# **A carbon and nitrogen isotopic investigation of a case of probable infantile scurvy (6th- 4th centuries BC, Slovenia)**

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**Abstract**

This paper presents a case study of a young infant, from a larger isotopic and osteological investigation of Bronze/Iron Age (14th-4th century BC) skeletal assemblages from Croatia and Slovenia. The osteological analysis of this infant identified pathological lesions including abnormal porosity and new bone formation consistent with malnutrition and phases of recovery. The distribution and appearance of these pathological lesions (i.e. diffuse micro-porosities and plaques of subperiosteal new bone formation on the skull and long bones) led to the conclusion that this infant probably suffered from scurvy (vitamin C deficiency). The diet and nitrogen balance of this individual were investigated by incremental dentine sampling and stable carbon and nitrogen isotope analysis. This sampling method provided a high resolution record of dietary and metabolic changes from pre-birth to around the time of death. The resulting isotope data exhibited unusually high δ13C values for this region and time period (between -11.3‰ and -12.6‰), while δ15N values were observed to be c. 3‰ above that of rib collagen sampled from contemporary adults recovered from the same site. The isotope profiles generated from the incremental dentine analysis show that δ13C and especially δ15N continue to increase until death. The evidence from the skeletal remains and high resolution isotopic data support the hypothesis that this infant suffered from severe malnutrition and an increasingly negative nitrogen balance. The paper discusses some scenarios which could have resulted in these unusual isotope ratios, whilst considering the diagnosis of possible metabolic disease. The paper also addresses the need for context when interpreting isotopic results. The isotope data should not be viewed in isolation, but rather as part of a multidisciplinary approach, considering the multiple causes of isotopic variability.

**Key words**

Incremental dentine, nitrogen balance, pathology, vitamin C deficiency, breastfeeding, Early Iron Age

**1. Introduction**



Figure 1. Map of south-eastern Europe with the site of Zagorje ob Savi marked in red (Sources: [www.freeworldmap.net](http://www.freeworldmap.net); [www.esri.com](http://www.esri.com)).

This paper provides a case study combining contextual, osteological and isotopic data to produce a biography for a single individual buried at the Early Iron Age cemetery at *Zagorje ob Savi*, (46°8'7.84" N, 14°59'36.04" E) Slovenia (6th-4th centuries BC). The study was drawn from a larger osteological and stable isotope investigation of skeletal material recovered from central and eastern Slovenia, and northern Croatia (Nicholls 2017). The infant presented in this paper was exceptional in both their osteological and isotopic results.

**1.1 *Zagorje ob Savi***

The Early Iron Age cemetery at *Zagorje ob Savi* (*Kidričeva cesta*) is located at the foot of the hill known as *Ocepkov hrib*, where the contemporary settlement is located. The location of this site is highlighted in Figure 1. It lies close to the *Medija* stream, a tributary of the *Sava*, one of the major rivers of the region. The site belongs to the *Dolenjska* Early Iron Age (Hallstatt) regional group, although it is located on the northern fringes of this group (Supplementary 1) and was active only in the Late Hallstatt period (6th-4th centuries BC), as was the case for a number of sites in this region.

The cemetery is unusual in containing flat inhumation graves, in a region where monumental burial mounds are much more common. It is one of fourteen such cemeteries in the broader region, which, where they can be dated, generally belong to the Late Hallstatt period. Most are located in the northern part of the *Dolenjska* cultural group area, i.e. the *Posavsko hribovje* hill range, close to the *Sava* River (Gabrovec 1966; Dular, Tecco Hvala 2007; Draksler 2007).

The cemetery at *Zagorje ob Savi* has been known since the end of the 19th century, when several graves were found during construction work. Some contained high status artefacts, including a belt plate featuring a hunting scene; however, contextual information on these burials is scarce (Gabrovec 1966; Draksler 2007). Recent development-led excavations provided the first opportunity to acquire high quality data from this site. This work revealed nine graves, six of which yielded human skeletal remains, set in three distinct groups. The central group comprised five graves including one with the infant skeletal remains that forms the basis of this paper.

It is impossible to date all the graves individually, as some were disturbed or lacked any grave goods (e.g. grave 1 and 4). Nevertheless, those containing datable finds seem without exception to date to the Late Hallstatt Period (Draksler 2011; Murko 2011; Draksler and Murko 2020 in press). The most informative finds are a serpentine type fibula with a saddle-shaped bow and a disc from grave 8 (variant IVb after Tecco Hvala 2014), dating to the 6th century BC (Tecco Hvala 2014), and a Certosa type fibula, from grave 2 (variant XII after Teržan 1976), which is characteristic of the 4th century BC (Teržan 1976). Radiocarbon dates for two of the graves (the infant burial discussed here, i.e. grave 7 and grave 8, with the serpentine fibula) fall within the so-called Hallstatt plateau, between approx. 750 and 400 BC.The infant grave (SUERC-69421, 2412 ± 29 BP) is weighted to the 6th and 5th centuries BC (550-401 cal BC (80.8 %); OxCal v4.2.4), while grave 8 (SUERC-69422, 2499 ± 28 BC) is weighted to the 8th to 6th centuries BC (781-538 cal BC (95.4 %)); OxCal v4.2.4; although, as mentioned above, the 6th century fibula from this grave would suggest that it falls towards the end of this distribution.

## **1.2 Evidence of diet in Iron Age Slovenia**

Iron Age Slovenia was inhabited by agricultural communities. Their diet probably consisted of domesticated plants and animals. Evidence of faunal remains obtained from large settlement centres, such as *Stična* and *Cvinger*, show a predominance of cattle, followed by caprine (sheep/goat) and pig (Dular and Tecco-Hvala 2007). Wild animals made up a relatively small proportion of faunal remains, the most common being red deer, wild boar and roe deer (Dular and Tecco-Hvala 2007).

The botanical material has provided evidence for a mixture of domesticated plants, including C3 cereals (barley, wheat, oats, and rye), legumes (vetch, faba bean, pea, lentils) and vegetables (cabbage, mustard, turnip and kohlrabi) (Dular and Tecco-Hvala 2007). The charred remains of millet (a C4 plant) grains found in over a dozen late prehistoric sites in south-east Slovenia and northern Croatia, are evidence that this crop was being exploited in this region by the Late Bronze Age, if not earlier (Dular and Tecco-Hvala 2007; Karavanić et al. 2015; Reed and Drnić 2016).

**1.3 Stable isotope analysis**

The analysis of stable carbon and nitrogen isotopes in human bone and dentine collagen has become commonplace for the investigation of diet in the past (Tykot 2004; Craig et al. 2009; Tafuri et al. 2009; Killgrove and Tykot 2013; Lightfoot et al. 2014). δ15N values have also been used for the investigation of health status, as this ratio is not only reflective of diet, but of an individual’s nitrogen balance (Fuller et al. 2004; Fuller et al. 2005; Beaumont et al. 2013; Beaumont et al. 2015). This balance is affected by dietary nitrogen, but also by metabolic processes and states, such as growth, malnutrition, and illness (Mekota et al. 2009; Reitsema 2013; Waters-Rist and Katzenberg 2010). It depends on the amount of nitrogen excreted in relation to the amount of dietary nitrogen ingested. The body falls into a negative nitrogen balance also known as catabolism, or a catabolic state, for example, during periods of nutritional stress or disease (Fuller et al. 2005; Mekota et al. 2009; Beaumont et al. 2013; Beaumont et al. 2015). Catabolism raises δ15N values as the body breaks down tissues to replace nitrogen missing from the diet, which results in an increase in the 15N in newly-formed tissues relative to the original, and can be misinterpreted as ingestion of higher trophic level foods. The body enters into a positive nitrogen balance when less nitrogen is excreted than ingested, commonly during times of rapid growth, such as infancy and puberty (Waters-Rist and Katzenberg 2010). This is also known as anabolism, or an anabolic state, and has been reported to lower δ15N values (Fuller et al. 2006, Waters-Rist and Katzenberg 2010). This current case study from Early Iron Age Slovenia aims to investigate temporal changes in nitrogen balance in the hard tissues of an infant exhibiting a range of pathological skeletal lesions.

This study illustrates the advantages of bringing together multiple strands of evidence to create more informed interpretations regarding life and death in prehistory. It also demonstrates the importance of context when interpreting stable carbon and nitrogen isotope data.

**2. Material and Methods**

**2.1 Osteological Analysis**

Poor preservation can limit the nature of information gleaned from osteological analysis (Bytheway and Ross 2010). The burial conditions within the current study area are predominantly detrimental to bone preservation, leading to severe cortical exfoliation, root etching, as well as a variable level of completeness, frequently 50% or less (Nicholls 2017).

Osteological and palaeopathological analysis was undertaken through macroscopic observation and using a magnifying hand lens. Standard analysis and recording methods were used for biological sex assessment (pelvis: (Walker 2005; Phenice 1969; Klales et al. 2012) skull: (Buikstra and Ubelaker 1994; Walker 2008; İşcan and Steyn 2013)) and age estimation (Buckberry and Chamberlain 2002; Brooks and Suchey 1990; Brothwell 1981). Tooth wear (Brothwell 1981) was used with caution due to a lack of published, calibrated, population-specific standards for the study area. Supporting evidence was taken from late fusing epiphyses (Belcastro et al. 2008; Webb and Suchey 1985). Due to the generally poor preservation of the skeletal material, broad ranges of young, middle and mature adult were considered more appropriate than numerical ages (after Buikstra and Ubelaker, 1994). Standard protocols and guidelines were used for recording the appearance and location of pathological lesions (Buikstra and Ubelaker 1994; Aufderheide and Rodríguez-Martín 1998; Brickley and Ives 2008). Although some radiographs were taken, they were of limited use due to the presence of taphonomic damage (including the loss of metaphases on all but the right femur of the infant) and soil within the medullary cavities and trabecular spaces. The degree of fragmentation did, however, allow for the observation of the internal structures of some of the long bones.

Age estimation for the infant was carried out using dental development (AlQahtani 2010; Moorrees et al. 1963). Due to the young age of the infant the remains were not assessed for sex.

**2.2 Isotope analysis**

Stable carbon and nitrogen isotope ratio analysis was carried out on rib collagen from three adults interred within the same site. A rib collagen sample was taken from the infant, in addition to incremental dentine collagen sampling. Care was taken not to sample from an area that exhibited any pathological lesions (Olsen et al. 2014). The study also included faunal material from three cows, two pigs and a deer. Animal bones were excavated from the Late Bronze/Early Iron Age settlement adjacent to the cemetery assemblage, and so represent a contemporary faunal baseline of stable carbon and nitrogen isotope ratios for the area.

All collagen extractions were carried out following the modified Longin method (Longin 1971; Brown et al. 1988). Rib fragments (c. 300mg) were demineralised in 0.5M HCl at 4°C, which took between two and four weeks. Once the production of CO2 had ceased and the reaction was complete, all samples were rinsed three times with deionised water and placed in an HCl solution of pH3 at 70°C for 48 hours to gelatinise. The solutions were filtered using Ezee filters, followed by centrifugal filtering using Millipore ultrafilters. The resulting liquid was then freeze-dried, weighed in duplicate and measured at the University of Bradford Stable Isotope Facility by combustion in a Thermo Flash EA 1112. Internal and external standards were run throughout, as well as separated N2 and CO2 references gases, using a Delta plus XL via a Conflo III interface. The analytical precision for both the δ13C and δ15N, based on instrumental error, is ± 0.2‰.

For the infant, seven 1mm incremental dentine samples were taken from the crown to the apex of the first right maxillary deciduous incisor, following method 2 from Beaumont et al. (2013). The first right maxillary incisor was selected for stable isotope analysis based on preservation and because this was the most developed of the available teeth. Because of the overlap of dentine layers in each 1mm sample (apart from the first and final sections as the tooth was forming at the time of death) the isotope results will represent a rolling average of dietary input (Beaumont et al. 2013). Temporal resolution is much better than that seen in bone collagen, but the results are discussed in terms of general trends of increasing and decreasing isotope values, rather than attempting to assign exact age to each section.

Isotopic data, together with associated chronological and other supporting information from this study, has been added in the IsoArcH database (Salesse et al. 2018; Salesse et al. 2019).

## **3. Results**

## **3.1 Osteological analysis**

The first and second right incisors had complete crowns and ¼ roots. The right canine and first molar had yet to erupt from the maxilla, while the crown of the second right molar (loose) was complete, but with no initiation of the root. The stage of development and tooth eruption is consistent with an age of 7.5 months +/- 3 months following AlQahtani (2010). The development of the second molar is consistent with an age of 6-9 months following Moorrees et al. (1963). It was concluded that the infant was 6-9 months old at the time of death based on dental development (AlQahtani 2010; Moorrees et al. 1963). The age and sex determinations of the adult skeletal remains are presented in Table 1.

* 1. The preservation and fragmentation of skeletal remains has had considerable consequences for the identification of pathological lesions. Subsequently, it cannot be said that pathological lesions were never present, only that there was no evidence upon examination.
  2. The middle aged adult female buried in grave 8 exhibited cribra orbitalia and diffuse porosity across the cranial vault fragments. Diffuse porosity and compact, striated bone was also identified on surviving fragments of the long bones. Probable systematic non-specific inflammation possibly due to infection or metabolic disease is suggested, however, further diagnosis was not possible due to the fragmented nature of the remains.
  3. No other palaeopathological lesions were observed on the other adults. However, the taphonomic damage to the skeletal remains was very severe, and the cortical surface of these remains was heavily impacted. There is a possibility that any pathological lesions were destroyed in the burial environment.

## **3.1.1 Palaeopathological analysis of the *Zagorje ob Savi* infant**

Some of the pathological lesions described here are depicted in Figure 2. The skeletal remains of the infant exhibited a high prevalence of abnormal porosity. In the cranium, this was noted in the roof of both eye orbits; diffusely across the ectocranial surface of the surviving fragment of frontal bone; and in small patches on both parietal bones around the parietal bosses and the coronal suture. The external areas of the right maxilla (Figure 3A), extending away from the alveolar bone towards the infra-orbital foramen, and the hard palate also exhibited extensive porosity.

Isolated small plaques of porous woven bone were observed on the endocranial surfaces of the parietal bones (Figure 3B) and the squamous portion of the temporal adjacent to the squamous suture. Layers of woven and compact new bone formation were also identified in the roof of the eye orbits (Figure 3C), the external (inferior) surface of the basilar portion (Figure 3D), and on the surface of the left pars lateralis of the occipital, adjacent to the occipital condyle and the hypoglossal canal.

Diffuse plaques of porous sub-periosteal new bone formation were identified on the surviving long bones, specifically; covering the middle third of both femoral and both humeral diaphysis (Figure 3E), the proximal and middle third of the right fibula and the entire surviving fragment of the right tibia (Figure 3F) (the left side being absent). Compact and striated bone was also noted along the linea aspera of the right femur (Figure 3G). The additional surviving fragments from the axial skeleton were too poorly preserved to detect the presence or absence of any pathological lesions.

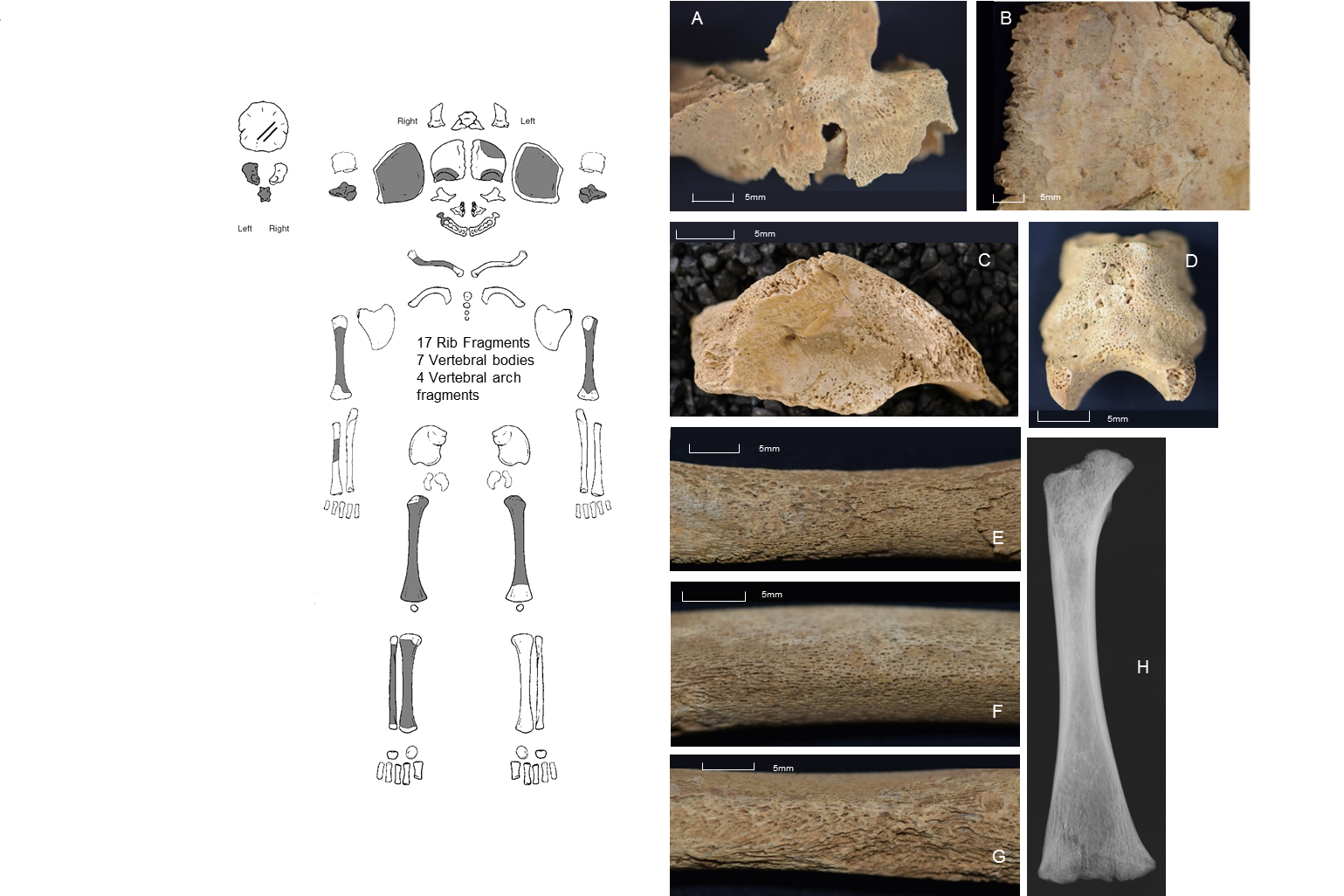


Figure 2. **Left:** Inventory of skeletal remains from the infant; **Right: A**. Right maxilla fragment showing porosity around maxillary sinus; **B**. Endocranial surface of parietal bone with plaques of compact bone; **C.** Right orbit with layers of porous woven bone; **D.** Basilar portion with abnormal porosity; **E.** Posterior right humerus showing plaque of porous woven and compact bone on top of original cortical surface **F**. Anterior right tibia showing plaque of porous woven and compact bone on top of original cortical surface; **G** Posterior right femur with porous compact bone on and around the linea aspera; **H**. Radiograph of right femur

### **3.2 Carbon and nitrogen Isotopes**

**3.2.1 Rib collagen samples of individuals buried at *Zagorje ob Savi***

The stable carbon and nitrogen isotope results of bulk rib collagen from four individuals buried at *Zagorje ob Savi* can be seen in Figure 3 and Table 1. All samples fell within the accepted C:N ratio and >1 % collagen yield, indicating good collagen preservation (Van-Klinken 1999).

Adult rib δ13C values range between -15.2 and -14.4‰, and δ15N values between 8.6 and 9‰. The collagen extracted from the rib of the infant has considerably higher δ13C (+2.9‰) and δ15N (+3‰) values when compared to the adult values.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| *Specimen* | *Element* | *Grave No.* | *Sex* | *δ***13***C‰* | *δ***15***N‰* | *C:N* | *Amt%N* | *Amt%C* |
| Middle adult\* | rib | 2 | M | -14.8 | 8.6 | 3.2 | 16.7 | 46.0 |
| Middle adult | rib | 8 | F | -15.2 | 8.9 | 3.2 | 17.4 | 47.6 |
| Middle adult | rib | 5 | F | -14.4 | 9.0 | 3.2 | 17.2 | 46.8 |
| Infant | rib | 2001 | N/A | -11.9 | 11.8 | 3.2 | 15.3 | 41.3 |
| Infant dentine (average of whole tooth root) | incisor | 2007 | N/A | -12.1 | 12.0 | 3.2 | 15.9 | 43.2 |
| Deer | metatarsal | 9 | N/A | -23.1 | 3.3 | 3.3 | 13.5 | 38.1 |
| Cow | scapula | 5 | N/A | -19.0 | 4.8 | 3.2 | 14.9 | 41.0 |
| Cow | metatarsal | 1 | N/A | -19.2 | 6.3 | 3.2 | 15.2 | 41.9 |
| Cow | metatarsal | 5 | N/A | -18.5 | 6.7 | 3.3 | 14.4 | 40.5 |
| Pig | mandible | 8 | N/A | -21.0 | 6.3 | 3.2 | 14.4 | 40.1 |
| Pig | scapula | 8 | N/A | -19.3 | 4.8 | 3.3 | 14.0 | 39.1 |

Table 1. Carbon and nitrogen ratios from human and faunal remains from *Zagorje ob Savi* (\*Middle adult age range 36-50yr, Buikstra and Ubelaker 1994)

**Figure 3. Plot of carbon and nitrogen isotope ratios obtained from the remains of individuals buried at *Zagorje ob Savi*. Samples include rib bone collagen from three adult individuals (black circles) and the rib (white circle) and mean dentine collagen extracted from the Infant (grey triangle). The plot also includes carbon and nitrogen isotope ratios of animal bone collagen sourced from graves in the same cemetery assemblage as the human remains. The error bars represent the analytical precision of carbon and nitrogen isotope analysis, based on instrumental error of ± 0.2‰.

**3.2.2. Incremental dentine collagen analysis of a first deciduous incisor from the infant**

The results of incremental dentine collagen analysis are shown in Table 2 and Figure 4. Figure 4 displays an increase in both δ13C (+1.2‰) and δ15N (+2.3‰) values throughout the development of the tooth root. If we accept that this tooth represents tissue from 2 months pre-birth through to 9 months post-birth, (Beaumont and Montgomery 2015) we can see that, in comparison to δ13C values, δ15N values are observed to increase more rapidly and by a higher magnitude, most notably from section 5 through to death (δ15N = +1.7‰; δ13C = + 0.8‰).

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| *Dentine increment* | *δ13C ‰* | *δ15N ‰* | *C:N* | *Amt%N* | *Amt%C* |
| 1 (crown) | -12.6 | 11.2 | 3.2 | 15.8 | 43.3 |
| 2 | -12.4 | 11.4 | 3.2 | 14.1 | 38.4 |
| 3 | -12.2 | 11.6 | 3.2 | 16.4 | 44.5 |
| 4 | -12.1 | 11.8 | 3.2 | 16.8 | 45.4 |
| 5 | -12.0 | 12.1 | 3.2 | 16.7 | 45.1 |
| 6 | -11.7 | 12.7 | 3.2 | 16.5 | 45.0 |
| 7 (apex) | -11.3 | 13.5 | 3.2 | 14.8 | 40.6 |

Table 2. Carbon and nitrogen isotope ratios of dentine increments sampled from the first deciduous incisor*, Zagorje ob Savi* infant

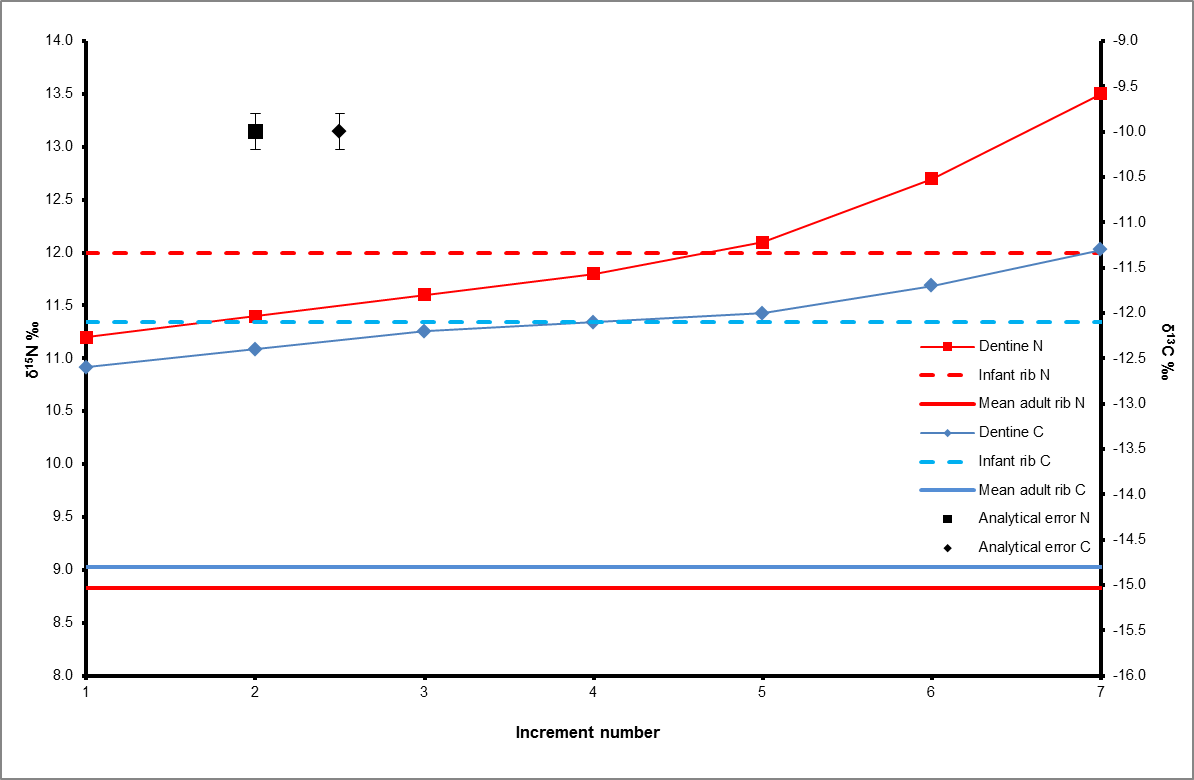


Figure 4. A plot of carbon and nitrogen isotope ratios of incremental dentine sections sampled from the first deciduous incisor of the *Zagorje ob Savi* infant. The red and blue trend lines represent changing metabolic and dietary conditions from pre-birth to around the time of death, reflected through isotopic variation. The red line represents the mean adult rib δ15N value of 8.8‰ and the blue line the mean adult rib δ13C value of -14.8‰. The red dashed line represents the infant rib δ15N value of 11.8‰ and the blue dashed line the infant rib δ13C value of -11.9‰. The error bars represent the analytical precision of carbon and nitrogen isotope analysis, based on instrumental error of ± 0.2‰.

## **4. Discussion**

### **4.1 Pathological lesions**

Normal growth and development of the infant skeleton causes areas of porosity because of increased vascularisation, particularly around the growth plates at the metaphyses of long bones (Cunningham et al. 2016). Ortner et al. (2001) suggest that porous lesions that extend 10 mm past the growing metaphyseal end of the diaphysis are less likely to be connected to normal bone growth. Sub-periosteal bone formation has been linked with several pathological conditions, including non-specific inflammation and metabolic disease, such as haemolytic (caused by the premature destruction of red blood cells) or megaloblastic (deficient in vitamin B12 and folic acid) anaemia, rickets (childhood vitamin D deficiency) and scurvy (vitamin C deficiency) (Ortner and Ericksen 1997; Ortner and Mays 1998; Ortner et al. 2001; Brickley and Ives 2006; Mays 2008a; Walker et al. 2009; Snoddy et al. 2018).

**4.2. Differential diagnoses**

Megaloblastic and haemolytic anaemias can be caused by a range of dietary and genetic factors (Walker et al. 2009). Porosity within the eye orbit and the cranial vault has been linked to anaemia, thought to occur as a response to increased red blood cell production. Expansion of the hematopoietic marrow within the diploë of the skull causes greater vascularisation at the expense of the outer and inner table of the cranial vault (Walker et al. 2009). Whilst porosity was observed at these locations in the current investigation, the absence of hyperplasia of the diploë and the presence of sub-periosteal new bone formation suggests these lesions were not primarily the result of anaemia (Ortner et al. 2001; Brickley and Ives 2006).

Rickets results from the inability to adequately mineralise bone due to the lack of vitamin D during growth. This produces pores in dry bone as the inadequately mineralised osteoid that once filled these spaces does not survive the burial environment (Ortner and Mays 1998; Mays et al. 2006). It is commonly associated with the deformation of long bones and sternal rib ends, as well as a roughened bone surface beneath epiphyseal joint plates. As some of these deformities are related to weight bearing whilst crawling or walking, it is unlikely they would be visible in the lower limbs of this infant. Bony changes have been recognised in infants as young as 3 months, affecting the upper limb, ribs, orbital roof, cranial vault, and growth plates (Ortner and Mays 1998). A pattern of struts and slits across major long bones were also observed by Ortner and Mays (1998), showing that rickets is diagnosable in infants who have not yet begun walking. In the *Zagorje ob Savi* case, macroscopic and radiographic (right femur; Figure 3H) observations did not identify deformation of the long bones, struts or slits near the metaphyses, and the one surviving metaphyseal surface (right, proximal femur) appeared normal. This makes a diagnosis of rickets unlikely.

Non-specific inflammation involving the periosteum can also result in abnormal bone porosity (Lewis 2004; Weston 2008). Plaques of new bone on top of the original cortical surface of long bones have also been associated with an inflammatory response to non-specific infection (Roberts and Manchester 2010). Whilst this type of lesion was observed in the *Zagorje ob Savi* infant, Ortner (2003) has cautioned that under the age of four, woven periosteal new bone at this location this cannot be differentiated from normal bone growth. Thus, whilst non-specific inflammation due to a systemic infection cannot be ruled out there is insufficient skeletal evidence to make this diagnosis.

The *Zagorje ob Savi* infant exhibits several features that have been associated with scurvy. Using criteria suggested from Snoddy et al’s (2018) synthesis of clinical and palaeopathological studies, we have highlighted skeletal lesions that have been described as ‘suggestive’ or ‘diagnostic’ (see Table 3).

|  |  |  |
| --- | --- | --- |
| Lesion Location | Lesion type | Proposed diagnostic strength |
| Frontal orbital roof | Bilateral sub-periosteal new bone | Diagnostic |
| Right maxilla: Anterior surface/ infra-orbital foramen | Abnormal cortical porosity | Diagnostic |
| Right maxilla: palatal surface | Abnormal cortical porosity | Diagnostic |
| Ectocranial parietal | Abnormal cortical porosity | Diagnostic |
| Endocranial parietal and squamous temporal | Islands of abnormal cortical porosity and sub-periosteal new bone | Suggestive |
| Occipital inferior surface: pars basilaris | Abnormal cortical porosity and sub-periosteal new bone | Suggestive |
| Femur: linea aspera | Sub-periosteal new bone | Suggestive |
| Appendicular skeleton: diaphysis | Sub-periosteal new bone (diffuse) | Diagnostic\* |

Table 3: Macroscopic lesions that have been attributed to scurvy (Snoddy et al. 2018) \*not exclusive to scurvy

There are only two additional features described by Snoddy et al. as ‘diagnostic’ and these are found on the sphenoid and scapulae bones. Unfortunately, in the current case, these elements were absent and, therefore, could not be assessed. However, we have identified five criteria thought to be diagnostic of scurvy and others that are suggestive (Table 3). Additionally, Ortner et al. (2001) have argued that scurvy can affect the skeleton without involving the greater wing of the sphenoid and Brickley and Ives (2006) have cautioned against the over-dependence on one skeletal element for the diagnosis of pathology.

Scurvy occurs when there is insufficient vitamin C intake. This micronutrient is responsible for the correct formation of collagen (Munday et al. 2005; Tsuchiya and Bates 2003). Collagen is a structural protein that is ubiquitous within the human body contributing to the formation of skin, muscle, tendon, the lining of vital organs, as well as bones and teeth. A lack of vitamin C can thus result in weakened connective tissues and vessels that are susceptible to haemorrhage (Hirschmann and Raugi 1999). Chronic bleeding of tissues adjacent to the bone surface can lead to an inflammatory response, i.e. porosity and sub-periosteal new bone formation (Mays 2008b). The pathological lesions consistent with scurvy have been argued to be more common in the skeletal remains of young children because of the high demands of rapid growth (Ortner and Ericksen 1997; Ortner et al. 1999). As the child develops, poor collagen formation leads to arrested osteoblastic activity and the collagenous bone matrix does not ossify.

The new bone growth identified across the skull and long bones was formed on top of the existing cortical surface, consistent with haemorrhage resulting from minor trauma, for instance lesions in the eye orbit may be caused by the movement of the ocular muscles (Lewis 2004; Mays 2008a; Moore and Koon 2017; Walker et al. 2009). The plaques of compact bone identified on the endocranial surface of the skull, in conjunction with the layers of woven and compact bone observed within the orbits and on the long bone diaphyses, suggest that there were phases of recovery and healing, as new bone could only have been laid down during a recovery phase following the reintroduction of vitamin C (Brickley and Ives 2006). The lesions are, therefore, consistent with chronic scurvy with some evidence of reintroduction of vitamin C prior to death (Brickley and Ives 2006; Mays 2008a; Walker et al. 2009).

It is important to note that a malnourished individual could suffer from more than one disease or condition. The occurrence of one can weaken the immune system, causing the individual to become more susceptible to other illnesses. The individual presented here may have been additionally affected by any of the disorders mentioned above, as well as other deficiencies and infections that do cause skeletal changes. Overall, it was concluded that this infant had suffered from severe and chronic malnutrition, resulting in vitamin C deficiency and probable scurvy.

Clinical symptoms associated with scurvy become identifiable after 1-3 months of insufficient vitamin C intake, depending on existing body stores (Hodges et al. 1969; Larralde et al. 2007) and haemorrhages only after around 6 months of total absence of vitamin C (Hodges et al. 1971). Given the presence of bony lesions on the infant skeleton it is likely they suffered from a chronic deficiency of vitamin C.

It is known from modern-day cases of infantile scurvy that abnormal collagen formation can lead to bleeding in the skin, mucous membranes, muscles, gastrointestinal tract and around the joints (Jackson and Park 1935; Larralde et al. 2007; Weinstein et al. 2001). Individuals suffering from scurvy may also have difficulties with coagulation of the blood, which can exacerbate the problem and lead to complications with healing wounds (Larralde et al. 2007). Clinical symptoms include fatigue, irritability, and delayed or stunted development. Haemorrhagic skin lesions are common, and haemorrhaging from the eyelids and gums has also been described (Larralde et al. 2007; Weinstein et al. 2001).

Swelling of the lower extremities has been observed, potentially with multiple causalities, including leaking capillaries and soft tissue haemorrhage (Larralde et al. 2007). Intense joint pain and muscle weakness, especially in the lower limbs, constricts movement and leads to children laying in an abduct or ‘frog-like’ position (Larralde et al. 2007). Therefore, the *Zagorje ob Savi* case of probable infantile scurvy would have resulted in particularly visible symptoms.

### **4.3 Carbon and nitrogen isotopes**

**4.3.1 Animal bone collagen**

There is a small shift in the carbon and nitrogen isotope ratios between wild (i.e. deer) and domesticated animals. This is probably due to the higher levels of controlled feeding in domesticated animal populations, including potential penning, which can raise δ15N values if animals are consuming vegetation grown in 15N enriched soil by their faeces, and foddering, where fodder may have similarly been grown on land enriched in 15N from manure (Bogaard et al. 2007; Fraser et al. 2011). Additionally, if this fodder included C4 plants (grain or crop waste), δ13C values would also be increased relative to non-foddered animals (Tieszen 1991; Tafuri et al. 2009). If the deer were living under heavily forested conditions, the δ13C values produced from their bone collagen could be reflecting the ‘canopy effect’, where plants become depleted in 13C because of either reduced light intensity or recycled CO2 (Bonafini et al. 2013; Noe-Nygaard et al. 2005; Van der Merwe and Medina 1991). This depletion is then transmitted up the food chain. If this were the case, the consumption of vegetation under forest conditions could mask the consumption of native C4 species.

When the human carbon and nitrogen isotope data is compared to the animal baseline, the predicted trophic level shift of c.2-5‰ between humans and most herbivores is present (Ambrose 1991; Hedges and Reynard 2007). This supports the interpretation that humans were ingesting herbivorous animal protein, either dairy or meat.

**4.3.2. Human bone and dentine collagen**

The δ13C and δ15N values produced as part of this study are indicative of a terrestrial-based diet, with a focus on a mix of C3 and C4 plants, probably millet, with the addition of herbivorous animal protein (Hedges and Reynard 2007; Richards 2003; Tieszen 1991; Tykot 2004). This is consistent with contemporary faunal and botanical evidence (Dular and Tecco-Hvala 2007).

Isotopically, there is a clear difference between the bone (and dentine) collagen of the infant and the three contemporaneous adults from *Zagorje ob Savi* (Figures 3 and 4). On first examination, and given the age of the infant (6-9 months), this elevated δ15N value is likely linked to breastfeeding. High δ15N values have been linked to breastfeeding practices in previous isotopic studies (Beaumont et al. 2013; Fuller et al. 2006; Jay et al. 2008; King et al. 2018; Millard 2000). Modern data (infant hair and nail) has demonstrated that breastfeeding results in a rapid trophic level shift in δ15N of 1.5‰ (de Luca et al. 2012) and up to 2-3‰, (Fuller et al. 2006) above birth values. This rise occurs as the infant ingests the proteins of their mother via breastmilk, which results in additional isotopic fractionation in comparison to the maternal δ15N values (Fogel et al. 1989). δ15N drops gradually to a value similar to the mother throughout the process of weaning (introduction of solid foods commonly combined with continued nursing) and the cessation of breastfeeding, provided the mother and child are sharing a similar diet (Fuller et al. 2006; Jay et al. 2008; Millard 2000) (see Supplementary 2) . A smaller trophic shift of 0.4 and 1‰ in δ13C values has similarly been observed (de Luca et al. 2012; Fuller et al. 2006; Tsutaya and Yoneda 2015).

The ~2‰ difference between the first infant dentine increment and the mean adult rib δ15N value fits this breastfeeding pattern. δ15N values continue to rise throughout the development of the dentine, a trend that similarly supports the interpretation of a trophic shift consistent with breastfeeding. The difference of 3‰ between the adult and infant rib collagen δ15N values is also consistent with a trophic shift related to breastfeeding. When viewed at face value, the infant isotope profiles presented here can reasonably be associated with the onset and continuation of exclusive breastfeeding. However, the first increment (which will include pre-natal dentine) is already comparatively high, and there is a gradual rise in the δ15N values rather than the rapid upward curve expected (see Supplementary 2). δ15N values for the infant then rise sharply through the final three increments prior to death. δ13C values follow a similar trend. The final increment of infant dentine (representing the time of death) has a δ15N value 4.7‰ above that of the mean adult rib value, while the δ13C value is 3.5‰ above the adults. Both of these differences in isotope ratios are greater than that anticipated from breastfeeding alone. Indeed, δ13C and δ15N from rib collagen (-11.9‰ δ13C; 11.8‰ δ15N) for this infant was notably higher than for other rib collagen samples from across Iron Age Slovenia, with mean δ13C and δ15N values of -15.3‰ and 9.1‰, respectively (Nicholls and Koon 2016; Nicholls 2017). With the additional context of chronic dietary deficiencies, the isotope data indicates a more nuanced interpretation.

**5. Interpretations**

The following presents some potential scenarios, which could explain the variation in carbon and nitrogen isotope ratios observed between the infant and associated adults.

**5.1 Dietary change**

The high δ13C and δ15N, particularly in the initial incremental dentine samples of the infant, suggest a possible change in maternal diet over time. The high δ13C values ranging between -12.6‰ and -11.3‰ indicate either a substantial marine or C4 dietary input and are considerably higher than that of the isotope ratios obtained from the three adults (+3 - 4‰). The ingestion of marine food is unlikely given the location of the cemetery, but also there is no isotopic or faunal evidence (to date) for the consumption of marine food during prehistory in Slovenia (Murray and Schoeninger 1988; Nicholls and Koon 2016). There is, however, substantial evidence for the consumption of C4 plants in the wider area during the Late Bronze and the Iron Age, namely millet (Dular and Tecco-Hvala 2007; Lightfoot et al. 2014; Murray and Schoeninger 1988; Nicholls and Koon 2016; Reed and Drnić 2016). The adult individuals buried at *Zagorje ob Savi* also provide evidence of C4 plant consumption. It is posited, then, that a substantial amount of the dietary carbon for the infant was obtained from millet, and that the amount of millet ingested increased with time, as indicated by the increase in δ13C values throughout the development of the tooth.

This is not to say that the infant was ingesting millet as a grain—given the age of the individual it is unlikely that they were eating solids—but that the C4 isotope signature was being transmitted indirectly into their diet through another route. This signal may have initially been introduced to the foetus in-utero via the mother. If the infant was nursing, the gradually increasing δ13C values could indicate that the mother was also ingesting increasing quantities of millet whilst breastfeeding. In addition, a small portion of the increase (i.e. 0.4-1.0 per mil) could be caused by the small carbon isotope breastfeeding trophic shift.

While this explains the carbon isotope values, it does not account the rapid increase in nitrogen isotope values observed in the latter three increments. The ingestion of millet would not result in elevated δ15N. Additionally, the increase in δ15N is beyond the trophic level shift associated with breastfeeding. This suggests that isotopic changes are not solely the result of a change in diet.

**5.2 Nutritionally stressed mother equals nutritionally stressed child**

Studies have shown that the nitrogen isotope ratios of children are complex, linking high δ15N values to changes in metabolism, maternal health, or falling into a negative nitrogen balance resulting in tissue catabolism (Beaumont et al. 2015; Reynard and Tuross 2015). When the body does not receive required dietary nitrogen (e.g. needed for protein synthesis), it instead obtains it from its own tissues. This breakdown of body tissues, known as catabolism, causes isotopic fractionation as molecules containing the lighter isotope, 14N, break down more readily. This results in an enrichment of the heavier isotope, 15N, in the tissues that are then sampled for isotopic analyses (Mekota et al. 2009). Increased δ15N values have been linked to malnutrition on several occasions (Hobson et al. 1993 (animal); Fuller et al. 2005; Mekota et al. 2009 (clinical); Beaumont et al. 2015 (archaeological)). If the mother of the infant was severely malnourished, this would have had a negative impact on their child. Studies into archaeological dentine and modern hair and breast milk samples have suggested that high δ15N values in infants are not exclusively the result of the ingestion of breastmilk (Beaumont et al. 2013; Beaumont et al. 2015; Beaumont and Montgomery 2016; de Luca et al. 2012; Romek et al. 2013). In the current study, δ15N values continue to rise post-birth, beyond the trophic shift associated with breastfeeding. This may reflect the mother’s continued metabolic status, in addition to her diet. It might equally reflect undernutrition in the infant to the extent that they are beginning to recycle their own tissues (Beaumont and Montgomery 2016).

When examining the earliest incremental dentine sections, probably including dentine formed in-utero, the high nitrogen isotope ratios could actually be reflecting a mother in the third trimester of pregnancy. The unborn foetus would have obtained its nutrients directly from the mother across the placenta, thereby transferring the maternal isotope ratios into the forming tissues of the unborn baby (Beaumont et al. 2015). δ15N values influenced by dentine formed prior to birth are notably higher than that of the three adult rib values (c.+2.4‰ between the mean adult rib δ15N value and the first dentine increment of the infant). As discussed above, this δ15N value may be indicative of the nitrogen balance of a physiologically stressed mother.

At this point, to understand more fully the comparatively high δ15N values (and comparatively smaller corresponding increase in δ13C values) from the bones and teeth of the infant, it is vital to consider the probable diagnosis of chronic scurvy. A lack of dietary vitamin C alone does not cause starvation or increased δ15N values associated with catabolism. In fact, clinical paediatric studies have shown normal weight gain in children with vitamin C deficiency, but an otherwise normal diet (Weinstein et al. 2001). Nevertheless, there are several reasons why scurvy may have been a factor in causing malnutrition. Symptoms of scurvy include painful and bleeding gums and irritability in infants; this may well have caused difficulties with feeding and suckling (Larralde et al. 2007). Also, vitamin C deficiencies are often linked to other nutrition deficiencies and co-morbidities (Schattmann et al. 2016), such as infections, that would exacerbate feeding problems and could themselves lead to malnutrition (Nguyen et al. 2013). A deficiency in vitamin C can also inhibit the absorption of iron from the small intestine, leading to iron deficiency anaemia (Brickley and Ives 2006; Weinstein et al. 2001) and leaving the infant more susceptible to infection and disease due to an under-developed immune system (Siegel, 1993; Thomas and Holt, 1978).

Even though there is evidence to suggest the mother was malnourished during pregnancy and breastfeeding, this should not normally have caused the skeletal manifestations associated with scurvy observed in these infant remains. It is known that breast milk usually contains enough nutrition for an infant to thrive without additions to their diet, including water, for the first 6 months (Larralde et al. 2007; WHO 2005). The minimum required intake of vitamin C for infants is unestablished; however, the recommended daily allowance for 0 to 12 months varies from 35-50mg (Byerly and Kirskey, 1985; NIH 2018; NRC 1989), which is well within the range of vitamin C available in breastmilk (Byerly and Kirskey, 1985) even in marginally nourished women with low serum vitamin C levels (Ahmed et al. 2004). During periods of short term stress, pregnant or lactating women will generally route necessary nutrients to the foetus or milk production at their own expense (Ahmed et al. 2004; Bates and Prentice 1994; Byerly and Kirskey 1985). For a breastfed infant of this age (6-9 months) to exhibit such severe nutritional deficiencies, the mother must also have suffered from severe malnutrition (Bates and Prentice 1994; Jackson and Park 1935). This has been demonstrated in very rare clinical cases of congenital scurvy, where it is likely that the mother herself suffered from the disease whilst pregnant or lactating (Hirsch et al. 1976; Jackson and Park 1935).

This interpretation of the mother’s health status should be treated with caution. The infant was buried in an individual grave and even had they been buried with an adult, it would be impossible to relate individuals without the use of aDNA, which could not be done in this instance. Furthermore, if it had been possible to identify the mother the slower rate of bone turnover in adults means that it is unlikely specific nutritional stress during pregnancy would be detectable in the rib collagen of the mother.

**5.3 Supplementation with animal milk?**

One possibility that could account for both the high δ13C and δ14N values (relative to adult rib collagen) is that they were the result of the ingestion of animal milk, as an emergency replacement or supplement for breastmilk, where the animal had been foddered on millet. As stated above, a successfully breastfed infant should not suffer from scurvy, as breast milk provides a good source of vitamin C. If a scenario arose where dietary supplementation was required, the ingestion of unmodified animal milk, for example bovine milk, by young infants (c. <12 months) is particularly unhealthy (Fleischer Michaelsen et al. 2000; Binns et al. 2007). Animal milk as a substitute for breastfeeding is a potential cause for this infant’s illness, as cow’s milk and goat’s milk are deficient in vital nutrients, including vitamin C. The consumption of cow’s milk by infants under the age of 12 months can cause blood loss from the gastrointestinal tract, also leading to iron deficiency, anaemia and diarrhoea (Binns et al. 2007; Wijndaele et al. 2009; Griebler et al. 2016). Although the ingestion of animal milk itself would not lead to raised δ15N values, the consequences to the health of an infant could result in a compromised immune system and illness, which, in turn could have resulted in tissue catabolism and 15N enrichment.

Overall, it is not possible to establish which of these scenarios was most likely. All, some or none of these scenarios could account for the isotopic and osteological evidence presented in this case study. It is clear that multiple factors can affect the interpretation of stable isotope data and palaeopathological lesions. It is, therefore, vital that any interpretations are augmented with all available contextual information.

**6. Conclusion**

The infant presented here suffered from severe and chronic malnutrition, probably scurvy. This has been supported by both the stable isotope data and osteological observations. The rib and dentine collagen from the infant suggest that the mother was malnourished during pregnancy passing on high δ13C and δ15N values to the tissues of the child. The isotope ratios of the adults presented in the current investigation are consistent with those found at a regional scale. The infant buried at *Zagorje ob Savi* had the highest rib collagen δ13C and δ15N values of the whole data set spanning central and eastern Slovenia, and northern Croatia, dating from the Bronze Age to the Late Iron Age. The combination of these findings and the palaeopathological status of this infant singled out this individual out as an interesting case study.

In the past, δ13C and δ15N values obtained from the tissues of infants have been linked to breastfeeding practices alone. Here we can see a pattern that could suggest breastfeeding, but is overlaid by higher δ13C and δ15N values than would be predicted by the rib samples from the adults. This case study offers several alternative scenarios and supports the argument that isotopic data are complex and should not be interpreted in isolation. Through the application of novel, high resolution sampling methods, in combination with osteological techniques, it has been possible to establish a more nuanced understanding of the complex life history of an infant buried in the Early Iron Age cemetery at *Zagorje ob Savi*, Slovenia.

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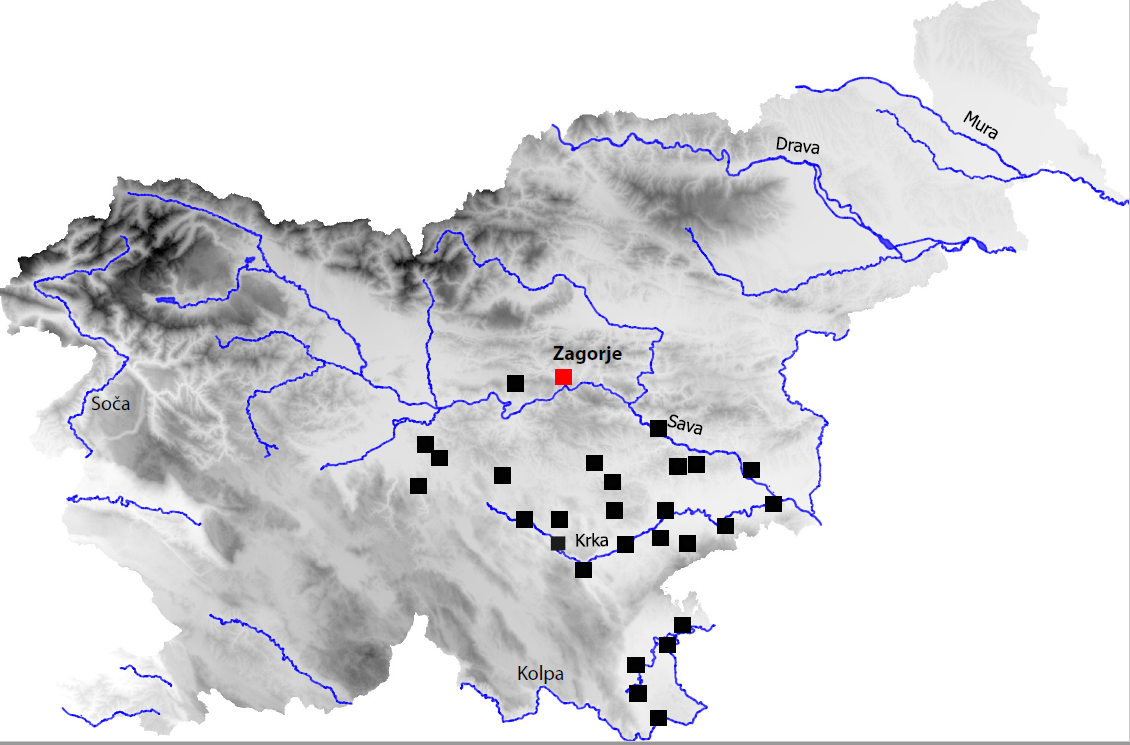
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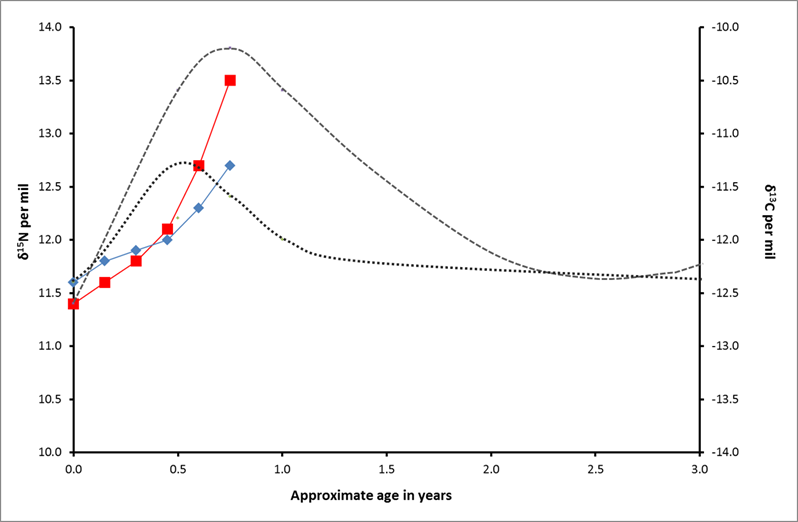
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Supplementary 1 : Map showing major Early Iron Age centres of the Dolenjska region of Slovenia (Zagorje ob Savi marked in red). Map: N Dolinar, Archive IPCHS.



Supplimentary 2 : A schematic model of change in C and N isotope ratios in collagen over time, related to breastfeeding and weaning adapted from King et al (2018) and Millard (2000). The dashed line represents an abrupt increase in δ15N values related to the onset of breastfeeding. The dotted line represents the smaller change in δ13C values, unless the diet of the mother changes to incorporate/reduce marine or C4 based foods. The incremental dentine data obtained from the Zagorje ob Savi infant presented in figure 5 is also included for comparison (δ15N values in red and δ13C values in blue).