1	A revised northern European Turonian (Upper Cretaceous)
2	dinoflagellate cyst biostratigraphy: integrating palynology and carbon
3	isotope events
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28 ABSTRACT

29 Organic walled dinoflagellate cyst (dinocyst) assemblage data are presented for a new 30 Turonian regional reference core (Bch-1) drilled at Běchary in the Bohemian Cretaceous 31 Basin, east-central Czech Republic. The detailed stratigraphic framework for the section is 32 summarised based on calcareous nannofossil and macrofossil biostratigraphy, regional e-log 33 correlation, sequence stratigraphy and carbon isotope chemostratigraphy. Dinocyst results 34 obtained for 196 samples from the 405 m long core offer the highest resolution (~22 kyr) 35 stratigraphically well-constrained data set available to date for the Turonian Stage, 93.9 – 89.8 36 Ma. A dinocyst biostratigraphic framework is presented based on the evolutionary first and 37 last occurrence, first common occurrence, and acmes of key species. Published dinocyst data 38 from English Turonian Chalk successions in East Sussex, Berkshire, Kent and Norfolk are 39 reviewed within a stratigraphic framework provided by macrofossil records and carbon 40 isotope event (CIE) chemostratigraphy. Critical analysis of existing published Turonian 41 dinocyst zonation schemes shows them to be untenable. Correlation of the English Chalk 42 data to Bch-1 provides a basis for defining a regional dinocyst event stratigraphy with 22 43 datum levels, and a revised dinocyst zonation scheme constrained within a chemostratigraphic 44 framework of 10 major CIEs. The new zones consist of a Cenomanian Litosphaeridium 45 siphoniphorum Zone, followed by the Cauveridinium membraniphorum Zone spanning the 46 uppermost Cenomanian to Lower Coniacian. This is subdivided into: Senoniasphaera 47 turonica (Lower - mid-Middle Turonian); and Raetiaedinium truncigerum (mid-Middle 48 Turonian – mid-Lower Coniacian) subzones. The *Oligosphaeridium pulcherrimum* Zone 49 (Senonisphaera rotundata Subzone) characterises the Lower Coniacian. The new 50 stratigraphy offers a basis for improved correlation and dating of Upper Cretaceous 51 successions.

52 **1. Introduction**

53 Over the last 40 years, a substantial amount of work has been undertaken on the 54 biostratigraphy of Turonian organic walled dinoflagellate cysts (dinocysts) from the English 55 and French Chalk and its correlatives in the Anglo-Paris Basin (Clarke and Verdier, 1967; 56 Foucher, 1974, 1975, 1976a, 1980, 1981, 1983; Tocher, 1984; Tocher and Jarvis, 1987, 1994, 57 1995; Jarvis et al., 1987, 1988a, b; FitzPatrick, 1992, 1995, 1996; Pearce, 2000, 2010; Pearce 58 et al., 2003, 2009; Prince et al., 2008; Lignum, 2009). However, to our knowledge, little has 59 been published previously on the Turonian – Coniacian dinocyst biostratigraphy of Central Europe, notable exceptions being the work of Prössl (1990), Kirsch (1991) and Svobodová et 60 61 al. (1998, 2002).

62 Here, new dinocyst data are presented from an expanded (> 300 m thick) Turonian hemipelagic succession in a fully cored research borehole drilled recently in the Bohemian 63 64 Cretaceous Basin at Běchary, east-central Czech Republic. Dinocyst records from four well-65 characterised English Chalk successions in East Sussex, Berkshire, Kent and Norfolk are reviewed and integrated using macrofossil biostratigraphy, lithostratigraphy and carbon 66 67 isotope chemostratigraphy (Jarvis et al., 2006; Pearce et al., 2009) to provide a regional 68 framework for comparison with results from the Czech Republic. Despite marked latitudinal 69 provinciality that occurred in the Late Cretaceous (see Lentin and Williams, 1980; Costa and 70 Davey, 1992), many of our marker taxa also occur to the north of the European Chalk 71 province, well into the siliciclastic-prone facies of the Shetland Group. It is a particular 72 strength of dinocysts to have a largely facies-independent preservation potential, 73 exemplifying their use as a powerful biostratigraphic tool. In this paper, a revised dinocyst zonation for the northern European Turonian tied to the carbon isotope event (CIE) 74

stratigraphy of Jarvis et al. (2006) is proposed, based on integrating the English Chalk data
with new results from Bch-1.

77 **2. Geological framework**

78 The Bohemian Cretaceous Basin was an intra-continental basin formed during the 79 Cenomanian – Santonian (100.5 - 83.6 Ma) as a seaway between the Boreal Sea and Alpine 80 Ocean (Fig. 1). The basin originated by the reactivation of a fault system in the Variscan 81 basement of the Bohemian Massif, and combined features of an epeiric sea formed during global transgression with those of a tectonically active setting that contains probably the 82 83 highest proportion of siliciclastics of all the European Cretaceous basins north of the Alps (Uličný et al., 1997, 2009). During Turonian – Coniacian times, sedimentation in the 84 85 Bohemian Basin was dominated by the repeated progradation of coarse-grained deltas and 86 adjoining shorefaces, and was affected by redistribution of siliciclastics by strong along-shore 87 tidal currents (Uličný, 2001). A maximum water depth of around 100 m is estimated for the 88 basin interior (Mitchell et al., 2010).

89 During 2010, a 405 m research core, Bch-1, was drilled through a representative Lower 90 Coniacian to Upper Cenomanian succession of offshore marine sediments in the Bohemian 91 Cretaceous Basin, to investigate the responses of multiple proxies to sea-level change (Uličný et al., 2014). The Bch-1 site (50.31506°N 15.29497°E), located in the village of Běchary, 92 93 east-central Czech Republic, is situated in the central basin between two depocentres (Fig. 1), 94 one adjacent to the Most-Teplice High and Western Sudetic Island in the northwest, the other 95 bordering the Bohemian Massif in the southeast. These source areas contributed varying 96 amounts of sediment through the Turonian, but with the Western Sudetic Island being by far 97 the most prominent source area.

98 The dominant lithofacies in the Bch-1 core consist of very dark grey marlstones and 99 calcareous mudstones with a varying proportion of quartz silt (Fig. 2). The mean percentage 100 of CaCO₃ through the core is ~ 35 %, and carbonate is generally represented by a micritic 101 component, some mm-scale bioclasts, and calcite spar in horizons with concretionary cement. 102 Total organic carbon contents (TOC) average 0.42%. Turonian lithofacies show abundant 103 bioturbation throughout the core, dominated by a distal *Cruziana* ichnofacies (sensu 104 MacEachern et al., 2010).

105 **3. Stratigraphic framework**

106 Deposition in the Bohemian Cretaceous Basin has been split into a number of genetic 107 sequences, termed TUR1 – TUR7, CON1 and CON2 (Fig. 2), which were detailed by Uličný 108 et al. (2009). The sequences are based on the analysis of a basin-scale correlation grid, 109 developed using well-log data (gamma-ray, resistivity, neutron porosity logs) from >700 110 boreholes, in most cases supplemented by outcrop and core data. These genetic sequences, 111 bounded by maximum transgressive surfaces, form a hierarchy of nested, composite 112 sequences, interpreted to record a response to relative sea-level changes. The sequences 113 record major cycles of regression and subsequent transgression, within which smaller scale 114 cycles also occur. Correlation of the sequence boundaries to Bch-1, as presented here, has been modified from Uličný et al. (2014) in the light of new stratigraphic data. 115

The macrofossil and calcareous nannofossil biostratigraphy, along with carbon isotope data for the Bch-1 core, were published by Uličný et al. (2014). Key biostratigraphic datum levels are plotted here against a lithological log and the carbon isotope profile for the core in Fig. 2.

119 *3.1. Litho- and biostratigraphy*

120 Sediments from the bottom of the Bch-1 core (404.6 - 402.4 m) yield a Cenomanian 121 nannofossil assemblage, including Lithraphidites acutus Verbeek & Manivit and Corollithion kennedyi Crux. The base of the Turonian (Sequence TUR1) is marked by a burrowed 122 123 omission surface at 402.35 m, overlain by a 70 cm thick zone of glauconite-rich marlstone 124 with phosphate concretions, interpreted as a hiatus, followed by condensed deposition (Valečka and Skoček, 1991; Uličný et al., 1993; Čech et al., 2005). A second, less prominent, 125 126 omission surface at 398.2 m, is overlain by a 20 cm thick, glauconite-rich, greenish-grey marlstone (Fig. 2). Calcareous nannofossil zones UC 5a – b (Burnett et al., 1998; correlative 127 128 to the uppermost Cenomanian upper Metoicoceras geslinianum and Neocardioceras juddii 129 ammonite zones) are absent, confirming the interpretation of Uličný et al. (1993) of a major 130 hiatus at this time in the central part of the basin. The first occurrence (FO) of Eprolithus 131 moratus (Stover) at 400.0 m depth indicates that at least nannofossil zones UC 5c - 6a are 132 contained in the lowermost 2.35 m of the Turonian succession in Bch-1. The prominent 133 earliest Turonian condensation event is attributed to the major flooding near the Cenomanian 134 - Turonian boundary (CTB) that established hemipelagic conditions over most of the basin (Klein et al., 1979; Uličný et al., 1997). 135

136 The first occurrence (FO) of the ammonite Collignoceras woollgari (Mantell), which 137 marks the base of the Middle Turonian, appears regionally together with Inoceramus cuvieri 138 (Sowerby) in the middle of Sequence TUR2. This datum level is placed at 374 m in Bch-1 139 (Fig. 2; Uličný et al., 2014), based on lithostratigraphic and e-log correlation to adjacent 140 sections. An acme of Inoceramus perplexus Whitfield occurs at the base of Sequence TUR5. 141 This level is distinctive due to a regional abundance of inoceramid prisms, identified at 246 m 142 in the core, and correlates to the "costellatus-plana Event" in NW Germany (Richardt and 143 Wilmsen, 2012). The base of the Upper Turonian, marked by the FO I. perplexus, is

correlated to the upper part of Sequence TUR4/3, at 252 m depth, immediately above the first
(rare) occurrence of the nannofossil *Marthasterites furcatus* (Deflandre) at 255 m. An
ammonite fauna typical of the *S. neptuni* Zone is recorded in the middle of TUR6, at 165 m.
This level is interpreted to represent the *Hyphantoceras* Event, a mid-Upper Turonian datum
level that can be recognised in England, parts of France, Germany, Poland and the Czech
Republic (e.g. Wiese et al., 2004).

150 The Upper Turonian – Lower Coniacian succession at Bch-1 is well constrained by 151 inoceramid bivalve and ammonite records from the core (Fig. 2), supplemented by e-log 152 correlation of FO datum levels from adjacent cores and outcrops. The uppermost Turonian 153 index taxon Mytiloides scupini (Heinz) is recorded at 134 m, while the position of its true FO 154 datum level is correlated to 140 m. Prionocyclus germari (Reuss) occurs at 121 m, with a 155 correlated FO at 130 m. The latest Turonian succession of three bivalve acme occurrences, Didymotis Events 0, I and II (Wood et al., 1984, 2004; Čech, 1989; Walaszczyk and Wood, 156 1998; Wiese, 1999; Walaszczyk et al., 2010) is placed between 115 – 95 m; Events I and II 157 158 are recorded directly in the core and the position of Event 0 is inferred by correlation. An 159 acme of Mytiloides herbichi (Atabekjan) occurs from 108 - 99 m depth, between Didymotis 160 Events I and II, as seen also on northern Germany (Wood et al., 2004).

The base of the Coniacian is placed at the correlated first occurrence of *Cremnoceramus deformis erectus* (Meek) at 94 m (cf. Walaszczyk et al., 2010), at the bottom of Sequence
CON1 and immediately above the top of *Didymotis* Event 2; *C. d. erectus* was recovered from
89 m in the core (Fig. 2). An acme of *M. furcatus* spans the Turonian – Coniacian boundary
interval.

The correlated FOs of *Cremnoceramus crassus inconstans* (Woods) at 45 m and *C. crassus crassus* (Petrascheck) at 37 m depth, and records of the latter species from the core at 33 m,

168 enable the placement of the standard European Lower Coniacian inoceramid zones

169 (Walaszczyk and Wood, 1998) in the core. The base of the Middle Coniacian is tentatively

170 placed at the FO Micula stauropora (nannofossil Zone UC10), at 16 m depth, a short distance

above the highest dinocyst sample at 17.5 m.

172 *3.2. Carbon isotope chemostratigraphy*

Turonian sediments display consistent secular variation in carbon isotope profiles 173 174 throughout northern and southern Europe (Jarvis et al., 2006; Voigt et al., 2007, 2008; 175 Richardt et al., 2013; Sprovieri et al., 2013), and remarkably similar trends have been documented from as far afield as Tibet and North America (e.g. Wendler, 2013; Joo and 176 Sageman, 2014). The Cenomanian – Coniacian carbonate carbon isotope ($\delta^{13}C_{carb}$) reference 177 curve for the English Chalk (Jarvis et al., 2006) and high-resolution $\delta^{13}C_{carb}$ curves from 178 179 Liencres, northern Spain (Wiese, 1999) and Saltzgitter-Salder (Voigt and Hilbrecht, 1997), Oerlinghausen and Halle (Voigt et al., 2007) in northern Germany, have been correlated to the 180 organic carbon isotope profile ($\delta^{13}C_{org}$) for Bch-1 by Uličný et al. (2014). A $\delta^{13}C_{org}$ profile 181 182 was favoured for Bch-1 due to concerns over possible diagenetic alteration in carbonate-183 carbon profiles generated from relatively low-carbonate mudrocks. Consistent relationships 184 were demonstrated between trends in the isotope profiles and the positions of key macrofossil 185 datum levels in the different sections (Uličný et al., 2014 fig. 3). A total of 20 peaks and 186 troughs in the profile from Bch-1 were correlated between the various sections, including the 187 major named CIEs of Jarvis et al. (2006).

The carbon isotope stratigraphy is consistent with an incomplete and attenuated Cenomanian – Turonian boundary succession at Bch-1, and the occurrence of an expanded Upper Turonian to Lower Coniacian section. The positions of the 9 most significant CIEs are shown for the Bch-1 section in Fig. 2, together with the primary $\delta^{13}C_{org}$ isotope data and a

- 192 smoothed chemostratigraphic profile. Placement of the CIEs follows Uličný et al. (2014),
- 193 with minor revision following the acquisition of new stratigraphic data.

194 **4.** Sampling and analysis

195 The 404.86 m Bch-1 core was drilled as a new Turonian regional reference section. Samples 196 of approximately 50 g were taken every 2 m for elemental, isotopic and palynological 197 analysis (196 samples). Based on an average compacted sedimentation rate for the Middle 198 and Upper Turonian of 9 cm/kyr (Uličný et al., 2014), sampling resolution was on the order of 199 22 kyr. Additionally, smaller samples (20 g, total 610) were obtained at 0.5 m (5.6 kyr resolution) intervals for carbon stable-isotope analysis of the organic fraction ($\delta^{13}C_{org}$). 200 Samples were cleaned, chipped to < 3 mm, and homogenised. Subsamples for isotopic and 201 202 elemental analysis were prepared and analysed following the methods described in Uličný et al. (2014). 203

204 It was necessary to process 10 g splits of chipped samples to yield representative 205 assemblages of palynomorphs for quantitative analysis. Palynomorphs >15 µm were 206 concentrated by a commercial processing company (PLS Ltd, Holvhead, UK) using the HF-207 HCl method of Lignum (2009) modified from Lignum et al. (2008, 'Company B' 208 methodology). All samples were spiked with tablets containing the modern spore 209 *Lycopodium* to allow statistically valid quantitative analysis of abundances (palynomorphs 210 per gram, ppg). Palynomorph identification and counting were undertaken using a light 211 microscope with a 400x objective. Three hundred dinocysts were identified per sample. 212 Broken or partial specimens were added to the count only if there was more than half of the 213 specimen present; unidentifiable specimens were recorded as 'indeterminate' and were not 214 included in the count of 300. Following this count, the remainder of the slide was scanned to

identify any additional species, which were marked as 'present', but in abundances too low tobe recorded among the 300.

217 5. Dinocyst biostratigraphy of the Bch-1 core

All samples yielded abundant and generally well-preserved dinoflagellate cysts, averaging
~4000 dinocysts per gram (~7000 dpg when corrected for carbonate-dilution, cf. Pearce et al.,
2009). The occurrence and ranges of 76 common dinocyst species identified in the Bch-1
core are plotted in Fig. 3, and the relative abundances of dominant species are shown in Fig.
4. A complete list of taxa (92 species) is provided in Appendix A.

223 Dinocyst biostratigraphic datum levels identified at Bch-1 include the sequential first

224 occurrences (FOs) of Cyclonephelium compactum – Cauveridinium membraniphorum in the

225 uppermost Cenomanian, of Senoniasphaera turonica in the Lower Turonian, and of

226 Oligosphaeridium poculum, Subtilisphaera pontis-mariae, Raetiaedinium truncigerum and

227 *Florentinia buspina* in the Middle Turonian (Figs. 2, 3; Plate I). The last occurrences (LOs)

228 of Stephodinium coronatum, Senoniasphaera turonica, C. compactum – C. membraniphorum

and *Kiokansium unituberculatum* occur in the Upper Turonian. The first common occurrence

230 (FCO) of Chatangiella ditissima and the FO of Surculosphaeridium belowii occur towards the

top of the stage. The FO of *Cribroperidinium wilsonii* in the lowermost Coniacian is

followed by the FCO of *Oligosphaeridium pulcherrimum*, above. With the exceptions of *K*.

233 *unituberculatum*, *S. pontis-mariae* and *C. ditissima*, most of these taxa occur sporadically

234 (Fig. 3) and in low numbers (<1% of assemblage), limiting stratigraphic resolution. The last

common occurrence (LCO) and reoccurrence (RO) datum levels of some species are also

considered to have stratigraphic potential (discussed further in Section 5, below).

237 A number of distinctive dinocysts range patterns are observed in the Bch-1 data (Figs. 3, 238 4). A generally inverse relationship between the relative abundance curves of 239 Palaeohystrichophora infusorioides and Spiniferites ramosus ramosus, with a marked 240 minimum of the former and an acme of the latter in Upper Turonian Sequences TUR5 - 6/1241 (Fig. 4), reflects the numerical dominance of these two taxa in the assemblages. A well-242 defined acme of Circulodinium distinctum distinctum occurs from 185 – 190 m in mid-Upper Turonian Sequence TUR6/1, with lesser floods of the species below, in the Middle Turonian 243 244 at 323.5 m (TUR3) and 271.5 m (TUR4/3).

Lower Turonian to basal Middle Turonian assemblages (Sequences TUR1 – 2) include
relatively high proportions of *Hystrichosphaeridium pulchrum* and *Surculosphaeridium longifurcatum* (Fig. 4). *Downiesphaeridium armatum* is most common in Middle Turonian
Sequences TUR3 – 4/1 and an acme of *Circulodinium latoaculeum* occurs in TUR4/3. Other
notable events include the LO *Microdinium distinctum* and LCOs of *Hystrichospheridium bowerbankii* and *Achomosphaera sagena* towards the top of the Middle Turonian (Sequence
TUR4/3).

252 A number of dinocyst species are more common within the regressive Upper Turonian 253 package of Sequences TUR5 – 6/1 (cf. Uličný et al., 2014), notably Downiesphaeridium aciculare, Oligosphaeridium prolixispinosum, Florentinia cooksoniae, Hystrichosphaeridium 254 tubiferum brevispinum, Kleithriasphaeridium loffrense, Kiokansium unituberculatum, S. 255 256 pontis-mariae, Tanyosphaeridium salpinx (Figs. 3, 4). Common species that temporarily disappear within this interval are *Florentinia mantellii* and *Downiesphaeridium armatum*. 257 258 The mutual exclusivity of a number of key taxa, points to palaeoenvironmental factors 259 limiting their stratigraphic distribution in the Bohemian Cretaceous Basin.

Several well-defined FOs of common species are apparent in the section (Figs. 3, 4),
notably, *Florentinia clavigera* towards the top of the Middle Turonian, and *Isabelidinium amphiatum* in the mid-Upper Turonian. *Surculosphaeridium belowii* and *Cribroperidinium wilsonii* first occur within the Turonian – Coniacian boundary interval, which from the upper
half part of Sequence TUR6/1 to mid-Sequence CON1, is characterised by assemblages
containing abundant *Sepispinula*? *ambigua*.

266 6. Turonian dinocyst biostratigraphy: review

267 Comparisons between dinocyst records from four key English Turonian Chalk sites (Fig. 268 1) and those obtained here from the Bohemian Cretaceous Basin (Figs. 3, 4) are illustrated in Fig. 5, based on macrofossil biostratigraphic correlation and further constrained by key CIEs. 269 270 Carbon isotope data (Jarvis et al., 2006) are available to more precisely place the stratigraphic 271 levels of dinocyst records (FitzPatrick, 1995; Pearce, 2000, 2010; Pearce et al., 2003; Prince 272 et al., 2008) from Berkshire (Banterwick Barn borehole). Kent (Dover composite section) and 273 Norfolk (Trunch borehole), but limited carbon isotope data are available for the Turonian of 274 East Sussex. Here, the positions of the CIEs have been placed using bio- and 275 lithostratigraphic criteria derived from other southern English Chalk sections where isotope 276 data are available (Jenkyns et al., 1994; Pearce et al., 2003; Jarvis et al., 2006). 277 6.1. Cenomanian – Turonian boundary interval

278 No Cenomanian dinocyst marker species were recognised in the Bch-1 core. For example,

279 Adnatosphaeridium tutulosum and Litosphaeridium siphoniphorum, which have last

280 occurrences in the Upper Cenomanian M. geslinianum – N. juddii zones (Foucher, 1980,

281 1981; Courtinat et al., 1991; Costa and Davey, 1992; Dodsworth, 2000; Pearce, 2000;

Lignum, 2009; Pearce et al., 2009), are absent. However, calcareous nannofossil records

place the base Turonian at 402 m, indicated by the FO *Eprolithus octopetalus* at 401.2 m and
the LOs of Upper Cenomanian species *Lithraphidites acutus* at 402.4 m and *Corollithion kennedyi* at 402.8.

286 Cauveridinium (formerly Cyclonephelium) membraniphorum and the Cyclonephelium 287 compactum – Cauveridinium membraniphorum 'complex' are important uppermost 288 Cenomanian and Turonian dinocyst markers in NW Europe (e.g. Clarke and Verdier, 1967; 289 Marshall and Batten, 1988; Pearce, 2000). The latter is found infrequently in Bch-1. The 290 taxon first occurs in the Upper Cenomanian near the core base at 403.6 m (Fig. 3), and is 291 recorded sporadically through the Lower and Middle Turonian, with a LO in the mid-Upper 292 Turonian S. neptuni Zone. Elsewhere, the LCO of C. membraniphorum is generally recorded 293 at or near the Turonian - Coniacian boundary (Prössl, 1990; Costa and Davey, 1992; Pearce, 294 2000; Pearce et al., 2003; Prince et al., 2008).

295 Cauveridinium membraniphorum ranges down into the Middle Albian (Foucher, 1981), 296 and has been recorded as high as the Santonian and Campanian (Foucher, 1979; Ioannides, 297 1986; Harker et al., 1990), but it is generally only a common component of assemblages in the top Cenomanian and Turonian. An acme of C. membraniphorum has been identified as a 298 299 useful dinocyst marker in the uppermost Cenomanian (Dodsworth, 2000; Lignum, 2009), but 300 this acme is absent at Bch-1 and likely correlates to a hiatus in the section. The species has a 301 LO or LCO near the top of the Upper Turonian in most English Chalk sections (Fig. 5), and is 302 rare in post-Turonian sediments. Prince et al. (1999) suggested that many non-figured 303 specimens attributed by previous workers to C. membraniphorum in Santonian or younger 304 sediments might be C. filoreticulatum, leaving a probable true LO of C. membraniphorum in 305 the Lower Coniacian (Pearce, 2000; Pearce et al., 2003; Prince et al., 2008). The relatively

low LO of the species in Bch-1 in the mid-Upper Turonian, reflects the low abundance of thespecies in the core.

308 Heterosphaeridium difficile and Florentinia buspina are biostratigraphic marker species 309 that have Lower Turonian bases in NW Europe (Davey and Verdier, 1976; Foucher, 1980, 1981; Tocher and Jarvis, 1987; Jarvis et al., 1988a; Costa and Davey, 1992; FitzPatrick, 1995; 310 311 Pearce et al., 2009). In the high northern latitudes, *H. difficile* has been recorded in the 312 Cenomanian (Bell and Selnes, 1997; Bloch et al., 1999) suggesting that it is a cold water 313 tolerant species that migrated southward with the predominant Late Cretaceous cooling. 314 Heterosphaeridium difficile is absent at Bch-1, and F. buspina first appears in the Middle Turonian (Fig. 3). A mid-Turonian FO or FCO of F. buspina is also noted in most English 315 316 Chalk sections (Fig. 5), although the exact stratigraphic position is variable, and sporadic 317 occurrences range down into the Cenomanian (FitzPatrick, 1995; Pearce, 2000; Pearce et al., 318 2009).

319 6.2. Senoniasphaera bioevents

320 The first occurrence of Senoniasphaera rotundata, originally described by Clarke and 321 Verdier (1967), has been widely regarded as being a marker for the Lower Turonian (Foucher, 322 1980, 1981; Tocher and Jarvis, 1987, 1994, 1995; Jarvis et al., 1988a, b; FitzPatrick, 1995; 323 Lamolda and Mao, 1999; Dodsworth, 2000; Pearce, 2000; Pearce et al., 2003, 2011). Recent 324 taxonomic revisions (Pearce et al., 2003, 2011), however, differentiate two main species of 325 Senoniasphaera in the Turonian: S. turonica (= S. rotundata alveolata of Pearce et al., 2003, 2009; Prince et al., 2008); and S. rotundata (= S. rotundata rotundata). Senoniasphaera spp. 326 327 are uncommon (Fig. 3) in Bch-1, but S. turonica first occurs within the Mammites nodosoides 328 Zone, between the Holywell and Lulworth CIEs.

329 Published FOs of S. 'rotundata' (interpreted here to be records of S. turonica; see Pearce et 330 al., 2011 for discussion) occur in the lower part of the Lower Turonian Mytiloides labiatus 331 Zone (equivalent to the W. devonense – M. nodosoides zones) in SE Devon (Jarvis et al., 332 1988b) and East Sussex (FitzPatrick, 1995; Pearce et al., 2009), at the top of the zone in Berkshire (Pearce et al., 2003), at the base of the Middle Turonian Terebratulina lata Zone 333 334 (=C. woollgari Zone) in Kent (FitzPatrick, 1995), and towards the top of that zone in Norfolk (Pearce, 2000), where the Lower to basal Middle Turonian is barren of dinocysts (Fig. 5). 335 336 The observed diachroneity of the FO S. turonica is likely facies controlled, dinocysts being 337 less abundant in, and the index species absent from, coarser-grained condensed nodular chalk 338 facies that characterise the Lower Turonian in many areas. 339 The Lower Turonian in Bch-1 yielded abundant dinocysts throughout, but S. turonica was 340 recorded in only two samples (Fig. 3). Its FO in the lower Mammites nodosoides Zone is 341 consistent with published lowest records of the species in the English Chalk (e.g. Tocher and 342 Jarvis, 1987; Pearce et al., 2009). However, this cannot be regarded as being definitive. 343 Sporadic occurrences of S. turonica consistently range down into the Middle Cenomanian, 344 upper Acanthoceras rhotomagense Zone, at Culver Cliff, Isle of Wight England, at Wunstorf 345 Quarry, northern Germany, and at Vergons, SE France, indicating that this is the true FO of 346 the species (Lignum, 2009; Pearce et al., 2011). The FCO of S. turonica, however, typically

347 lies towards the base of the Lower Turonian (Pearce et al., 2011).

Senoniasphaera abundances vary considerably between sections, but an acme of S. *turonica* occurs consistently in the Upper Turonian (*Sternotaxis plana* Zone; *Subprionocyclus neptuni* Zone) of the English Chalk successions (Fig. 5). *Senoniasphaera rotundata* s.s. first
appears in the Upper Turonian, with a FCO around the stage boundary and an acme in the
Lower Coniacian (*Micraster cortestudinarium* Zone). *Senoniasphaera rotundata* was not

353 recorded from Bch-1. However, the LO S. turonica occurs in the lower S. neptuni Zone,

above the Hitch Wood CIE, which is correlative with the uppermost acme of the species

355 elsewhere (Fig. 5).

356 6.3. Middle Turonian dinocyst events

Oligosphaeridium poculum first occurs towards the base of the C. woollgari Zone at Bch-1 357 358 (Fig. 3). Its FO is recorded at a similar stratigraphic level in Berkshire and Kent (FitzPatrick, 359 1995; Pearce et al., 2003), but occurs lower in East Sussex, at the base of the *M. nodosoides* 360 Zone (Pearce et al., 2009). The FO of O. poculum in the Trunch borehole of eastern England 361 (see Pearce, 2010 for locality details) lies within the basal Coniacian, albeit based on only two 362 records from the core (Pearce, 2000), and these may not be *in situ* as that interval shows 363 evidence of intermittent sediment reworking. The datum level appears to be a consistent 364 Turonian marker, although the species has been recorded from the Hauterivian of Germany (Prössl, 1990) and the Barremian – Albian, of Greenland, England and India (Jain, 1977; 365 366 Lister and Batten, 1988; Nøhr-Hansen, 1993), so the Turonian FO is in truth a reoccurrence (RO) datum level, as indicated by Pearce et al. (2009, fig. 3). 367

The FO *Subtilisphaera pontis-mariae* is found within the lower *C. woollgari* zone in Bch-1, marginally higher than the upper Lower Turonian placement of Hardenbol et al. (1998) for the Tethyan realm. However, the species has a long stratigraphic range, from at least Upper Albian (Davey, 1970; Lignum, 2009) to Lower Campanian (Prince et al., 1999; Pearce, 2000), so its stratigraphic utility is limited.

373 The FO *Raetiaedinium truncigerum* lies in the mid-*C. woollgari* Zone at Bch-1, slightly
374 below the Glynde CIE. The FO of *R. truncigerum* has been previously recorded also from the

upper Middle Turonian by Prössl (1990), or slightly higher in the Upper Turonian (Foucher,

376 1976a; Marshall, 1983; Williams and Bujak, 1985; Williams et al., 1993; Hardenbol et al.,

377	1998) or Lower Coniacian (Kirsch, 1991), and so the FO appears to be a potential Middle
378	Turonian marker. The species ranges into the Campanian in England and Germany (Kirsch,
379	1991; Williams et al., 1993; Prince et al., 1999; and by our personal observations).
380	Florentinia buspina also first occurs in the mid-C. woollgari Zone at Bch-1 (Fig. 3). A
381	mid-Turonian FO or FCO of <i>F. buspina</i> is noted in most English Chalk sections (Fig. 5),
382	although the exact stratigraphic position is variable. This is again a RO, since the species
383	extends down into the Lower Cenomanian at Trunch (Pearce, 2000), at Culver Cliff Isle of
384	Wight, and at Vergons SE France (Lignum, 2009). The species ranges upwards into the
385	Campanian in Germany (Davey and Verdier, 1976; Kirsch, 1991; Pearce, 2000).
386	FitzPatrick (1995), Pearce (2000) and Pearce et al. (2003) recognised 'acmes' of
387	<i>Circulodinium distinctum</i> within the Middle Turonian <i>T. lata</i> Zone of the English Chalk.
388	Three main peaks of <i>C. distinctum</i> are found in the Bch-1 core (Figs. 4, 5): the first, in the
389	mid-C. woollgari Zone, between the Round Down and Glynde CIEs, is the highest amplitude
390	peak but is confined to a single sample; the second broader peak is towards the top of the C .
391	woollgari Zone, at the level of the 'Pewsey' CIE; the third peak is located in the mid-S.
392	neptuni Zone mid-way between the Bridgewick and the Hitch Wood CIEs. The first and
393	second peaks correspond stratigraphically to floods and acme intervals of C. distinctum found
394	in the English Chalk (Fig. 5). The final higher peak in the Upper Turonian may be of local
395	significance, or may not have been recognised (or preserved) in the lower resolution sampling
396	of the more attenuated, potentially less complete, English Chalk Upper Turonian successions
397	(e.g. see discussion in Uličný et al., 2014).

398 6.4. Upper Turonian dinocyst events

The Upper Turonian at Bch-1 is marked by a series of last occurrence datum levels (Figs.
3, 5). The LO of *Stephodinium coronatum* occurs in the lower *S. neptuni* Zone at Bch-1. The

LO of the species has been recorded marginally higher elsewhere in the area, in the *M. scupini*Zone of the Úpohlavy section, located 25 km NNW of Prague (Svobodová et al., 2002). This
level correlates to a position just below the FO of *P. germari* in other Bohemian Cretaceous
Basin sections (cf. Fig. 2). This compares favourably to a top Upper Turonian LO of *Stephodinium coronatum* at Trunch (Pearce, 2000). The LO *S. coronatum* has also been
recorded from the Turonian – Coniacian boundary interval elsewhere (Foucher, 1976b; Costa
and Davey, 1992; Williams et al., 1993).

Bujak and Williams (1978) and Williams and Bujak (1985) stated that *S. coronatum* occurs
in the Santonian, but without distribution data it is unknown whether this observation might
be based on reworked specimens; Ioannides (1986) also recorded *S. coronatum* from
Santonian to ?Maastrichtian sediments, but in samples that clearly contained reworked taxa.
We therefore discount these records in the light of overwhelming evidence, at least for NW
Europe, that the LO of *S. coronatum* occurs in the uppermost Turonian, close to the Turonian
- Coniacian boundary.

The coincident LOs of S. turonica and C. compactum – C. membraniphorum occur in the 415 416 mid-S. neptuni Zone, below the Hitch Wood CIE; the former is somewhat lower than the last 417 occurrence of S. turonica in the English Chalk (Fig. 5). The FCO of Chatangiella ditissima occurs towards the base of the *M. scupini* Zone at Bch-1. The FO of *C. ditissima* has been 418 419 previously recorded from the lowermost Turonian (Costa and Davey, 1992), the Upper 420 Turonian (Sweet and McIntyre, 1988; Prössl, 1990; Scott, 2014) and the Lower Coniacian 421 (Williams et al., 1993). However, a high Upper Turonian (high S. plana Zone) FCO was 422 recorded at Trunch (Fig. 5) by Pearce (2000), at a comparable level to Bch-1, with lower sporadic occurrences in the Lower Cenomanian. The FCO of the species offers, therefore, a 423 424 potential biostratigraphic datum level. Chatangiella ditissima is a common component of

- 425 Santonian Campanian assemblages in many areas (e.g. McIntyre, 1975; Ioannides, 1986;
- 426 Costa and Davey, 1992; Skupien et al., 2009; Radmacher et al., 2014), with a likely LO in the
- 427 Maastrichtian (Kirsch, 1991; Williams et al., 1993; Lebedeva, 2006; Lebedeva et al., 2013).
- 428 6.5. Turonian Coniacian dinocyst boundary events

429 The Turonian – Coniacian boundary interval is marked by a series of benthic macrofossil 430 events, particularly affecting the inoceramid bivalves (Walaszczyk, 2000), that enable the 431 development of a refined biostratigraphy at Bch-1 (Fig. 5). A number of dinocyst events are 432 also apparent. The FO Surculosphaeridium belowii occurs immediately below the stage 433 boundary and the Navigation CIE at Bch-1 (Fig. 3). This species has been recorded 434 previously largely from Coniacian - Maastrichtian sediments in England, Germany and 435 Austria (Yun, 1981; Kirsch, 1991; Pearce, 2000; Soliman et al., 2009; Mohamed and 436 Wagreich, 2013), but it has also been identified from the Barremian of the Slovak Carpathians 437 (Skupien, 2003): however, its Turonian reoccurrence (RO) datum level may be of 438 stratigraphic value.

439 The FO Cribroperidinium wilsonii occurs immediately above the base Coniacian and the 440 Navigation CIE at Bch-1 (Figs. 3, 5). A Turonian – Coniacian FO is consistent with records 441 from Germany and Denmark (Kirsch, 1991; Schiøler, 1992), although the species ranges from 442 Upper Albian – Campanian in southern England (Prince et al., 1999, 2008; Pearce, 2000; 443 Lignum, 2009). However, the FCO C. wilsonii is observed in the high Upper Turonian mid-444 S. plana Zone above the Hitch Wood CIE at Trunch (Fig. 5), and here it becomes increasingly 445 common through the mid- to Upper Coniacian, confirming its importance as a significant 446 component of Coniacian dinocyst assemblages. Although Palaeohystrichophora 447 infusorioides is a major component of the dinocyst assemblages through most of the 448 succession at Bch-1, a prominent acme occurs in the lowest Coniacian C. d. erectus Zone

449	(Figs. 4, 5). A similar acme is recorded in the lowest Coniacian at Trunch (Pearce, 2000),
450	though it has not been identified in other Chalk sections studied to date.

451 The first common and consistent occurrence of Oligosphaeridium pulcherrimum occurs at 452 the top of the Cremnoceramus deformis erectus Zone at Bch-1, at the base of the Beeding 453 CIE; an acme of the species occurs a short distance above, at the base of the C. c. crassus 454 Zone. The FO of persistently occurring specimens, FCO and/or acme of the species are 455 observed around the Turonian/Coniacian boundary in Chalk sections throughout southern 456 England (Fig. 5). The oldest record of O. pulcherrimum is from the Upper Jurassic 457 (Brideaux, 1977), and it is a common component in the Lower Cretaceous (Prössl, 1990). 458 However, the species has been noted previously as becoming more common in the Coniacian 459 (Foucher, 1980; Pearce et al., 2003); in the zonation schemes of Williams (1975, 1977), O. 460 pulcherrimum is an important indicator species for the Coniacian O. pulcherrimum Zone. 461 *Pervosphaeridium truncatum* ranges through the Cenomanian – Coniacian succession 462 sampled at Bch-1. The species has a LO in the *M. coranguinum* Zone in Berkshire and 463 Norfolk (Marshall, 1983; Pearce et al., 2003), southern England (Fig. 5), indicating that the 464 top of Bch-1 probably lies below that stratigraphic level. However, records of the species 465 from the Campanian – Maastrichtian in Germany and Canada (Harker et al., 1990; Kirsch, 466 1991) may point to a younger true LO.

The FO *Heterosphaeridium verdieri* was proposed as a high Lower Coniacian marker by
Prince et al. (2008), based on records from Kent and the Isle of Wight. The species had
previously been reported from Santonian deposits in Germany (Yun, 1981; Kirsch, 1991).
The species has been recorded from the high-Middle Turonian in Berkshire (Pearce et al.,
2003) and as a very rare component (possibly misidentified?) from the Upper Cenomanian of
East Sussex (Pearce et al., 2009), but it has a FCO and acme in the low Middle Coniacian

- 473 (Fig. 5). The absence of *H. verdieri* and *Spinidinium echinoideum* (another Middle Coniacian
- 474 marker) at Bch-1 is consistent with the Early Coniacian age of the sampled section top
- 475 indicated by macrofossil and nannofossil records (*C. c. crassus* Zone, base UC10; Fig. 2).

476 **7. Turonian dinocyst zonation**

477 No generally accepted dinocyst zonation scheme exists for the Turonian Stage. Here,
478 previously published zonations are critically reviewed in the light of recent studies, including
479 the new data obtained from the Bch-1 core. The viability of individual zones is considered
480 and, following revision, a new composite scheme is proposed.

481 7.1. Clarke and Verdier (1967) zonation

482 The *Hystrichosphaeridium* (now *Litosphaeridium*) *siphoniphorum* Zone was created by 483 Clarke and Verdier (1967) to approximate the Cenomanian, from the stage base to a position 484 within the Plenus Marl of southern England, which at the time, was considered to be questionably Turonian (cf. Jefferies, 1963). The stage boundary is now placed higher, within 485 486 the overlying Ballard Cliff Member (Gale et al., 2005; Pearce et al., 2009). Litosphaeridium 487 siphoniphorum tends to be common between its range base in the Upper Albian and the 488 Upper Cenomanian. The LCO of L. siphoniphorum occurs consistently within the Upper 489 Cenomanian (*M. geslinianum* Zone) throughout Europe and in North America, although rare occurrences extend into the Lower Turonian (Davey, 1969; Foucher, 1979, 1980, 1982; 490 491 Marshall and Batten, 1988; Courtinat et al., 1991; Li Huan and Habib, 1996; Hardenbol et al., 492 1998; Lamolda and Mao, 1999; Dodsworth, 2000; Pearce, 2000; Lignum, 2009; Pearce et al., 493 2009). The absence of L. siphoniphorum at Bch-1 is consistent with an absence of low M. 494 geslinianum Zone and older Cenomanian sediments within the sampled interval.

495 Clarke and Verdier (1967) erected a Scriniodinium (previously Endoscrinium) campanula 496 Zone to represent the uppermost Cenomanian to the lower Turonian interval (Fig. 5), which 497 was characterised by the presence of *Xiphophoridium alatum* (now *Dinopterygium alatum*), 498 *Cometodinium obscurum* and *Cyclonephelium hughesii*. The base was defined by the top of 499 the L. siphoniphorum Zone, and the top by the LO of S. campanula. At Bch-1, S. campanula 500 was recorded only in two Lower Coniacian samples, and is absent from the Turonian. The 501 taxon occurs sporadically throughout the Cenomanian – Santonian at Trunch, with a LO here 502 and at Culver Cliff in the Upper Santonian, mid-U. socialis Zone (Prince et al., 1999, 2008; 503 Pearce, 2000), and it is never a common component of the assemblage. Furthermore, D. 504 alatum occurs through the Cenomanian to Upper Turonian (S. plana Zone), and C. obscurum 505 is only commonly recorded from the Upper Turonian (S. plana Zone), so the S. campanula 506 Zone as defined previously is not viable.

507 *Cauveridinium membraniphorum* is a very distinctive component of uppermost

508 Cenomanian and Turonian dinocyst assemblages throughout Europe (Section 6.1, above). A

509 Cyclonephelium (now Cauveridinium) membraniphorum Zone, of Turonian to Coniacian /

510 Santonian age, was erected by Clarke and Verdier (1967) based on limited sampling from the

511 Isle of Wight. The base of the zone was defined by the LO of *S. campanula*, and its top by

512 the LO of *C. membraniphorum* (Fig. 5). Subsequent records extending the range of *S.*

513 *campanula* into the Upper Santonian make the LO of this species untenable as a basal marker

514 for a largely Turonian zone. Accordingly, it is proposed here to redefine the base of the

515 *Cauveridinium membraniphorum* Zone by the LCO of *L. siphoniphorum*.

516 Clarke and Verdier (1967) erected a *Deflandrea echinoidea* (now *Spinidinium*

517 echinoideum) Zone to represent the ?lower – mid-Santonian, defined as the interval

518 immediately following the LO of *C. membraniphorum*, to immediately below the FO of

519 Hystrichosphaeropsis ovum, Dinogymnium albertii and D. heterocostatum. The FO of S.

520 echinoideum occurs in the Coniacian (Williams and Bujak, 1985); at Trunch and in Kent

521 (Pearce, 2000; Prince et al., 2008) its FO is in the mid-Coniacian, low *M. coranguinum* Zone

522 (Fig. 5). The index species has commonly low and varying abundance through the Coniacian

523 – Campanian in different sections, but it generally shows a FCO around the base Santonian,

524 making it a good Santonian marker (Foucher, 1976a; Heine, 1991; Kirsch, 1991; Prince et al.,

525 1999; Pearce, 2000).

526 Using the original definition of Clarke and Verdier (1967), beds above the LO of *C*.

527 *membraniphorum* at Bch-1 technically fall within their *S. echinoideum* Zone (Fig. 5).

However, in reality the FO of the zonal index species falls stratigraphically above the interval
represented in the Bch-1 core, and the zone is not considered to be relevant to the current
study.

531 7.2. Williams (1977) zonation

Williams (1977) 'global' Upper Cretaceous dinocyst zonation scheme, based largely on
assemblage data from southern England (Clarke and Verdier, 1967; Davey, 1969, 1970) and
offshore eastern Canada, employed *Kiokansium unituberculatum* (then referred to as *Cleistosphaeridium polypes*) as a Cenomanian index species (Fig. 5). Subsequently, the LO
of *Kiokansium unituberculatum* has been recorded as Lower Turonian (Foucher, 1981), and
sporadic occurrences extend into the basal Lower Coniacian at Trunch (Pearce, 2000) but in
an interval with significant Cenomanian reworking, so this latter record may be unreliable.

Williams (1977) Turonian zonal index, *Surculosphaeridium? longifurcatum* (Fig. 5), is a
long-ranging species, from Upper Barremian (Prössl, 1990) to Campanian (Williams and
Bujak, 1985; Kirsch, 1991; Williams et al., 1993; Pearce, 2000). It is abundant in the Lower
Turonian to basal Middle Turonian and in the Lower Coniacian at Bch-1, but its abundance

records show little consistency between Chalk sections and it is considered to be of littlebiostratigraphic value.

545 Williams (1977) recognised a Callaiosphaeridium asymmetricum – Oligiosphaeridium pulcherrimum Zone within the Coniacian - Lower Santonian (Fig. 5). The first species is 546 long ranging, from Hauterivian (Davey and Williams, 1966a, b; Warren, 1967) to Campanian 547 548 (Foucher, 1979; Kirsch, 1991). By contrast, O. pulcherrimum has a consistent FCO in the 549 Lower Coniacian (Pearce, 2000; Pearce et al., 2003; Prince et al., 2008), and it is proposed 550 here as a Coniacian zonal index species (Fig. 5). 551 7.3. Foucher (1981) zonation 552 Foucher (1981) erected a Senonisphaera rotundata Zone, defined by the FO S. rotundata, 553 for the Turonian (top not considered), with an Upper Cenomanian – Lower Turonian 554 Silicisphaera (now Florentinia) ferox Zone, below (Fig. 5). The latter is long-ranging, 555 Hauterivian (Gocht, 1959) to Maastrichtian (Clarke and Verdier, 1967; Foucher, 1975, 1976a; 556 Foucher and Robaszynski, 1977; Kirsch, 1991), and is therefore of limited stratigraphic value. By contrast, Senonisphaera rotundata sensu lato has proved to be a good biostratigraphic 557 558 marker (Section 5.2). The genus is used here to define a new Lower to Middle Turonian S. 559 turonica Subzone (C. membraniphorum Zone; Fig. 5) and a Lower Coniacian S. rotundata 560 sensu stricto Subzone (O. pulcherrimum Zone).

561 7.4. FitzPatrick (1995) zonation

562 A palynological zonation scheme for the uppermost Cenomanian to Turonian of the

southern English Chalk was proposed by FitzPatrick (1995). Three zones were erected:

564 Palynozones I to III. Palynozone I was considered to represent the Lower Turonian, with a

565 base defined by the FO of Heterosphaeridium difficile, Senoniasphaera rotundata sensu lato,

and *Florentinia buspina*. Another species thought to be an important index taxon was

567 *Litosphaeridium* sp. A of Marshall and Batten (1988), which was recorded as having a LO in

the lowest Turonian of East Sussex, immediately below the base of the *Mammites nodosoides*Zone.

Palynozone II was considered to represent the mid-Turonian (*T. lata* Zone), and was
defined by the FO of *Florentinia? torulosa* to the LO of *S. campanula*. Acmes of *H. difficile*and *C. distinctum* characterise the middle part of the zone. *Florentinia? torulosa* is absent in
the Turonian of Bch-1, and rare to absent in most English Chalk sections, so it is a rather poor
index species. More importantly, the LO of *S. campanula* is Santonian (Foucher, 1976a;
Kirsch, 1991; Costa and Davey, 1992; Prince et al., 1999; Pearce, 2000), so this zone cannot
be employed as a Turonian marker.

577 Palynozone III was erected to represent the Upper Turonian, and was defined as lying directly above Palynozone II (i.e. above the LO of *S. campanula*); the top of the zone was not 578 579 defined. An acme of Senoniasphaera 'rotundata' (considered here to be records of S. 580 *turonica*) occurring in the lower part of the zone was considered to be of correlative value. In 581 the light of the proven extended stratigraphic range of S. campanula in the English Chalk and 582 elsewhere, Palynozone III would represent uppermost Santonian or Campanian, not Turonian 583 strata. The Turonian palynozonation scheme of FitzPatrick (1995) is therefore fundamentally 584 flawed.

585 7.5. A revised Turonian zonation

586 Here, we propose a single *Cauveridinium membraniphorum* Interval Zone spanning the top

587 Cenomanian to basal Coniacian (Fig. 5). The base of the *C. membraniphorum* Zone is

588 marked by the LCO of *Litosphaeridium siphoniphorum* (top *L. siphoniphorum* Zone). The *C.*

589 membraniphorum Zone is subdivided into: Senoniasphaera turonica (Lower – Middle

- 590 Turonian) and *Raetiaedinium truncigerum* (Middle Turonian basal Coniacian) subzones.
- 591 The top of the *C. membraniphorum* Zone is defined by the FCO *Oligosphaeridium*
- 592 *pulcherrimum (S. rotundata* Subzone; base O. *pulcherrimum Zone)*.

593 The FO of *S. turonica* is lower Middle Cenomanian *Acanthoceras rhotomagense* Zone in

southern England, northern Germany and SE France (Pearce et al., 2011), but in northern

595 Europe the species is rare at its FO and it is commonly absent from the Upper Cenomanian.

596 However, it shows a widespread RO and FCO in the Lower Turonian (Foucher, 1980; Tocher

⁵⁹⁷ and Jarvis, 1987; Jarvis et al., 1988a; Lamolda and Mao, 1999; Pearce et al., 2003, 2009).

598 The FCO/RO S. turonica defines the base of our S. turonica Subzone; its top is placed at the

599 FO Raetiaedinium truncigerum.

The FO of *R. truncigerum* in Bch-1 occurs within the Middle Turonian *C. woollgari* Zone,
below the Glynde CIE, consistent with records of its lowest FO elsewhere (Section 5.3). It
defines the base of the *R. truncigerum* Subzone.

Despite ranging down into the Hauterivian, *O. pulcherrimum* shows a FCO and acme in
the basal Coniacian (*Cremnoceramus deformis erectus* Zone; Beeding CIE) of Bch-1, which
is consistent with records from the basal Coniacian of the English Chalk (Fig. 5). The FCO
(in reality, a reoccurrence datum level; Section 5.5) is used here to mark the base of the *O. pulcherrimum* Zone, the lower part of which includes the FCO and an acme of *S. rotundata*,
constituting the *S. rotundata* Subzone (top not defined).

609 8. Conclusions

610 An expanded Turonian cored succession (Bch-1) from Běchary in the Bohemian

611 Cretaceous Basin has yielded abundant and diverse assemblages of well-preserved dinocysts

612 throughout the Turonian – Lower Coniacian. Dinocyst records are constrained by calcareous

nannofossil, macrofossil and sequence stratigraphic data (Uličný et al., 2014). A highresolution (5.6 kyr) carbon isotope ($\delta^{13}C_{org}$) profile obtained from the core provides the basis for identifying the Turonian carbon isotope events (CIEs) of Jarvis et al. (2006) in the succession, and placing dinocyst datum levels directly within the CIE stratigraphy. To our knowledge, these records offer the highest resolution, fully stratigraphically constrained, Turonian dinocyst dataset published to date.

619 A succession of 18 dinocyst datum levels, considered to be of potential regional

620 biostratigraphic significance, are recognised in Bch-1. From bottom to top, these are (Fig. 5):

621 FOs of *C. compactum – C. membraniphorum* and then *S. turonica* in the Lower Turonian;

622 FOs of O. poculum, S. pontis-mariae, a flood of C. d. distinctum, the FOs of R. truncigerum

623 and *F. buspina*, and a second flood of *C. d. distinctum* in the Middle Turonian; LO S.

624 *coronatum*, a *C. d. distinctum* acme, the LOs *C. compactum – C. membraniphorum*, *S.*

625 turonica and K. unituberculatum, the FCO C. ditissima and FO S. belowii in the Upper

626 Turonian; FO C. wilsonii coincides with the base Coniacian, followed by an acme of P.

627 *infusorioides*, the FCO and then an acme of *O. pulcherrimum*. *Pervosphaeridium trunctatum*

628 ranges to the top of the sampled section (high Lower Coniacian).

629 A review of published Turonian records from English Chalk sections in East Sussex,

630 Berkshire, Kent and Norfolk demonstrates the presence of several consistent dinocysts datum

631 levels. However, detailed stratigraphic comparison is hampered by lateral thickness variation,

632 by differences in sampling resolution and stratigraphic completeness, and by the presence of

barren intervals and poor recovery in some sections. Consistent Chalk datum levels are

634 provided by the FO *S. turonica* low in the Lower Turonian, an acme of *C. d. distinctum* in the

635 high Middle Turonian, an acme of *S. turonica* in the mid-Upper Turonian with the FO *O*.

636	pulcherrimum towards the top of the substage. The Lower Coniacian is marked by the FCO		
637	or an acme of O. pulcherrimum, followed by an acme/FCO S. rotundata.		
638	Critical review of published Turonian dinocyst zonation schemes shows them to be		
639	untenable. A revised dinocyst zonation for the Turonian based on our review of the English		
640	Chalk data and new results from Bch-1, tied to the carbon isotope event stratigraphy of Jarvis		
641	et al. (2006), has been presented that is judged to be of likely wider geographic significance.		
642	Correlation of the English Chalk data to Bch-1 provides a basis for defining a regional		
643	dinocyst event stratigraphy with 22 datum levels, and a revised dinocyst zonation scheme		
644	constrained within a chemostratigraphic framework of 10 major CIEs.		
645	The proposed new dinocyst zones consist of a Cenomanian Litosphaeridium		
646	siphoniphorum Zone, followed by the Cauveridinium membraniphorum Zone spanning the		
647	uppermost Cenomanian to Lower Coniacian. This is subdivided into: Senoniasphaera		
648	turonica (Lower – mid-Middle Turonian); and Raetiaedinium truncigerum (mid-Middle		
649	Turonian – mid-Lower Coniacian) subzones. The Oligosphaeridium pulcherrimum Zone		
650	(Senonisphaera rotundata Subzone) characterises the Lower Coniacian.		
651	Key regional datum levels are (Fig. 6):		
652	(1) within the Cenomanian – Turonian boundary CIE: LCOs <i>L. siphoniphorum</i> and <i>W</i> .		
653	cassidata (Upper Cenomanian, M. geslinianum Zone), defining the base of the C.		
654	membraniphorum Zone;		
655	(2) FO <i>C. compactum</i> – <i>C. membraniphorum</i> and LO <i>A. tutulosum</i> followed by an acme o		
656	C. membraniphorum in the uppermost Cenomanian (Neocardioceras juddii Zone);		

- 657 (3) FO *S. turonica* below the Holywell CIE (Lower Turonian, *F. catinus* Zone), defining
- 658 the base of the *S. turonica* Subzone;
- 659 (4) RO *O. poculum* above the Holywell CIE;

- 660 (5) flood of *C. d. distinctum* in the mid-Middle Turonian *C. woollgari* Zone;
- 661 (6) FO *R. truncigerum* between the Round Down and Glynde CIEs (Middle Turonian, *C.*
- 662 *woollgari* Zone), defining the base of the *R. truncigerum* Subzone;
- 663 (7) acme of *H. difficile* coincident with the Glynde CIE;
- 664 (8) an acme interval of *C. d. distinctum* at the summit of the Middle Turonian *C. woollgari*665 Zone, spanning the 'Pewsey'CIE;
- 666 (9) acme interval of *S. turonica* spanning the Hitch Wood CIE *S. neptuni* Zone, following
- 667 the LO *C. compactum C. membraniphorum*;
- 668 (10) FCO C. ditissima in the higher Upper Turonian;
- 669 (11) RO S. belowii, then LCO C. membraniphorum immediately below the Navigation CIE
- 670 in the uppermost Turonian, with the LO *K. unituberculatum* immediately above in the671 base Coniacian;
- 672 (12) acme of *P. infusorioides* in the Lower Coniacian between the Navigation and Beeding
 673 CIEs;
- 674 (13) FCO O. pulcherrimum below the Beeding CIE (Lower Coniacian, F. (H.)
- 675 *petrocoriensis* Zone), defining the bases of the *O. pulcherrimum* Zone and *S. rotundata*676 Subzone.
- 677 (14) acmes of *O. pulcherrimum* and then *S. rotundata* in the mid-Lower Coniacian, with the
 678 FCO *H. verdieri* towards the top of the substage.
- 679 The new stratigraphy offers potential for improved correlation and dating of Upper
- 680 Cretaceous successions. However, additional high-resolution dinocyst datasets from multiple
- 681 sections are required to test and further refine the Turonian biostratigraphy.

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689	Appendix A. Species list		
690	Complete list of dinoflagellate cyst species recorded in the Bch-1 core. Numbers correspond		
691	to the orde	r of species plotted in Fig. 3. Other taxa, found less commonly, are also listed	
692	along with	the genetic sequence(s) from which they have been identified (i.e. $TUR1 - 7$,	
693	CON1 – 2,	Figs. 2, 3). Taxonomic references are given in Fensome et al. (2008) and Pearce	
694	et al. (2011).		
695	26	Achomosphaera ramulifera ramulifera (Deflandre 1937) Evitt 1963	
696	33	Achomosphaera regiensis Corradini 1973	
697	1	Achomosphaera sagena Davey and Williams 1966	
698	2	Callaiosphaeridium asymmetricum (Deflandre and Courteville 1939) Davey and	
699		Williams 1966	
700	41	Cassiculosphaeridia reticulata Davey 1969	
701	22	Cauveridinium membraniphorum – Cyclonephelium compactum Complex of	
702		Marshall and Batten 1988	
703	58	Chatangiella ditissima (McIntyre 1975) Lentin and Williams 1976	
704	3	Circulodinium distinctum distinctum (Deflandre and Cookson 1955) Jansonius	
705		1986	
706	39	Circulodinium latoaculeum (Yun Hyesu 1981) Islam 1993	
707	TUR3	Cometodinium obscurum Deflandre and Courteville 1939	
708	59	Cometodinium whitei (Deflandre and Courteville 1939) Stover and Evitt 1978	
709	CON1, 2	Conosphaeridium Cookson and Eisenack 1969	

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710	CON1	Conosphaeridium striatoconum (Deflandre and Cookson 1955) Cookson and
711		Eisenack 1969
712	4	Coronifera oceanica Cookson and Eisenack 1958
713	5	Cribroperidinium orthoceras (Eisenack 1958) Davey 1969
714	70	Cribroperidinium sp. Neale and Sarjeant 1962
715	76	Cribroperidinium wilsonii (Yun Hyesu 1981) Poulsen 1996
716	74	Cyclonephelium filoreticulatum (Slimani 1994) Prince et al. 1999
717	TUR6/1, C	CON2 <i>Cyclonephelium hughesii</i> Clarke and Verdier 1967
718	31	Dapsilidinium laminaspinosum (Davey and Williams 1966) Lentin and Williams
719		1981
720	72	Dinogymnium acuminatum Evitt et al. 1967
721	27	Dinopterygium cladoides Deflandre 1935
722	40	Disphaeria macropyla Eisenack and Cookson 1960
723	23	Downiesphaeridium aciculare (Davey 1969) Islam 1993
724	42	Downiesphaeridium armatum (Deflandre 1937) Islam 1993
725	37	Ellipsodinium rugulosum Clarke and Verdier 1967
726	24	Exochosphaeridium arnace Davey and Verdier 1973
727	43	Exochosphaeridium bifidum (Clarke and Verdier 1967) Clarke et al. 1968
728	60	Exochosphaeridium phragmites Davey et al. 1966
729	67	Florentinia buspina (Davey and Verdier 1976) Duxbury 1980
730	71	Florentinia clavigera (Deflandre 1937) Davey and Verdier 1973

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731	28	Florentinia cooksoniae (Singh 1971) Duxbury 1980
732	47	Florentinia laciniata Davey and Verdier 1973
733	7	Florentinia mantellii (Davey and Williams 1966) Davey and Verdier 1973
734	34	Florentinia sp. Davey and Verdier 1973
735	8	Hystrichodinium pulchrum Deflandre 1935
736	TUR4/1	Hystrichosphaeridium sp. Deflandre 1937
737	32	Hystrichosphaeridium bowerbankii Davey and Williams 1966
738	49	Hystrichosphaeridium conispiniferum Yun Hyesu 1981
739	9	Hystrichosphaeridium recurvatum (White 1842) Lejeune-Carpentier, 1940
740	50	Hystrichosphaeridium salpingophorum Deflandre 1935
741	10	Hystrichosphaeridium tubiferum tubiferum (Ehrenberg 1838) Deflandre 1937
742	63	Hystrichosphaeridium tubiferum brevispinum (Davey and Williams 1966) Lentin
743		and Williams 1993
744	52	Hystrichostrogylon membraniphorum Agelopoulos 1964
745	69	Impletosphaeridum clavulum (Davey 1969) Islam 1993
746	74	Isabelidinium? amphiatum (McIntyre 1975) Lentin and Williams 1977
747	TUR4/1	Kallosphaeridium? ringnesiorum (Manum and Cookson 1964) Helby 1987
748	54	Kiokansium unituberculatum (Tasch in Tasch et al. 1964) Stover & Evitt 1978
749	35	Kleithriasphaeridium loffrense Davey and Verdier 1976
750	11	Kleithriasphaeridium readei (Davey and Williams 1966) Davey and Verdier 1976

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751	TUR6/1, C	ON2 <i>Litosphaeridium arundum</i> (Eisenack and Cookson 1960) Davey 1979;
752		emend. Lucas-Clark 1984
753	36	Membranilarnacia polycladiata Cookson and Eisenack in Eisenack 1963
754	29	Microdinium distinctum Davey 1969
755	55	Nematosphaeropsis denseradiata (Cookson and Eisenack 1962) Stover and Evitt
756		1978
757	45	Odontochitina costata Alberti 1961; emend. Clarke and Verdier 1967
758	12	Odontochitina operculata (Wetzel 1933a) Deflandre and Cookson 1955
759	53	Odontochitina singhii Morgan 1980
760	TUR4/1	Odontochitinopsis molesta (Deflandre 1937) Eisenack 1961
761	13	Oligosphaeridium complex (White 1842) Davey and Williams 1966
762	48	Oligosphaeridium poculum Jain 1977
763	14	Oligosphaeridium prolixispinosum Davey and Williams 1966
764	46	Oligosphaeridium pulcherrimum (Deflandre and Cookson 1955) Davey and
765		Williams 1966
766	15	Palaeohystrichophora infusorioides Deflandre 1935
767	CON1	Pareodinia ceratophora Deflandre 1947
768	TUR2	Pervosphaeridium sp. Yun Hyesu 1981
769	TUR2 – 4/	1, 4/3 <i>Pervosphaeridium cenomaniense</i> (Norvick 1976) Below 1982
770	TUR3 – 4/	1, CON1 Pervosphaeridium monasteriense Yun Hyesu 1981
771	64	Pervosphaeridium pseudhystrichodinium (Deflandre 1937) Yun Hyesu 1981

772	16	Pervosphaeridium truncatum (Davey 1969) Below 1982
773	17	Prolixosphaeridium granulosum (Deflandre 1937) Davey et al. 1966
774	18	Pterodinium "crassimuratum" (Davey and Williams 1966) Thurow et al. 1988
775	68	Raetiaedinium truncigerum (Deflandre 1937) Kirsch 1991
776	CON1	Scriniodinium campanula Gocht 1959
777	44	Senoniasphaera turonica (Prössl, 1990 ex Prössl, 1992) Pearce et al. 2011
778	62	Sentusidinium sp. Sarjeant and Stover 1978
779	6	Sepispinula? ambigua (Deflandre 1937) Masure in Fauconnier and Masure 2004
780	38	Spiniferites membranaceus (Rossignol 1964) Sarjeant 1970
781	25	Spiniferites ramosus granosus (Davey and Williams 1966) Lentin and Williams
782		1973
783	19	Spiniferites ramosus ramosus (Ehrenberg 1838) Mantell 1854
784	51	Stephodinium coronatum Deflandre 1936
785	56	Subtilisphaera pontis-mariae (Deflandre 1936) Lentin and Williams 1976
786	TUR3, 4/1	Surculosphaeridium? basifurcatum Yun Hyesu 1981
787	75	Surculosphaeridium belowii Yun Hyesu 1981
788	20	Surculosphaeridium longifurcatum (Firtion 1952) Davey et al. 1966
789	21	Tanyosphaeridium salpinx Norvick 1976
790	61	Tanyosphaeridium variecalamum Davey and Williams 1966
791	65	Tenua hystrix Eisenack 1958
792	30	Trichodinium castanea Deflandre 1935

- 793 57 *Wrevittia cassidata* (Eisenack and Cookson 1960) Helenes and Lucas-Clark 1997
- 794 TUR4/1 Wrevittia helicoidea (Eisenack and Cookson 1960) Helenes and Lucas-Clark 1997
- 795 TUR4/3 6/2 *Xenascus ceratioides* (Deflandre 1937) Lentin and Williams 1973
- 79666Xiphophoridium asteriforme Yun Hyesu 1981

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1141

Figure and Plate Captions

1142	Fig. 1. Location and Turonian palaeogeography of the Bohemian Cretaceous Basin study
1143	section. (A) Simplified palaeogeography of the European epicontinental sea showing the
1144	location of the Bch-1 well. Adapted from Mitchell et al. (2010). (B) Detail of the Bohemian
1145	Cretaceous Basin, location shown by grey rectangle in A. Main siliciclastic source areas and
1146	sub-basins are shown: NW, Lužice-Jizera sub-basin; SE, Orlice-Ždár sub-basin. TUR2 –
1147	CON1 indicate regressive limits of nearshore strata in genetic sequences. Modified from
1148	Uličný et al. (2014)
1149	Fig. 2. Biostratigraphy and carbon isotope chemostratigraphy of the Bch-1 well. Calcareous

1149 1150 nannofossil, dinocyst, inoceramid bivalve and ammonite datum levels are plotted against lithology and carbon isotope ratios of bulk organic matter ($\delta^{13}C_{org}$), and regional genetic 1151 1152 sequences. Dinocyst datum levels from this study; other data from Uličný et al. (2014). Upper Cretaceous (UC) calcareous nannofossil zones after Burnett et al. (1998). Carbon 1153 1154 isotope events (CIE) of Jarvis et al. (2006) are identified following Uličný et al. (2014). 1155 Minor revision of datum levels, genetic sequences and the placement of CIEs incorporates 1156 results from new unpublished stratigraphic work. Abbreviations used here and in text: FO = 1157 first occurrence; LO = last occurrence; FCO = first common occurrence; LCO = last common 1158 occurrence; RO = reappearance datum level. *Cci* = *Cremnoceramus crassus inconstans*.

Fig. 3. Dinocyst ranges and datum levels plotted against the macrofossil biostratigraphy and lithology of the Bch-1 well. 'Floods' are short-term abundance spikes; 'acmes' are mediumto long-term abundance maxima. The positions of major CIEs and the location of regional genetic sequences are shown. A complete list of species identified and their taxonomic assignments is provided in Appendix A. The ranges of biostratigraphically significant species (in bold) are emphasised by the vertical yellow bars; examples of these are illustrated in Plate

1165 I. Species displaying intervals of consistent common occurrence that may be of

1166 palaeoenvironmental or local biostratigraphic significance are highlighted by the vertical blue

1167 bars. See Fig. 2 for biostratigraphy and lithological key.

1168 Fig. 4. Relative abundance of common dinocyst species in the Bch-1 well. Values derived

1169 from counts of 300 identified dinocysts per sample. Note the different scales used to enhance

1170 stratigraphic trends. Red fills indicate peridinioid (P-cysts) and blue fills gonyaulacoid (G-

1171 cysts) species. Assemblages are dominated by *P. infusorioides* and *S. ramosus ramosus*, and

1172 in the Turonian – Coniacian boundary interval, S. ambigua. See Fig. 2 for biostratigraphy and

1173 lithological key.

1174 **Fig. 5.** Biostratigraphic and chemostratigraphic correlation of Bch-1 with English Chalk

1175 Turonian sections, showing key dinocyst datum levels. Inset map (modified from Rawson,

1176 1992) shows locations of English sections. Traditional Chalk macrofossil zones based on

1177 inoceramid bivalves, brachiopods and irregular echinoids; ammonite zones are also shown for

1178 the East Sussex composite section. Carbon isotope events in East Sussex based on

1179 interpretation of lithostratigraphic and biostratigraphic data (Mortimore, 1986; FitzPatrick,

1180 1995; Gale, 1996; Mortimore et al., 2001; Jarvis et al., 2006), and Pearce et al. (2009) for the

1181 Cenomanian – Turonian boundary interval.

Fig. 6. Revised dinocyst zonation for the Turonian Stage, key dinocyst biostratigraphic

1183 markers, and their correlation to major carbon isotope events. Symbols as in Fig. 5. CTBE =

1184 Cenomanian – Turonian Boundary Event.

Plate I Biostratigraphic marker species identified in the Turonian – Coniacian of the Bch-1
core.

1187 1 Pervosphaeridium truncatum (Davey 1969) Below 1982, sample 335.5 m

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- 1188 2 *Oligosphaeridium pulcherrimum (*Deflandre and Cookson 1955) Davey and Williams
 1189 1966, sample 33.5 m
- 1190 3 Palaeohystrichophora infusorioides Deflandre 1935, sample 351.5 m
- 1191 4 Cribroperidinium wilsonii (Yun Hyesu 1981) Poulsen 1996, sample 57.5 m
- 1192 5 Surculosphaeridium belowii Yun Hyesu 1981, sample 51.5 m
- 1193 6 Chatangiella ditissima (McIntyre 1975) Lentin and Williams 1976, sample 33.5 m
- 1194 7 *Kiokansium unituberculatum* (Tasch in Tasch et al. 1964) Stover & Evitt 1978, sample
 1195 197.5 m
- 1196 8 Stephodinium coronatum Deflandre 1936, sample 215.5 m
- 1197 9 *Circulodinium distinctum distinctum* (Deflandre and Cookson 1955) Jansonius, 1986,
 1198 sample 187.5 m
- 1199 10 Florentinia buspina (Davey and Verdier 1976) Duxbury 1980, sample 285.5 m
- 1200 11 Raetiaedinium truncigerum (Deflandre 1937) Kirsch 1991, sample 55.5 m
- 12 Subtilisphaera pontis-mariae (Deflandre 1936) Lentin and Williams 1976, sample
 1202 219.5 m
- 1203 13 Oligosphaeridium poculum Jain 1977, sample 43.5 m
- 1204 14 Senoniasphaera turonica (Prössl, 1990 ex Prössl, 1992) Pearce et al. 2011, sample
 1205 389.5 m
- 1206 15 Cyclonephelium compactum complex of Marshall and Batten 1988, sample 398.6 m



Fig. 1



Fig. 2



Fig. 3





Fig. 5















25 μm

















