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Hydrogeology Journal DOI 10.1007/s10040-008-0339-5 Fine scale variability of hyporheic hydrochemistry in salmon spawning gravels with contrasting strandwater-surface water interactions • Tetzlaff

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35		Received	20 January 2008	

36 37	Schedule	Revised		
		Accepted 19 June 2008		
36 37 38	Schedule Abstract	Provided 19 June 2008 There is increasing realisation of the importance of groundwater-surface water (GW-SW) interactions in understanding freshwater ecology. A study that assessed the influence of local GW-SW interactions on shallow (<250 nm) proprietic water quality at two contrasting salmon spawning locations in Sottand, UK is reported. At a groundwater-dominated site, continuous logging renors revealed that hyporheic dissolved oxygen (DO) concentrations changed rapidy in response to changing hydrological conditions. Low volume (25 ml) spot samples revealed fine-scale spatial variability (<0.05 m) consistent with a delectrical conductivity values, characteristic of surface water. Small reductions in DO at this site are hypothesis to be associated with short residence hyporheic discharge. A comparison between in-stul (logging DO sensor data) and subtry and the upported context was typically characteristic of surface water. Small reductions in patial and temporal scales and that future studies need to design sampling programmes. This study demonstrates that hyporheic water quality arise over fine spatial and temporal scales and that future studies end to design sampling programmes. This study demonstrates that hyporheic water quality arise over fine spatial and temporal scales and that future studies over fine fractions of cales (S-GM) est de plus en plus reconnue. Une étude sur l'influence de fine fractions of cales (S-GM) est de plus en plus reconnue. Une étude sur l'andice end aparte superficielle (<50.50 mm) de leave hyporhéique à deux stations differentes of prayer à saumon localisées en Ecosse, Royaume Uni, est décrite ici, Sur un plus fechnice (S-GM) indiquent une variabilit ef a partie superficielle (<50.50 mm) de leave hyporhéique à deux stations differentes of chantilons portueis de faible volume (S-M) indiquent une variabilit ef estations offerentes et al. Pour leave desufrace (D-M) est de plus en plus reconne and tha trutte		
		hiporreicas de corto tiempo de residencia. Una comparación entre métodos in- situ (datos de sensores de monitoreo de OD) y ex-situ (muestreo de pequeños volúmenes) demuestra una buena concordancia, y potencialmente permite la utilización de los dos métodos en programas de muestreos estratificados. Este estudio demuestra que la calidad del agua hiporreica varía en escalas finas de espacio y tiempo, y que los estudios futuros necesitan diseñar estrategias de muestreo que consideren las escalas adecuadas tanto para los procesos ecológicos de interés como los hiporreicos.		
		Resumo: Existe uma percepção crescente da importância das interacções		

		águas subterrâneas-água superficial para a compreensão da ecologia dos cursos de água doce. Apresenta-se neste artigo um estudo de avaliação da influência daquelas interacções na qualidade da água de zonas hiporreicas a reduzida profundidade (<250 mm) em dois locais, com características contrastantes, de desova de salmão na Escócia, Reino Unido. Num primeiro local, em que predomina o fluxo de água subterrânea, a monitorização contínua revelou que a concentração de Oxigénio Dissolvido (OD) na zona hiporreica se alterava rapidamente em resposta a variações das condições hidrológicas. Amostras de água de volume reduzido (25 ml) mostram uma variabilidade espacial a escala reduzida (<0.05 mm) consistente com variações na posição vertical entre fontes de água (superficial e subterrânea). Num segundo local, em que predomina a influência das águas superficiais, a água da zona hiporreica era tipicamente caracterizada por valores elevados de Oxigénio Dissolvido (DO) e de condutividade eléctrica, característicos de águas superficiais. Pequenas reduções no valor de DO neste local são atribuídas a tempos de residência reduzidos das águas subterrâneas nas zonas hiporreicas. Uma comparação entre métodos in-situ (sensores de DO) e ex-situ (amostras de reduzido volume) demonstram uma boa concordância entre aquelas metodologias, potenciando a utilização de ambos os métodos em programas de amostragem em zonas estratificadas. Este estudo demonstra que a qualidade da água de zonas hiporreicas varia em escalas temporais e espaciais reduzidas e que estudos futuros devem considerar estratégias de amostragem adaptadas às escalas apropriadas para os processos ecológicos e para os processos da zona hiporreica a estudar.
39	Keywords separated by ' - '	Groundwater–surface-water relations - Hydrochemistry - Oxygen - Hyporheic - UK
40	Foot note information	

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Fine scale variability of hyporheic hydrochemistry in salmon spawning gravels with contrasting groundwater-surface water interactions

9 I. A. Malcolm · C. Soulsby · A. F. Youngson ·

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Abstract There is increasing realisation of the importance 12of groundwater-surface water (GW-SW) interactions in 13 14 understanding freshwater ecology. A study that assessed the influence of local GW-SW interactions on shallow 15(<250mm) hyporheic water quality at two contrasting 16 salmon spawning locations in Scotland, UK is reported. 17At a groundwater-dominated site, continuous logging 18 sensors revealed that hyporheic dissolved oxygen (DO) 19concentrations changed rapidly in response to changing 20hydrological conditions. Low volume (25ml) spot samples 21revealed fine-scale spatial variability (<0.05m) consistent 22with a vertically shifting boundary layer between source 2324 waters. At a surface-water-dominated location, hyporheic water was typically characterised by high DO and 25electrical conductivity values, characteristic of surface 26water. Small reductions in DO at this site are hypothesised 27to be associated with short residence hyporheic discharge. 28A comparison between in-situ (logging DO sensor data) 29and ex-situ (small volume sampling) methods revealed 30 good agreement, potentially allowing deployment of the 31two methods in stratified sampling programmes. This 32 study demonstrates that hyporheic water quality varies 33 over fine spatial and temporal scales and that future 34 studies need to design sampling strategies that consider 35the scales appropriate to both the ecology and the 36 hyporheic processes of interest. 37

- 38
- Keywords Groundwater–surface-water relations ·
 Hydrochemistry · Oxygen · Hyporheic · UK

Received: 20 January 2008 / Accepted: 19 June 2008

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Introduction

With increasing research focus on groundwater-surface 42 water (GW-SW) interactions, there is a growing realisation 43of the complex spatio-temporal dynamics exhibited by 44 physical, chemical and biological characteristics in the 45hyporheic zone (Dahm et al. 2006; Malcolm et al. 2008). In 46 particular, the chemical characteristics of the hyporheic 47 zone, as the important interface between groundwater and 48 surface water, are known to vary spatially at scales ranging 49from centimetres to kilometres (Wondzell and Swanson 501996: Brunke and Gonser 1997: Boulton et al. 1998: 51Soulsby et al. 2001; Malcolm et al. 2004; Malcolm et al. 522005; Poole et al. 2006) and temporally at scales ranging 53from storm event (sub-hourly) to inter-annual (Wondzell 54and Swanson 1996; Fraser and Williams 1998; Malcolm 55et al. 2004; Malcolm et al. 2006; Arntzen et al. 2006). 56

It is widely accepted that there is a need for improved 57characterisation of the hyporheic environment in order to 58enhance understanding of hyporheic ecology (Palmer 591993; Fowler and Death 2001; Brunke et al. 2003; 60 Boulton and Hancock 2006; Poole et al. 2006). Further-61 more, it has long been recognised that sampling of the 62 hyporheic zone poses particular problems in terms of 63 protocols and methodology (Palmer 1993). However, it is 64 also becoming increasingly clear that one of the central 65 challenges for hyporheic zone research is to sample at 66 temporal and spatial resolutions that are appropriate to 67 both the hyporheic processes of interest and the related 68 ecology (Palmer 1993; Youngson et al. 2005; Grimm et al. 69 2006; Malcolm et al. 2006). Previous studies of the 70 hyporheic zone have often employed sampling methods 71that operate at coarse temporal and spatial scales. Moreover, 72these often involve abstraction of large water samples that 73 integrate over an indeterminate volume of streambed, with 74 unknown recharge or equilibration times. This potentially 75risks failing to characterise important fine scale spatio-76 temporal variability and may result in a mis-match between 77 the (large) spatial scales characterised by hyporheic water 78quality sampling and the (smaller) scales often required to 79adequately characterise and understand the environment 80 experienced by the hyporheos (Palmer 1993; Malcolm et al. 81 2008). While the importance of hyporheic sampling 82 methodology has been highlighted for invertebrates (Fraser 83 and Williams 1997; Hunt and Stanley 2000; Scarsbrook 84

and Halliday 2002), the issue of water quality sampling has
not been addressed in a similar way. In fact, the issue has
been overlooked to the extent that in many cases the
important details of sampling and sample volumes are not
reported (e.g. Bernier-Bourgault and Magnan 2002; Bowen
and Nelson 2003; Greig et al. 2005) making interpretation
of data and comparison between studies difficult.

92Traditional hyporheic sampling methods typically in-93 volve water sampling under negative pressure from standpipes (Ringler and Hall 1975), piezometers (Curry and 94Noakes 1995; Baxter and Hauer 2003; Olsen and Town-95send 2003), incubators (Soulsby et al. 2001; Malcolm et al. 96 2003a, b) and temporary (Mermillod-Blondin et al. 2000) 97 or fixed (Youngson et al. 2005) sampling tubes, inserted to 98 specified depths in the streambed (ex-situ). These methods 99 have a number of potential problems, including direct 100connection between the streambed and surface water or 101 atmosphere, and the creation of preferential flow paths 102such that surface water is drawn down into the streambed 103104during sampling. However, these methods benefit from 105potentially high spatial coverage and relatively low cost. In-situ measurements (e.g. Malcolm et al. 2006), using 106 water quality probes, have the benefit of providing high-107 resolution temporal data with minimal sampling distur-108109 bance, but financial constraints often dictate that replicated sampling at fine spatial resolution is impractical. These 110applications are relatively scarce (few chemical determi-111 nants can be accurately measured this way) and individual 112probes are parameter specific. Furthermore, there is the 113potential that in-situ monitoring can reflect highly localised 114 conditions that are not more generally representative of the 115hyporheic zone at a given location and scale and that 116results are not comparable with traditional ex-situ methods. 117In the context of salmon embryo survival, previous work 118

by the authors has demonstrated that traditional sampling 119methods have often failed to adequately characterise both 120 the temporal dynamics (Malcolm et al. 2006) and spatial 121variability (Malcolm et al. 2005; Youngson et al. 2005) of 122123the hyporheic zone in a way that is biologically meaning-124ful. Salmon ova are deposited in open gravel structures 125called redds, constructed from streambed gravels during a 126process known as spawning. Egg burial depths are typically between 0.05 and 0.3 m beneath the streambed 127128(DeVries 1997). Survival is dependent on complex 129interactions of physical, chemical and biological processes 130which are reviewed in detail elsewhere (Malcolm et al. 2008). Critically, however, survival depends on the 131132delivery of adequate oxygen to meet the needs of developing embryos, and thus, is often influenced by the 133local nature of GW-SW interactions where groundwater is 134characterised by reducing conditions. 135

This paper examines the hydroecological importance of 136sampling at appropriate spatio-temporal scales and com-137pares the results of in-situ sampling with low volume, 138finely stratified, ex-situ sampling methods, using a case 139study of salmon embryo survival at two heavily utilised 140141 spawning locations with contrasting GW–SW interactions. Inter-site differences are discussed in the context of local 142hydrological controls. The importance of sampling meth-143

od and resolution are discussed with reference to previous 144 work investigating salmon embryo survival in field 145settings. Specifically this study aims to: (1) characterise 146hyporheic hydrochemistry at fine temporal and spatial 147 resolution during the period of time between salmon 148 spawning and embryo hatch; (2) use natural tracer 149 methods to infer the influence of local GW-SW inter-150actions on streambed DO; (3) assess the implications for 151embryo survival and (4) compare in-situ and ex-situ 152sampling methods and assess the implications for sam-153pling strategy in future studies of the hyporheic zone 154

155

Materials and methods

The work was carried out at the Girnock Burn catchment, 156a 31-km² sub-catchment of the River Dee in northeast 157Scotland, UK (Fig. 1). Detailed characteristics about the 158catchment are given elsewhere: Tetzlaff et al. (2007a) 159describe the general hydrology and dominant runoff 160 processes; Moir et al. (2002, 2004) describe the distribu-161 tion of salmon spawning sites and their hydraulic and 162 sedimentary characteristics; Soulsby et al. (2007) outline 163 the catchment scale GW-SW interactions, whilst Malcolm 164 et al. (2005) consider their implications for hyporheic 165water quality and salmon embryo survival. Briefly, the 166 catchment drains a montane area underlain by granitic and 167 metamorphic rocks. Groundwater drains through fractures 168 in these rocks and various glacial and paraglacial drifts, 169which cover much of the catchment, contributing 25-30%170of annual runoff. The catchment is largely dominated by 171heather (Calluna) moorland (ca. 95%), though the lower 172catchment has mixed forest cover of pine (Pinus) and 173birch (Betula). Rainfall is around 1,100 mm per annum, 174with a mean annual runoff of around 700 mm. 175

Two sites with contrasting GW-SW interactions and a 176long and documented history of salmon spawning were 177 selected for detailed monitoring of hyporheic chemistry and 178assessment of the mortality of salmon ova (Fig. 1). Both 179sites were previously included in catchment scale studies 180 of hyporheic hydrochemistry (Malcolm et al. 2005) and 181 embryo survival and performance (Youngson et al. 2005) 182using traditional broad scale ex-situ sampling procedures. 183Each site comprised a riffle ca. 10 m long. In the upper 184catchment, the reach containing site 7 (S7) was examined 185 in detail by Malcolm et al. (2004). The site is characterised 186 by strong groundwater upwelling which often results in 187 marked groundwater influence on the hyporheic chemistry. 188 The reach containing site 16 (S16) was investigated by 189 Malcolm et al. (2002, 2003b) using hydrometric, tracer and 190thermal data which indicated that the hyporheic zone was 191 dominated by surface water at this site. 192

At each site, novel methods for measuring hyporheic 193 water quality and embryo survival were employed. High 194 resolution DO and temperature data were obtained between 195 04 November 2005 and 11 April 2006, from the stream and 196 an artificially constructed redd at depths of 150 and 250 mm 197 in the hyporheic zone using Aandera 4175 shallow water 198 (rated to 300 m) DO optodes with analogue converters 199



Fig. 1 Location maps showing a the position of the River Dee catchment within the UK, b the position of the Girnock Burn within the River Dee catchment and c the location of study site 7 (7) and site 16 (16) within the Girnock Burn catchment

(Fig. 2). These were connected to Campbell dataloggers 200201programmed to sample DO (per cent saturation) and 202temperature at 30 second intervals and log average values over 15 min. Prior to installation, DO optodes were cross-203204calibrated in the laboratory at a range of O₂ concentrations 205and temperatures showing agreement to within $1\% O_2$ 206saturation and 0.1°C. Previous work in the same catchment 207 (Malcolm et al. 2006) had shown that in-situ installation for 208the period between spawning and egg hatch (ca. 5 months) 209without re-calibration provided excellent data quality. Data 210integrity was generally good, with the exception of two short periods early in the monitoring period at S16. 211

These high temporal resolution measurements were supplemented with high-spatial-resolution spot samples of DO, electrical conductivity and temperature from within vertically stratified incubation chambers (Fig. 2). The incubation chambers were adapted from those described by Youngson et al. (2005). Briefly, they comprised 217stacking 25-mm-high plastic containers, 42 mm in diam-218eter, regularly perforated with 6 mm holes. When screwed 219together, the containers formed a cylindrical column 220250 mm long. The top chamber was filled with stream 221gravel to exclude daylight. Each subsequent container was 222 lined with a 1-mm plastic mesh and contained 20 water-223 hardened salmon eggs taken from a single male and female 224mating to exclude parental effects. Fish were obtained 225from the Fisheries Research Services (FRS) Girnock trap 226facility. A control group of eggs was held in surface water 227 at the Girnock incubator facility. The control accounted for 228hyporheic affects on survival and performance by main-229taining oxygen concentrations near saturation for the entire 230incubation period between spawning and hatch. 231

In November 2005 (spawning time in the Girnock 232 Burn), the cylindrical arrays were placed into pre-prepared 233

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Fig. 2 Sampler design and installation within an artificially constructed redd

inserts within artificial redds, constructed at locations used 234 by spawners in previous years. The insert was then 235withdrawn from around the cylinder and any resulting gaps 236237were filled with surrounding gravel material (>4 mm). This 238resulted in egg chambers at depths of 25, 50, 75, 100, 125, 150, 175, 200, 225 and 250 mm beneath the streambed. A 239narrow diameter (4 mm i.d.) Nalgene tube led from each 240chamber to the streambed. During sampling, a volume 241 equivalent to that held in the sampling tube was discarded 242and a sample (25 ml) approximately equivalent to that held 243in the containers (container volume-ova volume) collected 244to characterise water quality in the immediate vicinity of the 245246ova. When not used for sampling a small plastic plug prevented direct connection between sampler and surface 247water. DO and temperature were measured using a 2-mm-248249diameter DO micro-sensor and thermistor connected to a 250Pre-Sens Fibox3 oxygen meter. The manufacturer stated reporting resolution for DO varies from 0.05% Sat. at 1% 251Sat. to 0.5% Sat. at 100% Sat. Accuracy is stated as $\pm 1\%$ 252Sat. at 100% Sat. to $\pm 0.15\%$ at 1% Sat. The reporting 253254resolution for temperature is 0.2° C with an accuracy of $\pm 1^{\circ}$ C. Electrical conductivity was measured using a Hannah HI 2559033 portable conductivity meter, reporting resolution 2560.1 μ S/cm, accuracy $\pm 2 \mu$ S/cm (0–200 μ S/cm range). Spot 257samples were collected at approximately fortnightly inter-258259vals where discharge and icing conditions permitted (n=7). 260Spot samples were compared with continuously logged 261data from the same depth to assess the comparability of 262methods. The chambers were excavated from the stream 263 bed on the day of the last sample collection on 11th April 2642006. Live and dead eggs were counted to provide 265percentage survival rates.

266 **Results**

267 **Temporal variability in hyporheic conditions**

268 Spawning-hatch (in-situ sampling)

269 The 2005–2006 spawning to hatch period (ca. November–

270 April in the Girnock catchment) was relatively dry with only

four moderate flow events over 3 m³/s (cumecs; Fig. 3). 271Stream temperatures at S7 and S16 were broadly similar. 272Early November was characterised by declining stream 273temperatures, with frequent icing events throughout the 274winter, before warming once more during March. The last 275icing period in early March corresponded to a prolonged 276period of late winter snowfall, whose subsequent melt 277resulted in a period of moderately elevated flows. 278

Previous hydrochemical and hydrometric work at the 279study sites indicated contrasting GW-SW interactions, with 280 the hyporheic zone of S7 being influenced by variable 281contributions of groundwater (Malcolm et al. 2004), while 282S16 was dominated by surface water (Malcolm et al. 2005). 283These differences in GW-SW interactions were reflected in 284different hyporheic temperature and DO characteristics 285between the sites. At S16, streambed temperatures were 286slightly moderated, showing less variable temperatures than 287surface water, with differences being most apparent at 288greater depths and during freezing periods (Fig. 3). At S7, 289 stream and shallow hyporheic water (150 mm) exhibited 290similar temperature characteristics. However, hyporheic 291water at 250 mm initially exhibited similar temperatures, 292 with moderation of temperature extremes increasing over 293time. This is consistent with increasing groundwater 294influence, where groundwater is typically characterised by 295more stable temperatures which are higher than surface 296water during winter months (Hannah et al. 2004). Differ-297ences in stream and hyporheic temperatures for the entire 298period where data were available at both sites are 299summarised in Figs. 4a and b. Only the 250 mm depth 300 sampler at S7 (S7-250) exhibited a notably different 301 thermal regime, showing some temperature moderation. 302

DO concentrations at S16 remained close to saturation in 303 both the stream and hyporheic water for the majority of the 304 study period, although small and short-lived gradients were 305 observed, particularly in the final months on the study 306 (Fig. 3). Between October and the end of February. DO at 307 150 mm was often lower than that at 250 mm. Much of this 308 variability can be explained by the moderated (generally 309 higher) temperatures in the streambed, which affect calcu-310 lated saturation values, i.e. there is no change in oxygen 311



Fig. 3 a Girnock Burn discharge; temperature at **b** S16 and **c** S7; and dissolved oxygen at **d** S16 and **e** S7, for the period between salmon spawning and embryo hatch. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm

concentration (mg/L), but small differences in temperature
change expected saturation values. Over the course of the
study, five periods of notable DO reductions were observed
where levels dropped below 70%. Four of these periods
were observed during the final month of the study.

At S7 DO concentrations in stream and shallow (150 mm) hyporheic water remained at or near saturation throughout the study. However, at 250 mm, concentrations were characterised by a dynamic response, varying between 0 and 100% saturation, often varying markedly over short periods in response to hydrological events. Typically, DO levels fell on the recession limb of storm hydrographs 323 shortly after peak discharge in agreement with observations 324 from previous years (Malcolm et al. 2004, 2006). DO 325 concentrations tended to recover in the aftermath of events. 326 Recovery times varied depending on event magnitude and 327 antecedent catchment wetness, which are thought to 328 influence water table elevation in the adjacent hillslopes 329 at this site (Malcolm et al. 2004, 2006). Between January 330 and the end of February, DO recoveries were only partial. 331In the final stage of the study from March onwards, DO 332 levels failed to exhibit any response recovery. Inter-site 333



Fig. 4 Temperature at a S16 and b S7, and dissolved oxygen at c S16 and d S7; duration curves for the period between spawning and embryo hatch. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm

differences in DO are summarised in the duration curves 334 335 shown in Fig. 4c and d. At S16 DO was always above saturation in surface water and at, or near saturation at 336 337 150 mm. DO at 250 mm was near to saturation for the 338 majority of the study dropping below 80% sat. for less than 3% of the time. At S7 DO concentrations were close to 339 saturation in surface water and at 150 mm for the entire 340 study period. However, at 250 mm DO concentrations 341 were near to saturation for only $\sim 30\%$ of the time, which is 342 comparable to the time spent at 0% saturation. 343

344 Event responses (in-situ sampling)

Event responses varied between sites, depending on event magnitude and antecedent conditions. Three contrasting event responses were identified: (1) DO response identi-

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fied only at S7, (2) DO response observed at both sites, 348 and (3) DO response observed at S16 with S7 characterised by constant low DO at 250 mm. 350

On the 10 November 2005, a complex double-peaked 351hydrograph was accompanied by mirrored declines in DO 352 concentrations at S7-250, punctuated by a short period of 353saturated DO at the main event peak (Fig. 5e). Falling DO 354concentrations on the recession limb were followed by 355 fairly rapid recovery. At S16, only a very slight decline in 356 DO was observed at 250 mm on the recession limb 357following the main event peak. Temperatures in the stream 358and hyporheic zone were similar at both sites, though small 359 differences at S7–250 were associated with the event peak. 360

Figure 6 shows a later event (12 January 2005) where 361 hyporheic DO concentrations declined at 250 mm at both 362 S7 and S16. On the rising limb of the hydrograph and at 363



Fig. 5 Event based (November 2005) oxygen and temperature responses showing: a discharge; temperature at b S16 and c S7; and dissolved oxygen at d S16 and e S7. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm

364the event peak, DO levels in stream water, S16 (150, 250 mm) and shallow hyporheic water at S7 (150 mm) 365 366 were close to saturation. In contrast, at S7–250, DO levels 367 declined on the rising and falling limb of the hydrograph, 368 with elevated DO levels during peak flow (Fig. 6e). At 369 S16-250, DO concentrations exhibited a small decrease in 370 DO on the recession, which was considerably lagged relative to that at S7. Following the event, DO concen-371 372 trations at both S7-250 and S16-250 recovered to near 373 saturation within 3 days. Temperature data from S16 (Fig. 6b), shows moderation of warmer pre-event and 374cooler post-event water. S7-250 exhibited distinct strati-375fication from surface and shallow hyporheic water on the 376 recession limb, while temperatures at S7-150 were 377 identical to those of surface water (Fig. 6c). 378

Towards the end of the monitoring period (22 March 2006), S7–250 was consistently characterised by near-zero

DO concentrations for almost a month (Fig. 3e). However, 381during this period, S16 exhibited a series of unusual, mode-382rate and occasionally prolonged reductions in DO that ap-383 peared to be associated with only minor hydrological events 384 in the Girnock Burn (Fig. 7d). Temperature data from S16 385 showed a moderated temperature gradient with depth that 386 also showed a lagged response (Fig. 7b). At S7, temperatures 387 in the stream and at 150 mm closely tracked, while temper-388 ature at 250 mm exhibited marked thermal moderation. 389

Fine scale spatial variability in hyporheic water quality (ex-situ sampling)

The continuous water quality monitors provide data of 392 excellent temporal resolution, but only provide relatively 393 coarse spatial information on hyporheic water quality. The 394 integrated embryo survival chambers and samplers facil- 395

390

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Fig. 6 Event based (January 2006) oxygen and temperature responses showing: **a** discharge; temperature at **b** S16 and **c** S7; and dissolved oxygen at **d** S16 and **e** S7. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm

396 itated collection of water samples at 25 mm vertical 397 resolution (25-250 mm) which revealed the fine-scale 398 spatial variability of hyporheic water quality with depth, 399 although at the expense of temporal resolution (Fig. 8). Additionally, spot samples allowed the collection of 400 electrical conductivity data (Fig. 8a, b) as an indicator of 401 402 source water provenance (Youngson et al. 2005). Electrical conductivity and DO saturation at S16 were relatively 403 uniform with depth over the entire study period indicating 404 a common source water. By contrast, depth-related 405stratification of both DO and conductivity was apparent 406 at S7 over much of the study and appeared to increase 407over time. Differences in conductivity and DO were 408409 consistent with an increasing groundwater influence with

depth (Malcolm et al. 2005). Higher conductivity values 410 indicative of longer residence water were generally 411 associated with lower DO. Stratification gradients at S7 412 were steep, with DO varying from nearly 100% saturation 413 to <10% over distances of only 50 mm. Gradients in DO 414appeared to be more consistent with depth than those 415 exhibited by electrical conductivity and it is possible that 416 this inconsistency reflected mixing between samples 417 collected from adjacent depths despite very low volumes. 418

A comparison of in- and ex-situ sampling methods 419 A comparison of the spot sample DO data, with 420 continuous data from the optodes located at approximately 421



Fig. 7 Event based (March 2006) oxygen and temperature responses showing: **a** discharge; temperature at **b** S16 and **c** S7; and dissolved oxygen at **d** S16 and **e** S7. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm

the same depth, reveals that the two methods were 422generally comparable for a given sampling occasion 423(Fig. 9). The two methods also produced broadly similar 424patterns of variability, although very few low DO spot 425samples were obtained due to the coarse sampling 426frequency. Given the fine-scale spatial variability of DO 427 428 revealed by the spot sampling, difficulties locating equipment with a high degree of spatial precision beneath 429430 the streambed and complexities associated with crosscalibration of seven independent measuring units, it is not 431 surprising that the two methods did not provide exactly 432the same DO values. However, a paired *t*-test (n=26)433 revealed that there was no significant difference between 434 the data obtained using the two methods (P=0.27). When 435436comparing the methods, it is clear that each has merit. The loss of temporal resolution is evident in the spot samples, 437 while the continuous data lacks potentially important fine 438439scale spatial resolution.

Embryo survival

In 2003–2004 and 2004–2005 embryo survival at controls 441 held at the Girnock Burn was 100%. During the 2005-442 2006 spawning season, unusually high mortality of 443fertilised ova occurred. The reasons for this mortality are 444 unclear, but appeared to affect many groups of ova 445 reflecting reduced viability in general or unknown 446 procedural problems during adult stripping or fertilisation. 447 Survival in the control group was 70%, although across 448 the incubator as a whole it was on average closer to 50%. 449Given this background, interpretation of ova survival at S7 450and S16 is difficult and it is possible that variability 451between sites and depths reflected random sampling from 452a variably impacted group of fertilised ova at the project 453outset. Embryo survival in the streambed incubators 454varied from 0-60% (Table 1) and for the most part did 455not show clear patterns that could be associated with 456 environmental variation. Nevertheless, at S7, complete 457

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Fig. 8 Temporal and spatial variability of electrical conductivity at a S16 and b S7 and dissolved oxygen at c S16 and d S7 in surface (S) and hyporheic water at depths ranging from 25–250 mm (see *legend*), separated at 25-mm intervals. Approximately fortnightly sampling occasions are shown as points

ova mortality was observed across the depth range 200–
250 mm. These mortalities are consistent with the sharp
DO concentration gradients observed at S7 for these
depths (Fig. 8). In contrast, survival at S16 was generally
higher than at S7, especially in the lower hyporheic zone,
though even here, survival at 250 mm was only 25%.

464 **Discussion**

Influence of local GW–SW interactions on fine scale spatio-temporal variability of hyporheic water quality At S7, DO concentrations varied spatially and temporally in a manner that was consistent with changing groundwater contributions to the hyporheic zone. Low DO was associated with higher electrical conductivities, thought to be associated with increased residence times. Groundwater influence was associated with steep DO gradients 472 (distances of ca. 0.05 m), which shifted vertically over 473time. This is contrary to the common conceptual under-474standing of a broad hyporheic mixing zone containing 475 groundwater and surface water (e.g. Malard et al. 2002) 476 and is more consistent with a temporally shifting sharp 477 boundary between groundwater and surface water, with 478limited mixing. DO concentrations changed rapidly in 479response to hydrological events (Malcolm et al. 2006). 480 The exact form of the response varied with antecedent 481 conditions and discharge magnitude. In general, dry, low 482 flow periods were characterised by high DO, while low 483 DO was observed during periods of wet antecedent 484 conditions, later in the winter, and the recession limb of 485hydrological events where water table levels are high. 486

At S16, where surface water dominated the hyporheic 487 zone, DO concentrations in the stream bed were compa-488



Fig. 9 A comparison of ex-situ spot sample data (ca. fortnightly) with in-situ continuous (15 min) data for comparable depths. Plots show: a S16 electrical conductivity spot samples, b S16 dissolved oxygen spot samples, c S16 continuous dissolved oxygen, d S7 electrical conductivity spot samples, e S7 dissolved oxygen spot samples, and f S7 continuous dissolved oxygen. *Black lines* show surface water, *green lines* show hyporheic water at 150 mm, *red lines* show hyporheic water at 250 mm. *Symbols* denote spot-sampling occasions

t1.1 **Table 1** Percentage embryo survival for depths ranging from 25–250 mm at site 7 (S7) and site 16 (S16)

t1.2	Depth (mm)	% survival	
t1.3		S7	S16
t1.4	25	40	35
t1.5	50	25	40
t1.6	75	55	50
t1.7	100	60	60
t1.8	125	40	40
t1.9	150	40	45
t1.10	175	40	55
t1.11	200	0	45
t1.12	225	0	45
t1.13	250	0	25

rable with stream water and consequently near saturation 489 for the majority of the monitoring period, although low 490DO episodes were observed towards the end of the study. 491 Reductions in DO were not associated with increased 492electrical conductivity and thus appear unlikely to be 493associated with intrusion of groundwater. On excavation, 494the incubation chambers were found to be entirely free of 495 sediment, and therefore it also seems very unlikely that 496 reductions in DO were associated with intrusion of fine 497 sediment to the redd environment. It is possible that 498 changing DO levels reflected changing short residence 499(hours to days) hyporheic dynamics at the site associated 500with changing bed morphology during the study period. 501 High flows over the winter led to the development of a 502substantial gravel bar immediately upstream of the 503

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504monitoring site. It is possible that hyporheic exchange 505passing through the bar feature, re-emerged on the downstream side under certain flow conditions (Tonina 506 507and Buffington 2007) and that DO could have been stripped from the water during transit (Claret et al. 1997). 508Under these circumstances the residence time would be 509too short for substantial changes to more conservative 510511water quality parameters but DO could be reduced. Alternatively, it is possible that low DO episodes reflected 512discharge of hyporheic water from the River Dee as S16 513lies at the bottom of the Girnock catchment, within course 514gravel sediments associated with the Dee floodplain. It is 515therefore possible that high flows from the River Dee, 516entering an abandoned channel adjacent to the Girnock 517Burn, could have altered local hyporheic dynamics 518resulting in discharge of Dee water or displacement of 519Girnock floodplain water through S16 (Rodgers et al. 5202004: Poole et al. 2006). 521

Implications for hyporheic sampling 522

Palmer (1993) identified a number of key challenges for 523hyporheic zone research. These included the need to 524conceptualise hyporheic zone boundaries through under-525standing of inter-site heterogeneity and the development 526of methods to sample the hyporheic environment at small 527spatial scales that could be calibrated and quantified in 528terms of spatial extent. This study combined adaptations 529of recently documented hyporheic sampling methodolo-530gies (Youngson et al. 2005; Malcolm et al. 2006) to 531

identify fine scale spatial and temporal differences in 532hyporheic chemistry and embryo survival at two salmon 533spawning locations with contrasting GW-SW interactions. 534

While in-situ methods revealed important temporal 535variability, the stratified incubators and ex-situ sampling 536method provided valuable information on the spatial 537 variability of water quality, embryo survival and also 538provided supporting hydrochemical data. For the depths 539and times for which data could be compared (150 and 540250 mm), the two methods showed good agreement (no 541significant difference between methods), indicating that 542in-situ measurements did not reflect unrepresentative 543micro-scale (mm's) conditions and, more importantly, that 544the two methods generated comparable data and therefore 545could be deployed in a stratified sampling programme to 546 give both high resolution spatial and temporal data in 547future expanded studies. 548

Implications for hydro-ecological studies of the hyporheic zone

Previous studies of the hyporheic environment have often 551used large or unspecified sample volumes and infrequent 552sampling intervals. These water quality data are then often 553related to hyporheic ecology such as invertebrate commu-554nities (Boulton et al. 1997; Mermillod-Blondin et al. 2000; 555Fowler and Death 2001) or salmonid embryo survival 556without fully considering the spatial and temporal scales of 557the water quality sampling, the variability of the hyporheic 558environment or the scales relevant to the ecology. Table 2 559

Table 2. Comparison of hypothetic oxygen sampling methods and frequencies for studies investigating salmonid embryo survival +9.1

	Tuble 2 Comparison of Appointee experimentation and nequenees for statutes investigating statistical entry of survival					
Authors		Water sampling method (DO measurement method)	Sample volume	Sample depth (m)	Sample frequency	
Malcolm et al	l. 2006 I	In-situ (Aanderaa DO optode)	NA	0.15, 0.3	30 s, averaged every 15 min	
Groves and C (2005)	handler I	Buried incubators with sampling tubes and piezometers. (flow-through cell and YSI DO electrode)	3× dead volume discarded sample volume for measurement unknown	0.25	Monthly	
Greig et al. (2	2005)	Standpipe (YSI 250 DO electrode)	Not stated	Not stated	Weekly to fortnightly	
Youngson et a	al. (2005)	Sealed flexible hyporheic sampling tubes (Hannah DO electrode)	Dead volume discarded 200 ml sample	0.2–0.3	Fortnightly	
Bernier-Bourg Magnan (2	gault and solution (2002)	Sampling pipe installed on sampling date (YSI 57 DO electrode)	Not stated	0.05-0.15	Not stated	
Bowen and N (2003)	lelson	Variable depth hyporheic sampling pipes (unspecified multi-parameter meter including DO electrode)	Not stated	0.3, 0.46	2 samples, 1 month apart	
Ingendahl (20	01) 1	Flexible sampling tube (portable DO electrode)	60 ml discarded 60 ml sample	0.1, 0.2, 0.3	Fortnightly	
Peterson and 0 (1996)	Quinn	Sampling tube (titration)	Dead volume discarded 185 ml sample	Variable, depending on egg burial depth	Weekly to fortnightly	
Sowden and 1 1 (1985)	Power 1	Mini-piezometer (YSI 54 DO electrode)	150 ml sample	0.15	Approximately monthly	
Ringler and H	Hall (1975)	Standpipe (titration)	60 ml	0.25	3 samples per week	
3 Coble (1961)	c.	Standpipe (not stated)	37 ml	0.25	Not stated	

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560shows a comparison of hyporheic studies, where the 561research focus was to understand salmonid embryo survival. It can be seen that sample depths generally 562563reflect the reported range of egg burial depths (DeVries 1997). However, the number of reported depths is typically 564only 1-3 (relatively coarse) and the sampling methods and 565566 volumes are highly variable or are not clearly specified. 567This effectively means that, depending on streambed characteristics and equilibration times, individual studies 568 will be reporting hyporheic water quality for highly 569variable, but generally poorly delineated volumes of 570extracted streambed water that are unlikely to reflect the 571environmental conditions experienced by the hyporheos, in 572this case salmonid embryos. If the temporal variability of 573hyporheic water quality and the general inadequacy of 574sampling frequency is also considered, then it is unsur-575prising that the results of field (Sowden and Power 1985; 576Rubin and Glimsater 1996: Ingendahl 2001) and laborato-577 ry (Alderdice et al. 1958; Silver et al. 1963) based studies 578 579of salmonid embryo survival are not in good agreement. 580Disparities in the apparent findings of these approaches probably reflect the controlled nature of laboratory experi-581ments and problems with adequately characterising an 582environment that is as temporally and spatially highly 583 584variable and inaccessible as the hyporheic zone.

Implications for salmonids 585

At S7, there was a sharp transitional gradient in hyporheic 586water quality over distances of <0.05 m which was reflected 587in the total mortality of embryos at greater depths. In recent 588 years there has been frequent discussion of the benefits of 589 greater burial depth to avoid washout or overcutting by later 590 arriving female fish (Steen and Quinn 1999). Since larger 591fish generally bury their eggs deeper (Crisp and Carling 5921989; DeVries 1997; Steen and Quinn 1999), there has been 593debate as to whether larger fish are favoured in locations 594where scour or super-imposition are likely to be problem-595atic. However, the results of this study show that burial 596597 depth can also impact on survival where reduced hyporheic 598water quality is associated with groundwater upwelling. 599Moreover, very small (0.025 m) differences in burial depth can have a potentially very large impact on survival. 600 Therefore, in terms of spawning, there may be a careful 601 602 tradeoff to be made between avoiding scour on the one hand and avoiding hypoxia of developing embryos on the other. 603

Much salmon-focussed research to date has focussed 604 605 on the sediment component of hyporheic dynamics. This has led to proposals for fine sediment water quality 606 standards under legislation such as the Water Framework 607 Directive and Habitats Directive (Naden et al. 2002) of the 608 European Union. It has also led to the development of 609 simplified tools (Alonso et al. 1996; Wu 2000) which do 610 not consider the full range of hyporheic processes relevant 611 to an understanding of embryo survival. This paper has 612 highlighted both the importance of appropriate sampling 613 614 methods and a holistic understanding of hyporheic processes, which includes understanding of local GW-615 SW interactions for understanding hyporheic ecology. 616

Future research

This study lasted only for 1 year, focussing on a particular 618 aspect of hyporheic ecology over a relatively short, but 619 ecologically relevant time period. The issues highlighted 620 in relation to the spatial and temporal scale of sampling 621 are clear. However, further work is required to assess the 622 influence of local GW-SW interactions on other aspects of 623 the ecology and to characterise and understand the 624 influence of antecedent conditions on catchment hydrolo-625 gy (e.g. Tetzlaff et al. 2007b) and the effect that this has 626 on GW-SW interactions at longer temporal scales. 627

Acknowledgements The authors would like to acknowledge staff 629 from the Environment and Resources Groups at FRS Freshwater 630 Laboratory for field assistance during this project and the Scottish 631 Environment Protection Agency (SEPA) for the provision of 632 discharge data. 633

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