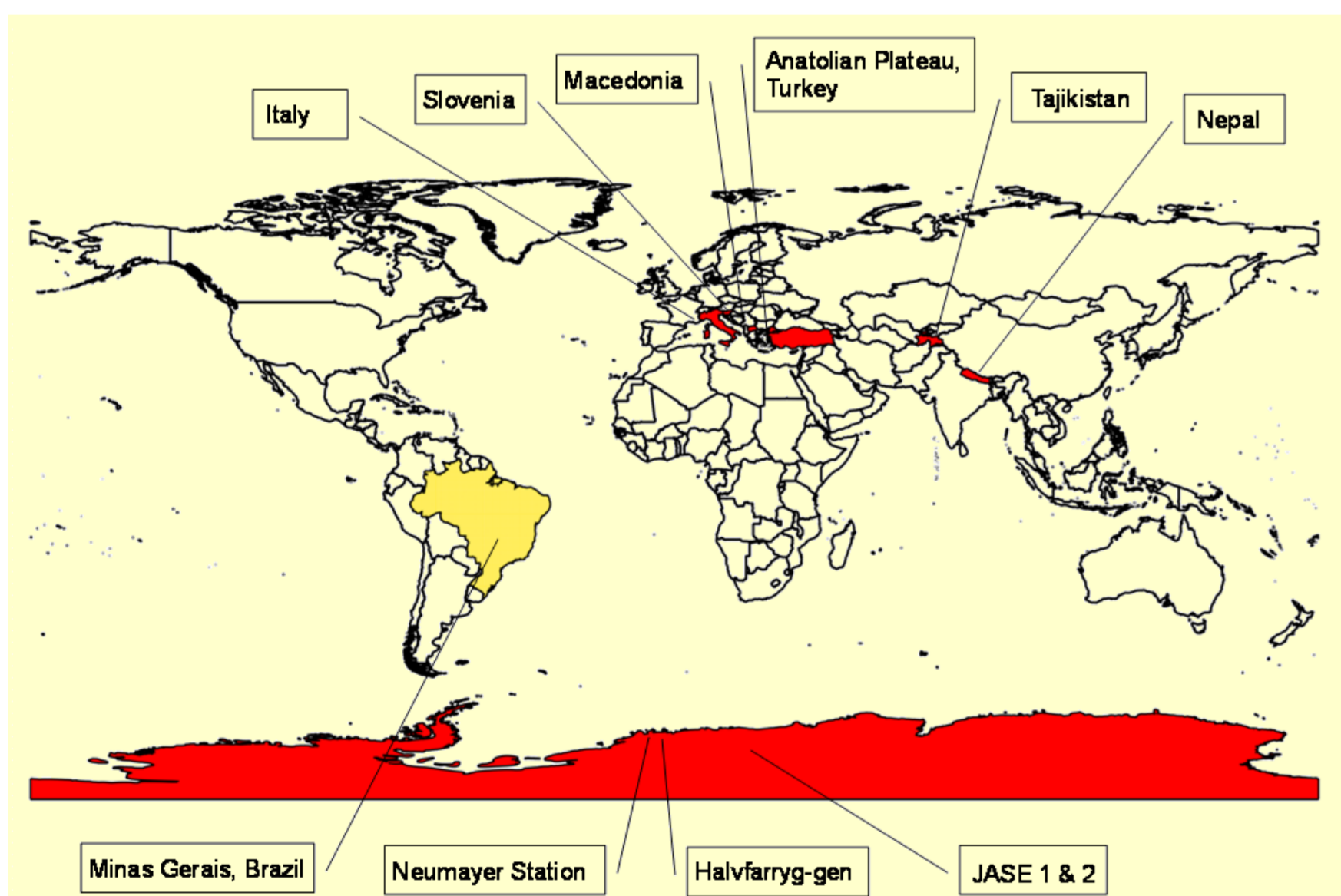


### Status @ DREAMS

- standards for  $^{10}\text{Be}$ ,  $^{26}\text{Al}$ ,  $^{36}\text{Cl}$ ,  $^{41}\text{Ca}$ ,  $^{129}\text{I}$  traceable via round-robins or cross-calibration elsewhere [ARN10, MER04, MER09, MER11]
  - special case  $^{10}\text{Be}$ : produced via  $^9\text{Be}(n_{th},\gamma)^{10}\text{Be}$  @ TRIGA, Atominstutut Vienna >>>  $(1.73 \pm 0.02) \cdot 10^{-12} \text{ }^{10}\text{Be}/^9\text{Be}$
  - $^{26}\text{Al}$ ,  $^{36}\text{Cl}$ ,  $^{41}\text{Ca}$  &  $^{129}\text{I}$  @ 3 different ratios each  $10^{-9} - 10^{-13}$
- "home-made"  $^{10}\text{Be}$  carrier & machine blank from shielded Be-containing mineral (phenakite -  $\text{Be}_2\text{SiO}_4$ ) [MER08]
- speeded-up  $^{10}\text{Be}$ -chemistry for ice core sample (5 d → 24 h / 10 samples)
- development of  $^{53}\text{Mn}$ -chemistry for "in-situ" samples (marcasite/pyrite/realgar)
- $^{10}\text{Be}$  &  $^{36}\text{Cl}$  targets from "in-situ" samples ( $\text{SiO}_2$  &  $\text{CaCO}_3$ ) measured @ ASTER & VERA >>> chemistry blank at least one order of magnitude lower than samples & high current
- chemistry training of external partners (non-chemists) for  $^{10}\text{Be}$  &  $^{36}\text{Cl}$  from "in-situ" samples satellite labs @ U Rennes & U Freiberg



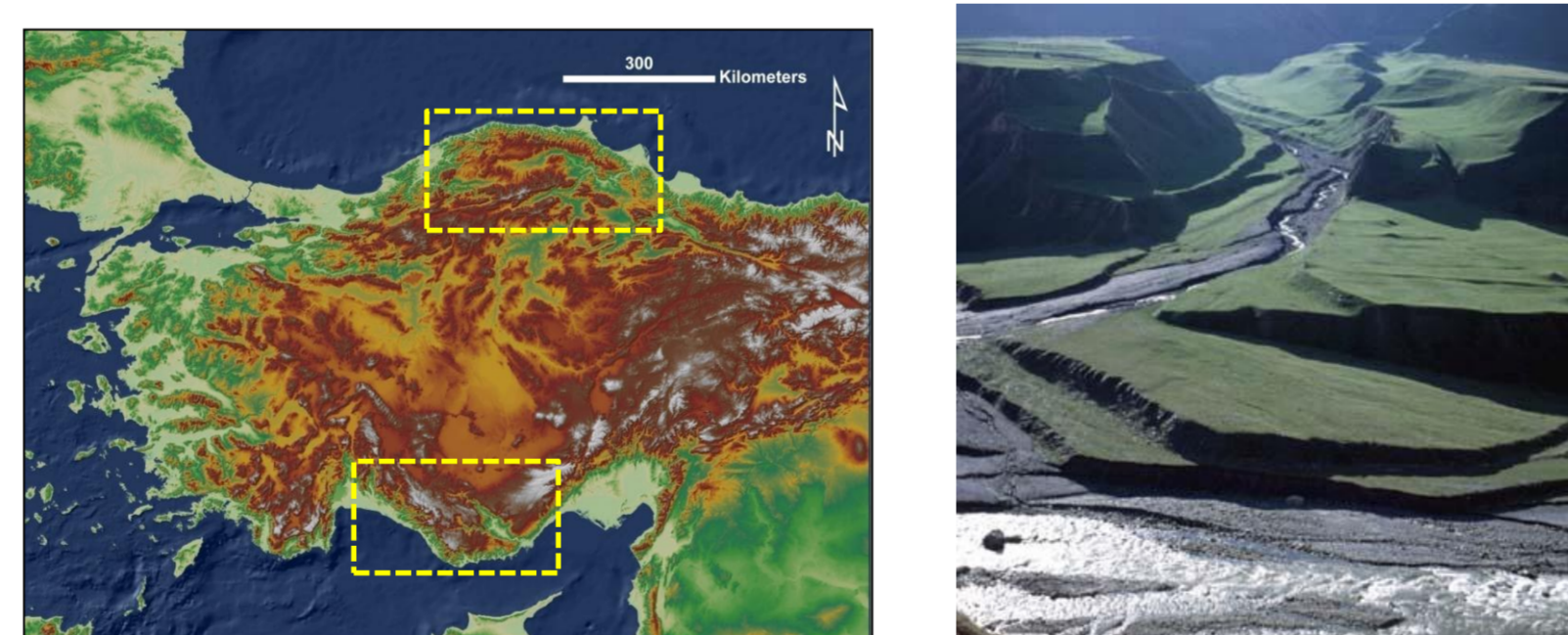
## Selection of sample origins & applications



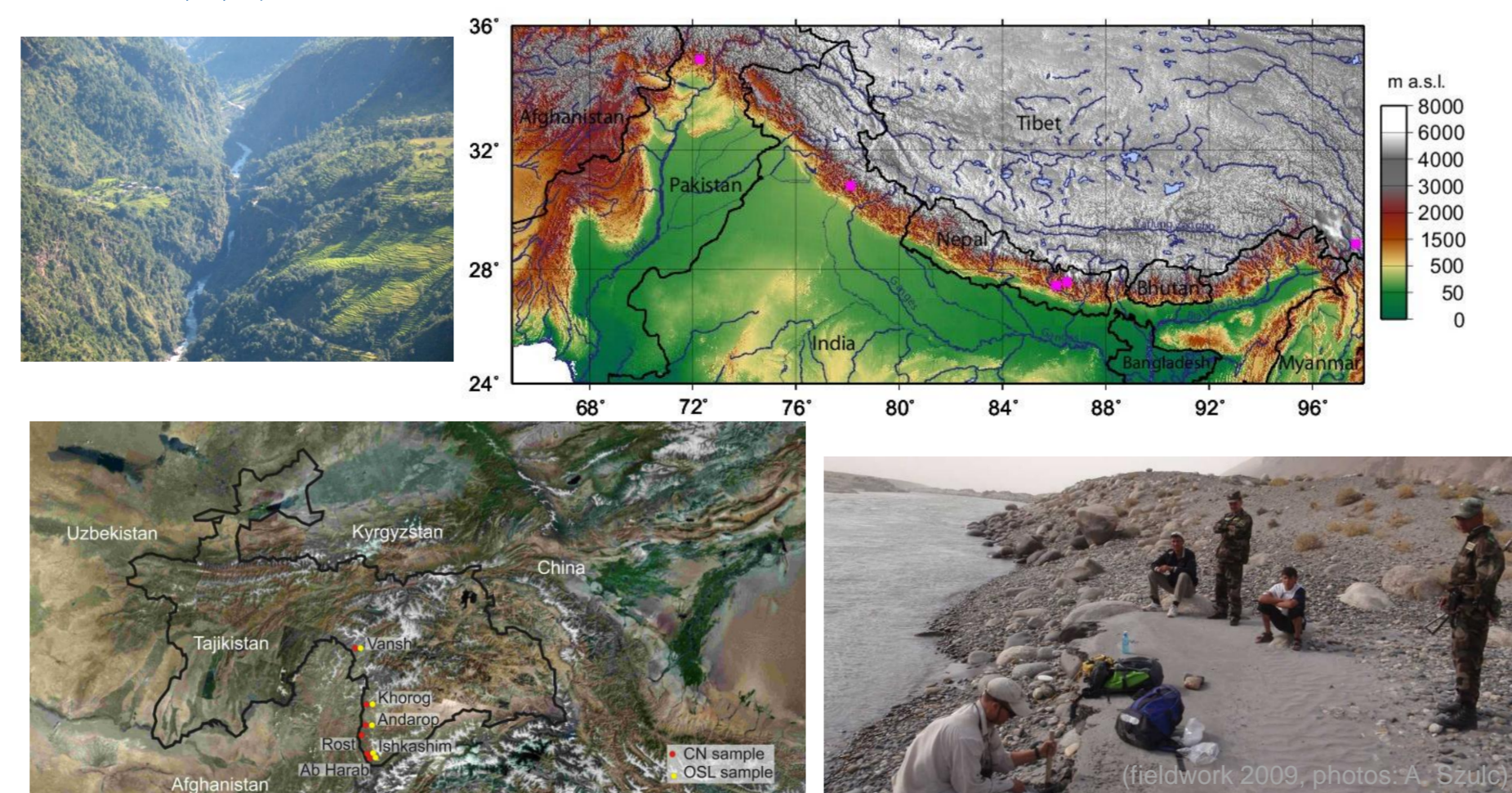
- Historical rock fall @ Veliki Vrh, Slovenia >>> Calibration site for production rate ( $^{36}\text{Cl}$ )



- Vertical Anatolian Movement Project >>> Uplift rates ( $^{36}\text{Cl}$ )



- Geomorphology @ Himalaya & Pamir >>> Basin-wide erosion rates ( $^{10}\text{Be}$ )



Detection limits of AMS can be as low as 20,000 atoms/g or  $10^{-9}$  Bq.

## Summary & Outlook

Routine radionuclides will be measured @ the DREsdEN AMS facility (DREAMS) first - except  $^{14}\text{C}$ . Generally, all long-lived radionuclides and stable nuclides can be measured by AMS. But due to the low precision of AMS (% to ‰), stable elements marked in yellow can only be measured if ratios are highly different from natural isotope ratios. Probably, other MS methods yield better results at normal isotope ratios.

H																	He
Li	Be	routine, sophisticated with long-lived radioactive nuclide(s), stable isotopes only										B	C	N	O	F	Ne
Na	Mg											Al	Si	P	S	Cl	Ar
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe
Cs	Ba	La	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn
Fr	Ra	Lr	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg	112	113	114	115	116	117	118
		La	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb		
		Ac	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No		

\* only one long-lived radioactive isotope / no stable isotope → very sophisticated set-up

## Acknowledgements

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