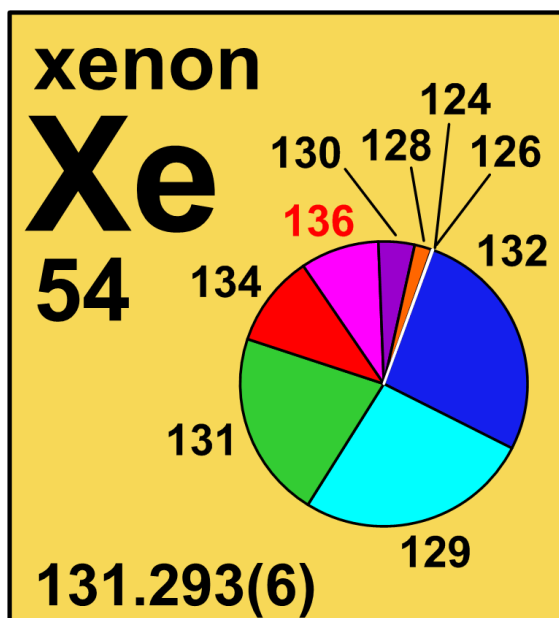





## 4.54 xenon

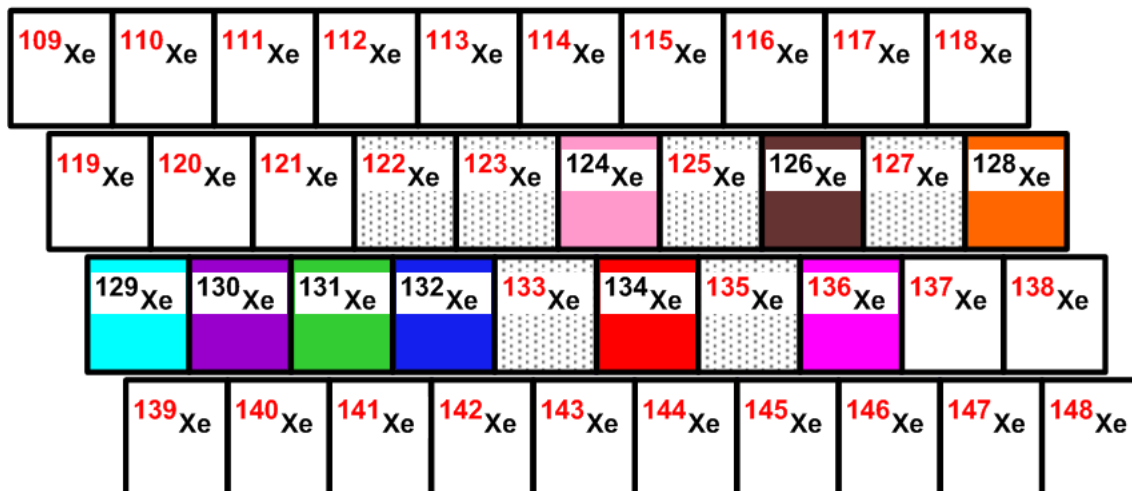


Stable isotope	Relative atomic mass	Mole fraction
$^{124}\text{Xe}$	123.905 89	0.000 95
$^{126}\text{Xe}$	125.904 30	0.000 89
$^{128}\text{Xe}$	127.903 531	0.019 10
$^{129}\text{Xe}$	128.904 780 86	0.264 01
$^{130}\text{Xe}$	129.903 5094	0.040 71
$^{131}\text{Xe}$	130.905 084	0.212 32
$^{132}\text{Xe}$	131.904 155 09	0.269 09
$^{134}\text{Xe}$	133.905 395	0.104 36
$^{136}\text{Xe}^\dagger$	135.907 214 48	0.088 57

<sup>†</sup> **Radioactive isotope** having a relatively long **half-life** ( $2.3 \times 10^{21}$  years) and a characteristic terrestrial **isotopic composition** that contributes significantly and reproducibly to the determination of the **standard atomic weight** of the element in **normal materials**.

## Half-life of radioactive isotope

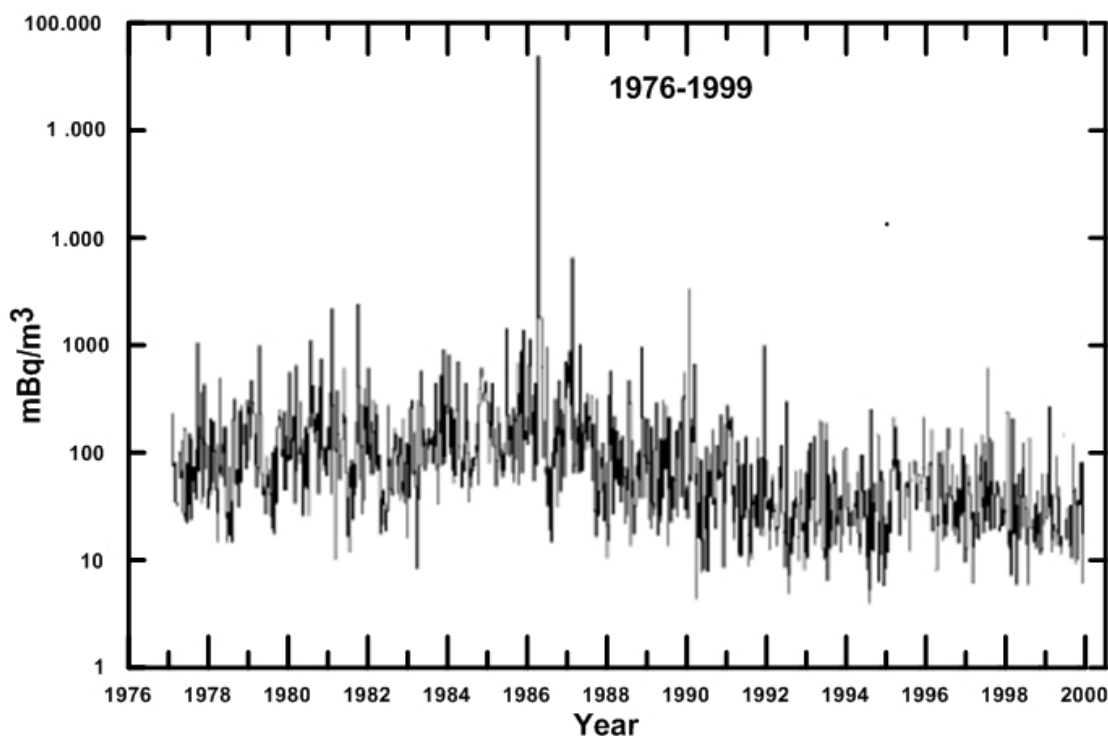
Less than 1 hour   
 Between 1 hour and 1 year   
 Greater than 1 year 



## 4.54.1 Xenon isotopes in forensic science and anthropology

**Radiogenic xenon isotopes** are produced by nuclear reactions in atomic bombs and nuclear reactors. For example,  $^{131}\text{Xe}$ ,  $^{133}\text{Xe}$ , and  $^{135}\text{Xe}$  are some of the **fission** products of  $^{235}\text{U}$  and  $^{239}\text{Pu}$ , and finding these isotopes would be evidence of a nuclear bomb reaction. Measurements of

1 xenon isotopes (e.g., in the atmosphere or the subsurface) have been used to identify  
 2 contamination from these sources, for example, to detect faults in nuclear reactors or to monitor  
 3 compliance with nuclear test bans (Figure 4.54.1) [393].  
 4



5  
 6 **Fig. 4.54.1:** The  $^{133}\text{Xe}$  time series in air at Freiburg, Germany, between 1976 and 1999. The  
 7 record indicates persistent low levels of **anthropogenic**  $^{133}\text{Xe}$  that are generally attributable to  
 8 normal acceptable releases from nuclear power plants with variability related in part to multiple  
 9 sources and changing wind patterns. A major spike occurred in 1986 during the Chernobyl  
 10 reactor accident in the Ukraine region of the former USSR. The **half-life** of  $^{133}\text{Xe}$  (5.2 days) is  
 11 sufficiently long for it to escape from its source and be distributed in air near the source, but  
 12 sufficiently short that long-term background levels are very low. Records such as this also can be  
 13 used to detect undocumented nuclear explosions. (Modified from [393]).  
 14  
 15

#### 16 4.54.2 Xenon isotopes in geochronology

17  
 18 The **stable isotopes** of xenon hold many clues about the formation of the **elements**, solar-system  
 19 history, and Earth processes [26, 98]. For example,  $^{129}\text{Xe}$  has been used as a detector of “extinct”  
 20 **radionuclides**. Some  $^{129}\text{Xe}$  is radiogenic as a result of being produced by the **radioactive decay**  
 21 of  $^{129}\text{I}$  (**half-life** =  $1.7 \times 10^7$  years). Because the half-life of  $^{129}\text{I}$  is much smaller than the age of  
 22 the Earth, **primordial**  $^{129}\text{I}$  (i.e., that which was present at the beginning of Earth’s history) is  
 23 essentially gone after it decayed to  $^{129}\text{Xe}$  over geologic time. This means that radiogenic  $^{129}\text{Xe}$   
 24 could be a marker of the former existence of the “extinct” isotope  $^{129}\text{I}$ . Because primordial  $^{129}\text{I}$

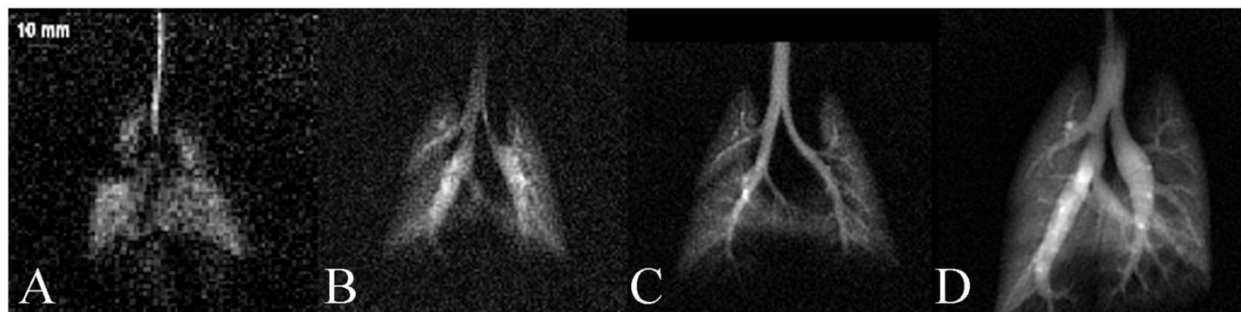
## IUPAC

1 was produced largely in **supernovae**, detection of radiogenic  $^{129}\text{Xe}$  in **meteorites** and terrestrial  
2 samples also implies that the time elapsed between  $^{129}\text{I}$  **supernova nucleosynthesis** and  
3 planetary condensation was short compared to the subsequent history of the Solar System. The  
4 many isotopes and reaction mechanisms of xenon have contributed numerous insights into Earth  
5 processes through the study of “xenology” (xenon isotopic variations used as geodynamic  
6 **tracers** to study the dynamics of the Earth) [394].  
7

### 8 **4.54.3 Xenon isotopes in medicine**

9

10 Xenon isotopes are used in numerous ways to investigate the movement of inhaled gases in lungs  
11 and other parts of the body. If **radioactive isotopes** of xenon [ $^{127}\text{Xe}$  (with a half-life of 0.1 year),  
12  $^{133}\text{Xe}$ , and hyperpolarized (having non-equilibrium alignment of nuclear spins, suitable for  
13 magnetic resonance)  $^{129}\text{Xe}$ ] are inhaled, they can be tracked throughout the body by externally  
14 monitoring their **decay products** using magnetic resonance microscopy (high resolution  
15 magnetic resonance imaging (MRI) at microscopic [nanometer] levels) (Figure 4.54.2). This  
16 imaging technique is used to assess how well oxygen is taken up and transported by the blood  
17 [395].  
18  
19



20  
21  
22 **Fig. 4.54.2:** Xenon ventilation imaging has progressed greatly since first being used in 1998.  
23 One of the first  $^{129}\text{Xe}$  images (A) has been enhanced by improving polarization, gas delivery  
24 technology, and magnetic resonance (MR) acquisition strategies. The standard  $^{129}\text{Xe}$  image (D)  
25 is further improved in image quality through improving polarization, gas delivery technology,  
26 and MR acquisition strategies. (Image source: Driehuys and Hedlund, 2007, © Sage  
27 Publications) [395].  
28

### 29 **4.54.4 Xenon isotopes used as a source of radioactive isotope(s)**

30

31  $^{124}\text{Xe}$  is used in the production of **radioisotopes**  $^{123}\text{I}$  and  $^{125}\text{I}$  (with half-lives of 0.55 day and  
32 59 days, respectively) via the reactions  $^{124}\text{Xe} (n, n p) ^{123}\text{I}$  and  $^{124}\text{Xe} (n, \gamma) ^{125}\text{I}$ , respectively,  
33 which are used in diagnostic procedures and cancer treatment, respectively [395].