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1 **THE GEOLOGICAL SIGNIFICANCE OF COSMOGENIC NUCLIDES**  
2 **IN LARGE LOWLAND RIVER BASINS**

3

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11

12 **ABSTRACT**

13 Measured from a bag of sand taken in large rivers, the concentration of cosmogenic  
14 nuclides such as *in situ*-produced <sup>10</sup>Be, <sup>26</sup>Al, and <sup>14</sup>C can be used to constrain the  
15 mean sediment flux of the headwaters and assess the duration of sediment storage  
16 from source to sink. We revisit these principles, with examples from the Amazon and  
17 Ganga basins.

18 We identify two end member cases controlling the concentration of cosmogenic  
19 nuclides in lowland river sediment: 1) if the time scale of floodplain sediment storage  
20 is short compared to the half-life of the nuclide, *in situ* cosmogenic nuclide  
21 concentrations are not significantly altered in lowland basins. In this case the  
22 concentration of e.g. *in situ*-produced <sup>10</sup>Be in the sediment taken in the lowland

23 basin equals in most cases that exported from the sediment source area, but the  
24 variability in nuclide concentrations between headwater streams is significantly  
25 averaged-out. Thus to convert the measured river sediments' *in situ* cosmogenic  
26 nuclide concentration into a catchment-wide denudation rate, production rates are  
27 scaled to include those of the sediment-producing mountainous areas only, rather  
28 than the entire catchment area. Nuclide production in the lowlands, where no  
29 sediment is being produced, is hence excluded. This correction is termed "floodplain  
30 correction". 2) If sediment buried for periods of the order of the nuclides' half-life is  
31 episodically re-entrained into the active river, paired nuclides, for example the ratio  
32 of  $^{26}\text{Al}/^{10}\text{Be}$ , through the differential decay of these isotopes, constrain the the  
33 storage duration and re-mixing of floodplain sediment.

34 As *in-situ* cosmogenic nuclides measure denudation rates over longer time scales,  
35 typically integrating between  $10^3$  and  $10^5$  yrs, their combination with modern  
36 estimates of sediment fluxes from river load gauging offers a rich potential of  
37 deciphering the controls of Earth surface fluxes across large basins. From this  
38 combination, we can assess how river systems react to external perturbations such  
39 as climate change or human interference, and how such signals are transmitted  
40 through the alluvial reaches of lowland basins. The most important results in both  
41 basins are that i) lowland Amazon and Ganga nuclide concentrations indeed broadly  
42 reflect the spatial average of Andean and Himalayan denudation, respectively, ii) the  
43 thus calculated sediment fluxes are within uncertainty of modern fluxes from  
44 sediment gauging, indicating complete sediment delivery to the sea without net loss  
45 into the basin, and iii) only the Amazon basin, given its size, contains a substantial  
46 fraction of sediment buried temporarily in the Quaternary.

47 Finally, we provide a short introduction to the use of meteoric cosmogenic nuclide  
48  $^{10}\text{Be}$  and its ratio to stable  $^9\text{Be}$ , released by rock weathering, in lowlands. In the  
49 Amazon basin we show that  $^{10}\text{Be}$  concentrations in fine-grained suspended loads,  
50 depth-integrated over the water column using Al/Si ratios, provide erosion rates  
51 whereas the  $^{10}\text{Be}/^9\text{Be}$  ratio on Be adsorbed to particles provides the denudation  
52 rate. Both are similar to denudation rates from *in situ*  $^{10}\text{Be}$ . We show that this  
53 system responds more sensitively to sediment storage than the *in situ* system, and  
54 both accumulation and decay of meteoric concentrations may thus be used to  
55 determine sediment residence time.

56

57 **Keywords:** Fluvial geomorphology, floodplains, denudation, erosion, *in situ* and  
58 meteoric cosmogenic nuclides, sediment budgets, Amazon, Ganga, Andes, Himalayas

59

## 60 **1. INTRODUCTION**

61 The role of lowland river basins in modifying source-specific sediment properties, for  
62 cycling of dissolved and particulate matter, and for setting the sensitivity of  
63 sediment transport to climate change and shifts in land use, has long been  
64 recognized. In the past years, cosmogenic nuclide methods have been developed for  
65 quantifying the mass transfer involved. Therefore, it is timely to highlight the recent  
66 methodological advances. These methods encompass *in situ* cosmogenic nuclides  
67 (produced within a mineral at the Earth's surface) or meteoric cosmogenic nuclides  
68 (produced within the atmosphere and bound to fine-grained material after  
69 precipitation). Cosmogenic nuclide methods are now well established as an integral

70 tool of Earth surface science. They serve to provide ages of Quaternary landforms  
71 and mass fluxes of eroding landscapes (see reviews of these applications by Gosse  
72 and Phillips (2001); Bierman and Nichols (2004); von Blanckenburg (2005); Granger  
73 and Riebe (2007); Ivy-Ochs and Kober (2008); Dunai (2010); Graly et al. (2010);  
74 Willenbring and von Blanckenburg (2010); Granger et al. (2013); von Blanckenburg  
75 and Willenbring (2014)).

76

77 Denudation rates from cosmogenic nuclides provide the sum of erosion and  
78 weathering in source areas that can be converted into sediment fluxes integrating  
79 over several kyr by multiplying with the sediment-supplying area. When compared  
80 with sediment gauging data, integrating over the past few decades, we can use the  
81 cosmogenic nuclide data to evaluate the temporal stability of sediment transport  
82 and thus source-sink connectivity. As recently reviewed by Romans et al. (2015) the  
83 stratigraphic record of source to sink sediment transfer records signals of external  
84 environmental forcings as well as internal sedimentary dynamics. To make use of  
85 this record it is of high importance to understand how rivers transmit signals of  
86 climate cycles and to estimate the fluvial preservation potential (Castelltort and Van  
87 Den Driessche, 2003; Phillips, 2003; Allen, 2008; Jerolmack and Paola, 2010;  
88 Armitage et al., 2011; Simpson and Castelltort, 2012; Armitage et al., 2013; Godard  
89 et al., 2013; Latrubesse, 2015).

90

91 But do lowland river cosmogenic nuclide concentrations, that during temporary  
92 storage in sedimentary deposits may both increase (by further irradiation) or  
93 decrease (by radioactive decay) indeed provide a new tool to quantitatively assess

94 these linkages? We show in this review how we can study the evolution of  
95 cosmogenic nuclide concentrations as sediment leaves the mountainous production  
96 zone and crosses large floodplain systems, by using both data and modeling  
97 approaches. We mainly review the principles of calculating denudation rates from *in*  
98 *situ* cosmogenic  $^{10}\text{Be}$  in lowland settings, as compared to steady state equations  
99 used to calculate denudation rates in the high-relief sediment production zone. We  
100 illustrate the application of these principles by summarizing existing denudation rate  
101 data for the Amazon and Ganga basins and their source areas, namely the Andes and  
102 the Nepalese Himalayas. In these areas an exceptional wealth of cosmogenic nuclide  
103 and modern river flux data has been generated in past years. We include a short  
104 review of mass balance models of cosmogenic nuclides in floodplains (Wittmann and  
105 von Blanckenburg, 2009; Lauer and Willenbring, 2010).

106

107 We also provide a short account of the state-of-the-art of the meteoric  $^{10}\text{Be}$  system  
108 and its ratio to stable  $^9\text{Be}$ , released by weathering, for lowland settings. We finally  
109 compare *in situ*- with meteoric-cosmogenic nuclide derived denudation rates in the  
110 Amazon basin and discuss whether these two methods are expected to agree with  
111 each other, or whether they provide divergent but complementary information.

112

### 113 ***1.1 Dates and rates: principles of cosmogenic nuclides as Earth surface*** 114 ***chronometers***

115 Cosmogenic nuclides are used to determine ***dates*** (exposure or burial ages) and  
116 ***rates*** (erosion, the removal of solids; and denudation, the sum of total solid and  
117 dissolved material removal). A fundamental prerequisite is that none of these

118 nuclides existed before Earth surface material was exposed to cosmic radiation, or  
119 fallout of meteoric nuclides. This is the case for most of the radioactive nuclides, of  
120 which the half-life is much shorter than the age of most geologic materials. Three  
121 types of chronometric information can be obtained in Earth surface materials.

122 *i) Exposure ages* are determined through measuring the buildup of nuclides. The  
123 concentration is a function of the nuclide production or fallout rate and the duration  
124 of exposure.

125 *ii) Burial ages* are determined when a sample is buried deeply enough to stop new  
126 nuclide production (like in thick floodplain sediment). Then the measurement of a  
127 shorter-lived radionuclide (e.g.,  $^{14}\text{C}$ ,  $^{26}\text{Al}$ ) relative to a longer-lived (e.g.,  $^{10}\text{Be}$ ) or stable  
128 (e.g.,  $^{21}\text{Ne}$ ) nuclide discloses the timing of burial from the radioactive decay of these  
129 coupled nuclides.

130 *iii) Denudation rates*, in contrast, can be determined if the build-up of cosmogenic  
131 nuclides at the surface is limited by the removal of mass from the surface that is  
132 continuously eroded, or dissolved, provided that such weathering occurs within the  
133 cosmic ray adsorption zone. In this case the concentration measured in a surface  
134 sample scales inversely with the rate of mass removal at the surface. If buried after  
135 erosion and shielded in sediment, decay-corrected concentrations provide a *paleo-*  
136 *denudation rate*, i.e. the rate at which the source area was eroding at the time of  
137 deposition. If the mass loss of elements from the weathering zone relative to parent  
138 bedrock by mineral dissolution can be constrained, both physical **erosion** and  
139 chemical **weathering** rates can be obtained.

140

141 Two varieties of nuclides have commonly been used in Earth surface research (von  
142 Blanckenburg and Willenbring (2014); see Fig. 1). Atmospherically produced  
143 nuclides, such as meteoric  $^{10}\text{Be}$ , were utilized for geomorphologic applications  
144 already in the late 1980's and early 1990's (Pavich et al., 1984; Brown et al., 1985;  
145 Brown, 1987; You et al., 1988; Brown et al., 1992; Monaghan et al., 1992; McKean et  
146 al., 1993). However, issues such as partial retentivity and the lack of accurate flux  
147 estimates impaired its routine application (Willenbring and von Blanckenburg, 2010).  
148 The meteoric  $^{10}\text{Be}$  nuclide was initially sidelined by the success of the *in situ* variety,  
149 produced in the uppermost layer of the Earth's surface (Lal, 1991; Gosse and Phillips,  
150 2001; Dunai, 2010).

151

152 The first principle explained in this review is the determination of denudation rates  
153 from the measurement of *in situ* cosmogenic nuclides in quartz from river sediment.  
154 It is less known that meteoric  $^{10}\text{Be}$  was first used to determine catchment-wide  
155 erosion (Brown, 1987; Brown et al., 1988; You et al., 1988). The first studies  
156 suggesting the *in situ*-produced nuclide variety as tool to determine catchment-wide  
157 denudation rates followed D. Lals (Lal, 1991) comprehensive framework and were  
158 published in the mid- to late 1990's (Brown et al., 1995; Bierman and Steig, 1996;  
159 Granger et al., 1996). The first systematic river basin scale studies appeared in 2001  
160 (Kirchner et al., 2001; Schaller et al., 2001). *In situ*-produced  $^{10}\text{Be}$ , measured in  
161 quartz (Fig. 1) became the most commonly used nuclide. The reasons for this success  
162 are that its half-life of 1.39 Myr (Chmeleff et al., 2010; Korschinek et al., 2010)  
163 ensures zero initial concentration for most materials first exposed at the Earth's  
164 surface, and it is in the range of many dating applications. Further, analytical efforts



165 and costs involved are manageable (Dunai, 2010), quartz is an abundant mineral that  
166 largely resists weathering (Kohl and Nishiizumi, 1992), and knowledge of production  
167 rates and their scaling for latitude and atmospheric scaling is constantly increasing  
168 (Lal, 1991; Masarik and Reedy, 1995; Stone, 2000; Balco et al., 2008; Lifton et al.,  
169 2014; Borchers et al., 2016; Phillips et al., 2016). Thus the nuclide can be used with  
170 high robustness and confidence for erosion studies in “sediment production”  
171 settings where the relief is such that sediment is readily exported from the  
172 catchment (see Fig. 1) (Bierman and Nichols, 2004; von Blanckenburg, 2005; Granger  
173 and Riebe, 2007).

174

175 The second principle explained in this review is the behavior of cosmogenic nuclides  
176 in sediment that has been stored in Quaternary deposits. Storage in sedimentary  
177 deposits may both increase (by further irradiation) and decrease (by radioactive  
178 decay), their cosmogenic nuclide concentrations thus depending on paleo-erosion  
179 rate, depositional age, and depositional depth (Granger and Muzikar, 2001; Schaller  
180 et al., 2004). Paleo-erosion rates were recently summarized in review articles (e.g.  
181 Granger and Schaller (2014); Dosseto and Schaller (2016)). Pairing cosmogenic  $^{10}\text{Be}$   
182 with stable noble gases such as  $^{21}\text{Ne}$  measured in quartz (Niedermann, 2000;  
183 Niedermann, 2002), or the longer-lived  $^{26}\text{Al}$  (half-life of 0.72 Myr; Samworth et al.  
184 (1972)), resulting in  $^{21}\text{Ne}/^{10}\text{Be}$  and  $^{26}\text{Al}/^{10}\text{Be}$  ratios (Granger and Muzikar, 2001;  
185 Granger, 2006; Balco and Shuster, 2009), is common to e.g. extend the time frame of  
186 investigation required for these applications (Granger and Muzikar, 2001; Schaller et  
187 al., 2002; Matmon et al., 2003; Partridge et al., 2003; Schaller et al., 2004; Balco and  
188 Stone, 2005; Stock et al., 2005; Haeuselmann et al., 2007; Kong et al., 2009; Hu et al.,

189 2011; Matmon et al., 2011; Wittmann et al., 2011b; Charreau et al., 2012; Bekaddour  
190 et al., 2014; McPhillips et al., 2016). Recently, the *in situ*-cosmogenic nuclide  $^{14}\text{C}$   
191 which has a much shorter half-life of 5730 yrs (Libby, 1955) was added to the family  
192 of cosmogenic isotopes measured in quartz (Lifton et al., 2001). When combined,  
193  $^{10}\text{Be}/^{14}\text{C}$  ratios can be used to constrain more recent Earth surface processes as this  
194 ratio is sensitive to kyr time scale variations in erosion rate, and sediment storage  
195 over kyrs (Hippe et al., 2012).

196

197 Studies exploring the transport and deposition zones of a river system (Fig. 1) with  
198 cosmogenic nuclides are still scarce. Granger et al. (1996) cautioned for the effect  
199 sediment storage can have on cosmogenic nuclide-derived denudation rates. These  
200 authors noted that *“during storage and transport, sediment can accumulate*  
201 *additional cosmogenic nuclides, or it can be shielded from cosmic ray exposure (...).*  
202 *However, the net effect on cosmogenic nuclide concentrations will be small as long as*  
203 *the mean residence time of storage and transport is much shorter than the erosional*  
204 *timescale”* (Granger et al., 1996). Wittmann and von Blanckenburg (2009) first  
205 explored this important prediction, by mapping-out the behavior and limitations of  
206 *in situ*-produced cosmogenic nuclides in global large lowland basins. In lowland  
207 settings, the mean residence time of storage is a function of the sediment  
208 accommodation space within the floodplain, and the rivers migration rate. We  
209 proceed to explain these systematics.

210

211

212 **2. PRINCIPLES OF *IN SITU* COSMOGENIC NUCLIDES IN THE SEDIMENT SOURCE**

213 **AREAS AND IN LOWLAND RIVER BASINS**

214 **2.1 Steady state denudation in the sediment production zone**

215 Within the sediment production zone of a basin (Fig. 1), sediment is generated by  
216 weathering and erosion processes on the hillslope, leaves the zone by riverine  
217 transport, and moves down the sediment cascade (Burt and Allison, 2010; Hinderer,  
218 2012). The concentration of an *in situ*-cosmogenic nuclide in the sediment leaving  
219 this zone is termed  $C_0$  (in at/g<sub>(Qtz)</sub>). A steady state  $C_0$  is attained if export by erosion  
220 equals the rate of its production (Lal, 1991; von Blanckenburg, 2005). In that case,  
221  $D_{insitu}$ , the denudation rate (in i.e. cm/yr) derived from *in situ*-produced <sup>10</sup>Be includes  
222 all mass losses from the surface by both physical erosion and chemical weathering  
223 reactions, where weathering loss is only accounted for if it occurs within the  
224 absorption depth scale (Lal, 1991; Riebe and Granger, 2013). The original equation  
225 derived to calculate steady state  $D_{insitu}$  (Lal, 1991) is as follows:

226

227 
$$D_{insitu} = \left( \frac{P_0}{C_0} - \lambda \right) \frac{\Lambda}{\rho} \quad (\text{Eq. 1})$$

228

229 However, since then the importance of additional nuclide production by slow and  
230 fast muons has been realized, such that the dependence of  $C_0$  on  $D_{insitu}$  is more  
231 complex:

232

233 
$$C_0 = \frac{\bar{P}_n}{\frac{\rho \times D_{insitu}}{\Lambda n} + \lambda} + \frac{\bar{P}_{\mu s}}{\frac{\rho \times D_{insitu}}{\Lambda \mu s} + \lambda} + \frac{\bar{P}_{\mu f}}{\frac{\rho \times D_{insitu}}{\Lambda \mu f} + \lambda} \quad (\text{Eq. 2})$$

234

235 where  $\bar{P}_n$ ,  $\bar{P}_{\mu s}$ ,  $\bar{P}_{\mu f}$  are basin-wide  $^{10}\text{Be}$  production rates (at/g<sub>(Qtz)</sub>/yr) for neutrons,  
236 slow muons, and fast muons, respectively. The terms  $\Lambda_n$ ,  $\Lambda_{\mu s}$ ,  $\Lambda_{\mu f}$  are effective  
237 attenuation lengths (g/cm<sup>2</sup>) of neutrons, slow muons, and fast muons, respectively  
238 (Braucher et al., 2011, see supplement A; note that these terms are often  
239 deconvolved into further subfractions denoting different exponential absorption  
240 paths). The density of removed rock or soil is given by  $\rho$  (g/cm<sup>3</sup>). The term  $\rho/\Lambda_n$  can  
241 be replaced by  $z^*$ , the absorption depth scale (cm), which is the distance over which  
242 the cosmic-ray flux decreases by a factor of  $1/e$ , or 63%. This vertical distance,  
243 divided by the denudation rate, gives the integration time scale of the method (von  
244 Blanckenburg, 2005). The time scale is a function of the denudation rate, and is the  
245 time required to erode a rock layer of a thickness of ca. 60 cm in rocks or ca. 100 cm  
246 in soils, representing the absorption depth scale of the cosmic rays producing these  
247 nuclides (Lal, 1991; von Blanckenburg, 2005). Note that for basin-scale studies,  
248 where denudation rates are not uniform, a requirement to solve equation (1) for  
249  $D_{\text{insitu}}$  is that the integration time scale is smaller than the time scale for radioactive  
250 decay, or  $D/z^* \gg \lambda$ . Therefore, the  $^{10}\text{Be}$  method is applicable in those settings where  
251  $D_{\text{insitu}} > 0.3$  mm/kyr, while for  $^{14}\text{C}$   $D_{\text{insitu}} > 80$ mm/kyr (von Blanckenburg, 2005). If  $\rho$  is  
252 omitted from equation 1, a denudational mass flux (g/cm<sup>2</sup>/yr) results. Denudation  
253 rates are typically reported in mm/yr or t/km<sup>2</sup>/yr.

254

## 255 ***2.2 Modifications of in situ nuclide concentrations in lowlands***

256 Equations 1 and 2 are only valid for a set of assumptions, the prime one being that  
257 steady state has been attained between cosmogenic nuclide production and their

258 removal by erosion. In the depositional zone of a river basin, however, there is no  
259 continuous sediment removal from the surface layer that equals its production;  
260 rather, sediment in the floodplain is locally and laterally eroded such as bank failure  
261 and avulsions, and sediment storage may lead to additional production or decay of  
262 nuclides. Thus sediment in the active channel comprises a mixture of fresh upstream  
263 sediment (nuclide concentration  $C_0$ ) with reworked floodplain material ( $C_{dep}$ ). The  
264 concentration of the active channel mixture  $C_{mix}$  depends on these concentrations  
265 and the relative sediment fluxes. In keeping with the terminology introduced by  
266 Wittmann and von Blanckenburg (2009), we define two floodplain end members  
267 (Fig. 2) that lead to characteristic cosmogenic nuclide patterns once the previously  
268 stored floodplain deposit is mixed with fresh river sediment from the source area.

269

270 The "**static floodplain**" is located away from the main stream. Reworking events by  
271 an irregular avulsion or channel diversion of the river (Fig. 2) are considered to be  
272 rare and episodic. As the main channel migrates away from the original sites of  
273 deposition the distance to the main stream and thus the probability of preserving an  
274 (increasingly older) deposit increase. These distal deposits are most likely of  
275 Quaternary or older age. The nuclide concentration in modern stream sediment  $C_{mix}$   
276 will encounter a transient change following a remobilization event.

277

278 The "**dynamic floodplain**" in contrast is limited to the active channel belt, a zone  
279 close to the main channel that is frequently reworked by the migrating river.  
280 Relatively young, mostly Holocene deposits are continuously reworked and mixed  
281 with fresh sediment. If at a given position within the main channel sediment storage

282 and sediment remobilization are balanced, the nuclide concentration at this site  $C_{mix}$   
283 will not change with time.

284

285 During storage in both cases the initial *in situ* cosmogenic nuclide concentration  $C_0$ ,  
286 established when the sediment was produced in the source area, is changed to a  
287 concentration in the lowland deposit  $C_{dep}$  (at/g( $Qtz$ )). At a given burial depth  $z$  (m) of a  
288 section of sediment stored in the floodplain,  $C_{dep,z}$  depends on the density of  
289 sediment ( $\rho_{dep}$ ; including pore water) in the floodplain, on the locally prevailing  
290 (most likely low) nuclide production rate at the sediment's surface  $P_{dep}$  (at/g( $Qtz$ )/yr)  
291 that contains all nucleogenic and muogenic production terms (for full equation  
292 including muonic production terms, we refer to equation 6 in Wittmann and von  
293 Blanckenburg (2009)), and the storage time  $t$  (yr) (Schaller et al., 2002):

294

$$295 \quad C_{dep,z} = C_0 \times e^{-\lambda t} + \frac{P_{dep}}{\lambda} \times e^{\left(-\frac{\rho_{dep} \times z}{\Lambda}\right)} (1 - e^{-\lambda t}) \quad (\text{Eq. 3})$$

296

297  $C_{dep}$ , the nuclide concentration contained in an integrated sediment column to the  
298 maximum mobilization depth ( $z_{max}$ ) is calculated by integrating  $C_{dep,z}$  from the  
299 surface ( $z_0$ ) over the depth of erodible floodplain sediment, providing a mean  $C_{dep}$ :

300

$$301 \quad C_{dep} = \frac{1}{z_{max}} \int_{z_0}^{z_{max}} C_{dep,z} dx \quad (\text{Eq. 4})$$

302

303 In the next two sections we illustrate how  $C_{\text{dep}}$  evolves in both the static and the  
304 dynamic floodplain end members, and we provide examples from the Amazon and  
305 the Ganga basin (see Fig. 3 for basin topography and sample locations).

306

307

### 308 ***2.3 Predicting nuclide concentrations in sediment re-entrained from static deposits***

309 We illustrate the effects of storage depth and time on  $C_{\text{dep}}$  for  $^{10}\text{Be}$  ( $^{10}\text{Be}_{\text{dep}}$ ), and the  
310  $^{26}\text{Al}/^{10}\text{Be}_{\text{dep}}$  ratio for a static deposit. In Fig. 4A we see that at the surface  $^{10}\text{Be}$   
311 concentrations almost always increase due to additional nuclide production, until  
312 steady state between production and decay is attained at  $>3$  Myr. We also see that if  
313 stored very deeply at ca. 50 m, where even deep muonic production is low,  $^{10}\text{Be}$   
314 concentrations decrease by radioactive decay. Fig. 4B shows that when measuring  
315 both *in situ*  $^{26}\text{Al}$  and *in situ*  $^{10}\text{Be}$  the resulting  $^{26}\text{Al}/^{10}\text{Be}$  ratio is sensitive to both  
316 storage time and depth.

317

318 The re-entrainment of such static sediment having a nuclide concentration  $C_{\text{dep}}$  and  
319 its mixing with fresh upstream sediment with  $C_0$  results in mixed nuclide  
320 concentrations  $C_{\text{mix}}$  (at/g<sub>(Qtz)</sub>). In the Amazon basin, such entrainment of old,  
321 formerly buried sediment has been traced in river sediment with paired *in situ*  $^{26}\text{Al}$   
322 and *in situ*  $^{10}\text{Be}$  in coarse-grained quartz (Wittmann et al., 2011b).

323

324 In an “erosion island” plot (Fig. 5), the  $^{26}\text{Al}/^{10}\text{Be}$  ratio is linked to  $^{10}\text{Be}_0$ .  $^{10}\text{Be}_0$  is a  
325 measure of the denudation rate of the sediment during initial erosion, called  
326 inheritance. Steadily eroding sediments with no pre-burial history plot on the steady

327 state erosion line (Fig. 5A). The ratio of  $^{26}\text{Al}$  and  $^{10}\text{Be}$  nuclide production rates at the  
 328 surface is 6.5 to 7 (Balco et al., 2008; Goethals et al., 2009; Hidy et al., 2014).  
 329 Samples that have been buried plot beneath the line, at lower  $^{26}\text{Al}/^{10}\text{Be}$  ratios. Upon  
 330 reactivation of buried sediment, the active channel will receive a fraction of buried  
 331 material, such that  $^{10}\text{Be}$  and  $^{26}\text{Al}$  concentrations, called  $^{10}\text{Be}_{\text{mix}}$  and  $^{26}\text{Al}_{\text{mix}}$ , and  
 332  $(^{26}\text{Al}/^{10}\text{Be})_{\text{mix}}$  emerge. In order to calculate the fraction of recycled material,  $f_{\text{buried}}$ ,  
 333 we must know  $^{10}\text{Be}_0$  from  $^{10}\text{Be}$  and  $^{26}\text{Al}_0$  from  $^{26}\text{Al}$ , such that the position on the  
 334 constant steady state erosion line is constrained (see Fig. 5B):

335

$$336 \quad \left(\frac{^{26}\text{Al}}{^{10}\text{Be}}\right)_{\text{mix}} = \frac{(^{26}\text{Al}_0 \times f_0 + ^{26}\text{Al}_{\text{buried}} \times f_{\text{buried}})}{(^{10}\text{Be}_0 \times f_0 + ^{10}\text{Be}_{\text{buried}} \times f_{\text{buried}})} \quad (\text{Eq. 5})$$

337

338 Since the sum of  $f_0$  (the fraction of sediment containing  $^{10}\text{Be}_0$  from steady state  
 339 erosion in the source area) and  $f_{\text{buried}}$  equals 1, we can solve equation 5 for  $f_{\text{buried}}$   
 340 (Wittmann et al., 2011b). As we can constrain  $^{10}\text{Be}_{\text{dep}}$  and  $^{26}\text{Al}_{\text{dep}}$  only from the  
 341 lowermost sample in the diagram, burial depths and ages are minimum estimates.  
 342 The fractions of formerly buried end member incorporated into the non-buried  
 343 channel material are thus maximum fractions.

344

345 At this stage it is important to point out that the “isochron method” was designed to  
 346 determine depositional ages of buried sediment in which nuclides are being  
 347 produced by cosmic rays not fully absorbed by the overlying sediment from linear  
 348 regression of  $^{26}\text{Al}$  versus  $^{10}\text{Be}$  concentrations (Balco and Rovey, 2008). This method  
 349 does not require assumptions on the inherited nuclide concentrations. However, the  
 350 method cannot be applied to determine burial ages on river sediment that



351 comprises a mixture of such old sediment and fresh source area-derived river  
352 sediment.

353

#### 354 ***2.4 Amazon nuclide concentrations in sediment re-entrained from static deposits***

355 To our knowledge, the Amazon data shown in Fig. 5 is the only currently available  
356 dataset suitable to estimate sediment reactivation by using the  $(^{26}\text{Al}/^{10}\text{Be})_{\text{mix}}$  ratio in  
357 bedload sediment.  $(^{26}\text{Al}/^{10}\text{Be})_{\text{mix}}$  ratios well below ca. 6.5, indicating shielding from  
358 cosmic ray production during deep burial (see Fig. 5), were detected in the  
359 headwaters of cratonic rivers and also downstream in isolated floodplain deposits.  
360 There, burial ages exceed 5 Myr. In contrast, Andean sediment evidently has not  
361 experienced burial showing  $^{26}\text{Al}/^{10}\text{Be}$  ratios of around 6.5 to 7 (see Fig. 5) (Wittmann  
362 et al., 2011b). Such old craton-derived sediment is admixed to the Amazon by the  
363 large Negro River, which joins the Amazon at Manaus, and the Madeira River, which  
364 drains the Bolivian Andes and parts of the Brazilian Craton, and joins the Amazon  
365 downstream of Manaus (Fig. 3).

366

#### 367 ***2.5 Predicting the nuclide concentrations in the dynamic floodplains of the Amazon*** 368 ***and Ganga basins***

369 We compiled sediment storage times  $t$  for dynamic floodplains (see supplement B ,  
370 Table A1). These storage times are significantly shorter than the half lives of  $^{10}\text{Be}$  and  
371  $^{26}\text{Al}$ . This means that storage times are too short to allow for significant radioactive  
372 decay of  $^{10}\text{Be}$  or  $^{26}\text{Al}$  even when shielded from cosmic irradiation (Fig. 4A). The  
373 question is whether for  $^{10}\text{Be}$  and  $^{26}\text{Al}$  the nuclide concentration  $C_{\text{mix}}$  still increases

374 during the repeated storage cycles. However, *in situ*  $^{14}\text{C}$  has a half-life exactly within  
375 the range of storage times and should quantify storage duration.

376

377 The change of nuclide concentrations first depends on the floodplain configuration,  
378 i.e. its width, depth (because the reservoir size governs the capacity of sediment  
379 storage), and the upstream sediment flux (which together with the reservoir size  
380 determines the storage duration). Nuclide concentrations secondly depend on  
381 nuclide-specific characteristics, i.e. the initial upstream concentration  $C_0$  (that  
382 depends on source area erosion rate), the production rate in the lowlands, the  
383 penetration depth of cosmic rays, and, mostly for  $^{14}\text{C}$ , on the half-life of the nuclide.  
384 Given this range in parameters, we need to revert to combined floodplain mass  
385 balance and nuclide production models that evaluate the dependence of nuclide  
386 accumulation and decay on floodplain parameters.

387

388 Two modeling studies have now mapped out the conditions under which nuclide  
389 concentrations of *in situ*-produced  $^{10}\text{Be}$ ,  $^{14}\text{C}$ , and  $^{26}\text{Al}$  are modified as sediment  
390 moves through dynamic floodplains. The “box” (compartment-based) model of  
391 Wittmann and von Blanckenburg (2009) used *a priori* choices of floodplain exchange  
392 parameters, sediment storage time, and sediment fluxes. These were explored for a  
393 wide range of global lowland river settings. The numerical model by Lauer and  
394 Willenbring (2010), later modified by Viparelli et al. (2013), uses numerically  
395 determined gain or loss of floodplain mass depending on overbank deposition and  
396 net bank erosion rates as input. The model was explored for the case of the local  
397 Neuse river (mid-USA) and also addresses the clay flux passing through the reach.

398 This model makes predictions on the behavior of meteoric  $^{10}\text{Be}$  in floodplains,  
399 although the way the isotope migrates with the water flux through sediment, its  
400 penetration depth, nor its adsorption behavior are specified. Both models have the  
401 *in situ* nuclide production mechanisms (spallation and muonic) in common and  
402 eventually derive the final steady state nuclide concentrations  $C_{\text{mix}}$  from mass  
403 balance. Although their mathematical complexity differs considerably, the  
404 predictions made by these two models are in good agreement. For comparison we  
405 have employed both models to the Amazon and the Ganga basin using a single-  
406 channel setup, i.e. assuming that the lowlands are fed by a single, source-area  
407 draining river (see a description of the setup in Table 1 and Fig. 6). In contrast, the  
408 numeric model as employed by Lupker et al. (2012a) for the Ganga basin uses a  
409 setup where multiple headwater channels feed the lower Ganga.

410 Both models predict very low increases of only 1.8% for the box model and 1.7% for  
411 the numerical model in long-lived (i.e.  $^{10}\text{Be}$ ) *in situ* nuclide concentrations relative to  
412  $C_0$  along the lowland reach from the Andes to the Amazon mouth (Table 1, Fig. 6).  
413 For the Ganga basin an increase of 1.5% is predicted by the box model and 6.5% by  
414 the numerical model. Lupker et al., (2012a) predicted an increase of 13% relative to  
415  $^{10}\text{Be}_0$  by notably more accurately addressing the variation in nuclide concentrations  
416 of multiple headwater rivers. Evidently, this range in model results that depends on  
417 each model configuration implies that a certain range in measured nuclide  
418 concentrations can be expected in real settings, too. For short-lived nuclides, a  
419 significant decrease in *in situ*-produced  $^{14}\text{C}$  nuclide concentrations is predicted (Fig.  
420 6, A-1,2).

421 The overall similar response of a few % increase despite differing sediment storage  
422 times (14 kyr in the Amazon, 1.4 kyr in the Ganga) are due to the different  
423 remobilization depths. For the much deeper Amazon channel,  $^{10}\text{Be}$  that is produced  
424 in the floodplain superficially is diluted by deeply sourced sediment that was  
425 shielded from further production. The shallower channel of the Ganga, in contrast, is  
426 more sensitive to irradiation, even at short storage times. In addition, the relatively  
427 high Amazon  $^{10}\text{Be}_0$  (stemming from the Andean erosion rate of ca. 0.2 mm/yr) is less  
428 sensitive to additional surficial nuclide production in the lowlands than the three  
429 times lower  $^{10}\text{Be}_0$  of the Ganga (corresponding to high mean Himalayan erosion of  
430 ca. 1.5 mm/yr; Fig. 5). Overall, however, changes in long-lived *in situ* nuclide  
431 concentrations are predicted to be minor as most lowland settings are located at low  
432 elevations, where production rates are much lower than those in the sediment-  
433 producing highlands.

434

435 For  $^{14}\text{C}$ , the significant predicted decrease from the headwaters to the floodplain  
436 (Fig. 6) is due to the much shorter half-life, which makes this system sensitive to  
437 storage times of a few thousand years. Due to this behavior, Wittmann and von  
438 Blanckenburg (2009) proposed this nuclide to be a good proxy for sediment  
439 residence time. Complicating factors for such use in floodplains however are: 1) The  
440 initial *in situ*- $^{14}\text{C}$  signal of the sediment eroded from the source area has to be stable  
441 over time and representative of the erosion process. Even short burial and decay of  
442  $^{14}\text{C}$  upstream of the floodplain might compromise this stability (Lupker et al., 2014).  
443 2) Sediment cannot be considered as being well-mixed with respect to *in situ*- $^{14}\text{C}$  in  
444 floodplain sediment. Sediment mixing in floodplains takes place over time scale of a

445 few kyr, which is the same time scale as the  $^{14}\text{C}$  half-life. Even small additions of  
446 distal sediment anywhere along the floodplain reach may introduce high variability  
447 into calculated residence times.

448

## 449 ***2.6 Identifying sediment sources from nuclide concentrations in lowland rivers***

450 To test the predictions made in the preceding section, namely that *in situ* nuclide  
451 concentrations imprinted in the source area are hardly modified in transit through  
452 the dynamic floodplain, requires tracing sediment from a single source. However,  
453 the nuclide concentration of large lowland rivers is also controlled by the influx of  
454 sediment from tributaries that drain source areas with different denudation rates.  
455 These will contain distinct initial nuclide concentrations.

456

457 In the Amazon lowlands, Wittmann et al. (2011b) were able to separate the  
458 entrained signature of the static craton-derived and other lowland deposits from  
459 Andean-sourced sediment. This separation was done based on  $^{26}\text{Al}/^{10}\text{Be}$  ratios,  
460 indicating entrainment of previously buried sediment from the cratons, and on  
461 different grain size distributions. A condition for use of grain size distributions as  
462 source indicator is that these are not obscured through downstream fining.  
463 Downstream fining occurs in larger sand-bed rivers through abrasion and selective  
464 transport, whereas selective transport dominates over abrasion due to durable  
465 lithologies, small grain sizes, and high degree of rounding (Frings, 2008). Evidence for  
466 preservation of the grain size distribution was provided by Nordin et al. (1980) and  
467 Mertes and Meade (1985) who showed that the particle size of the Amazon is  
468 uniform from Iquitos to Óbidos. The area-weighted  $^{10}\text{Be}$  nuclide concentration of

469 Andean sediment (comprising all grain sizes,  $n = 62$ , basins of  $>100 \text{ km}^2$ ) is ca.  
470  $5.6 \times 10^4 \text{ at/g}_{(\text{Qtz})}$  (Fig. 7). Note that we employed area-weighing of  $^{10}\text{Be}$   
471 concentrations by adapting equation 7 for nuclide concentrations (see section 3.1).  
472 An exception with respect to nuclide concentrations in the Andes is the Bolivian  
473 Grande basin, where  $^{10}\text{Be}$  nuclide concentrations are much higher at  $40 \pm 56 \times 10^4$   
474  $\text{at/g}_{(\text{Qtz})}$  ( $\pm 1\sigma$  std. dev.) and also highly variable depending on physiographic region  
475 (i.e. indicated by high std. dev.) (Kober et al., 2015). The  $^{10}\text{Be}$  nuclide concentrations  
476 of all other investigated regions are however very similar to the  $^{10}\text{Be}$  nuclide  
477 concentration detected downstream in fine-grained (125-500  $\mu\text{m}$ ) Amazon bedload  
478 ( $5.3 \times 10^4 \text{ at/g}_{(\text{Qtz})}$ ,  $n = 18$ , see Fig. 7; Wittmann et al. (2011a)). This grain size fraction  
479 is therefore derived from the Andes. Craton-derived sediment on the other hand  
480 mainly contributes coarse-grained sandy sediment (Nordin et al., 1980; Franzinelli  
481 and Potter, 1983) containing high nuclide concentrations (ca.  $26.2 \times 10^4 \text{ at/g}_{(\text{Qtz})}$ ) from  
482 slow erosion of the cratonic headwaters that was buried for Myr durations as shown  
483 by  $^{26}\text{Al}/^{10}\text{Be}$  ratios (section 2.3 and Wittmann et al. (2011b)).

484

485 In the lower Ganga basin Lupker et al. (2012a) measured mean  $^{10}\text{Be}$  nuclide  
486 concentrations in lowland sediment of  $1.75 \times 10^4 \text{ at/g}_{(\text{Qtz})}$  ( $n = 14$ , area-weighed).  
487 Notably, some variability in  $^{10}\text{Be}$  concentrations exists for smaller lowland rivers (i.e.  
488 Karnali, Kosi, Narayani, having between 17% and 37% lowland area at the sampled  
489 locations), but the area-weighed lowland  $^{10}\text{Be}$  nuclide concentration is very similar  
490 to an area-weighed concentration of  $1.5 \times 10^4 \text{ at/g}_{(\text{Qtz})}$  of the Himalayan source area  
491 (Fig. 7,  $n = 65$ ,  $>100 \text{ km}^2$ ). Lowland  $^{10}\text{Be}$  concentrations were corrected by Lupker et  
492 al. (2012a) for the sediment contribution from rivers draining the Indian craton (e.g.

493 the Chambal River, see Fig. 3), by using simple binary mixing equations.  $^{10}\text{Be}$  nuclide  
494 concentration there are very high with  $43 \times 10^4$  at/g<sub>(Qtz)</sub> (Lupker et al., 2012a).  
495 Although these areas are slowly eroding, the associated sediment flux leads to a ca.  
496 three times elevated  $^{10}\text{Be}$  nuclide concentration of the main stream below the  
497 confluence with the Chambal River. These cratonic rivers also influence the grain size  
498 of the lower Ganga. The overall grain size reduction is only 2% per 100 km of river  
499 length, negligibly affecting Himalayan nuclide concentrations, but a slight coarsening  
500 is observed downstream of Indian craton confluences (Singh et al., 2007). If similar  
501 to the Amazon basin, these areas might also deliver old, deeply stored sediment to  
502 the main Ganga. We cannot test this assumption because no  $^{26}\text{Al}$  measurements are  
503 available for the Ganga River.

504

505 These observations confirm that in both basins the erosion signal from the sediment  
506 source area imprinted in the form of  $^{10}\text{Be}$  nuclide concentrations is not significantly  
507 changed, or, in other words, nuclide concentrations are conserved during transfer to  
508 the sea. This is the case in both the Amazon and the Ganga Rivers. The models of  
509 channel-floodplain interactions (section 2.5) predict this absence of change in the  
510 concentration of long-lived *in situ* nuclides in lowland basins.

511

512

### 513 **3. CALCULATING DENUDATION RATES AND SEDIMENT FLUXES IN LOWLAND**

#### 514 **BASINS**

##### 515 ***3.1 Converting nuclide concentrations from lowland rivers into a denudation rate*** 516 ***of the sediment-delivering mountains***

517 We have seen in the preceding section that the concentrations of long-lived *in situ*  
518 nuclides that have been set in the eroding mountain hinterland are preserved in  
519 lowland areas. They can be extracted by grain size-specific analysis and their  
520 reduction by radioactive decay in re-entrained old sediment from static deposits can  
521 be identified by paired nuclide analysis. This preservation leads to the opportunity to  
522 determine the erosion signal of an entire mountain belt. If the denudation rate of  
523 the entire basin including the lowlands is to be determined (using equation 2), a  
524 floodplain-uncorrected denudation rate, termed “ $D_{insitu}$ ” results. As this denudation  
525 rate is based on the unreasonable assumption that the entire basin including the  
526 floodplain provides sediment by erosion from the whole basins’ surface, it is only an  
527 “apparent” denudation rate. A “floodplain-corrected” denudation rate, termed  
528 “ $D_{insituFC}$ ”, takes into account that the lowlands do not produce additional “primary”  
529 sediment eroded under cosmogenic steady state conditions (equation 2), but merely  
530 serve as a transfer route. Calculating a floodplain-corrected denudation rate is done  
531 by using the production rate of the source area, termed  $P_{FC}$ , containing all  
532 nucleogenic and muogenic production mechanisms, that is then scaled to the  
533 altitude and latitude of the hinterland (Eq. 2).

534

535 As illustrated in Figure 8, this correction removes the decrease in apparent  
536 denudation rates across lowland areas (see blue dashed curve in Fig. 8) that arises  
537 when using the total production rate at the sampling point including the low-  
538 elevation lowlands. Because all other areas that do not contribute sediment are not  
539 considered, the resulting floodplain-corrected denudation rates  $D_{insituFC}$  (mm/yr)  
540 correspond to the sum of the hinterlands sediment production and weathering



541 rates. In practice, this calculation involves the digital delineation of the basin and  
 542 clipping low-elevation floodplain areas by using a basin-specific cut-off elevation  
 543 (using Digital Elevation Models, i.e. DEMs). From this floodplain-corrected  
 544 denudation rate  $D_{insituFC}$ , we can calculate floodplain-corrected sediment and  
 545 dissolved fluxes  $QS_{insituFC}$  (t/yr) (that by definition also contain dissolved fluxes  
 546 because we use denudation rates, i.e. the sum of erosion and weathering rates). We  
 547 multiply  $D_{insituFC}$  by the area of the production zone  $A_{source}$  and the source rock's  
 548 density  $\rho_{bedrock}$  (here a value of 2600 kg/m<sup>3</sup> is used, to relate the mass flux to  
 549 bedrock lowering) and obtain:

550

$$551 \quad QS_{insituFC} = D_{insituFC} \times A_{source} \times \rho_{bedrock} \quad (\text{Eq. 6})$$

552

553 When considering regions, such as the entire Andes or the lowlands, an area-  
 554 weighed denudation rate is calculated because smaller basins contribute less  
 555 sediment than larger basins. This area-weighted denudation rate  $\overline{D_{insituFC}}$  is  
 556 calculated using equation (3) in Wittmann et al. (2009), in order to weigh each  
 557 tributary according to tributaries' area relative to the summed area of individual  
 558 subbasins  $i$ :

559

$$560 \quad \overline{D_{insituFC}} = \frac{\sum_{i=1}^n (D_{insituFC} \times A_{source})}{\sum_{i=1}^n A_{source}} \quad (\text{Eq. 7})$$

561

562 Equation 7 can also be used to calculate an area-weighted denudation rate of sub-  
 563 basins of the Andes, in this case employing  $D_{insitu}$ .

564

565 **3.2 Recent erosion rates from sediment gauging data and their correction in**  
566 **lowland basins**

567 The reconstruction of sediment dynamics over different temporal scales is  
568 fundamental to understanding source-sink connectivity, because the time scale of  
569 observation is crucial for signal analysis in sedimentary systems (Romans et al.,  
570 2015). To provide such comparison of sediment fluxes over two entirely different  
571 temporal scales we can compare cosmogenic nuclide-derived sediment fluxes  
572 (integrating over millenia) with estimates of present-day mean sediment fluxes  
573 resulting from instrumental monitoring of river loads (Wittmann et al., 2011a;  
574 Covault et al., 2013).

575

576 A measure of present-day sediment flux can be calculated from measurements of  
577 suspended sediment,  $QS_{\text{gaug}}$  (t/yr), derived from suspended sediment concentration  
578  $N_{\text{susp}}$  (mg/l) and water flux  $Q_{\text{water}}$  ( $\text{m}^3/\text{sec}$ ) measurements. If  $Q_{\text{water}}$  can be identified  
579 as the major control of  $N_{\text{susp}}$ , a sediment rating model can be established to calculate  
580 the suspended sediment discharge from the usually long river discharge gauging  
581 period and thus well-constrained  $Q_{\text{water}}$ . For the periods in which suspended  
582 sediment concentration time series are lacking, this rating curve can then be used to  
583 extrapolate  $QS_{\text{gaug}}$  by, e.g., using river discharge data alone (Asselman, 1999;  
584 Horowitz, 2003). For a catchment of higher stream order that contains several sub-  
585 catchments that all or partially have been gauged, a representative catchment-wide  
586  $QS_{\text{gaug}}$  can be calculated by normalizing the catchments' summed tributary  $QS_{\text{gaug}}$   
587 value to the entire catchments' area at the outlet (see Table 2).

588

589 Extrapolating gauging data, representing years to decades, to millennial time scales  
590 has some shortcomings, as the period of observation necessary to capture the full  
591 signal amplitude is required to cover the recurrence interval of high-magnitude, low-  
592 frequency discharge events. In settings where sediment influx to river systems is  
593 inherently stochastic and mostly driven by discrete events such as earthquakes and  
594 rainstorms (Benda and Dunne, 1997), finding ways to predict recurrence intervals of  
595 such extreme events is fundamental (e.g. Korup (2012)). However, after a large, e.g.,  
596 earthquake-triggered mass wasting event decades of suspended sediment  
597 measurements indicate that the increased erosion pulse may persist for several  
598 years until all sediment has been evacuated from the catchment (Lin et al., 2008;  
599 Hovius et al., 2011). Still the integration period of suspended sediment  
600 measurements is, compared to the time scale usually covered by the *in situ*  
601 cosmogenic method, relatively short, such that discrepancies related to different  
602 integration time scales are expected (Kirchner et al., 2001; Schaller et al., 2001;  
603 Dadson et al., 2003; Wittmann et al., 2009; Covault et al., 2013)

604

605 To make sediment gauging data comparable to *in situ*  $^{10}\text{Be}$ -derived denudation rates,  
606 one of two conversions is required. This need arises because sediment gauging  
607 measures sediment flux  $Q_{S_{\text{gaug}}}$  (t/yr) whereas  $D_{\text{insituFC}}$  is a measure of surface  
608 lowering (mm/yr) or mass flux from a given area (t/km<sup>2</sup>/yr). Thus either  $D_{\text{insituFC}}$   
609 needs to be converted into  $Q_{S_{\text{insituFC}}}$  (t/yr) by equation 6 (see previous section), or we  
610 need to convert measured sediment fluxes, termed  $Q_{S_{\text{gaug}}}$ , to erosion rates, termed  
611  $E_{\text{gaug}}$ . Note that unlike  $D_{\text{insituFC}}$  these data do not incorporate a weathering fraction

612 unless suspended sediment data is supplemented by dissolved data.  $QS_{\text{gaug}}$  are  
613 divided by total basin area ( $\text{km}^2$ ) to derive a specific sediment yield  $SSY_{\text{gaug}}$   
614 ( $\text{t}/\text{km}^2/\text{yr}$ ):

615

$$616 \quad SSY_{\text{gaug}} = \frac{QS_{\text{gaug}}}{A_{\text{total}}} \quad (\text{Eq. 8})$$

617

618 or by total basin area and bedrock density  $\rho_{\text{bedrock}}$  ( $2600 \text{ kg}/\text{m}^3$ ) to obtain an  
619 erosion rate ( $\text{mm}/\text{yr}$ ):

620

$$621 \quad E_{\text{gaug}} = \frac{QS_{\text{gaug}}}{A_{\text{total}} \times \rho} \quad (\text{Eq. 9})$$

622

623 For many basins, both gauging-derived erosion rates and specific sediment yields  
624 decrease with increasing basin area. This relationship can be described with a power  
625 law where the exponent globally varies between -0.06 and -0.85 and can even obtain  
626 positive values (Syvitski et al., 2005; de Vente et al., 2007). However the global mean  
627 exponent for basins world-wide is -0.5, which we attribute to reflect mostly the  
628 geometry-dictated increase in lowland area in equations 8 or 9 at a given upstream  
629 sediment flux  $QS_{\text{gaug}}$ . The variation around the global mean exponent arises as larger  
630 basins have proportionally more low relief area than do steep and rapidly eroding  
631 headwater basins (Milliman and Meade, 1983; Summerfield and Hulton, 1994;  
632 Hovius, 1998). Thus, sediment production decreases downstream, concurrently  
633 accompanied by an increase in deposition potential as accommodation space in  
634 larger valleys and floodplains increases (Hinderer, 2012). In order to be comparable

635 with the cosmogenic nuclide method and to provide meaningful erosion rates from  
636 gauging data that do not merely decrease across lowlands because basin area  
637 increases, we suggest a “floodplain correction” for lowland gauging data, too. A  
638 floodplain-corrected erosion rate  $E_{\text{gaugFP}}$  is calculated by simply replacing  $A_{\text{total}}$  in  
639 equation 9 with the area of the sediment-producing hinterland  $A_{\text{source}}$ . Similar to the  
640 cosmogenic data, we calculated an area-weighted  $\overline{E_{\text{gaug}}}$  for each sub-basin (Table 2)  
641 by adapting equation 7 to gauging data, yielding:

642

$$643 \quad \overline{E_{\text{gaug}}} = \frac{\sum_{i=1}^n (E_{\text{gaugFC}} \times A_{\text{source}})}{\sum_{i=1}^n A_{\text{source}}} \quad (\text{Eq. 10})$$

644

645 The floodplain correction allows evaluation of whether sediment is being lost due to  
646 deposition or added from erosion in lowland reaches. Thus, while long-lived *in situ*  
647 cosmogenic nuclides record the sediment production rate in the source area,  
648 suspended sediment data record sediment production *and* sediment transport,  
649 thereby spatially integrating over all processes occurring in the production and the  
650 deposition zones of a basin. For example, flooding caused by rainfall events in  
651 lowland parts of the basin or anthropogenic erosion (Syvitski et al., 2005; de Vente  
652 et al., 2007) may lead to erosion of temporarily stored floodplain sediment, resulting  
653 in elevated sediment concentrations. At the same time, cosmogenic nuclide  
654 concentrations will not vary, as long as no distal (old, partly shielded) sediment is  
655 incorporated into the channel during the flood event.

656

657 **3.3 Comparing fluxes from *in situ* cosmogenic nuclides with river loads in the**  
658 **Amazon and Ganga**

659 We calculated *in situ*-cosmogenic nuclide-derived denudation rates from the  $^{10}\text{Be}$   
660 cosmogenic nuclide dataset shown in Figure 7 (using basins  $> 100 \text{ km}^2$  in order to be  
661 spatially representative). We summarized these source area and lowland datasets in  
662 Table 2 and Figure 9, and compared these to erosion rates calculated from measured  
663 suspended sediment fluxes (equations 8, 9). The full cosmogenic and gauging  
664 datasets are given in supplement C. Note that all rates summarized here were area-  
665 weighed according to equations 7 or 10. The area-weighted analytical uncertainties  
666 for the cosmogenic denudation rates are 15% for the entire Andes/Amazon and 24%  
667 for the Himalayan/Ganga datasets, respectively. Quantifying uncertainties for the  
668 suspended sediment dataset is more complex. Although the long observation  
669 periods of several decades and bedload contribution of only a few percent result in  
670 high confidence in the floodplain datasets of the Amazon and Ganga, estimates from  
671 multiple observation periods differ by 50% (see Wittmann et al., 2011a, Amazon and  
672 Lupker et al., 2011, Ganga). Therefore, to compare both methods, we decided to  
673 present the  $1\sigma$  std. dev. of each dataset (Table 2) rather than uncertainties. The  
674 standard deviation gives a sense of natural variations present for each dataset and  
675 does not imply a precision that is inappropriate when comparing different methods.

676

677 For the lowlands of the Amazon basin at Óbidos (Fig. 3), we calculate an *in situ*  
678 cosmogenic nuclide-derived Amazon basin-wide denudation rate  $\overline{D_{in situ}}$  of 0.11  
679 mm/yr when no floodplain correction is employed. When the floodplain correction is  
680 employed, the  $\overline{D_{in situ FC}}$  is 0.18 mm/yr. This rate is similar to a  $\overline{D_{in situ}}$  of the entire

681 Andes of 0.19 mm/yr (see section 4.1) and thus can be considered to represent the  
682 combined sediment production and weathering rate of the source area. We  
683 converted this Andean denudation rate to sediment and weathering fluxes  $QS_{insitu}$  by  
684 using equation 6 and a sampled source area of  $7.58 \times 10^5 \text{ km}^2$  and obtained 380  
685 Mt/yr; conversion of the  $\overline{D_{insituFC}}$  of 0.18 mm/yr at Óbidos (using equation 6 and  
686 the same sampled source area) leads to a sediment flux of 350 Mt/yr (Figs. 9, 10;  
687 Table 2).

688

689 For the Ganga basin, a basin-wide  $\overline{D_{insitu}}$  of 0.33 mm/yr is calculated at Harding  
690 Bridge (see Table 2) when no floodplain correction is employed. A  $\overline{D_{insituFC}}$  of 0.82  
691 mm/yr (relating to  $QS_{insituFP}$  of 390 Mt/yr when using a sampled source area of  
692  $1.84 \times 10^5 \text{ km}^2$  in Eq. 6) compares to a  $\overline{D_{insitu}}$  of 1.14 mm/yr for the Himalayan source  
693 areas and an equivalent  $QS_{insitu}$  of 550 Mt/yr (see Figs. 9, 10).

694 The gauging-derived lowland dataset is similar to that measured in the source areas  
695 (see lower part of Table 2). For the Amazon basin at Óbidos, a basin-wide  $\overline{E_{gaug}}$  of  
696 0.07 mm/yr is calculated. After removing the lowland area from this rate by  
697 application of the floodplain correction (by using a sampled source area of  $6 \times 10^5 \text{ km}^2$   
698 in Eq. 6), the  $\overline{E_{gaugFC}}$  is 0.58 mm/yr, a value that compares to a source-area  $\overline{E_{gaug}}$  of  
699 0.47 mm/yr (see Table 2 and Fig. 9). In the Ganga basin, a non-floodplain corrected  
700  $\overline{E_{gaug}}$  is 0.17 mm/yr, and a  $\overline{E_{gaugFC}}$  denotes to 0.81 mm/yr (by using a sampled  
701 source area of  $1.85 \times 10^5 \text{ km}^2$  in Eq. 6), a value that compares to a source-area  $\overline{E_{gaug}}$   
702 of 0.80 mm/yr (see Figs. 9, 10).

703

704 By comparison, dissolved fluxes are minor. There is independent evidence that

705 dissolved fluxes as recorded in the Amazon lowlands comprise only ca. 10% of the  
706 suspended sediment flux (for summary on dissolved and suspended sediment fluxes  
707 see Table 2 in Wittmann et al. (2011a)). Silicate weathering rates determined from  
708 combined *in situ*- and gauging-derived estimates of D and W are similar for the  
709 Andes and the Amazon lowlands and are generally low at ca. 100-200 t/km<sup>2</sup>/yr  
710 (corresponding to 0.04-0.08 mm/yr, or to ca. 10% of total D (Bouchez et al., 2014)).  
711 Likewise, as explained in section 5, weathering intensities determined from the new  
712 weathering and erosion <sup>10</sup>Be(meteoric)/<sup>9</sup>Be proxy indicate similar weathering  
713 intensities in the Andes and in the lowlands, respectively (Wittmann et al., 2015).  
714 In the Himalayan source areas silicate weathering rates amount to only 17 t/km<sup>2</sup>/yr  
715 (corresponding to 0.007 mm/yr; Rai et al. (2010)). In such terrains, kinetically-limited  
716 conditions prevail, meaning that minerals are eroded before they weather to  
717 completeness (West et al., 2005; Gabet and Mudd, 2009; Dixon and von  
718 Blanckenburg, 2012; Dixon and Riebe, 2014). However, Lupker et al. (2012b) found  
719 that floodplain weathering dominated over mountain weathering in the Ganga  
720 basin. Yet these floodplain processes are associated with considerable partitioning of  
721 dissolved Si into the suspended load in clays that formed in the floodplain (Frings et  
722 al., 2015) such that net silicate weathering fluxes even decrease along the floodplain  
723 (Rai et al., 2010).

724

## 725 **4. THE GEOLOGICAL SIGNIFICANCE OF FLOODPLAIN DATA**

### 726 ***4.1 Across spatial scales: big brush or fine tip?***

727 Regarding the compiled denudation and erosion rate datasets we first discuss the



728 significance of denudation and erosion rates, their spatial averaging as catchment  
729 area increases, and evaluate sediment fluxes from the headwaters to the lowlands to  
730 the ocean. The following observations were made in the preceding section.

731 a) In the Amazon basin, D (from *in situ*  $^{10}\text{Be}$ ), differ by a factor of 2-3 to E (from  
732 gauging), both in the source area ( $\overline{D_{insitu}} = 0.19 \pm 0.18 \text{ mm/yr}$ ;  $\overline{E_{gaug}} = 0.47 \pm 0.90$   
733 mm/yr) and in the lowlands ( $\overline{D_{insituFC}} = 0.18 \pm 0.03 \text{ mm/yr}$ ;  $\overline{E_{gaugFC}} = 0.58 \pm 0.20$   
734 mm/yr; see Fig. 10 and Table 2).

735 b) In the Ganga basin, D and E show a good agreement, both in the source area  
736 ( $\overline{D_{insitu}} = 1.14 \pm 0.82 \text{ mm/yr}$ ;  $\overline{E_{gaug}} = 0.80 \pm 0.55 \text{ mm/yr}$ ; see Table 2) and in the  
737 Ganga lowlands ( $\overline{D_{insituFC}} = 0.82 \pm 0.27 \text{ mm/yr}$ ;  $\overline{E_{gaugFC}} = 0.81 \pm 0.27 \text{ mm/yr}$ ).

738 c) In both the Andes-Amazon and the Himalaya-Ganga systems, cosmogenic nuclide-  
739 derived D contains only a minor dissolved contribution. We can thus compare short-  
740 term erosion rates directly to millennial time scale denudation rates and ignore the  
741 weathering fluxes.

742

743 We summarize from our study the following first-order interpretations:

744 1) Anthropogenic perturbations, known to affect the Andean source areas by e.g.  
745 gold mining and changes in land use (Maurice et al., 1999; Cleary, 2000; Vanacker et  
746 al., 2007) can result in an increase in erosion towards modern times. The factor-of-2  
747 difference between  $D_{insitu}$  and  $E_{gaug}$  in the Andes supports this interpretation. For  
748 comparison, in the Highlands of Sri Lanka or the Paute basin in Ecuador, much higher  
749  $E_{gaug}$  than  $D_{insitu}$  are observed (see Fig. 11). For the Ganga data, where  $D_{insitu}$  and  $E_{gaug}$   
750 agree better, Lupker et al. (2012a) argued that human-impact in the Ganga basin  
751 have not increased sediment mobilization.

752

753 2)  $\overline{D_{insitu}}$  of the Himalayan source area might be overestimated due to several  
754 potential causes. Non-steady state conditions, due to the occurrence of episodic  
755 deep-seated (>5 m) landslides incorporating low  $^{10}\text{Be}$ -concentrated material into  
756 streams, may prevail. In catchments with frequent landslides, adequate spatial and  
757 temporal mixing of sediment is established for catchments larger than 100 km<sup>2</sup>  
758 (Niemi et al., 2005; Yanites et al., 2009). In our compilation (Table 2, Fig. 9), we have  
759 excluded catchments <100 km<sup>2</sup> for this reason. However, poor spatial mixing of  
760 sediment has also been attributed to large precipitation events, that locally  
761 remobilize large amounts of sediments (Lupker et al., 2012a). Further, in the  
762 Himalaya, the more rapidly eroding areas tend to be the most quartz-rich lithologies.  
763 Since the probability of non-ubiquitous quartz concentrations increases with  
764 increasing catchment size, higher  $D_{insitu}$  than  $E_{gaug}$  may be caused by not correcting  
765 for this effect.

766

767 3) We cannot exclude that  $E_{gaug}$  is underestimated if high magnitude, low-frequency  
768 events are not captured in decades-spanning records (see section 3.2). Considering  
769 the overall good agreement between cosmogenic nuclide-derived  $D$  and sediment  
770 gauging-derived  $E$ , and the potential of floodplains to buffer variations in fluxes (see  
771 next section), we attribute the gauging data a high likelihood to be representative.

772

773 4) The “sediment delivery ratio” (Walling, 1983; Syvitski et al., 2005; de Vente et al.,  
774 2007), describing the ratio of sediment entering a basin to that transmitted through

775 a basin, is unity within uncertainty for these two river basins. Apparently, all  
776 sediment is transmitted to the sea without intermittent loss.

777

778 5) We note that for both methods the variation contained in the data, estimated by  
779 the coefficient of variation,  $C_v$  (Table 2), decreases from the source to the lowlands  
780 in both basins. Source area- $C_v$  are on average a factor 3 higher than in the lowlands.  
781 Wittmann et al. (2009) showed for the Bolivian Beni basin that when sampling far  
782 away from the mountain front, fluvial transport and mixing processes average-out  
783 small-scale headwater variability, and a mean headwater denudation rate is  
784 preserved. We find a similar decrease in variability from source to lowlands for the  
785 Ganga basin. Thus a mean, representative orogenic denudation rate can be derived  
786 from a bag of sediment sampled downstream. This decrease in variation is most  
787 likely due to larger catchments being more representative of sediment transport  
788 processes than smaller catchments, especially if a transfer zone is present in a basin  
789 that is able to retain sediment for longer periods (Romans et al., 2015). Locally high  
790 sediment inputs from stochastic mass wasting are diluted downstream. The lowland  
791 floodplain has the ability to buffer changes in sediment supply and thus drives these  
792 averaging effects (see next section).

793

794 6) Good agreement is observed between sedimentary lowland fluxes and sediment  
795 volumes in offshore basins or deltas (Table 2): For the Amazon basin, Nittrouer and  
796 Kuehl (1995) estimated a mean sediment flux of 550 - 1030 Mt/yr from  $^{210}\text{Pb}$  activity  
797 profiles (integration time scale of ca. 1 kyr), an estimation that is slightly higher than  
798 the *in situ*- $^{10}\text{Be}$  based lowland sediment flux (370 Mt/yr) and also encompasses the

799 gauging-based lowland flux (990 Mt/yr) (Table 2). For the Ganga basin, Métivier and  
800 Gaudemer (1999) estimated a long-term deposition flux of ca. 510 Mt/yr for the  
801 Bengal fan over the Quaternary from the Ganga River alone. This estimation from  
802 the fan is at the higher end of gauging-derived estimates (ca. 200 - 550 Mt/yr, see  
803 Table 2), but broadly agrees with *in situ* cosmogenic-derived sediment fluxes (ca. 400  
804 Mt/yr).

805

#### 806 **4.2 Across temporal scales: Stability of lowland fluxes**

807 There are several potential causes for the major observations made in section 4.1,  
808 i.e. that sediment fluxes in the mountain sources and across floodplains are similar,  
809 that their variability decreases downstream, and that these sediment fluxes leaving  
810 the large floodplains are again similar to sediment volumes stored offshore.  
811 Constant source erosion rates over the time scales encompassed by the discussed  
812 methods is one explanation for the observed similarity.

813

814 Buffering of sediment transport through the floodplain is another explanation for the  
815 similarity over different time scales. Rivers react to external, high amplitude forcings  
816 in the source area by either incising their alluvial reservoir or by aggrading it to  
817 “buffer” against these changes. As a result sediment fluxes at the outlet are held  
818 constant (Métivier and Gaudemer, 1999; Phillips, 2003). However, it is unclear over  
819 which time scales these buffers act. Métivier and Gaudemer (1999) have calculated  
820  $4.7 \times 10^5$  yr as the time it takes to erode an alluvial reservoir of the size of the Ganga  
821 floodplain, called “reaction time”, and, based on their principles, Wittmann et al.  
822 (2011a) have calculated a reaction time of  $7 \times 10^5$  yr for the Amazon floodplain. These

823 time scales depend both on the size of the floodplain and on the sediment flux into  
824 the floodplain, and only signals that are longer than the reaction time should be able  
825 to pass the floodplain without alteration (Métivier and Gaudemer, 1999). The  
826 relation between floodplain reaction time and period of the external forcing is  
827 subject to an intense debate, and several models with differing outcomes have been  
828 put forward (Allen, 2008; Jerolmack and Paola, 2010; Simpson and Castelltort, 2012;  
829 Armitage et al., 2013; Godard et al., 2013). Romans et al. (2015) stress that signals  
830 may be preserved in the stratigraphic record, and thus “survive” fluvial transport if  
831 the period of signal forcing exceeds the reaction time of the alluvial reservoir and  
832 also exceeds the magnitude of internal oscillations, such that signals are not  
833 “shredded” by self-organized internal dynamics (Jerolmack and Paola, 2010).

834

835 In order to elucidate signal transmittance on an empirical basis, many authors use  
836 the comparison of longer-term (e.g. cosmogenic or infill-derived fluxes) to shorter-  
837 term (gauging) fluxes (Métivier and Gaudemer, 1999; Wittmann et al., 2011a;  
838 Covault et al., 2013). For example, Covault et al. (2013) note from their global  
839 compilation of modern gauging-based and longer-term cosmogenic nuclide-based  
840 sediment fluxes that 28 sites agree within a factor of 2, and even 81 of 103 sites  
841 exhibit a broad similarity by having within one order of magnitude similar fluxes.  
842 Covault et al. (2013) note that these sites are predominantly from larger basins (see  
843 Fig. 11). In Figure 11, we supplement Covault et al.’s (2013) compilation of longer-  
844 term cosmogenic-derived versus modern gauging-derived sediment loads with data  
845 from the European Alps (Wittmann et al., 2007; Norton et al., 2011; Hinderer et al.,  
846 2013), the Ecuadorian Intermontane Andes (Vanacker et al., 2007), the Bolivian

847 Andes (Safran et al., 2005; Aalto et al., 2006; Insel et al., 2010), and the Nepalese  
848 Himalayas (Andermann, 2011; Andermann et al., 2012). This added data supports  
849 the observation that it is mostly the larger basins exhibiting broad similarity in  
850 sediment discharge over time as recorded by the two methods. They show a broad  
851 balance between modern and longer-term fluxes for many large rivers of the world.  
852 If cosmogenic nuclides were added to the standard toolbox of methods applied to  
853 the stratigraphic record, more insight into the fluvial transport system and the role it  
854 is playing in transmitting source area and lowland signals could be gained.

855

## 856 **5. Meteoric and $^{10}\text{Be}$ and the $^{10}\text{Be}/^9\text{Be}$ ratio in floodplains**

### 857 ***5.1 General principles***

858 Meteoric cosmogenic  $^{10}\text{Be}$  atoms are produced in the atmosphere at high production  
859 rates. Their global flux amounts to ca.  $1 \times 10^6$  atoms/cm<sup>2</sup>/yr. After atmospheric  
860 redistribution and precipitation they adsorb onto fine particles with high reactive  
861 surface area (Willenbring and von Blanckenburg, 2010). Stable  $^9\text{Be}$ , present in silicate  
862 minerals in ppm concentrations, is released during weathering and mixes with  $^{10}\text{Be}$   
863 in soil or river solutions to form a characteristic  $^{10}\text{Be}/^9\text{Be}$  ratio. The potential of this  
864 new isotope ratio tracer lies in the versatility of its applications. In floodplains, the  
865  $^{10}\text{Be}$  concentration can be used to determine a) upland erosion rates and b)  
866 floodplain sediment storage duration. The  $^{10}\text{Be}/^9\text{Be}$  ratio provides c) denudation  
867 rates while the fraction of  $^9\text{Be}$  released during weathering from primary minerals  
868 discloses d) the degree of rock weathering. The principle framework for these  
869 applications is given in von Blanckenburg et al. (2012).

870

871 For the determination of erosion rates and exposure ages, sub-gram amounts of  
872 fine-grained samples can be measured, given the much higher meteoric  $^{10}\text{Be}$   
873 concentrations than *in situ* cosmogenic  $^{10}\text{Be}$ . While for determination of *in situ*  $^{10}\text{Be}$   
874 in quartz meteoric  $^{10}\text{Be}$  is removed by acid leaching (Kohl and Nishiizumi, 1992), the  
875 amounts of *in situ*  $^{10}\text{Be}$  are negligible when meteoric  $^{10}\text{Be}$  is measured. For the  
876 determination of denudation rates, reactive  $^{10}\text{Be}/^9\text{Be}$  ratios (reac) are determined  
877 from phases such as Mn-Fe-(oxy)-hydroxides. This reactive Be is separated from  
878 samples by sequential chemical extractions (Tessier et al., 1979; Bourlès et al., 1989;  
879 Wittmann et al., 2012), the mineral-bound  $^9\text{Be}$  retained in undissolved minerals is  
880 excluded, and the concentration is reported as  $^{10}\text{Be}_{\text{reac}}$  and  $^9\text{Be}_{\text{reac}}$  (in  $\text{at/g}_{(\text{solid})}$  or  
881  $\text{g/g}_{(\text{solid})}$ ), respectively.

882

## 883 **5.2 Atmospheric $^{10}\text{Be}$ deposition fluxes**

884 A first prerequisite to use meteoric  $^{10}\text{Be}$  for any quantitative information is  
885 knowledge of the  $^{10}\text{Be}$  flux reaching Earth surface  $^{10}\text{Be}$  flux  $F_{\text{met}}^{^{10}\text{Be}}$ , in units of  
886  $\text{at/cm}^2/\text{yr}$ . This parameter is still associated with substantial uncertainties; one  
887 approach to obtain  $F_{\text{met}}^{^{10}\text{Be}}$  that is most likely valid for large basin-wide studies  
888 (Wittmann et al., 2015) is by using a general atmospheric circulation model (GCM)  
889 coupled to an aerosol module (e.g. Heikkilä et al. (2013)) that also incorporates a  
890 physical model simulating processes of cosmic particle production and transport  
891 (Masarik and Beer, 1999) and variations in solar modulation and magnetic field  
892 strength. For smaller scales, measured inventories  $^{10}\text{Be}$  in soil for given latitudes and

893 precipitation rates can be used to determine local  $F_{met}^{10Be}$  (Reusser et al., 2010;  
894 Willenbring and von Blanckenburg, 2010; Graly et al., 2011; Ouimet et al., 2015). As  
895 >95% of meteoric  $^{10}Be$  production in the atmosphere takes place at elevations above  
896 3 km altitude, scaling for altitude is not required for most locations (Willenbring and  
897 von Blanckenburg, 2010).

898

### 899 ***5.3 Meteoric $^{10}Be$ as tracer of erosion rates and sediment storage duration***

900 The most simple application of meteoric  $^{10}Be$  is the determination of erosion rates  
901  $E_{met}$  (Willenbring and von Blanckenburg, 2010). To do so, we face two fundamental  
902 difficulties. First, only at near-neutral or higher pH is meteoric  $^{10}Be$  fully retained. We  
903 can account for this loss by using solid/fluid partition coefficients and runoff (von  
904 Blanckenburg et al., 2012) (see below, Eq. 11). Second,  $^{10}Be$  adsorption is grain size-  
905 dependent (Aldahan et al., 1999; Shen et al., 2004; Graly et al., 2010; Willenbring  
906 and von Blanckenburg, 2010) (Fig. 12). One means to remove this effect is to  
907 normalize over stable  $^9Be$ , where the  $^{10}Be/^9Be$  ratio does not depend on grain size  
908 (Figures 12B and 12D). If the  $^{10}Be/^9Be$  ratio is used, we need to find means to extract  
909 an erosion or denudation rate from that ratio, however. This procedure is explained  
910 in section 5.4. Another way to obtain a representative  $^{10}Be$  concentration from river  
911 sediment is to measure the Al/Si ratio in suspended sediment too, and correct the  
912 meteoric  $^{10}Be$  concentration by using depth-integrated Al/Si ratios (Fig. 12C).

913

914 Once the grain size dependence has been addressed, the reactive (or bulk sediment)  
915 meteoric  $^{10}Be$  concentration ( $^{10}Be_{reac}$ , in  $at/g_{solid}$ ) can be converted into an erosion  
916 rate  $E_{met}$  ( $g/cm^2/yr$ , convertible into  $mm/yr$  by using rock density  $\rho_{bedrock}$ ):



917

$$918 \quad E_{\text{met}} = \frac{F_{\text{met}}^{10\text{Be}}}{J_{\text{riv\_reac}}^{10\text{Be}}} - \frac{q}{K_d} \quad (\text{Eq. 11})$$

919

920 where  $q$  is water runoff ( $\text{cm}^3/\text{cm}^2/\text{yr}$ ) and  $K_d$  is the solid/fluid partition coefficient for  
921 Be ( $\text{ml/g}$ ). The  $q/K_d$  term takes into account the  $^{10}\text{Be}$  lost in solution (von  
922 Blanckenburg et al., 2012) and can be ignored where Be is fully retentive ( $\text{pH} > 6$ ).

923 A second application of meteoric  $^{10}\text{Be}$  is to evaluate the duration of static sediment  
924 storage in floodplains. This duration can be obtained by comparing the total mass

925 export rate of meteoric  $^{10}\text{Be}$  from a basin  $J_{\text{riv}}^{10\text{Be}}$  ( $\text{at/yr}$ ) to the atmospheric deposition

926 rate  $J_{\text{atm}}^{10\text{Be}}$ . While  $J_{\text{atm}}^{10\text{Be}}$  is the meteoric flux (i.e.  $F_{\text{met}}^{10\text{Be}}$  multiplied by the basins area),

927  $J_{\text{riv}}^{10\text{Be}}$  is the sum of the riverine solid reactive and dissolved fractions transported by

928 the river, called  $J_{\text{riv\_reac}}^{10\text{Be}}$  and  $J_{\text{riv\_diss}}^{10\text{Be}}$  respectively (all in  $\text{at/yr}$ ):

929

$$930 \quad J_{\text{riv}}^{10\text{Be}} = (J_{\text{riv\_reac}}^{10\text{Be}} + J_{\text{riv\_diss}}^{10\text{Be}}) \times (1 - \exp(-\lambda t)) \quad (\text{Eq. 12})$$

931

932 These  $J_{\text{riv}}^{10\text{Be}}$  are calculated by area-normalizing the reactive or dissolved  $^{10}\text{Be}$   
933 concentrations with the solid export rate (from e.g. *in situ*-derived erosion rates, in  
934  $\text{kg}/\text{m}^2/\text{yr}$ ) or water discharge (in  $\text{L}/\text{yr}$ ), respectively (Wittmann et al., 2015)).

935

936 If the  $^{10}\text{Be}$  inventory of the basin is at steady state,  $J_{\text{riv}}^{10\text{Be}} / J_{\text{atm}}^{10\text{Be}}$  equals 1 (Brown et

937 al., 1988). If this ratio  $< 1$ , the sediment storage duration can be calculated by solving

938 equation 12 for  $t$ . We illustrate this case for the riverine  $^{10}\text{Be}$  flux deficit observed in  
939 the Amazon basin in Figure 13A, i.e. the difference between  $J_{\text{riv}}^{10\text{Be}}$  and  $J_{\text{atm}}^{10\text{Be}}$ . The  
940 Amazon case provides a duration for storage of „static“ sediment and the decay of  
941  $^{10}\text{Be}$  contained in it of 3.7 Myr (Fig. 13A). An alternative explanation for the flux  
942 deficit is that no stored sediment (and the  $^{10}\text{Be}$  that is contained in it) reaches the  
943 active channel. For the Amazon lowlands, this area would be 64% (Wittmann et al.,  
944 2015).

945  
946 A third application is the calculation of sediment transfer times for the case of  
947 dynamic floodplains. Here we can make use of the fact that  $F_{\text{met}}^{10\text{Be}}$  does not change  
948 from the mountainous source areas to the lowlands of a basin (see Fig. 14). In  
949 contrast, *in situ*  $^{10}\text{Be}$  production rates decrease over this distance. Hence, meteoric  
950  $^{10}\text{Be}$  increases significantly during transfer along a floodplain, whereas *in situ*  $^{10}\text{Be}$   
951 mainly does not increase. We illustrate this distinct behavior schematically in Fig. 14.

952  
953 We can model the increase by a simple accumulation scenario where  $^{10}\text{Be}_{\text{reac}}$   
954 continuously accumulates in floodplain sediment during residence in the active  
955 floodplain (Fig. 13B). When using equation 6 in Willenbring and von Blanckenburg  
956 (2010) and assuming zero erosion of the stored lowland sediment, the sedimentary  
957  $^{10}\text{Be}$  inventory  $I$  ( $\text{at}/\text{cm}^2$ ) that is produced while the sediment is exposed to  
958 continuous atmospheric deposition  $F_{\text{met}}^{10\text{Be}}$ , can be calculated from equation 16 in  
959 Wittmann et al. (2015):

960

961 
$$I_{^{10}\text{Be}}(t) = \frac{F_{\text{met}}^{^{10}\text{Be}}}{\lambda} (1 - \exp^{-\lambda t}) + ^{10}\text{Be}_{\text{source}} \times \rho \times z \times \exp^{-\lambda t}$$

962 (Eq. 13)

963

964 In this case is  $t$  the sediment transfer time and the right-hand term in equation 13  
 965 reflects the decay-affected contribution of source area-derived  $^{10}\text{Be}_{\text{reac}}$  that is  
 966 provided to the lowlands from the Andes ( $^{10}\text{Be}_{\text{source}}$ , in  $\text{at/g}_{\text{solid}}$ ). As sediment density  
 967  $\rho$  we used a value of  $2000 \text{ kg/m}^3$  for wet, silty sand (see Balco et al. (2005)) and  $z$  (m)  
 968 is the remobilization depth of sediment. Note that the inventory of  $^{10}\text{Be}$  does not  
 969 depend on knowing the  $^{10}\text{Be}$  penetration depth (Willenbring and von Blanckenburg,  
 970 2010), as long as the entire inventory is captured by sampling. However, to convert  
 971 the  $^{10}\text{Be}$  inventory into the  $^{10}\text{Be}$  concentration contained in remobilized sediment,  
 972 the remobilization depth needs to be known. In the example shown in Fig. 13B we  
 973 use equation 13 with  $z = 1 \text{ m}$  for the case of very shallow floodplain remobilization  
 974 representing a topmost clay-rich layer ladden with meteoric  $^{10}\text{Be}$  that is deposited  
 975 during overbank spill and eroded during subsequent inundation. Using this depth, a  
 976 sediment transfer time in the floodplain of only ca. 1.6 kyr is predicted to cause the  
 977 additional accumulation of meteoric  $^{10}\text{Be}$  across the Amazon floodplain. A deeper  
 978 remobilization depth of several tens of meters down to the channel bottom (e.g. 20  
 979 m) would require longer sediment transfer times of ca. 29 kyr (not shown in Fig.  
 980 13B).

981

982

983 **5.4  $^{10}\text{Be}/^9\text{Be}$  as denudation rate and weathering degree proxy**

984 Within the mass balance framework of von Blanckenburg et al. (2012), meteoric  $^{10}\text{Be}$   
 985 is the atmospheric flux tracer, and  $^9\text{Be}$  is the tracer for the release of the trace metal  
 986 by weathering, and thus their combination can be used to simultaneously determine  
 987 denudation, erosion and weathering rates. In order to be able to calculate  
 988 denudation rates from equation 14 we need to know the fraction of a parent rock's  
 989  $^9\text{Be}$  that is released by weathering into the reactive and dissolved phases and the  
 990 initial "parent" bedrock concentration of stable  $^9\text{Be}$  ( $^9\text{Be}_{\text{parent}}$ , in  $\mu\text{g/g}$ ). For large scale  
 991 studies, a value of the average crustal composition of Be of  $2.5 \pm 0.5 \mu\text{g/g}$  (Rudnick  
 992 and Gao, 2004) can be used (von Blanckenburg et al., 2012) to calculate  $D_{\text{met}}$ :

993

$$994 \quad D_{\text{met}} = \frac{F_{\text{met}}^{10\text{Be}}}{\left(\frac{^{10}\text{Be}}{^9\text{Be}}\right)_{\text{reac}} \times [^9\text{Be}]_{\text{parent}} \times (f_{\text{reac}}^{9\text{Be}} + f_{\text{diss}}^{9\text{Be}})} \quad (\text{Eq. 14})$$

995  
 996  $(f_{\text{reac}}^{9\text{Be}} + f_{\text{diss}}^{9\text{Be}})$  can be determined in two ways (Wittmann et al., 2015): Either by  
 997 knowing both the riverine reactive and dissolved  $^9\text{Be}$  flux, or by sequential  
 998 extractions of the reactive and the residual mineral-bound concentration from a  
 999 given sediment sample. Then:

1000

$$1001 \quad f_{\text{reac}}^{9\text{Be}} + f_{\text{diss}}^{9\text{Be}} = \left(\frac{^9\text{Be}_{\text{min}}}{^9\text{Be}_{\text{reac}}} + 1\right)^{-1} \quad (\text{Eq. 15})$$

1002  
 1003  $(f_{\text{reac}}^{9\text{Be}} + f_{\text{diss}}^{9\text{Be}})$  describes the release of Be from primary minerals into the dissolved  
 1004 (diss) and particulate (reac) component and is thus a proxy for the degree of  
 1005 weathering. In the slow and complete weathering („supply-limited“) case, this

1006 fraction is close to 1, and all primary mineral-bound  $^9\text{Be}$  is released. In fast and  
1007 incomplete weathering („kinetically limited“) settings, this fraction is  $<1$ , and some  
1008  $^9\text{Be}$  remains in undissolved primary minerals. Note that this fraction may be biased  
1009 by grain size effects, as finer grains may yield higher  $^9\text{Be}_{\text{reac}}$  than  $^9\text{Be}_{\text{min}}$  inherited  
1010 from sorting between coarse grain sizes and clays in the river (von Blanckenburg et  
1011 al., 2012). This bias may then also affect denudation rates calculated from equation  
1012 14 where the weathering fraction is contained.

1013

1014 In the Amazon basin,  $(f_{\text{reac}}^{^9\text{Be}} + f_{\text{diss}}^{^9\text{Be}})$  have been calculated from  $^9\text{Be}$  data from  
1015 suspended sediment depth profiles; this data was corrected for grain size bias during  
1016 riverine sorting following the approach shown in Fig. 12C and is thus considered  
1017 representative for the Amazon River where the majority of sediment flux is  
1018 transported by suspended load (Wittmann et al., 2015). The  $(f_{\text{reac}}^{^9\text{Be}} + f_{\text{diss}}^{^9\text{Be}})$  values  
1019 are ca. 0.4 (i.e. 40% of Be mass has been released during weathering) for two  
1020 Andean and the two lowland depth profiles and thus the weathering degree does  
1021 not change from the Andes to the lowlands. One explanation for this stability in the  
1022 two regimes is that sediments contained in the Amazon lowlands are pre-weathered  
1023 in the Andes and potentially depleted of their  $^9\text{Be}$  (Wittmann et al., 2015). Other  
1024 studies (Bouchez et al., 2012; Bouchez et al., 2014) have found similar low  
1025 weathering during transfer of sediments through the Amazon lowlands (compare  
1026 section 3.3).

1027

1028 **5.5 Meteoric  $^{10}\text{Be}/^9\text{Be}$ -derived denudation rates compared to *in situ*-derived**

1029 **denudation rates in the Amazon**

1030 For a comparison between rates from *in situ*  $^{10}\text{Be}$  with those from the meteoric  $^{10}\text{Be}$   
1031 we first briefly repeat the types of rate information derived from both types, so that  
1032 true geologic causes can be distinguished from differences introduced by possible  
1033 methodological artefacts. Such comparison is possible for the Amazon basin, where  
1034 both methods have been applied. All meteoric Be data is from Wittmann et al.,  
1035 (2015), and the *in situ* rates from source areas and lowland areas, respectively, are  
1036 from Wittmann et al. (2009) and Wittmann et al. (2011a). We present

1037 –  $E_{\text{met}^*}$  that are meteoric-derived erosion rates from reactive  $^{10}\text{Be}$   
1038 concentrations (eq. 11) of high confidence that have been corrected for grain  
1039 size bias by means of depth integration using Al/Si in suspended loads  
1040 (section 5.3, Fig. 12C).

1041 –  $D_{\text{met}^*}$  that are meteoric  $^{10}\text{Be}/^9\text{Be}$ -derived denudation rates (Eq. 14) of high  
1042 confidence where the parameter  $f_{\text{reac}}^{9\text{Be}} + f_{\text{diss}}^{9\text{Be}}$  (Eq. 15) has been corrected for  
1043 grain size bias by means of depth integration using Al/Si in suspended loads  
1044 (section 5.4).

1045 –  $D_{\text{met}}$  that are meteoric  $^{10}\text{Be}/^9\text{Be}$ -derived denudation rates (Eq. 14) of lower  
1046 confidence that have not been corrected for grain size bias potentially  
1047 introduced by sorting that affects the  $^9\text{Be}_{\text{min}}/^9\text{Be}_{\text{reac}}$  ratio used in equation 14.

1048 –  $D_{\text{insitu}}$  that are *in situ*  $^{10}\text{Be}$ -derived denudation rates of high confidence from  
1049 source areas calculated from (Eq. 2) using the production rate scaling of the  
1050 source area.

1051 –  $D_{\text{insituFC}}$  that are *in situ*  $^{10}\text{Be}$ -derived denudation rates (Eq. 2) of high

1052 confidence from floodplain areas where the floodplain correction (section  
1053 3.1, Fig. 8) has been applied to account for zero additional *in situ* nuclide  
1054 production in the floodplain.

1055 – We further note that in floodplain samples all meteoric erosion ( $E_{\text{met}^*}$ ) and  
1056 denudation rates ( $D_{\text{met}^*}$ ,  $D_{\text{met}}$ ) are likely biased as meteoric  $^{10}\text{Be}$   
1057 concentrations are modified by both radioactive decay (Fig. 13A) or by  
1058 accumulation (Fig. 13B) of meteoric  $^{10}\text{Be}$  during floodplain storage.

1059

1060 We do not present  $E_{\text{met}}$  which are erosion rates (Eq. 11) based on reactive  $^{10}\text{Be}$   
1061 concentrations from either bedload or suspended sediment samples that are of low  
1062 confidence because they are not corrected for grain size bias introduced by particle  
1063 sorting.

1064

1065 Figure 15 shows the comparison of highest confidence rates where available.  $E_{\text{met}^*}$   
1066 and  $D_{\text{met}^*}$  are available at four locations within the Amazon basin, where river depth  
1067 profiles were sampled. At these sites,  $E_{\text{met}^*}$  and  $D_{\text{met}^*}$  are almost identical in three out  
1068 of four locations, testifying to the low fraction of chemical weathering from  
1069 dissolved loads in the basin that is discussed in section 3.3.

1070 For the Andes we calculated an area-weighted  $D_{\text{met}}$  of 0.44 mm/yr. The corresponding  
1071  $\overline{D_{\text{insitu}}}$  value is 0.19 mm/yr. Indeed, the same pattern, with  $D_{\text{met}}$  exceeding  $D_{\text{insitu}}$ ,  
1072 emerges for all large rivers draining the Andes (Upper Solimões, Madre de Dios,  
1073 Upper Madeira, Beni, Upper Mamoré). We can speculate this difference to reflect  
1074 true geologic causes: meteoric  $^{10}\text{Be}$  likely samples more rapidly eroding lithologies  
1075 such as shales that produce finer-grained sediment. Conversely, *in situ* denudation

1076 rates from coarse quartz reflect more erosion-resistant granitoid lithologies  
1077 (Wittmann et al, 2015).

1078 Rivers from the cratons (Negro, Branco, Brazilian Craton) yield low denudation rates  
1079 from both methods that are typical for slowly eroding cratonic landscapes.  $D_{\text{met}}$   
1080 range from 0.02 to 0.06 mm/yr. As in Andean rivers,  $D_{\text{met}}$  typically exceeds  $D_{\text{insitu}}$  by a  
1081 factor of ca. 2. Too-high  $D_{\text{met}}$  for the cratonic rivers are most likely due to a grain size  
1082 bias affecting the  ${}^9\text{Be}_{\text{min}}/{}^9\text{Be}_{\text{reac}}$  ratio (Eq. 14). However, denudation rates of both  
1083 methods are potentially compromised in these slowly eroding settings because  
1084 sediment storage may introduce another bias through radioactive decay (section 2.4  
1085 for *in situ*  ${}^{10}\text{Be}$ , Fig. 13A for meteoric  ${}^{10}\text{Be}$ ).

1086 In the central Amazon lowlands,  $D_{\text{met}^*}$  yields 0.17 mm/yr which compares to a  
1087  $\overline{D_{\text{insituFC}}}$  value of 0.18 mm/yr. This agreement contrasts with the observations from  
1088 the source areas, where  $D_{\text{met}}$  exceeds  $D_{\text{insitu}}$  by a factor of ca. 2. This shift in the  
1089 difference might be apparent: In the floodplain, meteoric  ${}^{10}\text{Be}$  concentrations can  
1090 be modified by both accumulation and decay during sediment storage (Figs. 13A,B).  
1091 Unlike  $D_{\text{insituFC}}$  that in the floodplain records the mean sediment production rate of  
1092 the source area, meteoric  ${}^{10}\text{Be}$  is more sensitive to storage.

1093

#### 1094 **5.6 Meteoric ${}^{10}\text{Be}/{}^9\text{Be}$ : outlook**

1095 We are only beginning to understand the exact mechanisms of how meteoric  ${}^{10}\text{Be}$   
1096 concentrations and  ${}^{10}\text{Be}/{}^9\text{Be}$  ratios are modified by floodplain processes and how  
1097 rivers transmit such signals to the sedimentary record. We have shown means to  
1098 correct for incomplete retentivity (by means of the  ${}^9\text{Be}$  normalization, Fig. 12B) and  
1099 grain size-dependent  ${}^{10}\text{Be}$  retention (by using depth-integrated Al/Si, Fig. 12C).



1100 Resolving the uncertainties associated with  $^{10}\text{Be}$  delivery from the atmosphere is  
1101 paramount. Potentially, however, this system can be applied to deriving paleo-  
1102 denudation rates in the terrestrial or marine sedimentary record, by using  $^{10}\text{Be}/^9\text{Be}$   
1103 in clays or reactive Be adsorbed onto e.g. Fe-Mn oxides. The latter approach can be  
1104 used to derive paleo-denudation rates from authigenic ocean sediments (von  
1105 Blanckenburg and Bouchez, 2014; von Blanckenburg et al., 2015). The resolution  
1106 with which changes in denudation as a function of climate change can be resolved  
1107 depends critically on the transport path of  $^9\text{Be}$  through basins in the dissolved and  
1108 the particulate forms and on the release of the latter in seawater (von Blanckenburg  
1109 and Bouchez, 2014; von Blanckenburg et al., 2015). Regardless, the entire, down to  
1110 upper Miocene sedimentary record can now be accessed for studying Earth surface  
1111 dynamics.

1112

## 1113 **6. SUMMARY AND SAMPLING GUIDELINES**

1114 We have analyzed the effects that sediment storage and deposition can have on the  
1115 *in situ* cosmogenic nuclide signal imprinted to sediment during erosion. When  
1116 sediment transport through lowland basins is “dynamic”, which is the case when  
1117 continuous exchange between channel sediment and sediment in the proximal  
1118 floodplain during lateral river movement takes place, geologically short floodplain  
1119 storage times (in the order of only several kyrs) result. These storage times are much  
1120 shorter than the about Myr half-life of long-lived cosmogenic nuclides such as  $^{10}\text{Be}$   
1121 and  $^{26}\text{Al}$ , and they are shielded from cosmic rays. Consequently, additional nuclide  
1122 production and also decay of *in situ*-produced nuclides during storage and

1123 deposition are limited, and a modification of the initial *in situ*-cosmogenic nuclide  
1124 signal inherited from the erosion process in the high-relief setting does not occur.  
1125 We find that large-scale sediment transport effectively averages-out source area  
1126 variability, and long-lived *in situ* nuclide concentrations, measured in lowland  
1127 sediment and treated using the nuclides' production rate in the source area, provide  
1128 a representative mean denudation rate of the source area. We show this for the  
1129 cases of the Amazon and Ganga basins. The conversion of these longer-term *in situ*-  
1130 cosmogenic nuclide-derived denudation rates to sediment fluxes (by using the  
1131 respective source areas and sediment density values) allows for the comparison to  
1132 modern, gauging-derived fluxes and to those derived from deltaic and off-shore  
1133 archives. From this comparison we find a remarkable temporal stability of sediment  
1134 fluxes over Holocene times to the present in these two basins.

1135

1136 This simple picture of non-changing long-lived *in situ*-produced nuclide  
1137 concentrations across floodplains is only challenged by 1) recycling of old, potentially  
1138 buried floodplain sediment during distal channel-floodplain interaction which is  
1139 coupled to 2) mixing of sediment with different provenance and grain sizes. For  
1140 example in the Amazon basin, the fine sand grain size originates in the Andes and is  
1141 not modified en route to the Amazon lowlands. Coarser sand, however, either  
1142 originates in the cratonic areas, where the slow erosion imprints a much higher  
1143 nuclide concentration, or is recycled from old lowland formations when the channel  
1144 avulses into distal, inherently older floodplain deposits. In this case the ratio of  
1145  $^{26}\text{Al}/^{10}\text{Be}$ , from its differential decay, may trace this admixing of recycled versus  
1146 "fresh" (in-channel source area-derived) material.

1147

1148 With the following *guidelines*, we aim to provide the reader with practical  
1149 recommendations for deriving *in situ*  $^{10}\text{Be}$ -based (or other long-lived cosmogenic  
1150 nuclides) denudation rates or sediment fluxes from lowland river sediment. These  
1151 guidelines summarize the findings from modeling approaches and field-based data:

1152 1) In lowland rivers with deep (10's of meters) actively migrating channels (e.g. >5  
1153 m/yr), the effect of nuclide production taking place at the floodplain surface is  
1154 minor compared to the vast majority of sediment that is shielded at depth before  
1155 remobilization, resulting in only small changes of the initial (source area-derived)  
1156 nuclide concentration. Sampling bedload from deep, laterally active rivers should  
1157 thus be preferred.

1158 2) Nuclide production during lowland transfer of sediment in shallow migrating  
1159 channels can be significant. But mostly such shallow floodplains are also  
1160 characterized by short sediment storage duration. Sampling bedload from rivers  
1161 with shallow floodplains is in order as long as sediment turnover is rapid. Thus,  
1162 we suggest estimating the ratio of ingoing to exported sediment fluxes, e.g. from  
1163 gauging data, before targeting specific areas for cosmogenic nuclide sampling.

1164 3) Rivers fed by rapidly eroding source areas are potentially more affected by  
1165 down-channel change as their initial source area-derived nuclide concentration is  
1166 low and thus sensitive to additional production. Unless sediment storage  
1167 duration is also short, more slowly eroding settings (0.1 to 0.5 mm/yr) should be  
1168 preferred.

1169 4) Entrainment of sediment from e.g. avulsions might reactivate old deeply stored  
1170 sediment in which long-lived nuclides might have decayed. In rivers in which

1171 lateral migration is limited to the active channel belt, modification of the source  
1172 area signal by such incorporation of deeply stored old floodplain material is less  
1173 likely.

1174 5) In order to assess the influence of old, deeply stored sediment deposits, another  
1175 long-lived nuclide, such as  $^{26}\text{Al}$ , should be monitored along with  $^{10}\text{Be}$ , as  $^{26}\text{Al}/^{10}\text{Be}$   
1176 ratios are indicative of potential admixing of formerly buried floodplain material.  
1177 If highly-weathered (i.e. cratonic) areas in the basin exist, a reconnaissance study  
1178 for the distribution of nuclide concentrations in different grain sizes might be  
1179 necessary.

1180 Finally, we point at the opportunities arising from new nuclide systems. *In situ*-  
1181 produced  $^{14}\text{C}$ , measured in quartz, may have the potential to estimate sediment  
1182 storage times over one to two of its half-life (ca. 5-10 kyrs). Meteoric  $^{10}\text{Be}$  can be  
1183 measured in fine-grained sediment where the presence of quartz is not a  
1184 requirement. The meteoric  $^{10}\text{Be}$  concentration alone yields an erosion rate, provided  
1185 a representative concentration can be inferred from suspended sediment depth  
1186 profiles and Al/Si. The flux balance of  $^{10}\text{Be}$  and its increase in floodplains can be used  
1187 to evaluate sediment residence times. Further, the adsorbed and reactive Be should  
1188 be extracted chemically from reactive phases and the stable  $^9\text{Be}$  measured, such that  
1189 erosion rates, denudation rates, and the degree of weathering can be simultaneously  
1190 determined from the meteoric  $^{10}\text{Be}/^9\text{Be}$  system.

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1195

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**Table 1: Comparison of model input parameters and results for the Amazon and Ganga basins for long-lived *in situ* <sup>10</sup>Be**

Model type <sup>a</sup> using each parameter		Parameters used for model calculation	Amazon basin <sup>b</sup>	Ganga basin <sup>c</sup>
Box	Numeric	Initial source-area derived <sup>10</sup> Be concentration	<sup>10</sup> Be <sub>(0)</sub> : 6.57x10 <sup>4</sup> at/g(Qtz)	<sup>10</sup> Be <sub>(0)</sub> : 2.21x10 <sup>4</sup> at/g(Qtz)
Box	Numeric	Initial sediment flux	QS <sub>insitu</sub> = 380 Mt/yr (A <sub>source</sub> = 8.5x10 <sup>5</sup> km <sup>2</sup> )	QS <sub>insitu</sub> = 550 Mt/yr (A <sub>source</sub> = 1.8x10 <sup>5</sup> km <sup>2</sup> )
Box	Numeric	Channel lateral migration rate	5 ± 2 m/yr	70 ± 35 m/yr
Box	Numeric	Active floodplain width	35 ± 15 km	40 ± 15 km
Box	Numeric	Active floodplain depth	23 ± 10 m	10 ± 5 m
Box	-	Sediment transfer time	14 kyr	1.4 kyr
Box	Numeric	Average transfer distance	3000 km	1000 km
-	Numeric	Floodplain density	1700 kg/m <sup>3</sup> (Dunne et al., 1998)	1500 kg/m <sup>3</sup>
-	Numeric	Channel sinuosity	1.15 ± 0.2 (avg. from Mertes et al., 1996)	1.3 ± 0.2
-	Numeric	Fraction of QS in channel in sand-sized fraction	0.15 ± 0.05 (Dunne et al., 1998)	0.35 ± 0.15
-	Numeric	Fraction of QS in floodplain in sand-sized fraction	0.1 ± 0.05 (Dunne et al., 1998)	0.1 ± 0.05
-	Numeric	Floodplain aggradation	steady-state (Mertes et al., 1996; Bouchez et al., 2012)	0.5 ± 0.25 mm/yr
Increase in <sup>10</sup> Be concentration relative to <sup>10</sup> Be <sub>(0)</sub> (%)			Box: 1.8	1.5
(also see Fig. 6)			Numeric: 1.7	6.3 <sup>d</sup>

<sup>a</sup>The box model is based on Wittmann and von Blanckenburg (2009) and the numeric model is based on Lauer and Willenbring (2010).

<sup>b</sup>If no reference is given next to the parameter, the reference used is Wittmann and von Blanckenburg (2009).

<sup>c</sup>If no reference is given next to the parameter, the reference used is Lupker et al. (2012a).

<sup>d</sup>Note that our result is lower than that obtained by Lupker et al. (2012a), because we used a single channel setup, whereas Lupker used four tributary input streams.

**Table 2: Summary of measured *in situ*-derived denudation rates (upper part) and measured gauging-derived sediment fluxes (lower part).**

AMAZON BASIN	Subbasin	Source or total basin area ( $\times 10^5$ km <sup>2</sup> )	<i>In situ</i> -derived cosmogenic data <sup>a</sup>				C <sub>v</sub> for denudation rates <sup>d</sup>
			n (>100 km <sup>2</sup> ) <sup>b</sup>	Area-weighted D <sub>insitu</sub> $\pm$ 1 $\sigma$ SD (mm/yr)	Measured D <sub>insitu</sub> at largest outlet (mm/yr) <sup>c</sup>	Area-weighted QS <sub>insitu</sub> $\pm$ 1 $\sigma$ SD (Mt/yr)	
<b>Andes:</b> Subbasins or largest outlets at mountain front	Beni	0.68	38	0.37 $\pm$ 0.20	0.4 - 0.86	66 $\pm$ 35	
	Grande	0.60	17	0.38 $\pm$ 0.14	0.10 - 0.57 <sup>g</sup>	59 $\pm$ 22	
	Solimões <sup>e</sup>	6.3	7	0.16 $\pm$ 0.11	0.27	257 $\pm$ 210	
entire sampled Andes draining to Amazon		7.58		<b>0.19 <math>\pm</math> 0.18</b>		<b>380 <math>\pm</math> 350</b>	<b>0.55</b>
<b>Lowlands</b> at Óbidos (using sampled source area for conversion from D to QS) <sup>f</sup>		7.58	17		<b>0.18 <math>\pm</math> 0.03</b>	<b>350 <math>\pm</math> 70</b>	<b>0.19</b>
<i>Lowlands at Óbidos (no floodpl. corr., using total basin area for conversion from D to QS)<sup>f</sup></i>		50.9	17		<b>0.11 <math>\pm</math> 0.03</b>	<b>1480 <math>\pm</math> 370</b>	
<b>GANGA BASIN</b>							
<b>Himalayas:</b> Subbasin or largest outlets at mountain front	Upper Ganga	0.32	3	1.22 $\pm$ 0.60	1.9	100 $\pm$ 50	
	Yamuna	0.096	18	0.76 $\pm$ 0.56	0.56	20 $\pm$ 14	
	Karnali	0.46	3	1.19 $\pm$ 0.75	0.8 - 1.9	140 $\pm$ 90	
	Narayani	0.34	25	1.54 $\pm$ 0.96	1.0 - 1.7	140 $\pm$ 70	
	Kosi	0.62	8	0.69 $\pm$ 0.78	0.4 - 0.8	110 $\pm$ 125	
entire sampled Himalayas draining to Ganga		1.84		<b>1.14 <math>\pm</math> 0.82</b>		<b>550 <math>\pm</math> 390</b>	<b>0.82</b>
<b>Lowlands</b> at Harding Bridge (using sampled source area for conversion from D to QS) <sup>h</sup>		1.84	8		<b>0.82 <math>\pm</math> 0.27</b>	<b>390 <math>\pm</math> 130</b>	<b>0.31</b>
<i>Lowlands at Harding Bridge (no floodpl. corr.; using total basin area for conversion from D to QS)<sup>h</sup></i>		8.73	8		<b>0.33 <math>\pm</math> 0.50</b>	<b>760 <math>\pm</math> 1140</b>	

			Gauging-derived suspended sediment data <sup>i</sup>					
AMAZON BASIN	Subbasin	Source or total basin area ( $\times 10^5$ km <sup>2</sup> )	n (>100 km <sup>2</sup> ) <sup>b</sup>	Catchment-wide QS <sub>gaug</sub> (Mt/yr) <sup>j</sup>	Measured QS <sub>gaug</sub> at largest outlet (Mt/yr)	Area-weighted E <sub>gaug</sub> (mm/yr)	C <sub>v</sub> for erosion rates <sup>d</sup>	
<b>Andes:</b> Subbasins or largest outlets at mountain front	Beni	0.68	13	210 ± 185	160 - 300	1.20 ± 1.06		
	Grande	0.60	15	210 ± 150	125 - 180	1.33 ± 0.97		
	Solimões <sup>e</sup>	4.7	14	630 ± 480	553	0.33 ± 0.25		
entire Andes draining to Amazon		6.0		<b>1050 ± 820</b>	<b>840 - 890</b>	<b>0.47 ± 0.90</b>	<b>0.98</b>	
<b>Lowlands</b> at Óbidos (using sampled source area for conversion from QS to E) <sup>f</sup>		6.0	6	<b>900 ± 310</b>	<b>560 - 1320</b>	<b>0.58 ± 0.20</b>	<b>0.34</b>	
<i>Lowlands at Óbidos (no floodplain corr.; using total area for conversion from QS to E)<sup>f</sup></i>		50.9	6			<b>0.07 ± 0.02</b>		
<b>Amazon Delta:</b> 550 - 1030 Mt/yr measured from <sup>210</sup> Pb activity profiles (Nittrouer and Kuehl, 1995).								
GANGA BASIN								
<b>Himalayas:</b> Subbasin or largest outlets at mountain front	Upper Ganga	0.32	3	12 ± 13	10.1	0.14 ± 0.15		
	Yamuna	0.096	2	18	18	0.73 ± 0.73		
	Karnali	0.46	8	110 ± 80	76 - 210	0.89 ± 0.66		
	Narayani	0.34	13	100 ± 50	97 - 120	1.09 ± 0.60		
	Kosi	0.62	8	130 ± 65	170 - 180	0.81 ± 0.40		
entire Himalayas draining to Ganga		1.84		<b>370 ± 230</b>		<b>0.80 ± 0.55</b>	<b>0.75</b>	
<b>Lowlands</b> at Harding Bridge (using sampled source area for conversion from QS to E) <sup>k</sup>		1.84	8	<b>390 ± 130</b>	<b>200 - 550</b>	<b>0.81 ± 0.27</b>	<b>0.33</b>	
<i>Lowlands at Harding Bridge (no floodplain corr.; using total area for conversion from QS to E)<sup>k</sup></i>		8.73	8			<b>0.17 ± 0.06</b>		



**Bengal Delta:** 320 - 920 Mt/yr measured from stratigraphic profiles (Wasson, 2003; Goodbred and Kuehl, 2000)<sup>l</sup>.

**Bengal Fan:** 510 Mt/yr from Métivier and Gaudemer (1999)<sup>l</sup>.

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*Note that uncertainty given gives the data spread ( $1\sigma$  std. dev. of the dataset). We chose to present standard deviations and not analytical uncertainties because the latter is not available for  $QS_{\text{gaug}}$ , they would not represent natural variations and would imply a precision that is not appropriate when comparing different methods. In situ-derived denudation rates were calculated using a total SLHL production rate of  $3.75 \text{ at/g}_{(\text{Qtz})}/\text{yr}$  and the scaling model of Dunai (2000) (see supplement A for details).*

<sup>a</sup>All values are area-weighted  $\pm 1\sigma$  std. dev., calculated according to Eq. 7. All lowland data is floodplain corrected (i.e. using the sampled source area for calculations), except where indicated. When no floodplain-correction was applied, the total basin area was used in Eq. 6 for conversion between D and QS.

<sup>b</sup>References for datasets used are displayed in Fig. 3. Grande dataset by Kober et al. (2015) not included (see text).

<sup>c</sup>A range is given if more than one measurement for the largest outlet is available to display variability.

<sup>d</sup> $C_v$  gives the coefficient of variation and is the ratio of the standard deviation to the mean of each dataset.

<sup>e</sup>In situ tributaries contained: Napo, Marañon, Ucayali; Gauging tributaries: also Huallaga (see Fig. 3).

<sup>f</sup>Comprises lowland data at Manacapuru, Iracema, Parintins, lower Madeira, Obidos, and Varzea data; for entire dataset see Tables 2,3 in Wittmann et al. (2011a).

<sup>g</sup>Two values measured for the Grande at Abapo deviate strongly (Wittmann et al., 2009; Kober et al., 2015). Note that a area-weighted  $D_{\text{insitu}}$  of the Grande basin is  $0.11 \pm 0.27 \text{ mm/yr}$  ( $n=71$ ).

<sup>h</sup>Comprises lowland data of lower Karnali, Kosi and Narayani and all data from lower Ganga below Varanasi to Harding Bridge; c.f. Table 1 in Lupker et al. (2012a).

<sup>i</sup>For measured  $QS_{\text{gaug}}$ , the range is given (for largest outlets) to display variability in data;  $E_{\text{gaug}}$  are area-weighted  $\pm 1\sigma$  std. dev. All lowland data is floodplain corrected (i.e. using the sampled source area for calculations), except where indicated (see Eqs. 8,9 for calculation of  $E_{\text{gaug}}$ ). No dissolved loads were added.

<sup>j</sup>Calculated using upscaling, i.e. the sum of all measured QS values was multiplied by the largest drainage area and then divided by the sum of all areas.

<sup>k</sup>Comprises measured QS data at Harding Bridge (cf. Lupker et al. (2012a)).

<sup>l</sup>Values are for Ganga only. Brahmaputra contribution was removed by assuming a 40% sediment flux contribution of the Ganga relative to the Brahmaputra.

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1213

FIGURE CAPTIONS

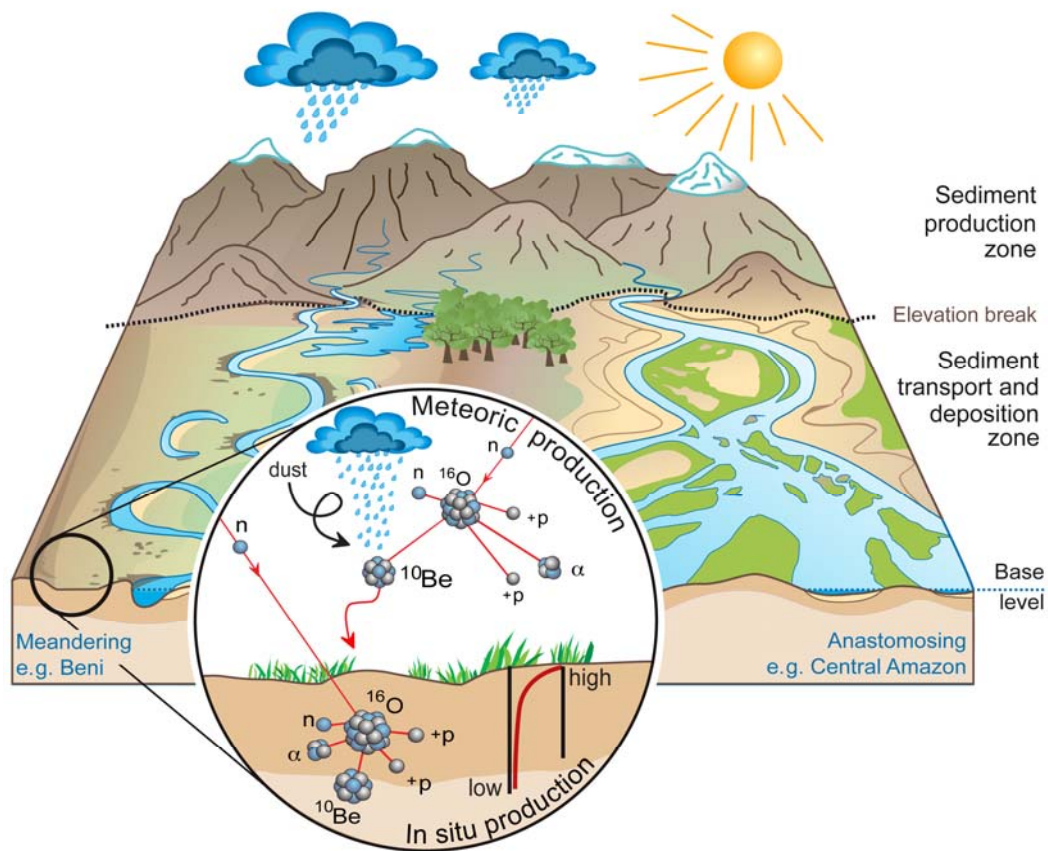


Fig. 1: Sediment routing from the zone of sediment production (in high mountain settings) to transport and storage in low-relief floodplain regions. The left-hand system is a lowland river that is migrating (e.g. the large Beni River in Bolivia), and the right-hand system is anastomosing (e.g. the Amazon River). Both types store and permanently release sediment that mixes the cosmogenic nuclides it carries with those contained in the main stream. The main production pathways of “meteoric” (i.e. within the atmosphere) and “*in situ*” (i.e. within the mineral grain) cosmogenic  $^{10}\text{Be}$  are shown in the inset. Both varieties of  $^{10}\text{Be}$  are mainly produced from spallation reactions of target atoms (e.g.  $^{16}\text{O}$ ) that are hit by high-energy cosmic rays (neutrons, protons). Secondary cosmic rays (protons, neutrons, and very light particles, such as muons) and cosmogenic nuclides are the result. *In situ* production is high at Earth’s surface and decreases with increasing depth through cosmic ray attenuation.

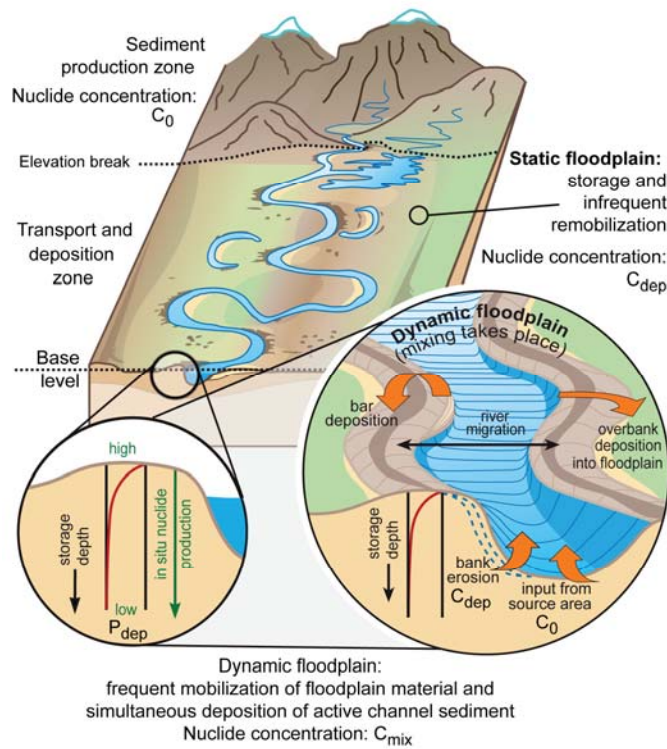


Fig. 2: Illustration of “dynamic floodplain” and “static” cases, and the nuclide concentrations in the participating compartments. In the static case, sediment is stored (nuclide concentration:  $C_{dep}$ ) and eventually eroded and re-incorporated into the main channel ( $C_{mix}$ ). In the case of the dynamic floodplain, channel- floodplain interaction leads to permanent remobilization and deposition due to the proximity to the main channel. The right-hand blow up shows parameters that influence the mixing between sediment in the channel and floodplain material. In the main channel, source area-derived material from close to the mountain front ( $C_0$ ) is mixed ( $C_{mix}$ ) with floodplain sediment eroded from bank deposits ( $C_{dep}$ ). At the same time, sediment is deposited in the floodplain by bar- and overbank deposition ( $C_{mix}$ ).

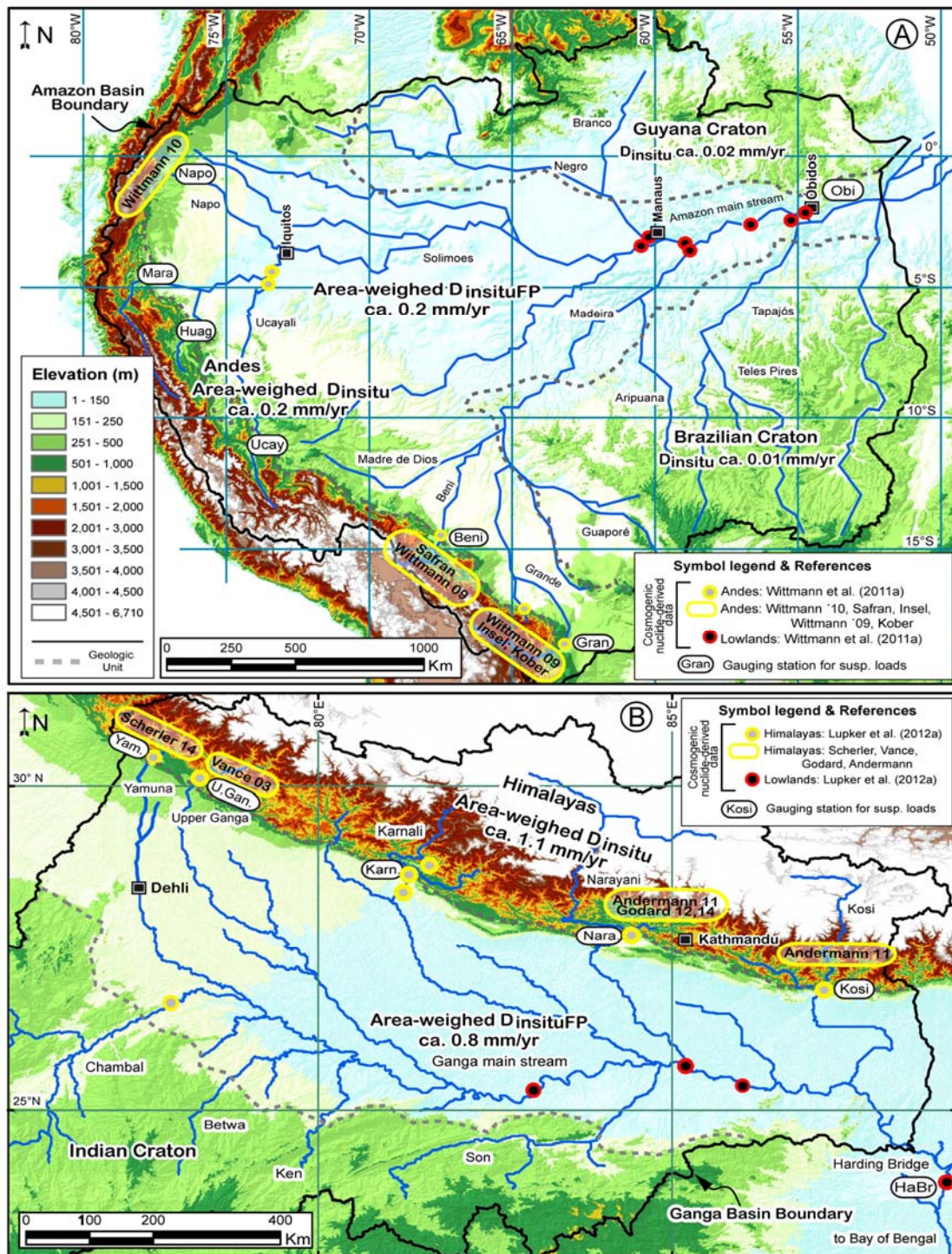


Fig. 3: Topographic and fluvial drainage map of the Amazon (A) and Ganga (B) basins. Elevation is scaled using identical color schemes in both basins. Yellow-rimmed symbols denote cosmogenic-nuclide sampling locations (only basins > 100 km<sup>2</sup> are shown) and first author's name of studies (or first authors name and year for ambiguous studies) that were used for compilation (Table 2). Mean source area  $D_{insitu}$  and floodplain-corrected (see section 3.1) lowland  $D_{insituFP}$  are given. White ellipses denote gauging station for suspended sediment (only basins >100 km<sup>2</sup> are shown; for compilation see Table 2). *Refs. for gauging data in*

Amazon basin are: Napo- Armijos et al. (2013); Mara = Marañon at Borja and Huag = Huallaga at Chazuta- Pepin et al. (2013); Ucay = Ucayali at Pucallpa- Santini et al. (2014); Beni = Beni at Rurrenabaque and larger stations within upper Beni basin- Aalto et al. (2006); Pepin et al. (2013); Gran = Grande at Abapo and larger stations within upper Grande basin- Guyot et al. (1996); Aalto et al. (2006); Pepin et al. (2013). In the Amazon lowlands at Obi = Óbidos, we compiled data from Meade (1985); Dunne et al. (1998); Guyot et al. (2005); Filizola and Guyot (2009); Martinez et al. (2009). Refs. for gauging data in Ganga basin are: Yam. = Yamuna- Jha et al. (1988); U. Gan. = Upper Ganga- Abbas and Subramanian (1984); Karn. = Karnali - Andermann et al. (2012); Lupker et al. (2012a); Nara. = Narayani- Sinha and Friend (1994); Gabet et al. (2008); Andermann et al. (2012); Lupker et al. (2012a); Kosi- Sinha and Friend (1994); Andermann et al. (2012); Lupker et al. (2012a). Note that the work by Lupker et al. (2012a) comprises a secondary data source for several gauging stations, including HaBr = Harding Bridge station that is located downstream of the Hooghly diversion.

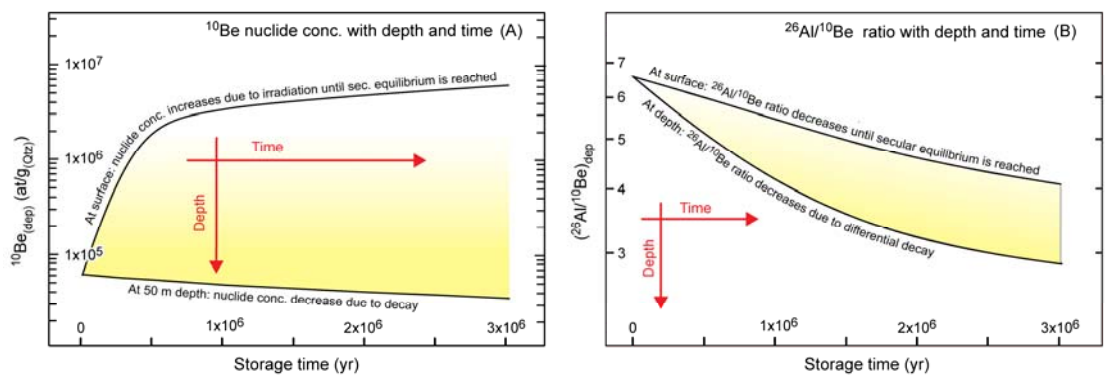


Fig. 4 A): Example of time- and depth dependent evolution of lowland *in situ*  $^{10}\text{Be}$  concentrations with storage time in a static deposit. The examples are calculated for an initial  $^{10}\text{Be}$  concentration  $^{10}\text{Be}_0$  of  $6 \times 10^4$  at/g<sub>(Qtz)</sub> and further irradiation at a total production rate of 3.75 at/g<sub>(Qtz)</sub>/yr and simultaneous decay. At the surface of the floodplain, ingrowth of the nuclide will occur with time, until a secular equilibrium between production and decay is attained. When stored deeply, decay becomes significant (lower curve calculated for a storage depth of 50 m, where further irradiation is virtually absent due to complete shielding). B): Example of the time- and depth-dependent evolution of the  $^{26}\text{Al}/^{10}\text{Be}$  ratio in this static case, from an initial  $^{26}\text{Al}/^{10}\text{Be}$  ratio of 6.5, which is the surface production ratio between the two nuclides. Due to the faster decay of  $^{26}\text{Al}$ , the ratio of the nuclides is more sensitive to storage than single nuclide concentrations.

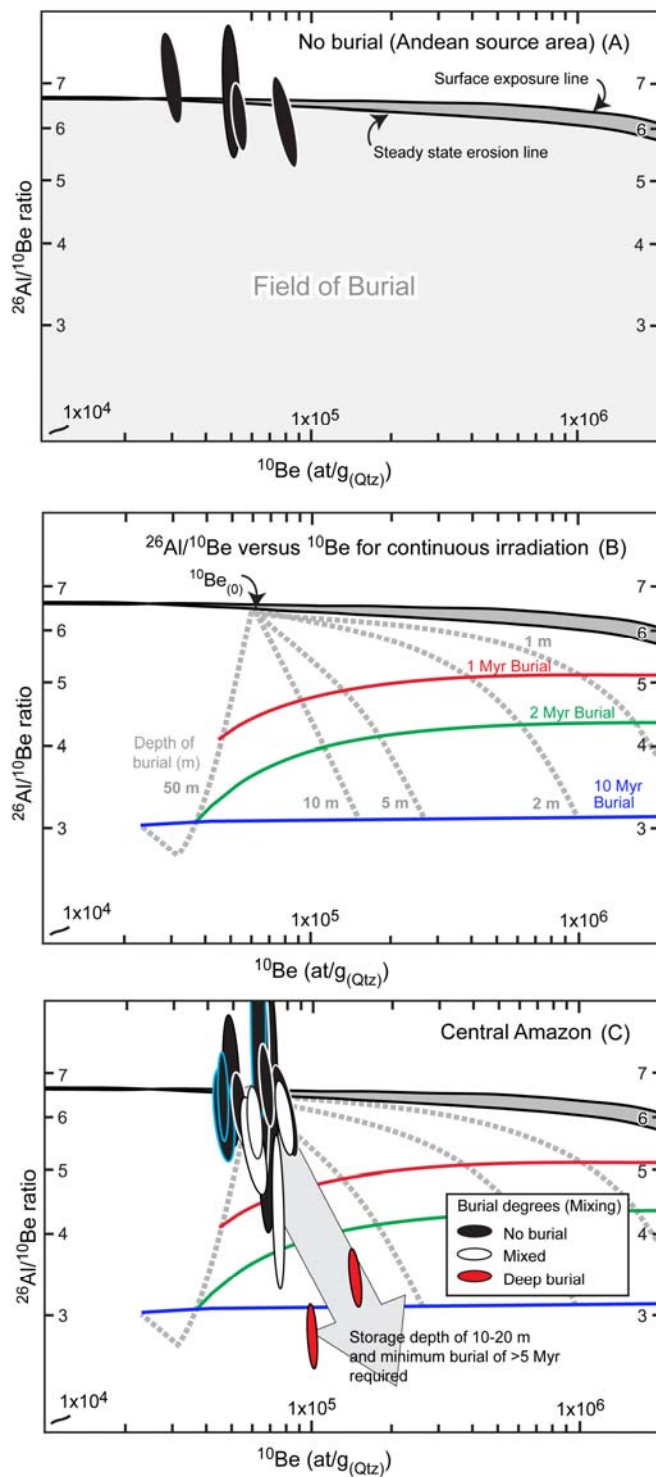


Fig. 5: The change in  $^{26}\text{Al}/^{10}\text{Be}$  ratio versus the  $^{10}\text{Be}$  concentration in quartz with duration and depth of burial (the “erosion island” plot, Lal (1991); showing the constant exposure and steady state erosion lines), with examples from the Amazon basin. A): Samples from the Andean source areas (black ellipses) have  $^{26}\text{Al}/^{10}\text{Be}$  ratios of around 6.5 and plot onto the steady state erosion line. These samples have never been buried. Samples that plot below the line would have been buried. B): When the overburden is not entirely shielding the sample from cosmic irradiation (e.g. in a shallow floodplain), the erosion island plot is calculated assuming simultaneous nuclide production and decay. A given  $^{26}\text{Al}/^{10}\text{Be}$  ratio then represents a burial age at a given depth. Contour lines of storage depth (stippled grey lines) are shown starting from a  $^{10}\text{Be}_0$  (representative for a given

family of samples). C): Burial-exposure history for the lower Amazon River (modified from Wittmann et al., 2011b). In the lower Amazon River, samples comprise mixtures of sediment with no previous burial (from Andean sources given in black or black with blue rim for visibility) and deeply buried sediment (red ellipses), resulting in mixed samples (open ellipses). If the initial  $^{10}\text{Be}$  concentration of  $^{10}\text{Be}_0$  is known or can be inferred, minimum storage depths and burial durations can be inferred.

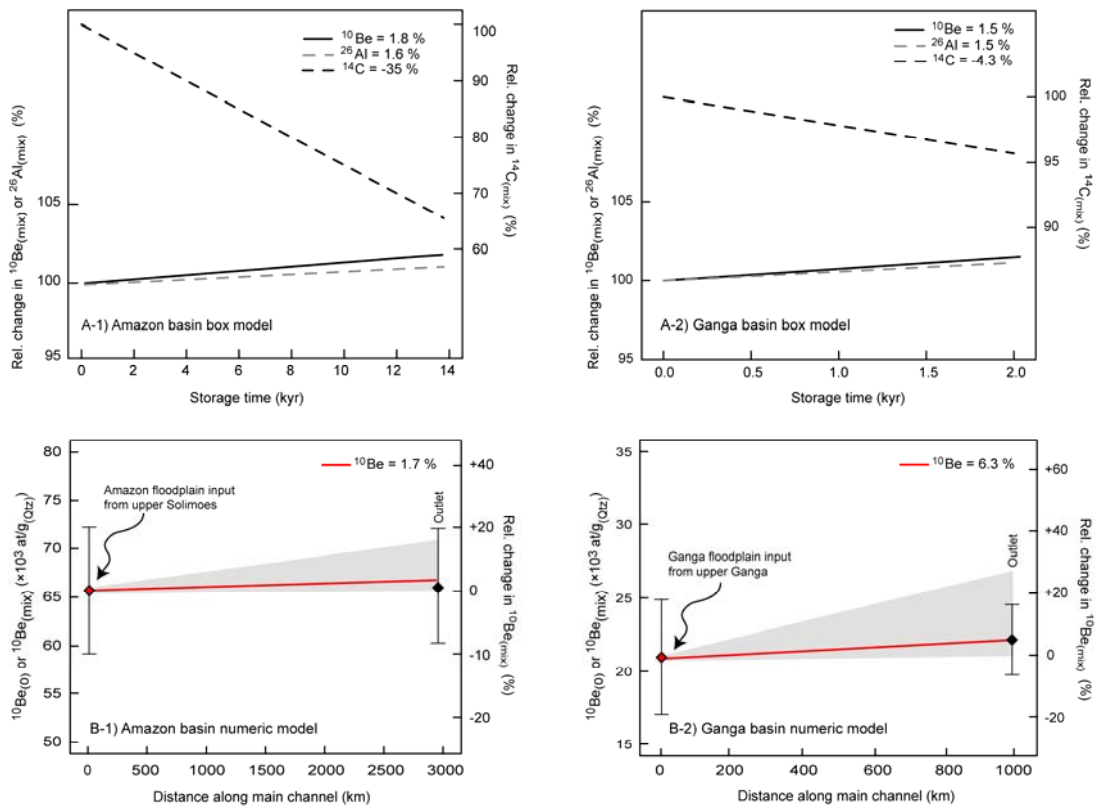


Fig. 6, Box (upper panels) and numeric (lower panels) models of relative changes in lowland river sediment nuclide concentrations  $C_{\text{mix}}$ . A-1,2): Results of the model of Wittmann and von Blanckenburg (2009) for the Amazon (A-1) and Ganga (A-2) rivers (using their eqs. 2-4). Long-lived *in situ* ( $^{10}\text{Be}$ ,  $^{26}\text{Al}$ ) nuclide concentration changes are shown on the left Y-axis, and *in situ*  $^{14}\text{C}$  nuclide concentration changes are shown on the right Y-axis. Relative changes in nuclide concentrations as a function of total storage time (kyr) in a dynamic floodplain were calculated for all three nuclides relative to the upstream concentration  $^{10}\text{Be}_0$  (see Table 1). The differences in the change of nuclide accumulation are mainly due to the different half-lives of the three nuclides. B-1,2): Results of the model by Lauer and Willenbring (2010), using their eq. 17, as a function of total distance along the main channel (km). Red line indicates change of  $^{10}\text{Be}$  signal along the floodplain within grey  $1\sigma$  uncertainty space, and red and black diamonds give the real measured  $^{10}\text{Be}_0$  and  $^{10}\text{Be}_{\text{mix}}$  (at/g<sub>(Qtz)</sub>) with their analytical uncertainties, respectively. Note that we used the same input parameters as Lupker et al. (2012) for the Ganga (Table 1), but using a single-channel system (i.e. having only one major tributary draining the source area). The Amazon system was also modeled as being a single channel from the Andes to the mouth. Note that both model types were (re-)calculated using the same total SLHL production rate of 3.75 at/g/yr for  $^{10}\text{Be}$  and muonic parametrization (Braucher et al., 2011) (for details see supplement A).

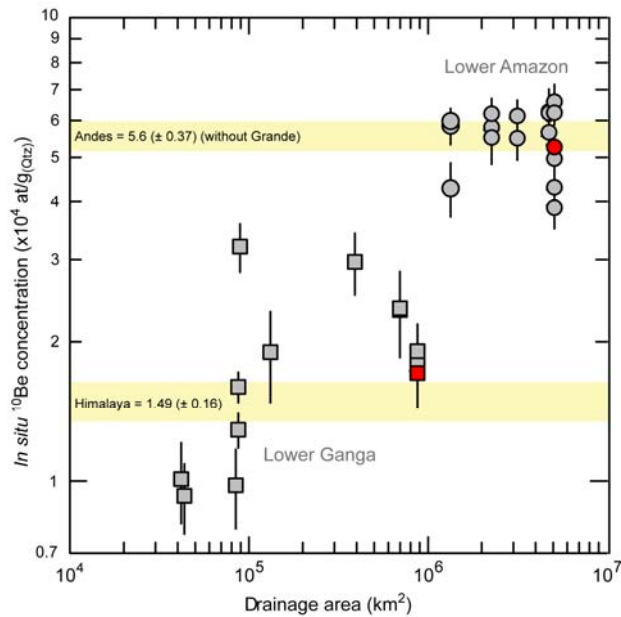


Fig. 7: *In situ*  $^{10}\text{Be}$  nuclide concentrations ( $\times 10^4$  at/g<sub>(Qtz)</sub>) and associated analytical  $1\sigma$  uncertainties measured in lowland river sediment of the Ganga and Amazon Rivers. The Amazon lowland dataset comprises only fine-grained sediment (n=18) and the Ganga lowland dataset (including all floodplain data, i.e. also data from lower Karnali, Narayani, and Kosi rivers, n = 14) is corrected for Indian craton input. Red symbols give area-weighted nuclide concentrations for the lowland areas; yellow bars give area-weighted ( $\pm 1\sigma$  analytical uncertainty) Andean and Himalayan source area data of catchments  $> 100$  km<sup>2</sup>. Andean data is from Safran et al. (2005); Wittmann et al. (2009); Insel et al. (2010); Wittmann et al. (2010); Wittmann et al. (2011a); the Grande dataset by Kober et al. (2015) is not shown (average is ca.  $40 \times 10^4$  at/g<sub>(Qtz)</sub>); upper Nepalese Himalaya data is from Vance et al. (2003); Wobus et al. (2005); Andermann (2011); Godard et al. (2012); Lupker et al. (2012a); Godard et al. (2014); Scherler et al. (2014).



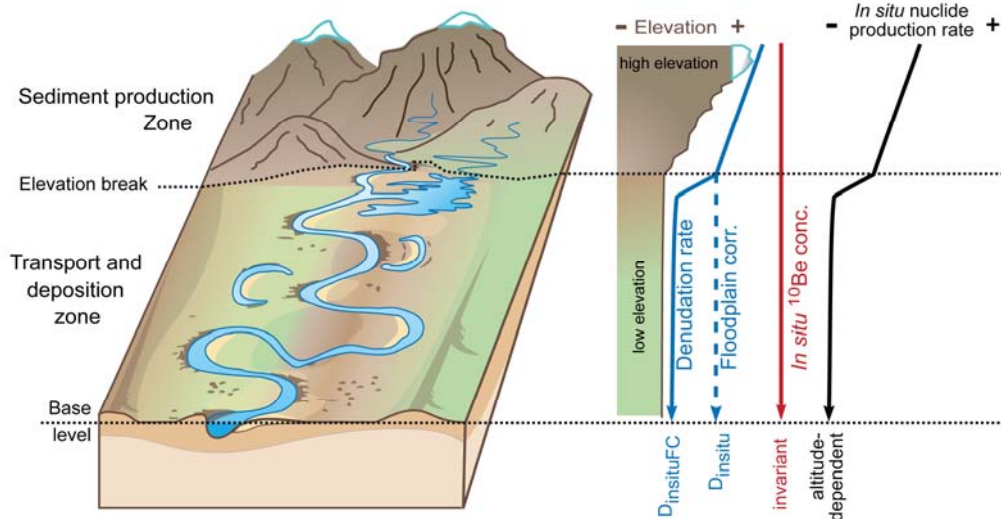


Fig.

8: Concept of the “floodplain correction” of denudation rates. The *in situ* nuclide production rate (black curve) is high in high-altitude mountain regions, and low in low-altitude floodplain regions. Long-lived *in situ* nuclide (e.g.  $^{10}\text{Be}$ ) concentrations of sediment transported along these domains are shown here as invariant (red curve). If the production rates  $P_{FC}$  of the source area are used, the denudation rate of the source area results ( $D_{in situ FC}$ ; blue dashed curve), whereas if the production rate of the whole basin above the sampling point is used, denudation rates integrate over the entire basin area and are thus apparent rates.

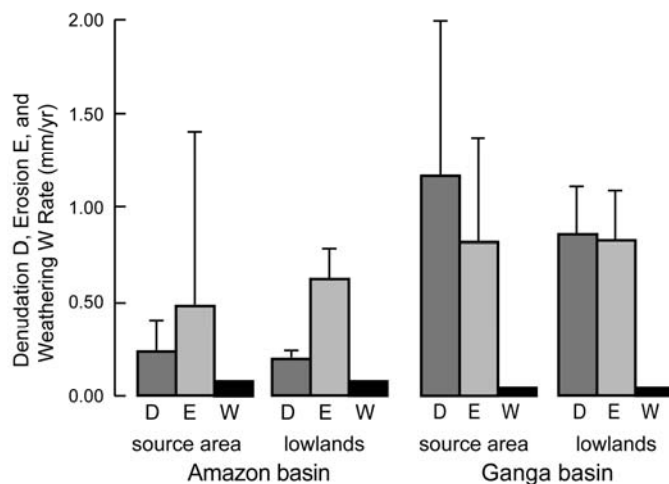


Fig. 9: Longer-term area-weighted denudation rates  $D$  from *in situ*  $^{10}\text{Be}$ , area-weighted modern erosion rates  $E$  from gauging, and published silicate weathering rates  $W$  (in mm/yr) across the Amazon and Ganga basins from source area to the lowlands. For references see text. Uncertainties denote the  $1\sigma$  std. dev. for the  $D$  and  $E$  datasets, and a 10% uncertainty was assumed for the  $W$  dataset.

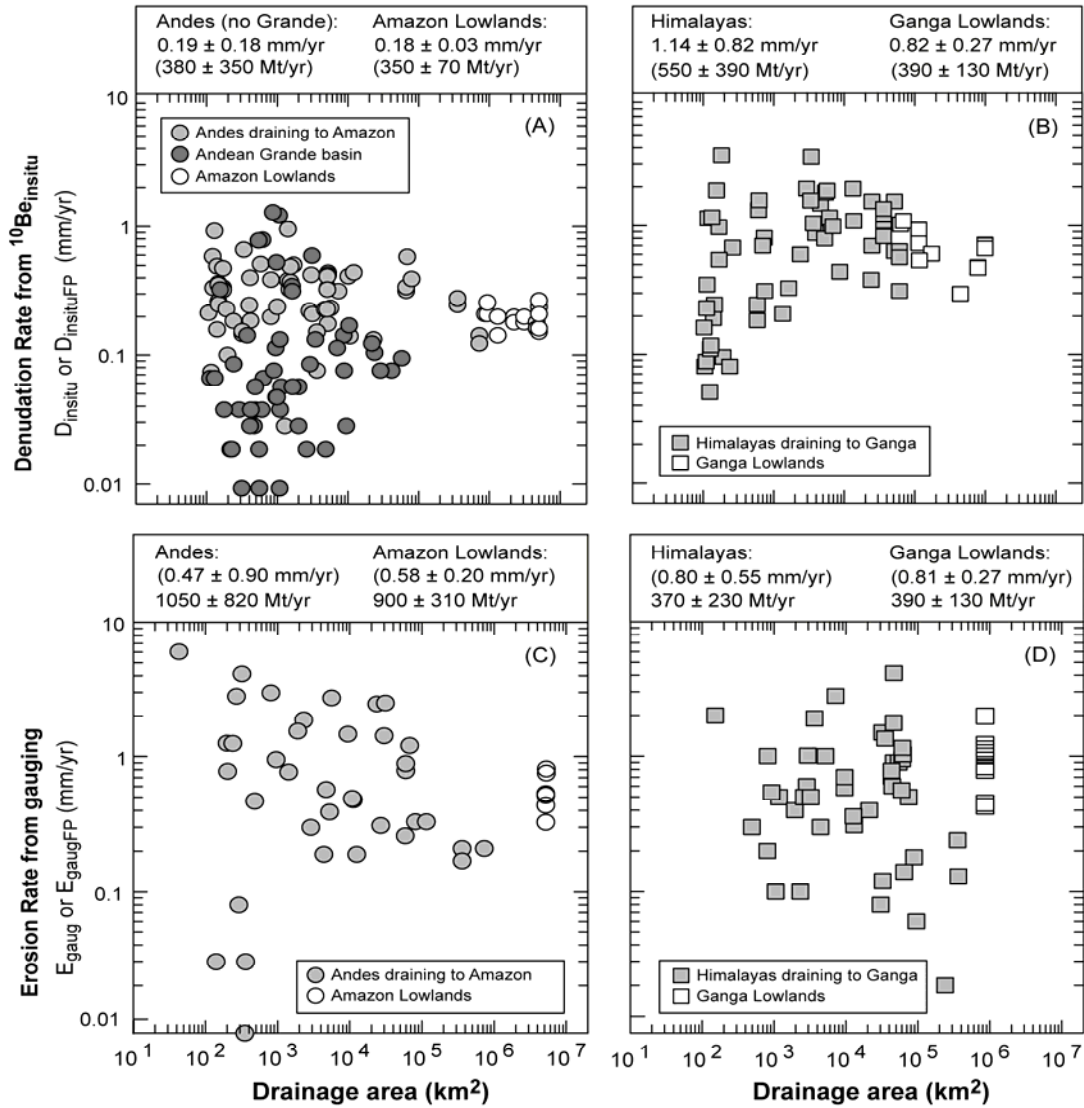


Fig. 10: Denudation from *in situ* cosmogenic <sup>10</sup>Be (A, B) and erosion rates from gauging (C, D) (all in mm/yr) from the source areas to the lowlands for the Amazon and the Ganga basins versus drainage area (km<sup>2</sup>, note that we omitted basins <100 km<sup>2</sup> to exclude small-scale effects). All lowland denudation and erosion rates are floodplain-corrected according to equations 6 and 10, respectively. Area-weighted denudation/erosion rates/sediment fluxes and the 1σ std. dev. of each dataset are shown above each panel. Converted values are shown in brackets, using the surface areas given in Table 2. For references see Fig. 3 and for details see Table 2.

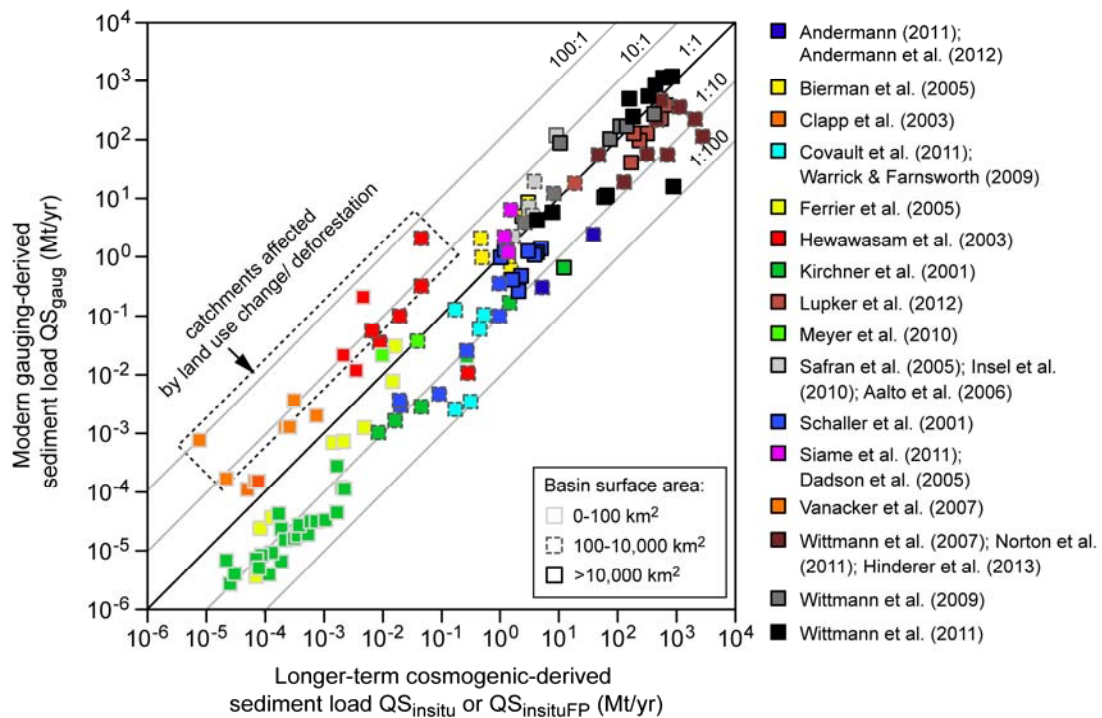


Fig. 11: Modern gauging-derived versus longer-term cosmogenic nuclide-derived sediment load data from approximately the same locations ( $n=129$ ), color-coded according to publication source (last author is always the reference for gauging dataset where applicable, i.e. if no second reference for gauging data is given, we rely on the compilation or measurement for gauging data in the given reference). Basin surface area ( $\text{km}^2$ ) is coded by rim color for small ( $0-100 \text{ km}^2$ ), medium ( $100-10000 \text{ km}^2$ ), and large catchments ( $>10000 \text{ km}^2$ ). For references on gauging data in Amazon and Ganga basins see Fig. 3. The datasets from the highly anthropogenized settings (Sri Lanka and Paute basin from Hewawasam et al. (2003); Vanacker et al. (2007), respectively) are highlighted by the dashed box. Note that the smaller Alpine basins, although having high sediment flux, display larger disagreement between  $Q_{S_{\text{gaug}}}$  and  $Q_{S_{\text{insitu}}}$ , whereas all other larger rivers agree reasonably well.

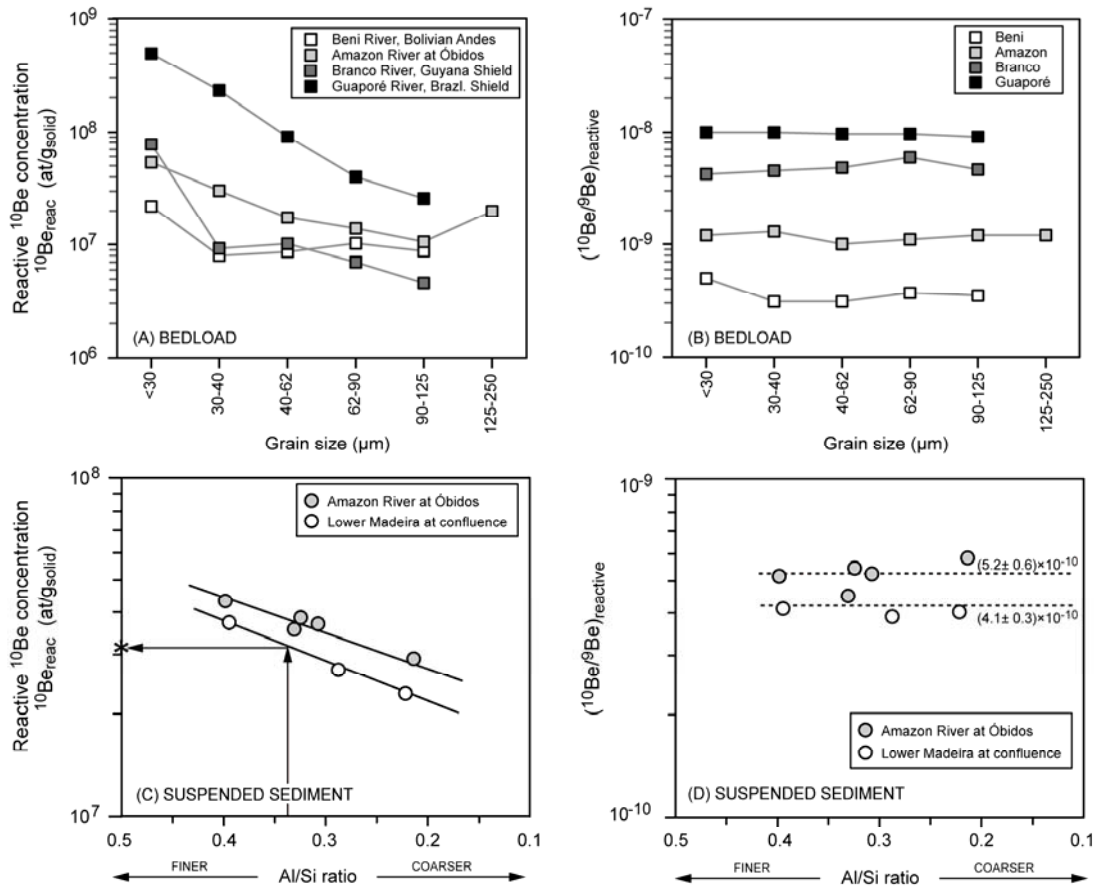


Fig. 12: The grain size-dependence of reactive (adsorbed and precipitated) meteoric  $^{10}\text{Be}$  concentrations on grain size (A, C) and the removal of this dependence by normalizing over reactive  $^9\text{Be}$  (B) or by linear regression using a representative Al/Si ratio as grain size indicator (C). A)  $^{10}\text{Be}_{\text{reac}}$  (at/g<sub>solid</sub>) and B)  $^{10}\text{Be}/^9\text{Be}$  ratio from bedload of different Amazon rivers (both modified from von Blanckenburg et al. (2012); data from Wittmann et al. (2012)), C)  $^{10}\text{Be}_{\text{reac}}$  (at/g<sub>solid</sub>) and D)  $^{10}\text{Be}/^9\text{Be}$  ratios (with mean and  $1\sigma$  std. dev. of each dataset) from suspended sediment in the lower Madeira and the Amazon at Óbidos, sampled at various depths in the river, versus Al/Si ratio (cosmogenic data from Wittmann et al. (2015) and Al/Si data from Bouchez et al. (2011b); both quantities measured on splits of the same sample). The Al/Si ratio is a proxy for grain size (Galy et al., 2008; Bouchez et al., 2011a). Note that when knowing a depth-integrated Al/Si ratio representative for the river's sediment flux, a representative  $^{10}\text{Be}$  concentration (black star) can be derived from linear regression (example shown in C), with arrows showing a depth-integrated Al/Si ratio of the Madeira of 0.34 from Bouchez et al. (2011b) in this example. From this  $^{10}\text{Be}$  concentration, an erosion rate can be calculated that is not affected by bias from riverine particle sorting. The same correction can be done with  $^9\text{Be}$  data in order to calculate a representative weathering fraction. Most uncertainties are within symbol size.

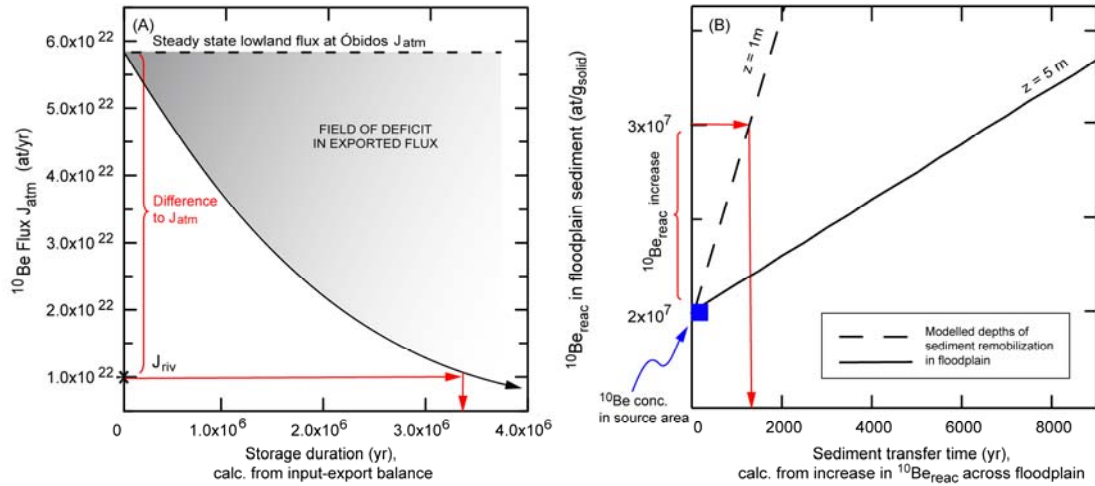


Fig. 13: A) The duration (in Myr) of static storage required to reduce the riverine flux of  $^{10}\text{Be}$  in the Amazon lowlands at Óbidos ( $J_{\text{riv}}$ ) relative to the depositional flux onto the floodplain upstream of Óbidos ( $J_{\text{atm}}$ , dotted black line). Here  $J_{\text{riv}}^{10\text{Be}}$  was calculated by using an *in situ*-derived sediment flux (Wittmann et al., 2015). The red horizontal arrow corresponds to the effective lowland riverine flux and shows potential  $^{10}\text{Be}$  radioactive decay during static storage the field of deficit representing 3.7 Myr of storage. B) Model of  $^{10}\text{Be}$  accumulation over time scales typical of dynamic floodplain-channel interaction in the Amazon basin. Note that for these short time scales, decay of  $^{10}\text{Be}$  is negligible. The observed increase in floodplain-derived  $^{10}\text{Be}_{\text{reac}}$  (ca.  $1 \times 10^7$  at/g) was added to the  $^{10}\text{Be}_{\text{reac}}$  of ca.  $2 \times 10^7$  at/g contained in Andean source area sediment (blue square). The predicted  $^{10}\text{Be}$  inventory was calculated using equation 12 and converted into  $^{10}\text{Be}_{\text{reac}}$  by assuming two different remobilization depths ( $z = 1, 5\text{ m}$ ). For a mean remobilization depth of ca. 1 m, which would correspond to the case where a clay layer deposited on top of the floodplain binds all meteoric  $^{10}\text{Be}$  deposited from the atmosphere, the observed increase in  $^{10}\text{Be}$  across the floodplain would correspond to a transfer time of ca. 1.6 kyr. Both Figures modified from Wittmann et al. (2015).

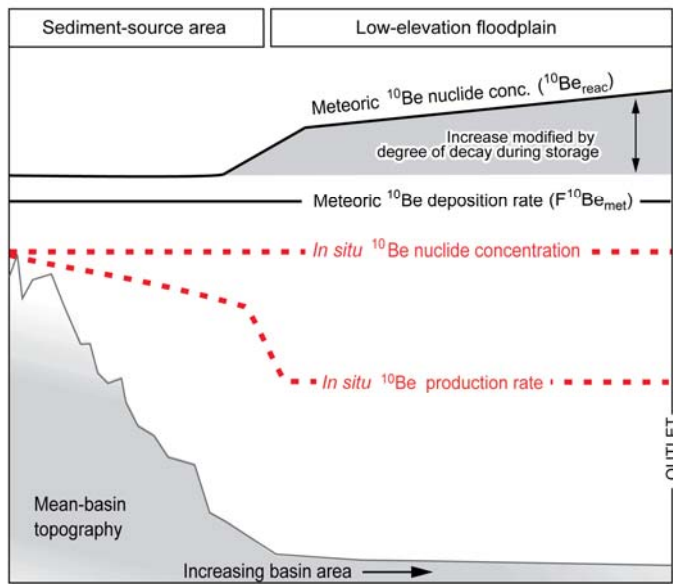


Fig. 14: Schematic illustration of the different behavior of meteoric versus *in situ*-cosmogenic nuclides in large lowland basins; principle is adapted from Wittmann et al. (2011a). *In situ* nuclide production decreases significantly across lowland areas, whereas meteoric depositional fluxes are not dependent on elevation (at elevations below 3 km). In dynamic floodplains, long-lived *in situ* nuclide concentrations are not modified during storage and reworking. Meteoric cosmogenic nuclide concentrations are thus much more sensitive to storage.

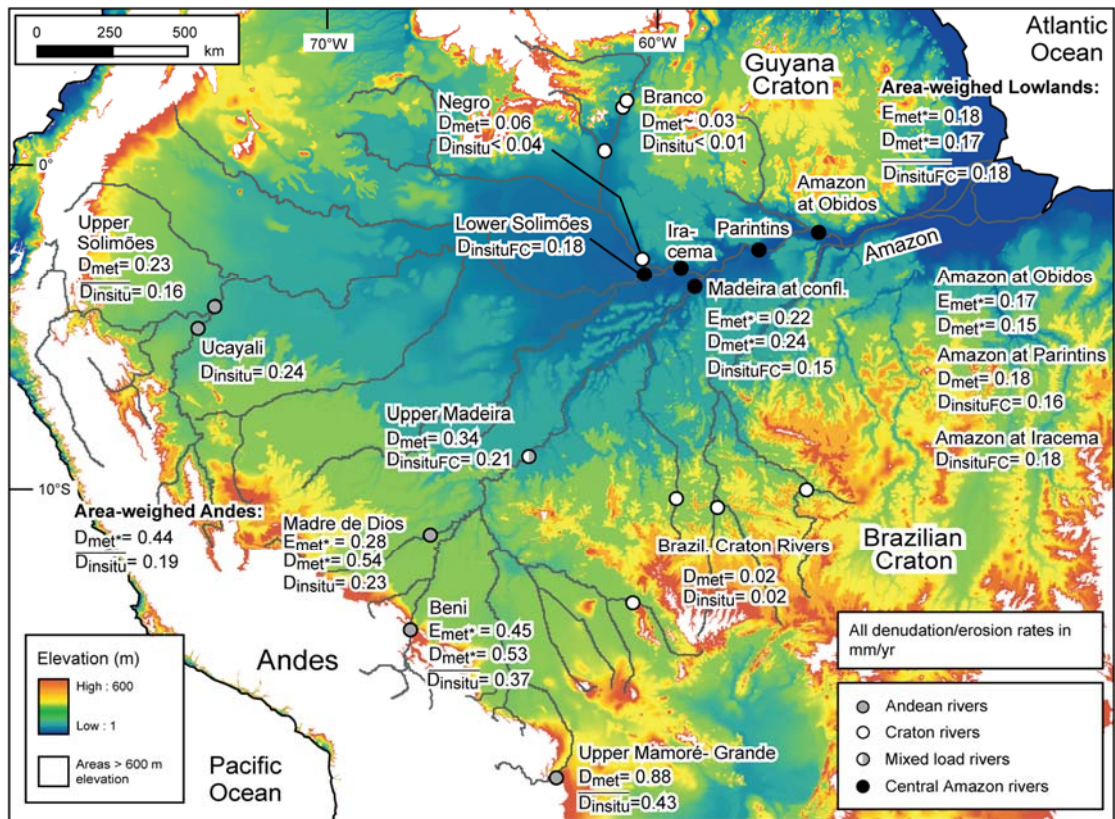


Fig. 15: Elevation map of the Amazon basin with comparison of denudation rates ( $D_{met}$ ;  $D_{met}^*$ ) and high-confidence erosion rates  $E_{met}^*$  with  $D_{insitu}$  and  $D_{insituFC}$  (from *in situ*  $^{10}\text{Be}$  concentrations in sand-sized quartz from bedload; Wittmann et al., 2009; 2011a;  $\overline{D_{insitu}}$  or  $\overline{D_{insituFC}}$  are area-weighted values from Table 2).  $E_{met}^*$  and  $D_{met}^*$  are from depth-integrated suspended sediment measured at four locations where depth profiles were available. Lower confidence rates are denudation rates from  $^{10}\text{Be}/^9\text{Be}$  from bedload ( $D_{met}$ ) where the  $^9\text{Be}$  fractions are potentially affected by grain size bias. We show area-weighted meteoric rates for the source areas (Andes) and the Amazon outlet; the area-weighted Andean rate was calculated using both  $D_{met}$  from bedload and  $D_{met}^*$ ; the area-weighted lowland rate was calculated using  $D_{met}^*$  from the two lowland depth profiles (Madeira and Amazon at Óbidos).

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## **Supplementary Information**

### Part A- TREATMENT OF COSMOGENIC NUCLIDE DATA

Please note that where necessary, we have recalculated published cosmogenic nuclide concentration data mentioned below relative to the secondary AMS standard “07KNSTD” ((Nishiizumi et al., 2007); measured relative to a half life of  $1.39 \pm 0.012 \times 10^6$  yrs by Chmeleff et al. (2010); Korschinek et al. (2010)) to make the data comparable. We then re-calculated all denudation rates relative to Dunai (2000) time-independent scaling scheme and a corresponding SLHL nucleogenic production rate (Borchers et al., 2016; Phillips et al., 2016) of 3.7 at/g<sub>(Qtz)</sub>/yr. Muonic contributions are ca. 1% at SLHL, yielding a total SLHL production rates of 3.75 at/g<sub>(Qtz)</sub>/yr used, where muonic coefficients were calculated following Braucher et al. (2011), using values of 157 g/cm<sup>2</sup> for neutrons, 1500 g/cm<sup>2</sup> for slow muons, and 4320 g/cm<sup>2</sup> for fast muons, respectively (Braucher et al., 2011). Note that the procedure for re-calculation of production rates and nuclide concentrations is summarized in the supplement table A1 of Wittmann et al. (2011). Production rates were calculated on a pixel-by-pixel basis using a 90 m SRTM DEM (except for large basins from Andermann (2011), and those by Lupker et al. (2012) which were

calculated using a 1 km-resolution DEM). Topographic shielding was carried out following Norton and Vanacker (2009) and the nucleogenic production rate was set to zero for pixels covered by ice using a digital glacier map for the Himalayas (National Snow and Ice Data Center, 1999). A snow shielding correction factor as well as a correction due to changes in Earth's magnetic field were not applied.

## Part B- COMPILATION OF SEDIMENT STORAGE TIMES IN FLOODPLAINS OF THE AMAZON AND GANGA BASINS

Basin & position in sediment cascade	Method	Estimated t (x10 <sup>3</sup> yr)	Mode of estimation	Reference	
Amazon	Headwaters	U-Series disequilibria	3	"lag time"	Dosseto et al., 2006
	Lowlands	Channel belt width and river migration rates	4	Beni tributary only	Wittmann and von Blanckenburg, 2009
		U-Series disequilibria	14	transfer time for reach Beni - Madeira confl.	Dosseto et al., 2006
		Size of entire floodplain reservoir and river migration rates	200	"floodplain response time" to remove entire floodplain reservoir during e.g. sea level drop <sup>a</sup>	Castelltort and Van Den Driessche, 2003
		Size of entire floodplain reservoir and river migration rates	700	"floodplain response time" to remove entire floodplain reservoir during e.g. sea level drop <sup>a</sup>	Wittmann et al., 2011 (based on Métivier & Gaudemer, 1999)
Ganga	Headwaters	Dating of valley fills	3-32; up to 100	"lag time"	Blöthe & Korup, 2013
	Lowlands	Channel belt width and river migration rates	1.4	time to remove sand reservoir in floodplain	Lupker et al., 2012a (based on Lauer & Willenbring, 2010)
		U-Series disequilibria	20-25	transfer time for susp. sediment	Granet et al., 2010
		U-Series disequilibria	<100	transfer time for bedload	Granet et al., 2007
		Size of entire floodplain reservoir and river migration rates	100	"floodplain response time" to remove entire floodplain reservoir during e.g. sea level drop <sup>a</sup>	Castelltort and Van Den Driessche, 2003
		Size of entire floodplain reservoir and river migration rates	470	"floodplain response time" to remove entire floodplain reservoir during e.g. sea level drop <sup>a</sup>	Métivier & Gaudemer, 1999

<sup>a</sup>Note that the "response times" estimated by Métivier and Gaudemer (1999) and Castelltort and Van Den Driessche (2003) are theoretical storage times. These denote the maximum time that would be needed to remove the entire floodplain reservoir (i.e. floodplain volumes are calculated from estimated storage depths and the width from entire fluvial valleys, by assuming continuous channel migration rates).

## Part C- COMPILED COSMOGENIC NUCLIDE AND RIVER-LOAD GAUGING DATA

### Cosmogenic nuclide data:

Reference	Sample ID	Basin/River	Denudation		Source area (km <sup>2</sup> ) (> 300m elevation)	Area-weighted value		Sediment flux (Mt/yr)	1σ SD of dataset	1σ SD of dataset
			Rate (mm/yr)	Total area (> ca.100 km <sup>2</sup> )		(mm/yr)	±			
<b>AMAZON BASIN</b>	Safran et al., 2005	BOL-31	Beni	0.21	111					
	Safran et al., 2005	BOL-16	Beni	0.073	120					
	Safran et al., 2005	BOL-28	Beni	0.57	127					
	Safran et al., 2005	BOL-01	Beni	0.90	134					
	Safran et al., 2005	BOL-25	Beni	0.16	146					
	Safran et al., 2005	BOL-27	Beni	0.48	146					
	Safran et al., 2005	BOL-40	Beni	0.26	152					
	Safran et al., 2005	BOL-24	Beni	0.25	154					
	Safran et al., 2005	BOL-34a	Beni	0.33	177					
	Safran et al., 2005	BOL-34b	Beni	0.46	177					
	Safran et al., 2005	BOL-34c	Beni	0.32	177					
	Safran et al., 2005	BOL-08	Beni	0.23	198					
	Safran et al., 2005	BOL-17	Beni	0.18	248					
	Safran et al., 2005	BOL-45	Beni	0.64	348					
	Safran et al., 2005	BOL-43	Beni	0.24	416					
	Safran et al., 2005	BOL-30	Beni	0.39	425					
	Safran et al., 2005	BOL-14	Beni	0.18	432					
	Safran et al., 2005	BOL-44	Beni	0.50	601					
	Safran et al., 2005	BOL-32	Beni	0.20	834					
	Safran et al., 2005	BOL-37	Beni	0.38	841					
	Safran et al., 2005	BOL-22	Beni	0.38	1460					
	Safran et al., 2005	BOL-41	Beni	0.93	1460					
	Safran et al., 2005	BOL-19	Beni	0.32	1540					
	Safran et al., 2005	BOL-20	Beni	0.49	1760					
	Safran et al., 2005	BOL-07	Beni	0.22	2930					
	Safran et al., 2005	BOL-13	Beni	0.21	3230					
	Safran et al., 2005	BOL-21	Beni	0.17	5360					
	Safran et al., 2005	BOL-38a	Beni	0.43	5410					
	Safran et al., 2005	BOL-38b	Beni	0.41	5410					
	Safran et al., 2005	BOL-15	Beni	0.23	5830					
	Safran et al., 2005	BOL-42	Beni	0.40	10400					
	Safran et al., 2005	BOL-18	Beni	0.14	10900					
	Safran et al., 2005	BOL-50	Beni	0.57	70200					
	Insel et al., 2010	N04	Beni	0.14	332					
	Insel et al., 2010	N04-2	Beni	0.15	332					
	Insel et al., 2010	N05	Beni	0.10	205					
	Wittmann et al., 2009	BE 1a	Beni	0.29	68000					
	Wittmann et al., 2009	BE 1b	Beni	0.31	68000					
						<b>0.37 ± 0.20</b>		<b>66 ± 35</b>		(using 68000 km <sup>2</sup> for conversion)
	Insel et al., 2010	S08	Grande	0.33	129					
	Insel et al., 2010	S09	Grande	0.35	152					
	Insel et al., 2010	S09-2	Grande	0.35	152					
	Insel et al., 2010	S12	Grande	0.24	1036					
	Insel et al., 2010	S06	Grande	0.028	1318					
	Wittmann et al., 2009	PIR 18b	Grande	0.36	1590					
	Wittmann et al., 2009	PIR 18c	Grande	0.48	1590					
	Insel et al., 2010	S15	Grande	0.42	3127					
	Insel et al., 2010	S04	Grande	0.08	3763					
	Insel et al., 2010	S04-2	Grande	0.15	3763					
	Wittmann et al., 2009	CHA 23a	Grande	0.22	4862					
	Insel et al., 2010	S10	Grande	0.23	5194					
	Wittmann et al., 2009	MAN 15b	Grande	0.33	5298					
	Wittmann et al., 2009	ICH 21a	Grande	0.31	7580					
	Wittmann et al., 2009	GR 25a	Grande	0.13	23500					
	Wittmann et al., 2009	GR 17b-Mts only	Grande	0.39	59800					
	Wittmann et al., 2009	GR 19b	Grande	0.54	59800					
						<b>0.38 ± 0.14</b>		<b>60 ± 22</b>		(using 59800 km <sup>2</sup> for conversion)
	Wittmann et al., 2010	Na 0a-1	Solimoes	0.37	5270					
	Wittmann et al., 2010	Na 0a-2	Solimoes	0.30	5270					
	Wittmann et al., 2010	Na 1b	Solimoes	0.40	12400					
	Wittmann et al., 2011	Pe 107a-1	Solimoes	0.23	238000					
	Wittmann et al., 2011	Pe 107a-2	Solimoes	0.25	238000					
	Wittmann et al., 2011	Pe 101a-1	Solimoes	0.13	630000					
	Wittmann et al., 2011	Pe 101a-2	Solimoes	0.12	630000					
						<b>0.16 ± 0.11</b>		<b>260 ± 180</b>		(using 630000 km <sup>2</sup> for conversion)
						<b>Entire Andes draining to Amazon:</b>				
						<b>0.19 ± 0.18</b>				
					<b>total area used for conversion: 757800</b>			<b>380 ± 350</b>		

Reference	Sample ID	Basin/River	Denudation		Source area (km <sup>2</sup> ) (> 300m elevation)	FC-corr D (mm/yr)	1σ SD of dataset	Sediment flux (Mt/yr)	1σ SD of dataset	
			Rate (mm/yr)	Total area (> ca.100 km <sup>2</sup> )						
<b>FLOODPLAIN</b>	Wittmann et al., 2009	MAR 18a-1	Marmoré	0.12	599400	282214	0.15			
	Wittmann et al., 2009	MAR 18a-2	Marmoré	0.10	599400	282214	0.13			
	Wittmann et al., 2009	MAD 19a	Madeira	0.16	881900	282214	0.19			
	Wittmann et al., 2009	MAD 20a-1	Madeira	0.16	954300	282214	0.19			
	Wittmann et al., 2009	MAD 20a-2	Madeira	0.20	954300	282214	0.24			
	Wittmann et al., 2011	(3)-Napo lower	Lower Napo	0.12	93171	17715	0.20			
	Wittmann et al., 2011	(4)-Man 0.2a	Manacapuru	0.13	2270010	500272	0.18			
	Wittmann et al., 2011	(4)-Man 1.1a	Manacapuru	0.13	2270010	500272	0.19			
	Wittmann et al., 2011	(4)-Man 2.85a	Manacapuru	0.12	2270010	500272	0.17			
	Wittmann et al., 2011	(7)-Ir 1.75a	Iracema	0.11	3154060	628165	0.17			
	Wittmann et al., 2011	(7)-Ir 1.75b	Iracema	0.12	3154060	628165	0.19			
	Wittmann et al., 2011	(8)-Mad 0.5a	Madeira	0.09	1336000	282214	0.14			
	Wittmann et al., 2011	(8)-Mad 1.8a	Madeira	0.13	1336000	282214	0.19			
	Wittmann et al., 2011	(8)-Mad 1.8b	Madeira	0.09	1336000	282214	0.13			
	Wittmann et al., 2011	Par 0.9a	Parintins	0.11	4736250	947182	0.17			
	Wittmann et al., 2011	Par 2.2a	Parintins	0.10	4736250	947182	0.15			
	Wittmann et al., 2011	Par 2.2b	Parintins	0.10	4736250	947182	0.15			
	Wittmann et al., 2011	(16)-Ama-a	Obidos	0.09	5087830	947182	0.14			
	Wittmann et al., 2011	Soc-b	Varzea	0.11	5087830	947182	0.22			
	Wittmann et al., 2011	Soc-c1	Varzea	0.11	5087830	947182	0.22			
	Wittmann et al., 2011	Soc-c2	Varzea	0.12	5087830	947182	0.25			
	Wittmann et al., 2011	Gran-b	Varzea	0.07	5087830	947182	0.15			
	Wittmann et al., 2011	Gran-c	Varzea	0.09	5087830	947182	0.19			
					<b>total area used for conversion: 757800</b>		<b>0.18 ± 0.03</b>	<b>350 ± 65</b>		

Reference	Sample ID	Basin/River	Denudation		Area-weighted value (mm/yr) ± dataset	1σ SD of dataset	Sediment flux (Mt/yr) ± dataset	1σ SD of dataset				
			Rate (mm/yr)	Area (> ca.100 km <sup>2</sup> )								
<b>GANGA BASIN</b>	Vance et al., 2003 AK43	Upper Ganga	0.87	21690	<b>1.22 ± 0.60</b>		<b>100 ± 50</b> (using 32400 km <sup>2</sup> for conversion)					
	Vance et al., 2003 AK36	Upper Ganga	0.87	21690								
	Lupker et al., 2012 BR924-Mts only	Upper Ganga	1.91	21690								
	Scherler et al., 2014 DS7-048/1	Yamuna	0.44	100								
	Scherler et al., 2014 DS6-080A/14	Yamuna	1.41	103								
	Scherler et al., 2014 DS7-088/16	Yamuna	0.25	123								
	Scherler et al., 2014 DS7-077/9	Yamuna	0.31	128								
	Scherler et al., 2014 DS7-049/2	Yamuna	0.13	168								
	Scherler et al., 2014 DS7-061/4	Yamuna	0.10	210								
	Scherler et al., 2014 DS6-088/18	Yamuna	0.84	231								
	Scherler et al., 2014 DS7-072/10	Yamuna	0.23	514								
	Scherler et al., 2014 DS6-094/P1	Yamuna	0.31	516								
	Scherler et al., 2014 DS6-081/13	Yamuna	1.63	536								
	Scherler et al., 2014 DS6-023C/15	Yamuna	1.94	548								
	Scherler et al., 2014 DS7-055/Y1	Yamuna	1.00	647								
	Scherler et al., 2014 DS6-018/Y2	Yamuna	0.26	1179								
	Scherler et al., 2014 DS6-085/P2	Yamuna	0.41	1440								
	Scherler et al., 2014 DS7-071/T1	Yamuna	1.09	3477								
	Scherler et al., 2014 DS7-076/T2	Yamuna	0.99	4670								
	Lupker et al., 2012 YAM Q3-Mts only	Yamuna	0.56	7620					<b>0.76 ± 0.56</b>		<b>20 ± 14</b> (using 9570 km <sup>2</sup> for conversion)	
	Lupker et al., 2012 CA10-5-Mts only	Karnali	0.48	21260					<b>1.19 ± 0.75</b>		<b>140 ± 90</b> (using 45970 km <sup>2</sup> for conversion)	
	Lupker et al., 2012 PB80-Mts only	Karnali	1.91	45430								
	Lupker et al., 2012 LO743-Mts only	Karnali	0.79	45430								
	Godard et al., 2012 NEP030	Narayani	0.06	110								
	Godard et al., 2012 NIB-975-02	Narayani	2.32	136								
	Andermann-2011 PhD NP-A5s	Narayani	0.69	148								
	Godard et al., 2012 NEP003	Narayani	0.87	614								
	Andermann-2011 PhD NP-A1s	Narayani	0.39	657								
	Andermann-2011 PhD NP081016A	Narayani	4.17	3000								
	Andermann-2011 PhD CAJ-7	Narayani	1.30	3207								
	Andermann-2011 PhD NP-A39s	Narayani	1.67	3981								
	Andermann-2011 PhD NP-A10s	Narayani	1.83	4086								
	Andermann-2011 PhD NP-A18s	Narayani	2.23	4798								
	Andermann-2011 PhD Arr-3A	Narayani	2.31	5048								
	Andermann-2011 PhD NP-A20s	Narayani	1.43	5469								
	Andermann-2011 PhD Arr-2	Narayani	1.23	6091								
	Andermann-2011 PhD Arr-3B	Narayani	2.38	11716								
	Andermann-2011 PhD Arr-4	Narayani	1.35	11997								
	Lupker et al., 2012 LO741-Mts only	Narayani	1.03	31770								
	Lupker et al., 2012 CA10-10-Mts only	Narayani	1.35	31770								
	Lupker et al., 2012 LO1001-Mts only	Narayani	1.51	31770								
	Lupker et al., 2012 LO1001-Mts only	Narayani	1.67	31770								
	Godard et al., 2014 PO-150311-05	Narayani	0.21	91								
	Godard et al., 2014 KP-160311-09	Narayani	0.10	93								
	Godard et al., 2014 KP-160311-10	Narayani	0.11	97								
	Godard et al., 2014 KP-090311-05	Narayani	0.29	99								
	Godard et al., 2014 KP-090311-01	Narayani	0.14	111								
	Godard et al., 2014 TR-170311-03	Narayani	1.21	147					<b>1.54 ± 0.96</b>		<b>140 ± 70</b> (using 34450 km <sup>2</sup> for conversion)	
	Godard et al., 2014 EK-180311-05	Kosi	1.43	116.5								
	Andermann-2011 PhD NP080912A	Kosi	0.75	2105								
	Andermann-2011 PhD NP080913A	Kosi	2.39	2572								
	Andermann-2011 PhD NP080913B	Kosi	1.94	2962								
	Andermann-2011 PhD NP080924A	Kosi	0.15	113								
	Lupker et al., 2012 LO763-Mts only	Kosi	0.79	53610								
	Lupker et al., 2012 CA987-Mts only	Kosi	0.40	53610								
	Lupker et al., 2012 CA987-Mts only	Kosi	0.72	53610								
									<b>0.69 ± 0.78</b>		<b>110 ± 125</b> (using 62000 km <sup>2</sup> for conversion)	
									<b>Entire Himalayas draining to Ganga: 1.14 ± 0.82</b>		<b>550 ± 390</b>	
		total area used for conversion:		<b>184389</b>								

Reference	Sample ID	Basin/River	Denudation		Source area (km <sup>2</sup> ) (> 300m elevation)	FC-corr D (mm/yr) ± dataset	1σ SD of dataset	Sediment flux (Mt/yr) ± dataset	1σ SD of dataset
			Rate (mm/yr)	Total area (> ca.100 km <sup>2</sup> )					
<b>FLOODPLAIN</b>	Lupker, 2012 BR355-1 Fc	Karnali	1.10	87040	64940	1.15			
	Lupker, 2012 BR355-2 Fc	Karnali	0.88	87040	64940	0.91			
	Lupker, 2012 BR343 Fc	Karnali	0.45	131840	72010	0.76			
	Lupker, 2012 BR336 Fc	Narayani	1.50	41800	34450	1.27			
	Lupker, 2012 BR310 Fc	Narayani	1.49	43500	34450	1.35			
	Lupker, 2012 BR330 Fc	Kosi	0.51	89110	56210	0.68			
	Lupker, 2012 BR8135 Sc	Ganga	0.04	390410	45540	0.37			
	Lupker, 2012 BR315 Sc	Ganga	0.15	697540	153270	0.60			
	Lupker, 2012 BR8138 Sc	Ganga	0.15	697540	153270	0.59			
	Lupker, 2012 BR418 Sc	Ganga	0.23	873240	220150	0.84			
	Lupker, 2012 BR717 Sc	Ganga	0.23	873240	220150	0.89			
	Lupker, 2012 BR8221 Sc	Ganga	0.23	873240	220150	0.84			
	Lupker, 2012 BR8252-1 Sc	Ganga	0.23	873240	220150	0.88			
	Lupker, 2012 BR8252-2 Sc	Ganga	0.23	873240	220150	0.84			
		total area used for conversion:		<b>184389</b>		<b>0.821 ± 0.27</b>		<b>390 ± 130</b>	

## Gauging data:

	River	Station Code	Total area (> ca.100 km2)	Sampled source area (km2)	Gauging-derived Load (Mt/yr)	Calculated erosion rate (mm/yr)	Area-weighted erosion value (mm/yr)	±	1σ SD of dataset	
<b>AMAZON BASIN</b>	Pepin et al., 2013	Beni	AB	67500		161.8	0.92			
	Aalto et al., 2006	Beni	AIN	29900		115.0	1.48			
	Aalto et al., 2006	Beni	AQM	9400		36.8	1.51			
	Aalto et al., 2006	Beni	COT	5600		40.6	2.79			
	Aalto et al., 2006	Beni	LUR	810		6.4	3.04			
	Aalto et al., 2006	Beni	MIG	360		0.047	0.050			
	Aalto et al., 2006	Beni	MIS	350		0.012	0.013			
	Aalto et al., 2006	Beni	POR	240		0.80	1.28			
	Aalto et al., 2006	Beni	SIR	270		2.0	2.85			
	Aalto et al., 2006	Beni	SRC	4700		7.1	0.58			
	Aalto et al., 2006	Beni	TAM	950		2.4	0.97			
	Aalto et al., 2006	Beni	VBA	1900		7.8	1.58			
	Aalto et al., 2006	Beni	VER	140		0.011	0.030			
				<b>sum area = 122120</b>		<b>sum load = 380.8</b>		<b>1.20</b>	<b>± 1.06</b>	
				<b>largest basin= 67500</b>		<b>"Upscaled load"= 210.5</b>				
	Aalto et al., 2006	Grande	ARC	23700		154.3	2.50			
	Aalto et al., 2006	Grande	HUR	11200		14.1	0.48			
Aalto et al., 2006	Grande	MIZ	10800		14.1	0.50				
Aalto et al., 2006	Grande	PAZ	4360		2.2	0.19				
Aalto et al., 2006	Grande	PNA	31200		296.9	3.66				
Aalto et al., 2006	Grande	AP	59800		125.0	0.80				
Aalto et al., 2006	Pirai	ANG	1420		2.9	0.79				
Aalto et al., 2006	Pirai	BER	480		0.60	0.48				
Aalto et al., 2006	Pirai	EPS	203		0.42	0.80				
Aalto et al., 2006	Pirai	LBE	2880		2.3	0.31				
Aalto et al., 2006	Pirai	PEI	4160		1.0	0.09				
Aalto et al., 2006	Mamoré	ICO	2300		11.4	1.91				
Aalto et al., 2006	Mamoré	LOC	200		0.67	1.29				
Wittmann et al., 2009	Grande	AP	59800		138.0	0.89				
Pepin et al., 2013	Grande	ABA	59480		178.2	1.15				
			<b>sum area = 271983</b>		<b>sum load = 942.1</b>		<b>1.33</b>	<b>± 0.97</b>		
			<b>largest basin= 59480</b>		<b>"Upscaled load"= 206.0</b>					
Pepin et al., 2013	upper Marañon	BOR	115410		136.6	0.46				
Pepin et al., 2013	upper Ucayali	CHA	70270		70.3	0.38				
Pepin et al., 2013	middle Ucayali	ATA	191020		200.0	0.40				
Santini et al., 2014	middle Ucayali	LAG	191180		445.0	0.90				
Santini et al., 2014	middle Ucayali	PIN	22300		60.0	1.03				
Santini et al., 2014	middle Ucayali	PUC	261070		300.0	0.44				
Armijos et al., 2013	Napo	SEB	5272		6.2	0.46				
Armijos et al., 2013	Napo	FDO	12443		8.2	0.25				
Armijos et al., 2013	Napo	ROC	26861		20.8	0.30				
Armijos et al., 2013	Pastaza	PAS	12700		8.7	0.26				
Armijos et al., 2013	Santiago River	SAN	23880		17.3	0.28				
Santini et al., 2014	lower Ucayali	REQ	360490		205.0	0.22				
Guyot et al., 96	lower Marañon	SRE	360550		168.0	0.18				
Wittmann et al., 2011	upper Solimoes	TAM	733470		413.0	0.22				
			<b>sum area = 2386916</b>		<b>sum load = 2059.1</b>		<b>0.33</b>	<b>± 0.25</b>		
			<b>largest basin= 733470</b>		<b>"Upscaled load"= 632.7</b>					
							<b>Area weighted total erosion rate:</b>			
<b>sum upscaled=</b>						<b>1050.0</b>	<b>0.47</b>	<b>± 0.9</b>		
<b>Floodplain</b>	Fillizola & Guyot 2009	Amazon	OBI	5087830	(used this value for conversion)	556.0	0.36			
	Dunne et al., 1998	Amazon	OBI	5087830	600000	1239.0	0.79			
	Guyot et al., 2005	Amazon	OBI	5087830	600000	810.0	0.52			
	Meade, 1985 (USGS)	Amazon	OBI	5087830	600000	1321.5	0.85			
	Martinez et al., 09	Amazon	OBI	5087830	600000	688.0	0.44			
	Martinez et al., 09	Amazon	OBI	5087830	600000	801.0	0.51			
				<b>total measured load=</b>		<b>902.6</b>		<b>0.58</b>	<b>± 0.2</b>	

	River	Station Code	Total area (> ca.100 km <sup>2</sup> )	Sampled source area (km <sup>2</sup> )	Gauging-derived Load (Mt/yr)	Calculated erosion rate (mm/yr)	Area-weighted erosion value (mm/yr)	±	1σ SD of dataset	
<b>GANGA BASIN</b>	Andermann et al., 2012	Upper Karnali	#240	21121		10.4	0.19			
	Andermann et al., 2012	Karnali	#280	45967		75.9	0.64			
	Lupker et al., 2012	Karnali	-	42890		87.0	0.78			
	Lupker et al., 2012	Karnali	-	45967		210.3	1.76			
	Lupker et al., 2012	Karnali	-	44000		68.6	0.60			
	Andermann et al., 2012	Saradha	#286	808		0.30	0.14			
	Andermann et al., 2012	Upper Rapti	#350	3648		18.3	1.93			
	Andermann et al., 2012	Rapti	#360	5197		13.2	0.98			
				<b>sum area =</b>	<b>209598</b>	<b>sum load =</b>	<b>484.1</b>			
				<b>largest basin=</b>	<b>45967</b>	<b>"Upscaled load"=</b>	<b>106.2</b>			<b>0.89 ± 0.66</b>
	Jha et al., 1988	Yamuna	-	9572		18.08	0.73			0.73 ± 0.73
	Gabet et al., 2008	Koto	#1	812		2.1	1.00			
	Gabet et al., 2008	Nar Khola	#2	1052		0.27	0.10			
	Gabet et al., 2008	Upper Dharapani	#5	1946		2.0	0.40			
	Gabet et al., 2008	Dudh Khola	#6	491		0.38	0.30			
	Gabet et al., 2008	Lower Dharapani	#8	2605		3.4	0.50			
	Gabet et al., 2008	Bhulbule	#9	3217		4.2	0.50			
	Gabet et al., 2008	Khudi Khola	#10	152		0.79	2.00			
	Andermann et al., 2012	Narayani	#450	32002		97.2	1.17			
	Andermann et al., 2012	Trishuli	#447	4428		2.4	0.21			
Andermann et al., 2012	Kali Gandaki	#410	7170		31.9	1.71				
Lupker et al., 2012	Gandak	-	34988		122.8	1.35				
Sinha & Friend, 1994	Burhi Gandak	-	9580		14.4	0.58				
Andermann et al., 2012	Bagmati	#589	2849		4.2	0.57				
			<b>sum area =</b>	<b>101292</b>	<b>sum load =</b>	<b>286.1</b>				
			<b>largest basin=</b>	<b>34988</b>	<b>"Upscaled load"=</b>	<b>98.8</b>			<b>1.09 ± 0.60</b>	
Andermann et al., 2012	Bhote Koshi	#0	2308		0.3	0.050				
Andermann et al., 2012	Sapta Koshi	#695	54024		62.7	0.45				
Lupker et al., 2012	Kosi	-	60400		149.2	0.95				
Lupker et al., 2012	Kosi	-	62000		166.0	1.03				
Lupker et al., 2012	Kosi	-	61000		182.4	1.15				
Lupker et al., 2012	Kosi	-	59280		89.6	0.56				
Sinha & Friend, 1994	Baghmati	-	12973		10.5	0.31				
Sinha & Friend, 1994	Kamala-Balan	-	2945		7.7	1.01				
			<b>sum area =</b>	<b>314930</b>	<b>sum load =</b>	<b>668.4</b>				
			<b>largest basin=</b>	<b>62000</b>	<b>"Upscaled load"=</b>	<b>131.6</b>			<b>0.81 ± 0.40</b>	
Abbas et al., 1984	Ramganga		32400		10.1	0.12				
Abbas et al., 1984	Gomti		30400		6.3	0.080				
Abbas et al., 1984	Ghaghra		12700		11.9	0.36				
			<b>sum area =</b>	<b>75500</b>	<b>sum load =</b>	<b>28.3</b>				
			<b>largest basin=</b>	<b>32400</b>	<b>"Upscaled load"=</b>	<b>12.2</b>			<b>0.14 ± 0.15</b>	
<b>Area weighted total erosion rate:</b>										
<b>sum upscaled=</b>						<b>370.0</b>	<b>0.8</b>	<b>±</b>	<b>0.55</b>	
(used this value for conversion)										
<b>Floodplain</b>	Lupker et al., 2012	Ganga	Harding Bridge1	873240	184927	480.0	1.00			
	Lupker et al., 2012	Ganga	Harding Bridge2	873240	184927	448.0	0.93			
	Lupker et al., 2012	Ganga	Harding Bridge3	873240	184927	210.0	0.44			
	Lupker et al., 2012	Ganga	Harding Bridge4	873240	184927	340.0	0.71			
	Lupker et al., 2012	Ganga	Harding Bridge5	873240	184927	200.0	0.42			
	Lupker et al., 2012	Ganga	Harding Bridge6	873240	184927	480.0	1.00			
	Lupker et al., 2012	Ganga	Harding Bridge7	873240	184927	548.0	1.14			
	Lupker et al., 2012	Ganga	Harding Bridge8	873240	184927	390.0	0.81			
<b>total measured load=</b>						<b>390.0</b>	<b>0.82</b>	<b>±</b>	<b>0.27</b>	

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