Correlation between the base of Drumian Stage and the base of middle Caesaraugustan Stage in the Iberian Chains (NE Spain)

Rodolfo Gozalo, Juan Bautista Chirivella Martorell, Jorge Esteve & Eladio Liñán



Inter-regional Cambrian correlations are difficult amongst different palaeogeographic provinces, because of a lack of shared genera and species between different palaeogeographic provinces. The Drumian Stage shows this difficulty because it was defined by the FAD of the agnostoid *Ptychagnostus atavus*, which species has not yet been found in the Mediterranean subprovince. Thus it is necessary to establish a correlation between the ISCS scale and the local scale using other trilobites. New data from the Iberian Chains and a review of previous works suggest that the bases of *Pardailhania hispida* and *Ptychagnostus atavus* zones are approximately equivalent. Therefore *Pardailhania hispida* FAD could be used to mark the beginning of the Drumian Stage in the Mediterranean region. This level is easily identifiable in several places: Montagne Noire (France), Cantabrian Mountains, Demanda Mountains and Iberian Chains (Spain), Sardinia (Italy), Amanos and Sultan Mountains (Turkey), and, probably, Antiatlas (Morocco). • Key words: trilobites, biostratigraphy, Cambrian Stage 5 – Drumian boundary, middle Cambrian / Cambrian Series 3, Mediterranean region.

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During recent years the International Subcommission on Cambrian Stratigraphy (ISCS) has been engaged in defining new series and stages for the Cambrian System (see Peng et al. 2009). One of the most important problems facing inter-regional correlation in the Cambrian is the lack of genera and species shared between different palaeogeographic provinces. For example, many of the zonal trilobite species for Cambrian Series 3 are representatives of the family Ptychagnostidae, and the Drumian Stage was defined by Babcock et al. (2007) at the first occurrence of Ptychagnostus atavus, whereas localities in the Mediterranean subprovince (sensu Sdzuy et al. 1999; Fig. 1) do not yield any ptychagnostids. Nevertheless, a correlation between the official chronostratigraphy ISCS scale and the Mediterranean scale is needed. Therefore this correlation has to be carried out using polymeroid trilobites. Our aim is to recognize the base of Drumian Stage in the Iberian Chains (NE Spain).

Hitherto, the base of *Ptychagnostus atavus* Zone has been correlated with an undetermined level in the upper part of the lower Caesaraugustan (Sdzuy 1971, 1972; the Iberian Peninsula was developed originally by Sdzuy (1968). However each zone in Caesaraugustan was used with different criteria (*e.g.* Acme zones, assemblage zones, range zones or phylozones, see Álvaro & Vizaïno 1998). These disagreements as to criteria make any international correlation difficult. We have worked on the revision of the Caesaraugustan zones in various sections of the Iberian Chains (northeast Spain). The new detailed samplings allow us to propose interval phylozones throughout the Caesaraugustan Stage. The sections studied are close to the villages of Murero, Jarque, Villafeliche, and Purujosa, which are located about 100 km west and south-west of Zaragoza, in the Aragonese branch of the Iberian Chains (Fig. 2). From a geological point of view, these localities belong to the Badules Unit (see Gozalo & Liñán 1988).

Shergold & Geyer 2003). Caesaraugustan biozonation of

We have studied the trilobites of the transition from lower to middle Caesaraugustan succession from the basal part of the Murero Formation in five sections: Rambla de Valdemiedes 1 and 2 (RV1 and RV2: Liñán & Gozalo 1986; Gozalo *et al.* 1996, 2003a; Chirivella Martorell



2008; Liñán *et al.* 2008), Jarque 1 (J1: Liñán *et al.* 1996, 2008, Chirivella Martorell 2008), Villafeliche 1 (Vi1: Gozalo *et al.* 1993, 1996; Chirivella Martorell 2008) and Purujosa 6 (Pur6, unpublished data).

Biostratigraphic framework

The classical Spanish middle Cambrian biozonation was established by Sdzuy (1968, 1971, 1972) and has been used with minor modifications until now (*i.e.* Liñán & Gozalo 1986; Sdzuy *et al.* 1996; Álvaro & Vizcaïno 1998; Liñán *et al.* 2002, 2004; Gozalo *et al.* 2003b, 2008; Esteve *et al.* 2008; Fig. 3). From a historical point of view the Mediter-

ranean zones that are involved in the base of Drumian Stage seem to correspond with the *Badulesia granieri* and *Pardailhania hispida* zones (*sensu* Sdzuy 1968, 1971, 1972). According to Liñán *et al.* (1993) the boundary between these zones also marks the lower-middle Caesaraugustan boundary.

Sdzuy (1968, 1971, 1972) put the base of *Pardailhania hispida* level at the FAD of *Pardailhania hispida hispida*; however the older specimens of *Pardailhania hispida* n. spp. were recorded together with *Badulesia granieri* and *Badulesia paschi* in the previous level. Therefore the FAD of *Pardailhania hispida* lies lower than the former use of the *P. hispida* Zone that was used as acme zone by Sdzuy (1968, 1971, 1972) and considered as an interval zone by

Sdz	Sdzuy 1968		Sdzuy 1971, 1972			Liñán & Gozalo 1986			Sdzuy <i>et al.</i> 1996, 1999; Liñán <i>et al.</i> 2002, 2004; Gozalo <i>et al.</i> 2003b			Álvaro & Vizcaïno 1998, Esteve <i>et al.</i> 2008, Gozalo <i>et al.</i> 2008		New proposal		
HANIA (pars)	Pardailhania hispanica		ars)	nia (pars)	2	ars)	nia (pars)	hispanica		Parda hisp	ilhania anica		Parda multi:	uilhania spinosa		Pardailhania multispinosa
PARDAILH	Parda his	ilhania pida	DPSIDAE (p	Pardailha	1	DPSIDAE (p	Pardailha	hispida	STAN (pars)	Parda his	ilhania pida	STAN (pars)	Parda his	iilhania pida	STAN (pars)	Pardailhania hispida
oars)	ılesia nieri	B.p.	ERUC		4	ERUC		g. + p.	AUGU	ılesia nieri	B. p.	AUGU	ılesia nieri	B.p.	AUGU	Dadulasia
ESIA (J	Badı gra	Badule julivei	ENOPI	sia (pars	3	ENOPI	sia (pars	granie) julive	ESAR	Badi gra	Badule julivei	ESAR	Badı gra	Badule julivei	ESAR	granieri
BADULI	Badulesia tenera		SOL	Badule	2	SOL	Badule	tenera	CA	Badı ten	ilesia vera	CA	Badi ten	ilesia vera	CA	Badulesia tenera

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Figure 3. Sketch showing the main biostratigraphic subdivisions of the level studied in the Middle Cambrian of the Spain and France. *B.p.: Badulesia paschi, g. + p.: granieri + paschi*

Liñán & Gozalo (1986), Álvaro & Vizcaïno (1998) and others. Nevertheless they did not change the position of the lower boundary of the P. hispida Zone. The problem has been that all the researchers until now accepted the facto Sdzuy's idea about the existence of two subspecies of Pardailhania hispida; although Pardailhania hispida n. spp. never was defined and/or figured (Sdzuy 1971, table 1; 1972, table 2). Currently we accept that Pardailhania hispida has a wide and continuous morphological variation within the features of cranidia (Chirivella Martorell 2008, Esteve et al. 2008) and it is not possible to recognise two subspecies, as Sdzuy had proposed. Thus, the lower boundary of the Pardailhania hispida interval Zone should be placed at the P. hispida FAD, slightly below Sdzuy's original definition for the P. hispida Zone (Figs 3, 4). This modification of the systematic interpretation of P. hispida makes it necessary to redefine the Badulesia granieri and Pardailhania hispida zones.

Badulesia granieri interval Zone

Synonymy: "Nivel de *B. granieri*" (part.), sublevel "*B. juliverti*" (Sdzuy 1968). "Nivel de *B. juliverti* y *B. granieri*" (Sdzuy 1971). "Biozona *Badulesia granieri* + *Badulesia juliverti*" (Liñán & Gozalo 1986). "Superbiozone à *B. granieri*" (part.) and "phylozone à *B. juliverti*" (Álvaro & Vizcaïno 1998). See Fig. 3.

Our proposal is to consider this zone as an interval phylozone, the base of which coincides with the *Badulesia* granieri FAD and the top is the level immediately below the *Pardailhania hispida* FAD.

The trilobites recorded in the Iberian Chains in this time interval (see Chirivella Martorell 2008; Fig. 5) are: *Pero*nopsis acadica, Condylopyge sp., Eccaparadoxides asturianus, E. sulcatus, E. sequeirosi, E. acadicus (Fig. 6K, L), Badulesia granieri, B. tenera, Parabailiella schmidti, P. languedocensis, Bailiella cf. tenuicincta and Ctenocephalus (Hartella) cf. terranovicus.

So far *Badulesia granieri* has been recorded in France (Montagne Noire: Courtessole 1973, Álvaro & Vizcaïno 1998); in Spain (Sdzuy 1961, 1968, 1969; Liñán & Gozalo 1986; Gámez *et al.* 1991; Gozalo *et al.* 1993; Liñán *et al.* 1995) and Morocco (Geyer *et al.* 1995, Geyer & Landing 2006). Furthermore, it may be possible to recognise equivalent levels of this zone in Turkey and Italy, where the index taxa for the previous and/or subsequent zones have been recognized.

Pardailhania hispida interval Zone

Synonymy: "Nivel de *B. granieri*" (part), sublevel "*B. paschi*" and "Nivel de *P. hispida*" (Sdzuy 1968). "Nivel de *B. paschi* y *B. granieri*" and "Nivel de *P. hispida*" (Sdzuy 1971). "Biozona de *Badulesia granieri* + *Badulesia juliverti*" and biozona "*Pardailhania hispida*" (Liñán & Gozalo 1986). "Superbiozone *B. granieri*", "phylozone à *B. paschi*" and "phylozone à *P. hispida*" (Álvaro & Vizcaïno 1998). See Fig. 3.

We consider the *Pardailhania hispida* Zone as an interval phylozone, the base of which lies at the FAD of *Pardailhania hispida* and the top immediately below the FAD of *Pardailhania multispinosa* (see Esteve *et al.* 2008).

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Figure 4. Middle Cambrian chronostratigraphic units for the Mediterranean Subprovince (modified from Liñán *et al.* 2002) and the correlation with the Cambrian Stages and Series proposed by ISCS, with a subdivision into sequences and systems tracts (from Gámez *et al.* 2009). Abbreviations: HST – highstand system tract; LST – lowstand system tract; TST – transgressive system tract.

Our proposal to define the *P. hispida* Zone as an interval phylozone implies that the levels with Badulesia paschi (former B. granieri + B. paschi Subzone) that occur in the top of *B. granieri* Zone (lower Caesaraugustan) currently are the base of P. hispida Zone (middle Caesaraugustan). The P. hispida Zone seems to bear the highest diversity and relative abundance of trilobites in the Iberian Chains (see Chirivella Martorell 2008; Fig. 5). At least eighteen species have been collected in the lower part: Peronopsis acadica, P. segmenta (Figs 6A-C, 7), Peronopsella westergardi, Diplagnostus planicauda (Figs 6G-I, 8), Condylopyge rex (Fig. 6D-F), Pardailhania hispida (Fig. 6M), Badulesia granieri, B. paschi, Eccaparadoxides asturianus, E. sequeirosi, E. acadicus (Fig. 6K, L), E. rouvillei (Fig. 6J), Parabailiella schmidti, P. languedocensis, Ctenocephalus (Hartella) antiquus, Agraulos longicephalus, A. arenosus and Skreiaspis miqueli. The diversity of agnostids, five species, may indicate a transgressive pulse.

So far *Pardailhania hispida* has been recorded in France (Montagne Noire: Thoral 1935, Courtessole 1973, Álvaro & Vizcaïno 1997), in Spain (Sdzuy 1961, 1968; Palacios 1982; Liñán & Gozalo 1986), in Italy (Sardinia: Rasetti 1972, Loi *et al.* 1995) and in Turkey (Shergold & Sdzuy 1984; Dean 2005, 2006). In addition, Geyer *et al.* (1995) and Geyer & Landing (2006) cited the presence of the genus *Pardailhania* in Morocco.

Correlation between the base of the Drumian Stage and the base of the middle Caesaraugustan

The classical correlation between Spain and Baltica (Sdzuy 1971, 1972) placed the base of Pardailhania within the Tomagnostus fissus and Ptychagnostus atavus Zone without further qualification, and slightly above of the Paradoxides hicksii Zone in Eastern Newfoundland. These correlations have been followed by later workers (e.g. Shergold & Geyer 2003, Liñán et al. 2002, Gozalo et al. 2008). Recently, Fletcher (2006, p. 68, 2007) and Elicki & Geyer (2010) moved this boundary below the previous proposal. They have considered that the correlation of the base of Ptychagnostus atavus Zone would be with a level just below the base of the Badulesia granieri (see Fletcher 2006, fig. 18) or with the base of Caesaraugustan (base of Badulesia tenera Zone) (see Elicki & Geyer 2010, fig. 1). On the other hand, the redefinition of the Pardailhania hispida Zone, the lower boundary of which lies below the old Pardailhania hispida Zone (sensu Sdzuy 1971, 1972). Therefore, it is likely that the correlation of the base of Pardailhania hispida would be roughly equivalent to the base of Tomagnostus fissus and Ptychagnostus atavus Zone. This roughly corresponds to the base of Drumian Stage.

The available data reinforce this correlation. Although many of trilobite species are endemic to the Mediterranean

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Spanish Stages	Zones	Agnostids	Polymeroids	I.S.C.S. Stages
middle CAESAR-	Pardailhania multispinosa		sturianus dti — Badulesia tenera — Badulesia tenera	DRUMIAN
AUGUSTAN (part.)	Pardailhania hispida		Eccaparadoxides a Eccaparadoxides a aus Parabalitella schmi Cenocephalus (Ha Cenocephalus (Ha and and and and and conoco Hydrocephal	
lower	Badulesia granieri	 Condylopyge sp. Peronopsella westergardi Condylopyge rex Diplagnostus planicanda Peronopsis segmenta 	oxides sdztyji oxides sdztyji sei as andutina a matutina a matutina a matutina a matutina a matutina a matutina a segnensis doxides sequeirosi loxides sequeirosi doxides sequeirosi a sequeirosi fordes sequeirosi fordes areadicut a streiaspis miqueli Agrataspis miqueli Agrataspis miqueli Agrataspis miqueli Agrataspis miqueli Agradovida Ctenocephalus (Harella b	STAGE 5
CAESAR- AUGUSTAN	Badulesia tenera	Peromopsis acadica	Eccapara - Hydrocephalus brau - Acadolenus sp. - Arasolenpis unpeuro on - Arasolenpis unpeuro on - Holocephalina? leve - Bailiaspis tuberci Badulesia gramieri Eccapara Eccapara	STACE 3

Figure 5. Stratigraphic distribution of trilobite species in the lower and middle Caesaraugustan of the Iberian Chains showing the proposed correlation with ISCS subdivision. Names of species comment in the correlation section are highlighted.

region (e.g. Badulesia granieri, Badulesia paschi, Pardailhania hispida, Eccaparadoxides sequeirosi, Parabailiella languedocensis, etc.), there are some species that have been recorded also in other palaeogeographic regions: Eccaparadoxides acadicus, Agraulos longicephalus, Diplagnostus planicauda, Peronopsis segmenta and Condylopyge rex. We discuss the biostratigraphical information of these species below.

a) *Eccaparadoxides acadicus* (Avalonian species) (Fig. 6K, L) whose stratigraphic range is in the *Eccaparadoxides eteminicus* and *Hartella* zones in New Brunswick and Newfoundland (Kim *et al.* 2002) and which was recorded from the top of the *Ptychagnostus gibbus* or base of the *Tomagnostus fissus* Zone in England (Rushton *et al.* 2007); thus its presence in the *Pardailhania hispida* Zone indicates that the youngest age for this zone would be the base of *Ptychagnostus atavus* Zone, which is the base of Drumian Stage.

b) Agraulos longicephalus has been recorded in Great Britain, eastern Newfoundland, Spain and France. The range in Wales is *Tomagnostus fissus* and *Hypagnostus parvifrons* zones (Thomas *et al.* 1984); eastern Newfoundland, *Paradoxides hicksii* Zone (Martin & Dean 1988; Fletcher 2006, 2007); Spain, *Pardailhania hispida* to *Solenopleuropsis ribeiroi* zones (Liñán & Gozalo 1986); France, levels A1, A2 and B? (Courtessole 1973), which is equivalent to middle Caesaraugustan and upper Caesaraugustan (Álvaro & Vizcaïno 1998). The biostratigraphical range is equivalent to *Ptychagnostus atavus* Zone in the sense of Peng & Robison (2000).

c) *Diplagnostus planicauda* (Figs 6G–I, 8) has a wide geographic and stratigraphic distribution. Its FAD is recorded in the base of *Ptychagnostus atavus* Zone (Peng & Robison 2000), thus its presence in the base of *Pardailhania hispida* Zone indicates that the oldest age for this zone would be Drumian.

d) *Peronopsis segmenta* (Figs 6A–C, 7) has been recorded only in the eastern Great Basin (Robison 1964, 1982, 1984) and the observed range is from middle *Ptychagnostus gibbus* to *Ptychagnostus punctuosus* Zone or upper part of the Cambrian Stage 5 and Drumian Stage.

e) *Condylopyge rex* (see Appendix 1; Fig. 6D–F) has been recorded beyond the Mediterranean region in Bohemia, Baltica, England and eastern Newfoundland where shows a wide stratigraphic range. The range in Bohemia is *Eccaparadoxides pusillus* to *Onymagnostus hybridus*



Figure 6. Trilobites of the *Pardailhania hispida* Zone. • A–C – *Peronopsis segmenta* Robison, 1964; A – cephalon, internal mould, MPZ 2007/350; B, C – pygidium, internal mould (B) and latex cast (C) of the same specimen, MPZ 2007/348. • D–F – *Condylopyge rex* (Barrande, 1846); D – pygidium, internal mould, MPZ 2007/456; E – cephalon, latex cast, MPZ 2007/447; F –cephalon, internal mould, MPZ 2007/456; E – cephalon, latex cast, MPZ 2007/422; H – latex cast, MPZ 2007/428; I – internal mould, MPZ 2007/423. • J – *Eccaparadoxides rouvillei* (Miquel, 1905), cranidium, latex cast MPZ 2011/78. • K, L – *Eccaparadoxides acadicus* (Matthew, 1883); K – cranidium, latex cast, MPZ 2007/501; L – thoracic segments and pygidium, latex cast, MPZ 2007/516. • M – cranidia of *Pardailhania hispida* (Thoral, 1935), latex cast, MPZ 3078 and 3079 (specimens figured by Liñán & Gozalo 1986, pl. 24, figs 9, 10). All the specimens are housed in Museo de Paleontología de la Universidad de Zaragoza (MPZ). Scale bars: figures A–I 1 mm, figures J–M 5 mm.

zones (Fatka *et al.* 2004); in Sweden and Norway, *Hypa*gnostus parvifrons Zone (Westergård 1946) or top of *Ptychagnostus atavus* Zone sensu Høyberget & Bruton (2008); in Warwickshire, England, it occurs in the *hicksii* Zone (Illing 1916; Fletcher 2007, fig. 2) in beds equivalent to the *Ptychagnostus gibbus* and *Tomagnostus fissus* zones (Rushton 1979, Thomas *et al.* 1984); in Spain, *Pardailhania hispida* to *Solenopleuropsis thorali* zones (Sdzuy 1961, Liñán & Gozalo 1986, Chirivella Martorell 2008); in France, level B (Courtessole 1973) equivalent to *Solenopleuropsis ribeiroi* Zone (Álvaro & Vizcaïno 1998); in Eastern Newfoundland, *Paradoxides hicksii* and *P. davidis* zones (Howell 1925; Hutchinson 1962; Fletcher 2006, 2007, fig. 3), and in Germany, upper part of Caesaraugustan (Sdzuy 2000). In summary the total range of this species would be from the upper part of the *Ptychagnostus gibbus* to the *Ptychagnostus atavus* zones or the upper part of the Cambrian Stage 5 and Drumian Stage.

We do not know currently whether the base of Drumian lies at a lower level. However, a comparison of the biostratigraphic ranges of these species seems to show that the best correlation of the basal level of *Pardailhania hispida* is likely with the basal part of the *Ptychagnostus atavus* Zone. Thus, the FAD of *Pardailhania hispida* in the Mediterranean subprovince can be used as the local index trilobite for the Drumian Stage in this region.

Other data

The biofacies analysis and the sequence stratigraphy established for the Iberian Chains (see Gámez Vintaned *et al.* 2009; Fig. 4) show that the basal beds of *Pardailhania hispida* Zone was deposited during the Transgressive System Tract of the Sequence 6. The relative abundance of agnostids and diversity of polymeroids trilobites could be indicative of a relative maximum transgressive for these levels. Likewise the basal levels of the *Ptychagnostus atavus* Zone in Drum Mountains [(USA Babcock *et al.* 2007)] are associated with a relative maximum transgressive event also supports our proposal of correlation between the bases of *Pardailhania hispida* and *Ptychagnostus atavus* zones.

Recently, Álvaro et al. (2008) on the basis of the carbon isotopic studies suggested a different correlation for the Mediterranean subprovince. They considered that a negative $\delta^{13}C_{org}$ shift from a background of -22% to a negative peak of -24.5% in the Campelou section Montagne Noire, France) belong to the Drumian Carbon Isotope Excursion (DICE), more or less equivalent to the base of Drumian (see Ahlberg et al. 2009, Howley & Jiang 2010). The DICE falls in the base of middle Languedocian (of the Mediterranean scale) in the Campelou section Álvaro et al. (2008). Because the Montagne Noire and Iberian Chains share trilobite faunas, this correlation suggested that all the species listed above in the Iberian Chains have an age of Cambrian Stage 5. However the middle Languedocian in the Campelou section does not share any trilobites with areas where the DICE excursion has been detected. This disagreement between the biostratigraphic data and isotopic data could be due to various issues. Firstly, this disagreement is likely the result of low resolution, 27 samples in a section of 200 m thick with different intervals between samples and meter scale close to chemostratigraphic anomalies. Despite of fine sampling close to the chemostratigraphic anomalies, many negative anomalies



Figure 7. Pygidium of *Peronopsis segmenta* Robison, 1964. • A – internal mould, B – external mould of the same specimen, also figured in Fig. 6B, C, MPZ 2007/348. Scale bar 1 mm.



Figure 8. Pygidium of *Diplagnostus planicauda* (Angelin, 1851). • A – internal mould, B – external mould of the same specimen, also figured in Fig. 6G, MPZ 2007/422. Scale bar 1 mm.

are detected in other researches close to SPICE (Montañez *et al.* 2000; Zhu *et al.* 2004, 2006). Therefore a lack of a biostratigraphical control with trilobites in the Campelou section prevents us from knowing whether this negative anomaly belongs to DICE or to a negative excursion closer to the SPICE peek. For the moment, the biostratigraphic data of Montagne Noire and Iberian Chains and their trilobitic correlation with other regions show that the negative isotope excursion considered as DICE by Álvaro *et al.* (2008) in the Montagne Noire really correspond with a high stratigraphic level within Drumian, not below.

Conclusion

The new data and the previous work on biostratigraphy and sequence stratigraphy demonstrate that the bases of the *Pardailhania hispida* and *Ptychagnostus atavus* zones are roughly equivalent. The FAD of *Pardailhania hispida* could be used in the Mediterranean region as a marker for the beginning of the Drumian Stage. This level is easily identifiable in several places: the Montagne Noire (France), Cantabrian Mountains, Demanda Mountains and Iberian Chains (Spain), Sardinia (Italy), Amanos and Sultan Mountains (Turkey), and, probably, the Antiatlas (Morocco).

Systematic appendix

Condylopyge rex (Barrande, 1846) Figure 6D–F

- v 1986 Condylopyge aff. regia (Sjögren, 1872). Liñán & Gozalo, pp. 39, 40, pl. 1, fig. 9.
- v 1986 *Condylopyge rex rex* (Barrande, 1846). Liñán & Gozalo, p. 40 (with previous synonym).
- v 1986 *Condylopyge rex hispanica* n. ssp. Liñán & Gozalo, pp. 40, 41, pl. 1, figs 10–20 (with previous synonym).
- 2002 *Condylopyge rex* (Barrande, 1846). Valíček & Szabad, pp. 75, 76, pl. 1, fig. 1 (with previous synonym).
- 2004 *Condylopyge rex* (Barrande, 1846). Buchholz, p. 516, pl. 3, figs 10, 11.
- 2005 *Condylopyge rex* (Barrande). Cotton & Fortey, fig. 1A.
- 2006 *Condylopyge rex* (Barrande, 1846). Valíček, pl. 1, figs 7–9, fig. 2.
- v 2008 *Condylopyge rex* (Barrande, 1846). Chirivella Martorell, pp. 69–72, pl. 3, fig. 12, pl. 4, figs 1–11 (with previous synonym).
 - 2008 *Condylopyge rex* (Barrande, 1846). Høyberget & Bruton, pp. 71, 72, pl. 12, figs K, M (with previous synonym).
 - 2009 *Condylopyge rex.* Ahlberg, Axheimer, Babcock, Eriksson, Schmitz & Terfelt, fig. 3G.
 - 2009 Condylopyge rex. Fatka & Mergl, fig. 9e.

Comments. – Liñán & Gozalo (1986) subdivided the species *C. rex* in two subspecies. They defined *C. rex hispanica* from Spain and France and suggested the possibility of another Scandinavian subspecies. Their main criteria were geographic and the different morphological combinations of the pygidium and cephalon borders. The Bohemian specimens have both convex borders (see Šnajdr 1958). The Mediterranean specimens have the plane cephalic border and convex pygidial border (Sdzuy 1961, Liñán & Gozalo 1986), and the Scandinavian forms have a plane cephalic border and convex pygidial border (Westergård 1946). Also the age of the three subspecies seemed different.

The material classified as *C. rex hispanica* by Liñán & Gozalo (1986) have been recorded of the upper Caesaraugustan levels. They found a similar cephalon from the base of *Pardailhania hispida* Zone that classified as *Condylopyge* aff. *regia* (Sjögren, 1872) because the cephalon morphology is close to this species, but the lack of pygium prevent to classify in open nomenclature. The new samplings on the level where had been found *Condylopyge* aff. *regia* by Liñán & Gozalo (1986) allow us to find several cephala and pygidia of *Condylopyge*. The pygidium morphology is *C. rex* type, thus we assign this material to this species.

New specimens of *C. rex* have been published from Bohemian region (Valíček & Szabad 2002, Valíček 2006) that show an important morphological variability in the cephalic and pygidial borders and increase the stratigraphic range for this species (see Fatka *et al.* 2004). Now the age range of Spanish specimens is similar to the Bohemian and Scandinavian specimens, and the morphological variability does not permit to recognise the different subspecies proposed by Liñán & Gozalo (1986).

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