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Petrographic and rare earth elemental characteristics of Cambrian Girvanella oncoids exposed in the North China Platform: Constraints on forming mechanism, REE sources, and paleoenvironments

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Abstract

Oncoid-bearing Mioalingian strata in the North China Platform represent late highstand/forced regressive systems tract(s). In the current study, sequence stratigraphy, sedimentology, and microscopic studies were carried out to interpret the forming mechanism and paleoenvironment of these oncoids. Microscopic studies show the typical distribution of filamentous cyanobacteria (Girvanella) in the Cambrian oncoids. Based on these observations, the oncoids can be classified into three kinds, i.e., Girvanella-core oncoids, Girvanella-cortex oncoids, and Girvanella-full oncoids. Furthermore, the studied oncoids hold pyrite crystals and dolomite minerals that also indicate the influence of sulfate-reducing bacteria on their formation. The presence of Girvanella, pyrite, and dolomite in the cores and cortices of these distinctive oncoids depicts its microbial origin that flourished under relatively high-energy settings. The present work offers a reference example for the involvement of microbes, particularly Girvanella, in the development of microbial carbonates in the North China carbonate platform during the Cambrian period. Geochemistry of the Cambrian oncoids is interpreted in order to determine the involvement of detrital fractions, sources of rare earth elements (REEs), and paleoenvironmental settings. The results of slightly negative Eu anomalies, higher La/Yb (6.8–15.1), and the larger Y/Ho variations (30.17–42.08) in the studied oncoids advocate that these oncoids have been subjected to terrigenous input during their formation. Considering the absence of Ce anomaly, Gd/Gd^* ratios (< 1), and lesser values of Y/Ho (30.17–33.93), we propose that these oncoids have been influenced by diagenetic activities. Relatively greater values of Er/Nd (0.09–0.23) suggest seawater signature preserved by the marine sediments. Meanwhile, observed weak negative Eu anomalies within the oncoids point out the retention of the original marine water characteristics, whereas lower values of the Y/Ho propose freshwater participation from the rivers in the seawater during precipitation of oncoids. Based on the ratios Er/Nd , $(Nd/Yb)_{N}$, $La/$ Yb, and depletion and variation of REEs, it can be confirmed that the sources of these REEs within the Cambrian oncoids are mainly from terrigenous input. However, the higher Er/Nd ratios are sourced from marine carbonates. Relatively lower values of Mo/U (1.10–2.46), V/Cr (< 2), V/(V + Ni) ratios (0.24–0.33), and deficiency of Ce anomaly in studied oncoids suggest their development under oxic environments. In addition, the variation of Sr/Cu (42–54) and Sr/Ba (35–79) indicates their growth under arid climatic conditions and classic marine settings. Considering these geochemical signatures, it can be concluded that the oncoids of the North China Platform flourished under oxic, arid, and classic marine conditions.

Keywords $Girvanella \cdot Microbial \, mat \cdot Trace$ and rare earth elements \cdot Cambrian (Miaolingian) oncoids \cdot China

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Introduction

Oncoids $(> 2$ mm in diameter) are generally considered as rounded forms that comprise of nuclei of varying grain types and cortices composed of entirely or a combination of calcified microbes (Riding [1991](#page-13-0); Hanken et al. [2015](#page-12-0); Mei et al. [2019a](#page-12-0), [b](#page-12-0); Xiao et al. [2020a](#page-13-0), [b](#page-13-0)). The oncoid usually exhibits dual nature: (1) oncoids possess the mobile attribute of coated grain that has an irregular and uneven surface and are generally considered microbial in origin (Liu and Zhang [2012](#page-12-0); Xiao et al. [2020a,](#page-13-0) [b\)](#page-13-0), which is distinctive from ooids whose origin is debatable (Duguid et al. [2010](#page-12-0); Diaz and Eberli [2019\)](#page-12-0), and (2) oncoids also represent biosedimentary structures, such as stromatolite, which has successive laminations around the core (Riding [2011a](#page-13-0), [b\)](#page-13-0).

Oncoids are the major component of carbonates that flourished in various environments in the entire chronostratigraphic column such as (1) lacustrine and shallow water environments in the early Cambrian (Schaefer et al. [2001](#page-13-0)), (2) fluviatile to lacustrine environments in the middle Cambrian to Jurassic (Olivier et al. [2004](#page-12-0); Brigaud et al. [2009](#page-11-0); Zhang et al. [2014a](#page-13-0), [b;](#page-13-0) Xiao et al. [2020a,](#page-13-0) [b](#page-13-0)), and (3) lacustrine, riverine, and tidal flat environments of the shallow sea in the recent (Jones [1992](#page-12-0); Zhang et al. [2013](#page-13-0)). According to various depositional environments, the origin of oncoids can be interpreted as (1) sophisticated metabolic mechanisms within microbial mats, (2) microbially mediated calcium carbonate precipitation, and (3) formation of particles via trapping and binding in a marine environment (Dahanayake [1977;](#page-11-0) Peryt 1981; Flügel and Munnecke [2010;](#page-12-0) Jones and Renaut [2010;](#page-12-0) Jones [2011;](#page-12-0) Zhang et al. [2014a](#page-13-0), [b;](#page-13-0) Han et al. [2015](#page-12-0); Zhang et al. [2015a](#page-14-0), [b;](#page-14-0) Qi et al. [2016](#page-13-0); Mei et al. [2019a,](#page-12-0) [b\)](#page-12-0). The evolutionary study of ancient oncoids is significant to interpret fluctuations in depositional conditions as well as control of microbial activities on the forming mechanism of oncoids. Hence, the studies of oncoids from the ancient strata are important clues to restore paleoclimate, paleoenvironment, sea-level fluctuations, and sedimentary environment (Vedrine et al. [2007;](#page-13-0) Zhou et al. [2017](#page-14-0); Xiao et al. [2020a,](#page-13-0) [b\)](#page-13-0).

Several researchers studied the Cambrian oncoids of the North and South China to highlight its various aspects such as classification, forming environment, textural composition, growth mechanism and controlling factors (Yang et al. [2011\)](#page-13-0), mass-occurrence (anoxic) event of oncoids (Zhang et al. [2014b](#page-13-0), [2015a\)](#page-14-0), the ultra-fabrics and biomineralization of oncoids (Zhang et al. [2014a\)](#page-13-0), forming mechanism of oncoids (Han et al. [2015](#page-12-0)), coupling variation of oncoids (Zhang et al. [2015b](#page-14-0)), formation of oncoids via microbial activities (Wang and Xiao [2018;](#page-13-0) Mei et al. [2019a,](#page-12-0) [b](#page-12-0)); and the various shape of oncoids and their paleoenvironmental conditions (Xiao et al. [2020a,](#page-13-0) [b\)](#page-13-0). However, a study on the involvement of microbes particularly Girvanella in different parts of Cambrian oncoids is quite insufficient. Moreover, geochemistry of the Girvanella oncoids regarding the involvement of detrital

fractions, sources of rare earth elements (REEs) in these oncoids, and paleoenvironmental settings are still unresolved. The present study describes the Girvanella oncoids, its different forms, and the influence of cyanobacteria in the formation of oncoids in the Miaolingian strata of the North China Platform (NCP). In addition, current research highlights that these oncoids have been subjected to terrigenous input and diagenetic activities during their formation.

Geological background

The tectonic history of China delineates the development of main tectonic blocks including Tarim, North China, and South China (Zheng et al. [2013;](#page-14-0) Myrow et al. [2015\)](#page-12-0) (Fig. [1a, b](#page-2-0)). The studied NCP was developed in the tropical to subtropical regions of the Sino-Korean Block, close to Australia, in the early Paleozoic time (Wotte et al. [2007\)](#page-13-0) (Fig. [1a\)](#page-2-0). It is restricted by main suture zones on the north and south by Hinggan Fold Belt and Qinling Dabieshan Belt, respectively (Myrow et al. [2015](#page-12-0)) (Fig. [1b\)](#page-2-0), whereas it is bounded by east and west with Tanlu Fault and Helan aulacogen, accordingly (Zheng et al. [2013;](#page-14-0) Myrow et al. [2015\)](#page-12-0) (Fig. [1b\)](#page-2-0). Deposition of the Cambrian sediments in the NCP initiated approximately in late Series 2 with a sea-level rise that deposited Cambrian strata on it as shown in Fig. [1c](#page-2-0) (see Mei [1996](#page-12-0), [2011](#page-12-0); Riaz et al. [2019a,](#page-13-0) [b](#page-13-0)). The Cambrian strata overlap with the Precambrian strata in this widespread carbonate platform in a similar pattern as described in North America (e.g., Peters and Gaines [2012](#page-13-0)). Consequently, Cambrian succession with the thickness of 700 m was developed on the NCP, which can approximately be divided into three successions in ascending order (Meng et al. [1997;](#page-12-0) Mei [1996,](#page-12-0) [2011](#page-12-0); Riaz et al. [2019a,](#page-13-0) [b\)](#page-13-0) (Fig. [1c\)](#page-2-0): (1) a mixed rock unit of the Series 2, which is composed of red beds (i.e., dolostone) and carbonate rocks (i.e., limestone), (2) carbonate strata of the Miaolingian predominated by oolitic grainstones along with the minor amount of oncolites at some places, and (3) carbonate succession of the Furongian that mainly comprises carbonate muds. These rock units delineate characteristic cyclicity that helps in further classifying them into nine third-order sequences (from DS_1 to DS_9) as shown in Fig. [1c.](#page-2-0) The sequence boundaries of the Cambrian strata can be marked by punctuated surfaces and drowning unconformities (Fig. [1c\)](#page-2-0).

According to the new chronostratigraphic division (Peng and Zhao [2018](#page-13-0); Mei et al. [2020a](#page-12-0)), the studied Miaolingian strata consist of the Maozhuang, Xuzhuang, Zhangxia, and Gushan formations that deposited in four third-order depositional sequences (DS_3 to DS_6 in Fig. [1c\)](#page-2-0). The Maozhuang Formation and the lower part of the Xuzhuang Formation comprise a succession of a tidal-flat depositional system marked by carbonate rocks with the intergrowth of red beds (Fig. [1c\)](#page-2-0). Thick-bedded massive oolitic grainstones along with the minor amount of oncolites are commonly

Fig. 1 a Global location of the North China Platform (NCP). b Map showing the geological framework of the NCP (modified after Riaz et al. [2019a](#page-13-0), [b](#page-13-0)). Gray regions show the exposed Cambrian strata. Inset

satellite image (lower right corner) showing the NCP in a regional perspective. c The sedimentary succession of the Cambrian strata (from $DS₁$ to $DS₉$) in the NCP (modified after Mei et al. $2020a$)

concentrated in the upper parts of the Xuzhuang, Zhangxia, and Gushan formations $(DS_4$ to DS_6 in Fig. 1c), which constitute the time-specific facies (Brett et al. [2012](#page-11-0); Schinder [2012\)](#page-13-0). These deposits develop in both highstand and forced regressive systems that fulfill the forced regressive model proposed in several previous studies (i.e., Helland-Hansen and Gjelberg [1994;](#page-12-0) Schlager and Warrlich [2009](#page-13-0); Mei [2010](#page-12-0); Samanta et al. [2016](#page-13-0)). During each depositional period of FRSTs of the Miaolingian strata $(DS_4$ to DS_6), a broad oolitic shoal developed in the NCP that represents a carbonate platform dominated by ooidal sands (Pratt et al. [2012\)](#page-13-0).

Materials and methods

Extensive geological fieldwork has been carried out in the NCP. Macroscopic features associated with oncoids are examined in the upper part of the Cambrian (Miaolingian) Xuzhuang and Zhangxia formations (Fig. [2a](#page-3-0)–h). The oncoids of the Xuzhuang Formation are irregular and rounded in shape without clear separation of cores and cortices (Fig. [2b, d\)](#page-3-0), whereas the Zhangxia Formation comprises of of elongated and elliptical to spheroidal shape oncoids with definite cores and successively repeated laminations in their cortices similar to stromatolites (Fig. [2f, h](#page-3-0)). These formations have conformable upper and lower contacts with Cambrian Miaolingian strata; however, at places, the Xuzhuang Formation has unconformable lower contact with Precambrian strata (Fig. [2a,](#page-3-0) [g](#page-3-0)). A petrographic analysis of 360 carbonates samples from the targeted bed was carried out. Detailed investigation of microfacies was conducted in order to understand the microscopic composition and fabric of the formation, the structure of oncoids, its various types, and the presence of microbes particularly cyanobacteria as well as grains other than oncoids. Thin sections were observed under a high-magnification petrographic microscope at the China University of Geosciences, Beijing (CUGB). Photomicrographs for the high-resolution examination of oncoids were captured with a camera attached

Fig. 2 Important sequence boundaries, sedimentary features of the Miaolingian strata and locations of oncoids in the various region of the North China Platform (NCP). a The Xuzhuang Formation having massive oolitic limestone along with oncolite, conformably overlain by the Zhangxia Formation. b Irregular oncolite in the Xuzhunag Formation. c Drowning unconformity separates the Miaolingian strata. d Rounded oncolite in the Xuzhunag Formation. e The contact between Zhangxia and Gushan formations. f Elongated oncolite in the Zhangxia Formation. g Division of entire Cambrian Miaolingian strata. h Elliptical to spheroidal oncolite with dominant core in the Zhangxia Formation

to the petrographic microscope. CorelDraw graphic software was used to construct different figures and collage of different photomicrographs used in the current study.

Additionally, geochemical data, i.e., inductively coupled plasma–mass spectrometry (ICP–MS), was adopted from Xiao et al. ([2020a\)](#page-13-0) to determine the exact sources of REEs and paleoenvironmental conditions of Girvanella oncoids. The results of the geochemical data are presented in Supplementary Table 1.

Petrography of oncoids

Oncoids are mostly elliptical to spheroidal in shape under the microscope. The shape of oncoids is controlled by the shape of core and concentric laminae. These oncoids were classified into three varieties: Girvanella-core oncoids, Girvanella-cortex oncoids, and Girvanella-full oncoids based on texture and outline of the cortex. The detail descriptions of these oncoids are given below.

Girvanella-core oncoids

Description These oncoids are typically > 1.5 mm in diameter and surrounded by sparite cement (Fig. $3a-c$ $3a-c$). These oncoids have a well-developed core of Girvanella and cortex of dolomite (Fig. $3a-c$ $3a-c$). These oncoids (Fig. [3a, b](#page-4-0)) are similar to superficial ooids that have a thick core and thin cortex. The cortices of the oncoids are associated with crystals of dolomite that range in size from 100 to 150 μ m (Fig. [3a](#page-4-0)–c). Moreover,

pyrite grains are observed within and the surrounding areas of the oncoids. Further observation shows the minor amount of bioclasts (brachiopods and/or trilobites) and patches of micrite (possibly Girvanella) in the surrounding region of the oncoids (Fig. 3a).

Interpretation Sparite cement and presence of brachiopods and/or trilobites indicate the high-energy setting of these oncoids (Flügel [2004\)](#page-12-0). The superficial shape of oncoids indicates its migration from low- to high-energy conditions (Riaz et al. [2019b\)](#page-13-0). The cortices of these oncoids are thick indicating more life span in a high-energy environment (Riaz [2019](#page-13-0)). Furthermore, the cortex is associated with crystals of dolomite that possibly formed by capillary condensation due to fluctuation of relative sea-level (Guo et al. [2020](#page-12-0)). The dolomite is genetically associated with extracellular polymeric substance (EPS) degradation that forms cyanobacteria-dominated microbial mat (Decho and Gutierrez [2017](#page-12-0)). In the process, the elimination of EPSs occurred through both active and passive mineralization that produces Mg^{2+} ions, which is captured by in vivo EPSs. Further pyrite crystals in the surrounding area of the oncoids represent the metabolize products of sulfatereducing bacteria in cyanobacterial microbial mat (Baumgartner et al. [2006](#page-11-0)).

Girvanella-cortex oncoids

Description These oncoids ranges in size from 1 to 1.5 mm in diameter, with uneven, irregular, and nonsmooth cortices (Fig. [4a](#page-5-0)–c). The cortices of the oncoids associated with preserved fossils of Girvanella with either core of bioclasts (trilobite and/or brachiopods) (Fig. $4a-a'$ $4a-a'$) or microbial envelop/ bacterial biofilm (Fig. [4b](#page-5-0)–c). Further, observation shows the various shapes of oncoids that linked with the typical structure of cortex such as ellipsoidal in shape with rough and irregular laminations (Fig. $4a-a'$ $4a-a'$), irregular shape with unevenly

distribution of *Girvanella* in the cortex (Fig. [4b](#page-5-0)–b'), and rounded shape with an abundance of Girvanella in the outermost part of the cortex (Fig. $4c-c'$ $4c-c'$). In addition, pyrite crystals are also observed in all these oncoids (Fig. [4a](#page-5-0)′, [b](#page-5-0)′, [c](#page-5-0)′). These oncoids are surrounded by sparite cement (Fig. [4a](#page-5-0)–c) that is relatively different from cementing material that observed in Girvanella-core oncoids (Fig. 3a–c).

Interpretation The Girvanella in the cortex of oncoid depicts the characteristics of laminated stromatolite (i.e., Riding [2011a,](#page-13-0) [b](#page-13-0)). The oncoids with the core of bacterial biofilm (cyanobacteria and sulfate-reducing bacteria) show the similar features of the high-energy modern Bahamian coated grains that nucleus formed by the combination of cyanobacteria, sulfate-reducing bacteria, and diatoms (Brehm et al. [2003,](#page-11-0) [2006\)](#page-11-0). These microorganisms excrete EPS in the microbial mat to form multiple biofilms (Decho [2010](#page-12-0)). Further EPS provides a substrate for precipitation of calcite in the Cambrian calcite sea (Stanley [2006;](#page-13-0) Dupraz et al. [2009](#page-12-0); Mei et al. [2019c](#page-12-0), [2020a,](#page-12-0) [b](#page-12-0)). The oncoids with cores of bioclasts also depict the relatively high-energy setting of these deposits (Hanken et al. [2015;](#page-12-0) Riaz [2019](#page-13-0)). Moreover, the cementing material of these oncoids indicates the relatively low-energy setting as compared with the Girvanella-core oncoids that flourished in relatively high-energy setting above the normal wave base (Wang and Xiao [2018](#page-13-0); Riaz [2019;](#page-13-0) Xiao et al. [2020a,](#page-13-0) [b\)](#page-13-0).

Girvanella-full oncoids

Description The petrographic observation shows the same composition from center to outermost cortices of the oncoids that mostly range in size from 1.5 to 2 mm in diameter (Fig. [5a](#page-6-0)–c). These oncoids are associated in variable shape, i.e., elliptical (Fig. $5a-a'$ $5a-a'$), sub-angular (Fig. $5b-b'$ $5b-b'$), and asymmetric (Fig. [5c](#page-6-0)–c′) that show the abundance of Girvanella in their

Fig. 3 Photomicrographs depict the Cambrian Miaolingian Girvanella-core oncoids in the NCP. a Oncoids showing rounded shapes that are associated with Girvanella in the core and crystals of dolomite in the cortex. The yellow arrows show the pyrite crystals. Red and green arrows indicate the bioclasts and patches of micrite, respectively. b Elliptical to the spheroidal shape of oncoids having an elliptical arrangement of Girvanella in the core. c Rounded shape of oncoid is due to the rounded shape of Girvanella core

Fig. 4 Photomicrographs showing Girvanella-cortex oncoids surrounded by sparite cement. a An ellipsoidal oncoid with a core of trilobite/ brachiopods/other bioclasts and a clotted cortex. a′ Under crosspolarized light depicts the pyrite grains. b Irregular oncoid with a clotted and roughly laminated texture, comprised Girvanella fossils. b′ Under cross-polarized light showing the irregularshaped oncoids and pyrite grains. c Rounded oncoid grain, having roughly laminated cortices, with laminae thick upward and thin downward. c′ Under crosspolarized light portray pyrite crystals.

cores and cortices (Fig. [5a](#page-6-0)–c). Further observations reveal the pyrite crystals in these oncoids (Fig. [5a](#page-6-0)′, b′). These oncoids are surrounded by sparite cement (Fig. $5a-c$ $5a-c$) similar to the cement observed in Girvanella-cortex oncoids (Fig. 4a–c).

Interpretation These oncoids are associated with the cortex of variable shape indicate the high-energy turbulence condition (i.e., Chow and James [1987;](#page-11-0) Heller et al. 1990). Further core and cortex predominantly comprise Girvanella fossils that indicate the involvement of filamentous cyanobacteria in the formation of Cambrian oncoids (i.e., Elliott [1975;](#page-12-0) Rees et al. [1989](#page-13-0); Riding [1991;](#page-13-0) Mei et al. [2019a](#page-12-0), [b\)](#page-12-0). Furthermore, pyrite crystals indicate a close connection with the sulfate reduction and may show the complex microbial precipitation inside the microbial mat during the formation of the oncoids (e.g., Dupraz et al. [2009](#page-12-0); Riding [2011b](#page-13-0)). The cementing material of these oncoids is similar to Girvanella-cortex oncoids that indicate a relatively low-energy setting for their formation (Riaz [2019\)](#page-13-0).

Geochemistry of oncoids

Trace elements

Table [1](#page-7-0) lists the contents of the trace elements of the studied oncoids samples. Within the large ion lithophile elements (LILEs), Ba and Rb represent depletion (Table [1\)](#page-7-0). In comparison, Rb is extremely depleted in studied oncoids of the NCP. Among LILEs, the Sr concentration within the oncoids is relatively higher (Table [1\)](#page-7-0). In the case of Pb concentration, most of the oncoid samples reveal slight enrichment (Table [1\)](#page-7-0). Among the ferromagnesian trace elements, Ni and Co indicate concentrations of 7.2–9.3 ppm and 0.74–1.91 ppm while Cr and V show contents of 1.2–2.4 ppm and 2.7–3.7 ppm respectively (Table [1\)](#page-7-0). Within all of the high field strength elements (HFSEs) (Nb, Hf, Zr, Y, Th, and U), Nb, as well as Hf, is extremely depleted (Table [1](#page-7-0)).

Fig. 5 Photomicrographs show Girvanella-full oncoids. a–c Girvanella growth in all stages of oncoids development. a Elliptical shape of oncoid without a nucleus is surrounded by sparite cement. a ′ Under cross-polarized light showing the pyrite crystals. b Sub-angular shape of oncoid associated with the same composition from center to outermost cortex. b′ Under crosspolarized light filamentous Girvanella along with pyrite minerals. c Asymmetric shape of oncoid without a nucleus. c′ Under cross-polarized light same feature observed and close view of Girvanella also shown

The studied oncoids indicate lower values of Mo/U ratios (1.10–2.46; average 1.76; Table [1](#page-7-0)). Most of the V/Cr ratios of oncoids are < 2 , and $V/(V + Ni)$ ratios vary from 0.24 to 0.33. Moreover, the variation of Sr/Cu between 42 and 54 and Sr/Ba (35–79) is presented in Table [1.](#page-7-0)

REEs

Table [1](#page-7-0) presents the rare earth element contents (REEs) of the studied oncoids. On NASC-normalized REE pattern, these oncoids are observed with slightly weak negative Eu anomalies and indicates the weak enrichment of LREEs (Fig. [6](#page-8-0)). The Eu/Eu* ratios reveal a range of $0.34-1.03$ $0.34-1.03$ $0.34-1.03$ (Table 1). Meanwhile, the variation of Ce/Ce* ratios in Cambrian oncoids ranges from 0.48 to 1.41 along with the absence of Ce anomaly (Table [1;](#page-7-0) Fig. [6](#page-8-0)). The oncoids are represented by higher Er/Nd and La/Yb ratios while most of the Gd/Gd*

ratios of the Cambrian oncoids are < 1. The Y/Ho ratios of the Cambrian studied oncoids indicate a range of 30.17 and 42.08 (Table [1\)](#page-7-0). The REE contents within the studied oncoids indicate a narrow range of 5.71 to 7.69 ppm with an average of 6.69 ppm (Table [1\)](#page-7-0).

Discussion

Forming mechanism

Girvanella oncoids deposited in the Miaolingian strata of the NCP. These oncoids indicate high-energy depositional environment formed during FRST (forced regressive systems tract) of the third-order levels that represents a typical example of drowning unconformity (Schlager [1989](#page-13-0), [1998](#page-13-0), [1999](#page-13-0)). These oncoids are depicting dual characteristics, i.e.,

WH= Wuhai Section-Inner Mongolia Province

DQ= Diaoquan Section-Shanxi Province

XWD= Xiaweidian Section-Beijing

SDG= Sandaogou Section-Liaoning Province

XZ= Xuzhuang Formation

ZX= Zhangxia Formation

laminated stromatolites and mobile coated grains that represent a spectacular sedimentary phenomenon (Bosak et al. [2013;](#page-11-0) Pederson et al. [2015](#page-13-0); Peters et al. [2017](#page-13-0)). The abundantly preserved Girvanella in these oncoids shows its involvement at different growth stages that significantly alter the microstructure and consequently, different types of Girvanella

Fig. 6 NASC-normalized REE patterns of oncoids indicating slightly weak negative Eu anomalies and weak enrichment of LREEs

oncoids are developed (Fig. 7). In Girvanella-core oncoids, Girvanella along with few heterotrophic organisms make a dark micrite in the center of the oncoids (Figs. [3a](#page-4-0)–c and 7). These microorganisms particularly cyanobacteria excrete gellike material called extracellular polymeric substance (EPS) that provides the substrate for endorsing precipitation of $CaCO₃$ (Dupraz et al. [2009](#page-12-0); Bosak [2011](#page-11-0); Decho and Gutierrez [2017](#page-12-0)). In the *Girvanella*-cortex oncoids, Girvanella filaments were involved in the development of the cortices. Consequent calcification caused the development of Girvanella-rich laminae, which served as textural elements of these oncoids. Similarly, Girvanella-full oncoids experienced all stages of Girvanella growth as compared with the first two forms of Girvanella oncoids (Fig. 7). The presence of Girvanella in all stages of oncoids demonstrates credible confirmation regarding the contribution of Girvanella in the development of these carbonate grains. Furthermore, petrographic observations clarify that Girvanella-core oncoids flourished under high-energy conditions above the normal wave base, whereas Girvanella-cortex and Girvanella-full oncoids deposited in both high and relatively low-energy settings above and below the normal wave base.

The cores and cortices of these oncoids are composed of Girvanella and sulfate-reducing bacteria (Figs. [3,](#page-4-0) [4,](#page-5-0) [5,](#page-6-0) and 7) that are a possible demonstration of the microbial origin of the Cambrian oncoids of the NCP. These Girvanella fossils are equivalent to either the modern Plectonema (Riding [1991](#page-13-0)) or the modern Tychonema (De los Ríos et al. [2015](#page-12-0)). The evenly outlined walls of the Girvanella fossil can be interpreted as a

residue of *in vivo* calcification that represents an ecophysiological mechanism, which is genetically related to the atmospheric carbon dioxide concentration mechanisms (CCMs; Riding [2000;](#page-13-0) Kah and Riding [2007\)](#page-12-0). Moreover, the in vivo calcification may coordinate with calcification of EPSs containing extracellular sunscreen scytonemin (Soule et al. [2009](#page-13-0); Mei et al. [2020a](#page-12-0)), which is different from the calcification of common EPSs forming microbial mats. The interior parts of filaments are filled by microspars that may be the residues of cyanobacterial cells. Further, the filamentous fossils in the cores (Fig. [3\)](#page-4-0) and cortices of oncoids (Figs. [4](#page-5-0) and [5\)](#page-6-0) strongly reflect the sophisticated calcification of photosynthetic biofilms that occurred after the in vivo calcification of cyanobacterial sheaths (Riding [2000](#page-13-0); Reitner [2011a](#page-13-0), [b](#page-13-0)). These credible pieces of evidence provide certain clues of the microbial origin of Cambrian oncoids of the NCP. However, it is difficult to reach certain conclusions whether Girvanella fossils are directly involved in the formation of oncoids or not; nevertheless, these fossils provide certain evidences for the microbial origin of oncoids.

Furthermore, the presence of pyrite crystal in the Cambrian oncoids indicates microbial precipitation which is related to sulfate reduction (Riding [2000;](#page-13-0) Baumgartner et al. [2006](#page-11-0)). Moreover, pyrite crystals describe two important factors, i.e., alkalinity engine (Kempe and Kazmierczak [1994](#page-12-0); Gallagher et al. [2012\)](#page-12-0) and EPS (Tourney and Ngwenya [2014](#page-13-0); Decho and Gutierrez [2017\)](#page-12-0), which actively participate in carbonate precipitation in the cortex and core of these oncoids. In other words, cyanobacteria (Girvanella) abundantly and successively excrete a mess of EPSs to form biofilm in the microbial mat, and its degradation via heterotrophic organisms like sulfate-reducing bacteria caused the precipitation. The exact mechanism of precipitation in microbial mats dominated by Girvanella is still an important subject that needs to be investigated in the future.

Geochemical characteristics and source of REEs in Cambrian oncoids

Rare earth elements (REEs) in carbonate bodies are mainly inherited from ancient seawater. But, in several cases, the concentrations of REEs in ancient carbonates such as oncoids were also affected by detrital involvement as well as diagenesis. Thus, REEs of oncoids can be significant to reveal the participation of terrigenous input in addition to diagenetic influence. The rare earth elements (REEs) ratios such as Eu/Eu* can be valuable to infer the involvement of terrigenous material during the carbonates formation. The Eu/Eu* ratios of the Cambrian oncoid samples indicate a range of 0.34–1.03 (Table [1\)](#page-7-0). In general, the slightly positive Eu anomalies in REE patterns which are rare in seawater possibly caused by the diagenetic variations in the carbonates (Abiding and Calagari [2015](#page-11-0)) and enhanced through processes of hydrothermal activities (Madhavaraju and Lee [2009](#page-12-0)). On the contrary, the slightly negative anomalies of Eu are indicative of the terrigenous input in carbonate samples (Dai et al. [2016\)](#page-11-0). The slightly negative Eu anomalies in the studied oncoids suggest that these oncoids have been subjected to terrigenous participation during their formation (Fig. [6](#page-8-0)). Therefore, based on these findings, we propose that the slightly negative Eu anomalies in oncoids are indicative of terrigenous involvement.

Several research works have been conducted on the variations of Ce in the marine settings for defining paleoenvironments (e.g., Nath et al. [1997\)](#page-12-0). Also, it is documented that the scarcity of Ce in oceanic water is due to redox variations of Ce (Elderfield [1988](#page-12-0)). The variation of Ce/Ce* ratios in Cambrian oncoids ranges between 0.48 and 1.41 (Table [1\)](#page-7-0). Both Ce contents and the lack of Ce anomaly in studied oncoids can be governed by lithological input, and diagenesis may impart an important character in the incorporation of REEs (particularly Ce). According to Armstrong-Altrin et al. [\(2001\)](#page-11-0), the diagenesis can indicate a worthy character in removing Ce anomaly in the samples of oncoids. Considering these views, it is suggested that the absence of Ce anomaly in Cambrian oncoids can be attributed to diagenetic activities. Further, the impact of diagenetic conditions is most likely defined by Er/Nd ratios (Abiding and Calagari [2015\)](#page-11-0). Their higher ratios in carbonates show the seawater signature conserved by the sediments. It is widely reported that the ratio (Er/Nd) is nearly 0.27 in normal seawater (De Baar et al. [1988\)](#page-12-0). Both diagenetic processes and terrigenous materials can be responsible for discretional amounts of Nd comparative to Er and may decrease the Er/Nd values to < 0.1 (Abiding and Calagari [2015](#page-11-0)). However, these ratios in the oncoids vary between 0.09 and 0.23. Fairly higher values of Er/Nd (Table [1\)](#page-7-0) point out that the impact of terrigenous materials and diagenetic activities was insignificant. Relatively greater values of Er/Nd in oncoids of the NCP indicate seawater signature conserved by the marine sediments.

The Gd/Gd* ratios of the carbonates of modern shallow seawater range from 1.05 to 1.30, as indicated by De Baar et al. ([1985](#page-12-0)). However, most of the Gd/Gd* ratios of the Cambrian oncoids are < 1 (Table [1\)](#page-7-0), suggesting the influence of diagenetic processes on the oncoids of the NCP. The super chondritic Y/Ho ratios of the studied oncoids are less than the values of seawater $($ \sim 44–74) as defined by Bau ([1996](#page-11-0)). Relatively lesser values of oncoids in comparison to sea water suggest the impact of diagenetic activities on the carbonates of the studied area. Meanwhile, the higher La/Yb values (Table [1\)](#page-7-0) of oncoids (6.8–15.1) suggest the involvement of limited detrital material during the genesis of Cambrian oncoids (Sholkovitz [1990;](#page-13-0) Condie [1991\)](#page-11-0). Considering these views, it can be concluded that the oncoids of the NCP have been influenced by diagenetic activities and limited terrigenous input during their formation.

The REE patterns in carbonates are frequently significant to typify source rocks (Tobia [2018](#page-13-0)). NASC-normalized REEs values of the studied oncoids are mostly depleted and this points out the contribution of detrital material within the Cambrian oncoids (Fig. [6](#page-8-0); Sen and Mishra [2015](#page-13-0)). The studied samples reflect a narrow range of discrepancy in ΣREE contents (5.71 to 7.69, average 6.69; Table [1\)](#page-7-0). These narrow ranges in contents of ΣREE are also a consequence of variation in detrital materials within the Cambrian oncoids (Chen et al. [2014](#page-11-0)).

The $(Nd/Yb)_{N}$ ratios of the oncoids differ from 0.64 to 2.29 (Table [1\)](#page-7-0) and are not comparable to the seawater-like REE patterns. Therefore, oncoids with higher $(Nd/Yb)_N$ ratios confirm that the source of these REE ratios is from terrigenous input within the Cambrian oncoids. Meanwhile, it is also widely accepted that higher ratios of Er/Nd indicate that they are sourced and preserved by marine carbonates (De Baar et al. [1988\)](#page-12-0). On the contrary, the ratios of Er/Nd having values less than 0.1 are suggestive of terrigenous input and diagenetic activities within the oncoids (Tobia [2018](#page-13-0)). Therefore, the ratios Er/Nd, $(Nd/Yb)_{N}$, La/Yb, and depletion and variation of REEs point out the occurrence of a detrital fraction which is the most credible source of REEs within the oncoids. The Er/ Nd ratios of the oncoids are within the range of 0.09 to 0.23 (Table [1](#page-7-0)), indicating that these higher rare earth elemental ratios are sourced and preserved by marine carbonates (De Baar et al. [1988\)](#page-12-0).

Implications on paleoenvironments of Cambrian oncoids

Several researches have reported that the positive Eu anomalies are owing to hydrothermal activities (Armstrong-Altrin et al. [2003\)](#page-11-0) or because of diagenetic alteration (Tobia [2018](#page-13-0)). The studied oncoids are flourished in shallow carbonate platform (Xiao et al. [2020a\)](#page-13-0); thus, local enhancement of feldspar as well as diagenetic variations can produce positive Eu anomaly. However, on the REE pattern, the observed oncoids of the NCP show weak negative Eu anomalies (Fig. [6\)](#page-8-0). These observed negative anomalies point out the preservation of the original marine water signatures within the oncoids.

Y/Ho ratio is reflected as one of the consistent tool for the marine environment (Xiao et al. [2020a\)](#page-13-0). In modern marine settings, this ratio differs between 40 and 90 (Bau [1999](#page-11-0)). The Y/Ho ratios of the Cambrian studied oncoids indicate a range of 30.17 and 42.08 (Table [1](#page-7-0)) with an average value of 34.53. This implies that the oncoids have thrived in a marine setting; however owing to the detrital input or because of the diagenesis, the average Y/Ho ratio of the oncoids is declined (Tobia [2018\)](#page-13-0). Further, this ratio is analogous to other Mesoproterozoic carbonates such as Rohtas carbonates (Banerjee and Jeevankumar [2007\)](#page-11-0). The observed larger

variations in the Y/Ho ratios of the oncoids exhibit that the studied oncoids are contaminated by the impact of detrital materials. The lower values of the Y/Ho ratio such as 30.17 to 33.93 in the studied oncoids also propose freshwater participation from the rivers in the seawater during precipitation of these Cambrian oncoids.

Modern studies report that the geochemical ratios such as Mo/U are significant pointers of redox conditions of carbonates (McManus et al. [2006\)](#page-12-0). Therefore, these conditions of oncoids can be rebuilt by trace elemental ratios present in the studied oncoids. Algeo and Tribovillard ([2009](#page-11-0)) confirmed that Mo/U ratios higher than 7.9 are considered as a pointer of anoxic/sulfidic conditions and can provide significant information by studying sediments from modern marine settings. However, the studied oncoids indicate lower values of Mo/U ratios (1.10–2.46; average 1.76; Table [1\)](#page-7-0). Lower values of Mo/U ratios of studied oncoids suggest that oncoids thrived in oxic environments.

Several researches reported the application of Ce anomaly in the marine environments for highlighting the paleo conditions (Hua et al. 2013; Khelen et al. [2017\)](#page-12-0). To decide the redox condition, Ce anomalies are consistent and more reliable since Ce valences and solubility differ in redox condition. Under the oxidized environment, a negative or lack of Ce anomaly promotes (Hua et al. 2013). However, under anoxic condition, positive Ce anomaly results. In this study, the oncoids are displaying a deficiency of Ce anomaly (Fig. [6\)](#page-8-0), which suggests deposition of oncoids under the oxic environment.

In addition, $V/(V + Ni)$ and V/Cr are important to define the paleoenvironmental conditions of carbonates (Xiao et al. [2020a\)](#page-13-0). Low V/Cr ratio (< 2) is suggestive of oxic conditions, and $V/(V + Ni) < 0.6$ also represents the oxidizing condition (Rimmer [2004;](#page-13-0) Xiao et al. [2020a](#page-13-0)). Similarly, most of the V/Cr ratios of oncoids are < 2 , and $V/(V + Ni)$ ratios vary from 0.24 to 0.33 (Table [1\)](#page-7-0), suggesting the flourishment of Cambrian oncoids under oxic conditions.

The Sr/Cu and Sr/Ba ratios are measured as significant proxies to delineate paleoclimatic conditions of carbonates (Fouke et al. [2005](#page-12-0)). It is widely reported that the range of Sr/Cu (1.3 to 5.0) shows humid climate, whereas higher than 5.0 displays arid climate (Ni et al. [2010\)](#page-12-0). The oncoids of this study represent a variation of Sr/Cu between 42 and 54 (Table [1\)](#page-7-0), signifying arid climatic environments. Moreover, if the Sr/Ba value is < 1 , it is representative of terrestrial condition; nonetheless, greater than 1 value of Sr/Ba is suggestive of marine condition (Tripati et al. [2009\)](#page-13-0). The studied oncoids reveal the variation of Sr/Ba (35–79), signifying classic marine conditions.

Considering the above discussion regarding paleoenvironmental analyses of Cambrian oncoids, it can be concluded that the oncoids of the North China Platform (NCP) flourished under oxic, arid, and classic marine conditions.

Conclusions

- 1) Girvanella oncoids of the North China Platform (NCP) were distinguished into Girvanella-core, Girvanella-cortex, and Girvanella-full oncoids based on the morphology of the cortices. Girvanella-core oncoids indicate both high and relatively low-energy depositional settings based on the cementing material and the thickness of the cortex, while Girvanella-cortex oncoids and Girvanellafull oncoids are formed under relatively low-energy settings. These oncoids are associated with EPSs, which form a number of biofilms in a relatively thick microbial mat dominated by cyanobacteria, where they grow and roll over it. Moreover, abundantly preserved filamentous fossils, i.e., Girvanella in the cortex and nuclei of oncoids, strongly suggest the independent growth of these oncoids over microbial mats. The presence of pyrite grains and dolomite crystals in these oncoids also clarifies the calcification and decomposition of EPSs in cyanobacteria-dominated microbial mats.
- 2) The slightly negative Eu anomalies, higher La/Yb values, and the observed larger Y/Ho variations in the Cambrian oncoids of NCP suggest that these oncoids have been subjected to terrigenous participation during their formation. Considering the absence of Ce anomaly, Gd/Gd* ratios $(< 1$) and lesser values of Y/Ho, it can be concluded that the oncoids of the NCP have been influenced by diagenetic activities. Relatively greater values of Er/Nd suggest seawater signature preserved by the marine sediments. In addition, observed weak negative Eu anomalies within the oncoids point out the retention of the original marine water characteristics while lower values of the Y/Ho propose freshwater participation from the rivers in the seawater during precipitation of oncoids.
- 3) Based on the ratios Er/Nd, $(Nd/Yb)_N$, La/Yb, and depletion and variation of REEs, it can be confirmed that the source of these REEs within the oncoids is mainly from terrigenous input, which is the most credible source of REEs. However, the higher Er/Nd ratios are sourced and preserved by marine carbonates.
- 4) Lower values of Mo/U, V/Cr, and V/(V + Ni) ratios and deficiency of Ce anomaly in the studied oncoids suggest that these oncoids were flourished under oxic environments. These oncoids represent a variation of Sr/Cu between 42 and 54 and Sr/Ba (35–79), suggesting their development under arid climatic conditions and classic marine settings.

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Compliance with ethical standards

Conflict of interest The authors declare that there is no conflict of interest.

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