Technical note: Accelerate coccolith size separation via repeated centrifugation

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Abstract:

Coccolithophore play a key role in the marine carbon cycle and ecosystem. The carbonate shells produced by coccolithophore, named as coccolith, could be well preserved in the marine sediment for million years and become an excellent archive for paleoclimate studies. The micro filtering and sinking-decanting method have been successfully designed for coccolith separation and promoted the development of geochemistry studies on coccolith, such as the stable isotopes and Sr/Ca ratio. However, these two methods are still not efficient enough for the sample-consuming methods. In this study, the trajectory of coccoliths movement during a centrifugation process was calculated in theory and carefully tested by separations in practice. We offer a matlab code to estimate the appropriate parameter, angular velocity at a fixed centrifugation duration, for separating certain coccolith size fractions from bulk sediment. This work could improve the efficiency of coccolith separation, especially for the finest size fraction and make it possible to carry the clumped isotope and radio carbon analysis on coccolith in sediment.

1. Introduction

Coccolithophores are a group of marine calcifying eukaryotic phytoplankton., whose calcite exoskeletons (i.e. coccolith) contribute significantly to the particulate inorganic carbon (PIC) export from the euphotic zone into the deep ocean (Young and Ziveri, 2000). Coccoliths preserved in marine sediment are also excellent archive for paleo-productivity reconstruction (Beaufort et al., 1997). The element ratio, Sr/Ca, in coccolith is correlated with the growth rate of calcite crystal (Stoll et al., 2002) thereby becoming a competitive candidate for coccolithophore growth rate which is an essential parameter in the paleo-CO_2 reconstruction by alkenone carbon isotope. However, the coccolith geochemical analyses are limited by the difficulty of separating coccolith from bulk sediment. To solve this problem, different separating methods have been proposed in the past a few decades (Paull and Thierstein, 1987; Stoll and Ziveri, 2002; Minoletti et al., 2008).

Most of them, in general, could be categorized into two groups: the first one is micro-filtering and the second is sinking-decanting technique. The micro-filtering method relies heavily on the specifications of micro filter membrane (such as 3μm, 5μm and 8μm pore size), which is highly effective in separation of the larger size coccoliths, but useless for coccolith smaller than 2μm. The sinking-decanting method, on the other hand, could offer more freedom in coccolith size by adjusting the sinking durations, thereby separating both small and large coccoliths. However, because of the slow sinking speed, a single separation of particles smaller than 2 μm may take more than 10 hours in settling. Moreover, about 6–8 times operations should be repeated, which means a full separation may takes at most one week. Hence, it is necessary to improve this method by reducing the time cost in coccolith separation.

Based on the Stokes sinking equation, the sinking rate of a certain particle increases with the increase of density difference between particle and liquid, decrease of the liquid viscosity and the increase of gravity. Changing the physical property of liquid often leads to the organic and toxic solvent which could lead to potential

52 contaminations for the further geochemistry analyses. A better way to accelerate

53 coccolith sinking speed is changing the gravity, or the acceleration speed of the

54 reference system, which can be easily achieved by centrifugation. One study has

55 mentioned the usage of centrifugation in coccolith separation, but only centrifugation

56 settings for a special case were provided (Hermoso et al., 2015). Here in this study, the

57 method of separation coccolith by centrifugation is introduced systemically. We first

58 calculate the trajectory of coccolith movement in a centrifugation processes and show

59 how to estimate the centrifugation parameters in different situations. After that, two

60 tests are performed to confirm the robustness of our calculations. Ultimately, a sample

61 containing coccoliths ranging from 2 μm to 12 μm is selected for a separation case in

62 practice.

63 **2. Trajectory of coccoliths during centrifugation**

64 The movement of coccolith in centrifugation is similar to that under the gravity.

65 Previously, we have calculated the separation ratio variation with time during the

66 settling (Zhang et al., 2018). All calculations in this study are with an assumption that

67 the coccolith is in the force balance all the time during both settlings and centrifugations

68 for a convenience of calculation. Here we offer a brief proof for this assumption based

69 on sphere particles (the sink speeds of sphere particles are ~30% high than that of

70 coccolith in same size) and do a quick review of derivation we did before.

71 Based on the Newton second law, the force balance of a sphere object during

72 sinking can be described by the following equation:

$$F = \frac{4}{3}\pi r^3 \rho_p g \ - \ \frac{4}{3}\pi r^3 \rho_l g - 6\pi\eta r v = \frac{4}{3}\pi r^3 \rho_p \frac{dv}{dt} \qquad (\text{Eq. 1.})$$

74 where F is the join force of particle, which is equal to zero in force balance, r is

75 the radium of sphere, ρ_p and ρ_l are the density of particle and liquid, respectively, η is

76 the velocity of liquid and v is the particle sinking speed, dv/dt is the particle acceleration

77 speed, which can be also marked as a. On the right side of the first equal mark, the first

78 term is the gravity force, the second term is buoyancy and the third term is the dragging

79 force from liquid. Transform Eq. 1, we can obtain the expression of accelerated speed

80 (a = F/m) of sphere as Eq. 2:

$$a = \frac{dv}{dt} = -\frac{9\eta}{2r^2}v + \frac{g}{\rho_{cal}}(\rho_p - \rho_l) \qquad (\text{Eq. 2.})$$

82 Given the initial value of sinking speed is equal with zero at the initial time (t = 0),

83 we can solve the differential equation Eq. 2 and obtain the variation of velocity with

84 time as following equation:

$$v = \frac{-e^{\left[-\frac{9\eta}{2r^2}t + \ln(-\frac{g}{\rho_{cal}}(\rho_p - \rho_l))\right]} + \frac{g}{\rho_{cal}}(\rho_p - \rho_l)}{\frac{9\eta}{2r^2}} \qquad (\text{Eq. 3.})$$

86 when the value of t is large enough, the first term of numerator in Eq. 3 is close to

87 zero, which represents the sinking velocity is close to the termination sinking velocity

88 described in Stocks equation (Eq. 4).

$$\lim_{t \to \infty} v = \frac{2(\rho_p - \rho_l)gr^2}{9\eta} \qquad (\text{Eq. 4.})$$

90 Given the particle as a 5 μm in radium calcite carbonate sphere with a density of

91 2.7 g cm^{-3} and the density of water is equal to 1.0 g cm^{-3}, when the t is equal to 10^{-7} s,

92 the first term of numerator is 3.7×10^{-44} m s^{-2} and small enough to be neglected compared

93 with the second term, which is 6.3 m s^{-2}. The time scale in coccolith separation is minute

94 for centrifugation and hour for settling, therebefore we suggest that it is reasonable to

95 assume the coccolith sinks with the 'terminal speed' from the very beginning.

96 The only difference between the terminal speed in centrifugation and under gravity

97 is the acceleration speed. If the g in Eq. 1 – 4 is adapted by a, which is the acceleration

98 speed of coccolith during centrifugation, these four equations above can also describe

99 the sphere movement in the centrifugation if we adapt the gravity to centripetal

100 acceleration (ca). Here we define a new parameter named as Sinking Parameter (*sp*):

101
$$sp = \frac{v}{g} \qquad (\text{Eq. 5.})$$

102 The physical meaning of sp is the influence of coccolith shape and liquid property

103 (density and viscosity) on sinking velocity without considering the effect of gravity (or

104 the acceleration rate of reference system). The sinking speed of coccolith in water

105 during a centrifugation (v') can be described as following:

106
$$v' = sp \times ca = sp \times \omega^2 \times (L + D) \qquad (\text{Eq. 6.})$$

107 where the ca is centripetal acceleration during centrifugation, ω is angular velocity

108 of centrifuge, the (L+D) is the rotation radium as illustrated in **Figure 1**. The L is a

109 fixed value for a certain type of centrifuge and the D depended on the position of

110 coccolith in the tube. Here we should notice two issues: the first one is that the rotation

111 radius is varying when coccolith is moving in the centrifuge tube, in other words, D is

112 always changing. This effect could be ignored when the L is much larger than D, but,

113 unfortunately, most of centrifuge employed in geochemistry laboratory is not large

114 enough. The second one is the angular velocity is dynamic during when the centrifuge

115 is accelerating and decreasing. To solve these two dynamic parameters, Eq.6 was

116 transformed into a form of differential equation as Eq. 7 for the convenience of

117 integration in the next step.

118
$$dt = \frac{dD}{v} = \frac{dD}{sp \times \omega^2 \times (L + D)} \qquad (\text{Eq. 7.})$$

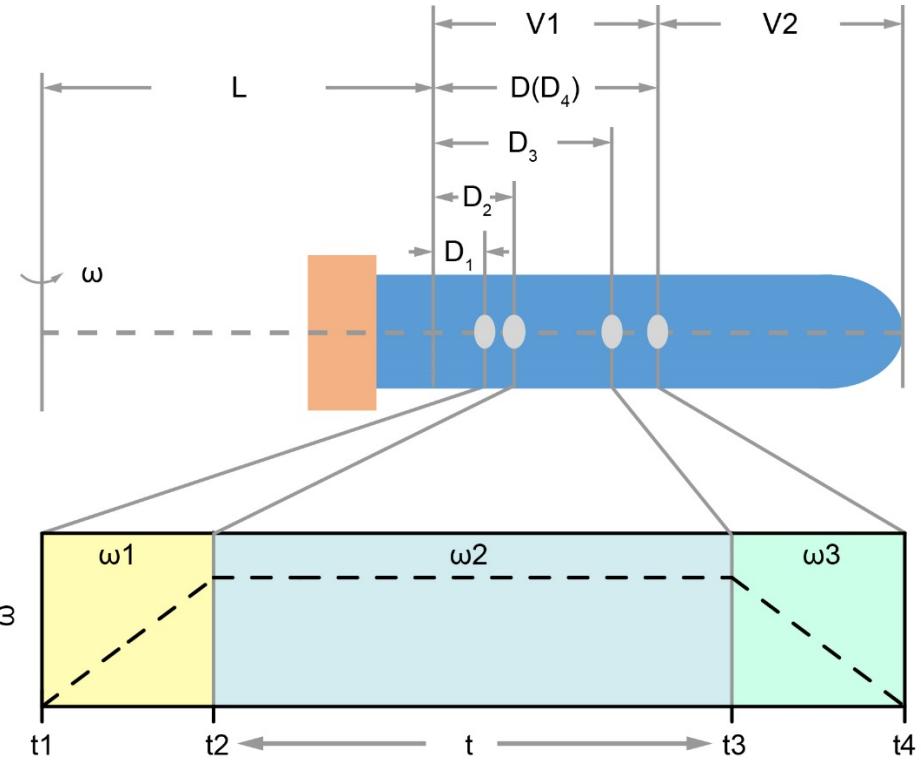
119 For all centrifugations there are three stages: the acceleration stage (t_1 to t_2 in

120 Figure 1), the constant angular velocity stage (t_2 to t_3 in Figure 1) and the deceleration

121 stage (t_3 to t_4 in Figure 1). The duration of acceleration stage and deceleration stage can

122 usually be controlled and the angular velocity is changing with a constant speed. For

123 those machines which the angular velocity dynamic ($\omega=f(t)$) is unknow we should

124 measure it manually.

125

126   **Figure 1.** The position of coccolith and the variation of ω in the three centrifuging stages: L

127   represents the minimum rotation radium, the V_1 and V_2 represent the volume of two parts; in the

128   first stage, the angular velocity increases from zero to ω1 (it could be linear or cubic, which

129   depends on the machine). Meanwhile the coccolith moves a distance of D_2-D_1.; similarly, the

130   coccolith moves a distance of D_3-D_2 in the second stage and it march a distance of D_4-D_3 in the

131   last stage.

132   After knowing the angular velocity curve, integrate the D over t in the Eq. 7 by

133   three steps from t_1 to t_4:

134
$$sp \times \int_{t_1}^{t_2} \omega_1{}^2 dt = ln^{(L+D_2)} - ln^{(L+D_1)} \qquad (\text{Eq. 8.})$$

135
$$sp \times \int_{t_2}^{t_3} \omega_2{}^2 dt = ln^{(L+D_3)} - ln^{(L+D_2)} \qquad (\text{Eq. 9.})$$

136
$$sp \times \int_{t_3}^{t_4} \omega_3{}^2 dt = ln^{(L+D_4)} - ln^{(L+D_3)} \qquad (\text{Eq. 10.})$$

137   Add the Eq. 8 – 10 together gives:

$$sp \times \left(\int_{t_1}^{t_2} \omega_1{}^2 \mathrm{dt} + \int_{t_2}^{t_3} \omega_2{}^2 \mathrm{dt} + \int_{t_3}^{t_4} \omega_3{}^2 \mathrm{dt} \right) = ln^{(L+D_4)} - ln^{(L+D_1)} \qquad (\text{Eq. 11.})$$

139   Set D_4 equal to D, which represents the maximum distance that a coccolith can

140   move in the upper suspension V_1. Now we can use the coccolith sinking property, *sp*,

141   and centrifugation settings to describe the coccolith position after centrifugation D_1:

$$D_1 = \frac{L + D}{e^{\left[sp \times \left(\int_{t_1}^{t_2} \omega_1{}^2 \mathrm{dt} + \int_{t_2}^{t_3} \omega_2{}^2 \mathrm{dt} + \int_{t_3}^{t_4} \omega_3{}^2 \mathrm{dt} \right) \right]}} - L \qquad (\text{Eq. 12.})$$

143   The meaning of D_1 is all coccolith with an initial position on the right side of D_1

144   in **Figure 1** will move to the right side of D_4 and then be kept in the suspension after

145   pumping, while the coccolith on the left side of D_1 will be removed by pumping.

146   In our previous publication (Zhang et al., 2018), we defined a parameter named as

147   separation ratio (R), which represents the percentage of coccolith removed in one

148   separation if we pump the upper V_1 volume suspension out of $(V_1 + V_2)$ suspension in

149   total.

$$R = \frac{V_1 \times \frac{D_1}{D}}{V_1 + V_2} \qquad (\text{Eq. 13})$$

151   Replacing the D_1 in Eq. 15 with Eq. 12 gives the separation ratio (R) as a function

152   of centrifugation settings:

$$R = \frac{V_1}{V_1 + V_2} \times \frac{1}{D} \times \left(\frac{L + D}{e^{\left[sp \times \left(\int_{t_1}^{t_2} \omega_1{}^2 \mathrm{dt} + \int_{t_2}^{t_3} \omega_2{}^2 \mathrm{dt} + \int_{t_3}^{t_4} \omega_3{}^2 \mathrm{dt} \right) \right]}} - L \right) \qquad (\text{Eq. 14})$$

154   The R can be employed in estimating the centrifugation parameters for separating

155   one type of coccoliths from another. For example, if we want to separate a group of

156   coccolith (marked as Coccolith$_A$, with sinking parameter sp$_A$) from another group of

157   coccolith (marked as Coccolith$_B$, with sinking parameter of sp$_B$ and sp$_A$ < sp$_B$), the R of

158   CoccolithB should be set as zero, which means all Coccolith$_B$ in the section V_1 have

159   sunk into V_2 after centrifugation and therefore all coccolith pumped out should be

160     $Coccolith_A$. To solve the angular velocity (ω_2) and centrifugation duration ($t = t_3-t_2$) in

161   Eq.14, we need to fix at least one of them. Usually the duration could be safely set as 1

162   min or 2 min, then solve the suitable angular velocity with known parameters V_1, V_2,

163   D and L. The matlab code for the parameter estimation is in attachment. After repeating

164   these 'centrifugation-pumping' routines several times, the $Coccolith_A$ could be fully

165   separated from $Coccolith_B$.

166 3.Test of the correctness of calculations

167 3.1 Experimental design

168   To test the robustness of our estimation in the last section, we performed two

169   groups of experiments comparing the observed with predicted separation ratio. Here we

170   select two different coccoliths, *F. profunda* and small *Gephyrocapsa*, with small size

171   and thereby slow sinking speed sampled from ODP 807 and IODP U1304, respectively.

172   Most of small *Gephyrocapsa* employed in this study are smaller than 3 μm with a

173   mixture of *G. muellerae* less than 10%. Two centrifuges from Anting Company, TDL–

174   40B and DL–5B, were selected to perform the tests. The angular velocity of DL-5B can

175   be set as linear increased or decreased with time in the acceleration or deceleration

176   stages, while the angular velocity of TDL–40B was measured manually by reading the

177   number on the instrument panel. The centrifugation duration can only be adapted by a

178   step of one minute on both of these two machines. The slowest angular velocities of

179   these two machines are 500 revolutions per minute (rpm). If we selected the water as

180   dispersion agent, most of the coccolith we used will sink to the tube bottom after two

181   minutes even with the slowest angular velocity. Hence, to slow down the coccolith

182   sinking speed in these tests, glycerol solution was employed in this equation test, which

183   can be dissolved with water in any proportion and washed away from carbonate calcite

184   particles conveniently. The density and viscosity data can be found in **Table 1**.

185   All calculations above are for the situation that particles sinking in the water or the

186   diluted solution, the physical property of which is close to water. However, in this case,

187 the property of glycerol is significant different with water. Here we define a new

188 parameter, τ, to transform the sinking speed in water to that in different liquid. The

189 physical meaning of τ is a ratio turning the sinking velocity in water (v) to the velocity

190 in any liquid with different density and viscosity (v'):

$$v' = v \times \tau \qquad \text{(Eq. 15)}$$

191

192 Based on the definition of Stokes equation, the term τ can be calculated as

193 following:

$$\tau = \frac{(\rho_p - \rho_l)}{(\rho_p - \rho_w)} \times \frac{\eta_w}{\eta_l} \qquad \text{(Eq. 16)}$$

194

195 where the ρ_p, ρ_l and ρ_w are density of particle, liquid (in this study is glycerol

196 solution) and water; the η_l and η_w are the viscosity of liquid and water.

197 Combine the Eq. 14–16 forming the separation ratio as a function of centrifugation

198 settings in different liquid:

$$R = \frac{V_1}{V_1 + V_2} \times \frac{1}{D} \times \left(\frac{L + D}{e^{\left[\frac{v}{g} \times \frac{(\rho_p - \rho_l)}{(\rho_p - \rho_w)} \times \frac{\eta_w}{\eta_l} \times \left(\int_{t_1}^{t_2} \omega_1{}^2 \mathrm{dt} + \int_{t_2}^{t_3} \omega_2{}^2 \mathrm{dt} + \int_{t_3}^{t_4} \omega_3{}^2 \mathrm{dt} \right) \right]}} - L \right) \quad \text{(Eq. 17)}$$

199

200 In this test, the calculated R by Eq. 17 will be compared with measured one. To

201 perform these tests, about 100 mg bulk sediments were scattered into 30 ml 0.5%

202 ammonia and, after that, particles larger than 20 μm particles were removed by mesh.

203 In this test, we should obtain suspensions with nearly monospecific coccolith. To

204 achieve it, in the test with *F. profunda*, coccoliths larger than 3 μm were removed by

205 the sinking method described in Zhang et al. (2018) and coccoliths larger than 5 μm

206 were removed by the same method in the test with small *Gephyrocapsa*. Briefly, the

207 suspension was (1) set in a 100 ml Reagent bottle sinking freely for a few hours, and

208 then (2) pumped out the upper 2cm. Repeat these two steps for 5–8 times until

209 coccoliths were purified. The sinking duration was 2 hours for *F. profunda* sample and

210 1.25 hours for small *Gephyrocapsa* sample, respectively.

211   Then 50 ml tubes with 45 ml coccolith suspensions were mounted in the centrifuge

212   and run with the settings shown in Table 1. After centrifugation, the upper 30 ml

213   supernatant was pumped out by pipette and then filtered onto 0.4 μm polycarbonate

214   member with a vacuum pump. The coccoliths on polycarbonate membrane were

215   resuspended into 20 ml diluted ammonia again and coccoliths number in the suspension

216   was measured with the same method described in our previous work (Zhang et al.,

217   2018). Finally, the separation ratio, R, was calculated by the coccolith number in the

218   upper 30 ml suspension divided by the total coccolith number. All the centrifuging

219   experiments were carried out in laboratory with temperature controlled around 20 (±1) °C

220   to avoid the variation of physical properties, especially the viscosity, with temperature.

221   **Table 1.** The settings of two tests: the density and viscosity of glycerol in 20°C, data from

222   Dorsey (1940); the parameters of centrifuge employed in this study: Fp and G60 represent the

223   experiment carried out with *F. profunda* in 70% glycerol and small *Gephyrocapsa* (<3 μ m) in

224   60% glycerol, respectively; L represents the minima rotation radius of centrifugation, which

225   represents the distance between the shaft and top of suspension as illustrated in Figure 1; A, B and

226   C are the terms on the left side of equal mark in Eq. 8–10.

	glycerol (%)	η (mPa s)	ρ (g cm^{-3})	τ	Centrifuge	L (cm)	A (s^{-1})	B (s^{-1})	C (s^{-1})
Fp	70%	22.5	1.16	0.040	TDL40B	6.2	1.060×10^6	$9.867 \times 10^4 \times t$	1.937×10^6
G60	60%	10.8	1.14	0.084	DL-5B	8.37	7.457×10^5	$9.867 \times 10^4 \times t$	2.193×10^6

227   **3.2 Result of experiments**

228   In the test, 30 ml suspension was pumped out from 45 ml suspension leading to

229   the initial R should be 60%. However, the intercept of calculated R is smaller than 60%

230   as the gravity settling in Zhang et al. (2018), because the time in the x-axis of Figure 2

231   is the period in which angular velocity remains constant. In other words, even the time

232   is set as zero, the centrifuge will still do the acceleration and deceleration processes and

233   coccolith will move toward the bottom. The results of observed R (dots in **Figure 2**)

234   are close to the theoretical values (dash lines in **Figure 2**), though a few measured

results are lower than prediction. We suggested that this difference may be caused by coccolith loss during harvesting of the coccolith from glycerol solutions into ammonia solution.

So far, we have obtained the coccolith movement equation in the centrifugation and prove its correctness. In the next section, a case of coccolith separation by centrifuging method will be carried out giving an example of separation.

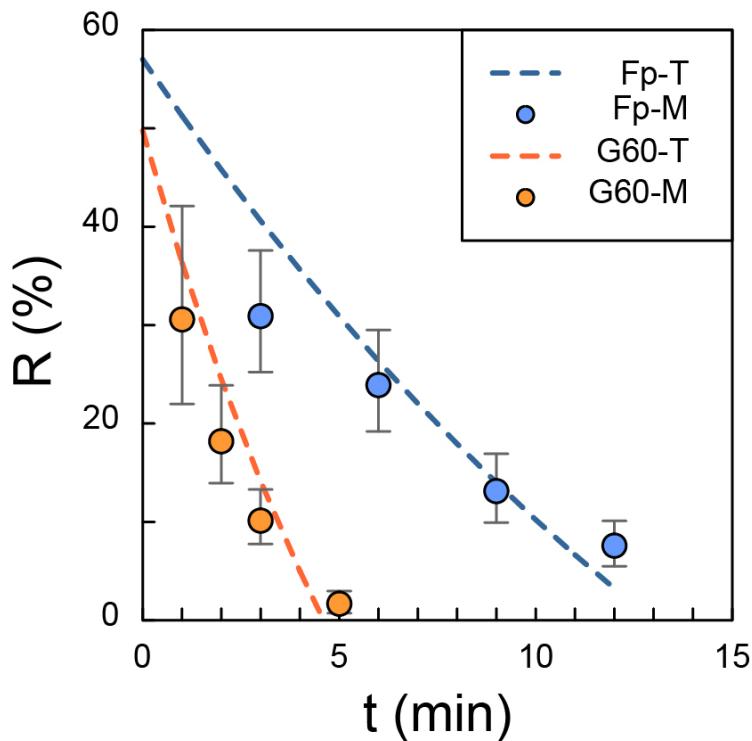



Figure 2. The comparison of theoretical and measured separation ratio (R): the dots represent the measured values and dish lines are theoretical calculations. The error bars represent 95% error based on the assumption that the error of counting coccolith follows the Poisson distribution. The orange dots represent the measured R in small *Gephyrocapsa* with 60% glycerol test (G60-M) and the blue ones represent the measured R in *F. profunda* with 70% glycerol test (Fp-M). The orange dashed line is the theoretical values for small *Gephyrocapsa* with 60% glycerol test (G60-T) and the blue one is the theoretical values for *F. profunda* with 70% glycerol test (Fp-T). The raw pictures for coccolith counting were shown in **Figure S1** and **S2**.

4. Separation of coccoliths in practice

4.1 Separation steps

The aim of this section is using the centrifugation method to separate a sample in practice. A sample form ODP 982B (56X Section 5 5-9cm) dated around mid-Miocene (nannofossil zoon NN4) was selected in this test. The coccolithophore *Reticulofenestra* spp. dominated in the assemblage, with long axis length ranging from 2 μm to more than 12 μm, offering an ideal sample to test the coccolith separation method. *Calcidiscus* spp. (4–10 μm), *Helicosphaera* spp. (5–10 μm) and *Coccolithus* spp. (6–8 μm) were also found in this sample, which contributed less than 10% of all coccoliths together. The preservation of fossil was moderate with many coccolith fragments but no evidence of dissolution in the raw sample. The detailed operations are as following:

Step 1: weigh about 40 mg bulk sediment, scatter with 45ml 0.5% ammonia solution and transfer the suspension into a 50 ml centrifuging tube;

Step 2: Calculate the centrifugation parameters (angular velocity and duration). Here we did not measure coccolith sinking velocities, but employ the length-velocity relationship in the previous study directly: sinking rate at $25°C= 0.0982×length^2$ (Zhang et al., 2018). Based on this length-velocity equation and the centrifuge properties listed in **Table 1**, we estimated that the angular velocity and duration for separating coccolith with a length of 2 μm, 3 μm, 5μm, 8 μm and 10 μm should be 1850 rpm 2 min, 2250 rpm 1 min, 1400 rpm 1min, 1000 rpm 1min and 600 rpm 1min, respectively. The Matlab code for calculating the angular velocity at fixed centrifugation duration (1 or 2 minutes) are in the supplementary.

Step 3: Mount the tube into the centrifuge and balance weight, set the angular velocity as 1850 rpm and the duration as 2 minutes and start the machine;

Step 4: Pump out the upper 30 ml suspensions and remove them into a beaker (500 ml or larger beaker, depends how many times repeating this step) and drop about 100 μl onto a glass cover. Dry the suspension on glass cover and mount the cover on slider. The details in this step follow Bordiga et al. (2015);

Step 5: Repeat Step 2–5 with different centrifugation parameters listed in Table 2;

279   **Step 6:** Take pictures of coccoliths in each slider on microscope and measure the

280   coccolith size on computer with the method described by Fuertes et al. (2014).

281

282   **Table 2**. Centrifugation parameters in the Miocene coccolith separations

	<2 μm	2–3 μm	3–5 μm	5–8 μm	8–10μm
Angular velocity (ω_2, rpm)	1850	2250	1400	1000	600
Duration ($t = t_3-t_2$, s)	120	60	60	60	60

283   **4.2 Coccolith length in each fraction**

284   The coccolith size distribution harvested from different centrifugation settings are

285   shown in **Figure 3** (the coccolith size was measured in circular polarizing microscope

286   and coccoliths under cross polarizing microscope were shown in **Figure S3-S9** for

287   species identification). The results show that the separated coccolith size increased with

288   the decrease of angular velocity and the differences of mean coccolith lengths are

289   significant between each size fractions. However, we should also notice that there is

290   still overlap of coccolith sizes between two neighbouring fractions. With the

291   centrifugation parameters set as 2250 rpm and 2 min, the coccoliths harvested have

292   long axis lengths around 2–4 μm and when the centrifugation parameters was varied to

293   1400 rpm and 1 min, the coccolith long axis size ranges from 3 μm to 7 μm, which

294   means coccoliths with a length between 3–4 μm appear in two fractions. Such situations

295   may also happen in both settling and micro filtering methods, but the range of overlap

296   seems to be larger for the centrifugation method compared with the size fractions

297   harvested by other methods.

298

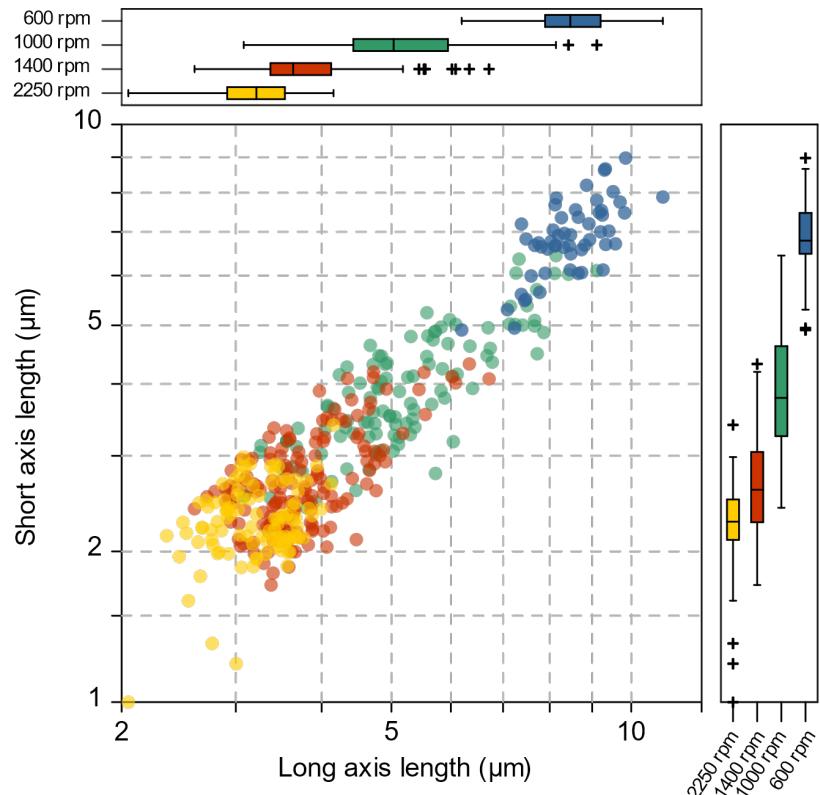



Figure 3. The coccolith size in different fraction after centrifuging separation: the yellow, red, green and blue dots represent 2250 rpm-2min, 1400 rpm-1min, 1000 rpm-1min and 600 rpm-1min, respectively.

302

4.3 Troubleshooting

The first potential reason leading to overlap may be the repeating times are not enough. This could be the main problem for settling under gravity, since the time costs for separation under gravity is much larger than the centrifugation method. Bolton et al. (2012) suggested that 4–6 times separations are enough for fossil extraction and in our separations, we repeated more than 8 times for a certain centrifugation setting. Considering these facts, we suggest that this overlapping was not caused by the separation times.

Another reason could be that larger coccoliths, which are supposed to sink into the lower suspension, are pumped out after centrifugation. When the upper suspension was

313   pumped out, the pumping speed could be too fast drawing up larger coccolith from the

314   lower suspension. This problem could be solved by reducing the pumping speed.

315   Hoverer, in practice, the pumping speed of pipette is difficult to control. Here we

316   recommend to modify the tips of pipette as following steps: (1) suck a drop of glue into

317   the top of pipette tips (the Norland optical adhesive 74 was employed in this study); (2)

318   solidify the glue with ultraviolet ray to seal the top of tips; (3) dill holes above the glue

319   horizontally. After this modification, the suspension will go into tips horizontally

320   instead of vertically (**Figure 4a**) to avoid mixing larger coccoliths with smaller ones.

321   The size overlapping could also be caused by the centrifugation tube not remaining

322   perfectly horizontal during centrifugation. In our calculations, the tubes are assumed to

323   be perfectly horizontal during all centrifugation processes and, thereby it was assumed

324   that there should be no collisions between coccoliths and tube wall nor among

325   coccoliths. However, in practice, the tubes in centrifuge are not always horizontal and

326   even a few degrees slope of the tubes can lead some coccoliths to knock and stick on

327   the tube wall forming a significant coccolith layer on one side of tube wall as illustrated

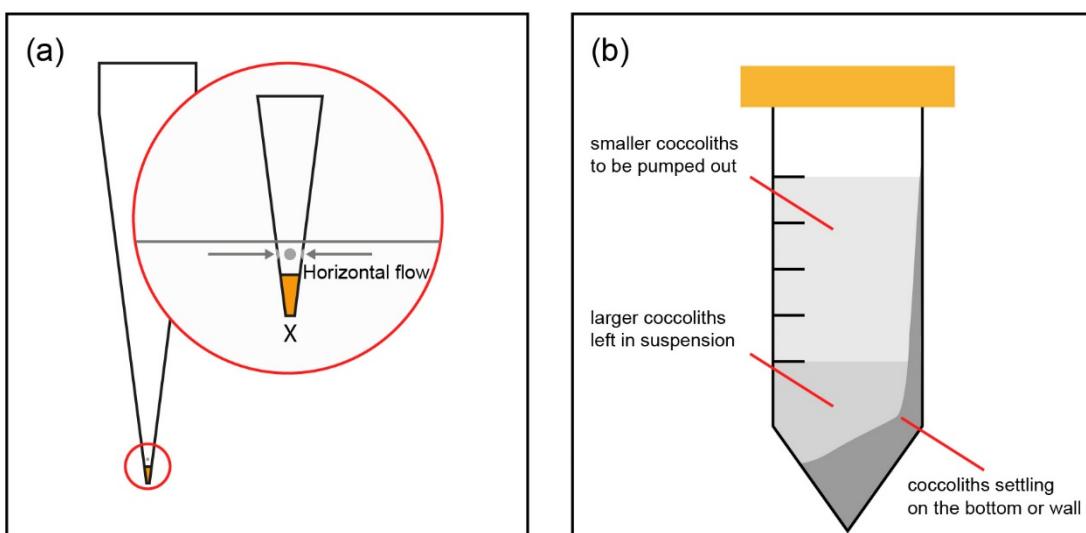
328   in **Figure 4b**. These coccoliths on tube wall will be pumped out after centrifugation

329   causing the coccolith length overlapping among two fractions. To avoid this problem,

330   before the step of pumping out suspension, we should observe the tube carefully. If a

331   coccolith layer can be found on the tube wall, the pipette tip should be placed on the

332   opposite of the coccolith layer to reduce the size overlapping.

333

334   **Figure 4.** Two methods to reduce the coccolith size overlapping. (a) Adaption of pipette tip:

335   the orange part on tip represents sealed by solidified glue and the gray parts mean that small holes

336   should be drilled allowing the suspension flowing in horizontally; (b) Choose a property pumping

337   position to avoid extracting the coccolith on tube wall: the lightest gray part in the tube represents

338   the suspension in which the smaller coccolith floats, most of the larger coccoliths are in the lower

339   part of the suspension and the tube bottom.

5. Summary

341   In this study, we described the method of separating coccolith from bulk sediment

342   by centrifuge. The rotation speed for separating coccoliths within a certain range of

343   length could be solved after measuring the rotations radium (property of centrifuge)

344   and fixing the centrifugation duration.

345   The centrifugation method is not perfect accurate and could still mix different

346   species of coccolith as other traditional separating methods. The size overlapping of

347   this method could be reduced by adapting the pipette tips and avoiding pumping the

348   coccolith on tube well out. However, this method is more efficient in separating the

349   finest particle (smaller than 3 μm) out of bulk sediment, which is always the time-

350   consuming step in micro-filtering and sinking method. Thereby, this method can be

351   widely used in the sample preparation for analyses needing a large amount of material,

352   such as coccolith clumped isotope and radioactive carbon isotope measurement.

353   Moreover, the centrifugation method can be combined with other separation steps, for

354   example using the centrifugation method to remove the finest particles followed by

355   micro filtering with different size of membrane. This method could largely reduce the

356   time cost in sample preparation for coccolith geochemistry analyses and have the

357   potential for wide use in the future.

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Author contributions.

405   This study was conceived by H.Z. and C.L. Measurements and calculations were
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