

Predicting the nationwide outmigration timing of Atlantic salmon (*Salmo salar*) smolts along 12 degrees of latitude in Norway

Knut Wiik Vollset¹  | Robert J. Lennox¹  | Anders Lamberg² | Øystein Skaala³ | Anne Dagrund Sandvik³ | Harald Sægrov⁴ | Eli Kvingedal⁵ | Torstein Kristensen⁶ | Arne Johan Jensen⁵ | Tormod Haraldstad⁷ | Bjørn Torgeir Barlaup¹ | Ola Ugedal⁵

¹Laboratory for Freshwater Ecology and Inland Fisheries at NORCE Norwegian Research Centre, Bergen, Norway

²Scandinavian Nature Surveillance, Ranheim, Norway

³Institute of Marine Research, Bergen, Norway

⁴Rådgivende Biologer AS, Bergen, Norway

⁵Norwegian Institute for Nature Research (NINA), Trondheim, Norway

⁶Faculty of Biosciences and Aquaculture, Nord University, Bodø, Norway

⁷Norwegian Institute for Water Research, Grimstad, Norway

Correspondence

Knut Wiik Vollset, Laboratory for Freshwater Ecology and Inland Fisheries at NORCE Norwegian Research Centre, Bergen, Norway.
Email: knvo@norceresearch.no

Funding information

Ministry of Trade, Industry and Fisheries

Editor: April Blakeslee

Abstract

Aim: Accurate predictions about transition timing of salmon smolts between freshwater and marine environments are key to effective management. We aimed to use available data on Atlantic salmon smolt migration to predict the emigration timing in rivers throughout Norway.

Location: In this study, we used outmigration timing data of Atlantic salmon (*Salmo salar*) smolts from 41 rivers collected from 1984 to 2018 to make a predictive model for the timing of out-migrating salmon smolts along 12 degrees latitude.

Methods: Smolt migration data were collated from rivers where smolts are monitored with traps, video and tagging and matched to river-specific metadata. Using a mixed-effects generalized additive model, we tested for the effects of spatial location, air temperature, river flow and counting methods on the timing of 25% smolt emigration from rivers.

Results: After accounting for spatial effects and repeated measures (across years and among rivers), air temperature and counting method were significant drivers of the estimated timing of smolt emigration. In-sample predictions yielded strong correlation with observed values, as did 10-fold cross-validation. Out-of-sample predictions suggested that the previous national estimates underestimated the migration timing in southern populations (linear model intercept = 39.73 days).

Conclusion: Model-derived estimates of run timing can be used to more accurately predict the timing of outmigration in order to better calibrate environmental flows and regulate management of marine industries such as aquaculture that may affect migration success at this particularly sensitive life stage.

KEYWORDS

biodiversity, migration, phenology, restoration, Salmonidae, telemetry

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

© 2021 The Authors. *Diversity and Distributions* published by John Wiley & Sons Ltd.

1 | INTRODUCTION

Atlantic salmon born in freshwater move to the ocean following smoltification to exploit the productivity of marine waters (Klemetsen et al., 2003; McCormick et al., 1998). The post-smolt life stage when salmon are exiting the rivers and transitioning to life in the marine environment is a critical bottleneck in their survival (Lothian et al., 2018; Stich et al., 2015; Thorstad et al., 2012). The timing of migrations has been adapted by evolutionary processes to match optimal conditions for survival (Lennox et al., 2016); for Atlantic salmon smolts, this window of opportunity is narrow in order to match physiological preparedness for migration with environmental conditions (McCormick et al., 1998). Optimal conditions are not directly observable by animals; therefore, they rely on secondary cues to synchronize their migration timing (e.g. Duston & Saunders, 1990). Deviations from the optimum time frame can yield high mortality when anadromous species enter the ocean to encounter suboptimal conditions (Scheuerell et al., 2009). For example, Hansen and Jonsson (1989) released smolts from the River Imsa throughout the year and observed returning adults came from groups released in the spring, illustrating the sensitive time window that smolts must initiate their seaward migration for their long-term success.

Evidently, there is strong pressure for Atlantic salmon smolts to migrate at appropriate times (Otero et al., 2014). Anthropogenic effects that have negative impacts on post-smolts during this window are of great concern to salmon conservation (McCormick et al., 1998; Thorstad et al., 2012). Consequently, management must have the appropriate tools available to them in order to predict the timing of the smolt migration to evaluate how such effects may overlap with the smolt migration and give advice on when mitigation efforts are most effective. Yet, the most frequently used tool for this is presently the designation of index rivers, nearby watersheds that are monitored and used to predict the timing of the smolt migration in all other proximate systems (e.g. Johnsen et al., 2021; Kristoffersen et al., 2018). Models have been developed to predict the timing of smolt migrations in single rivers and generally show that temperature and discharge are important to explaining the timing of smolt migration in rivers (Hansen & Jonsson, 1989; Whalen et al., 1999). Models across several rivers that attempt to predict general patterns of smolt migration have also been made, but with variable overarching goals. For example, Hvidsten et al., (1998) used data from five rivers to postulate that salmon migrate to sea when marine temperature exceeds 8°C. More recently, Otero et al., (2014) synthesized data throughout the range of Atlantic salmon showing a temporal shift in the timing of outmigration, and a correlation with both sea surface temperature and freshwater temperature.

A model that can be used to accurately predict outmigration timing on relevant management scales (such as aquaculture production zones in Norway) would clearly be of great importance for the conservation of Atlantic salmon. Generating predictions for the timing of smolt migration can provide a tool for monitoring the status of Atlantic salmon rivers and observing changes across time. An effective predictive model could be used to make predictions for rivers

that are not monitored due to access or financing challenges (i.e. out-of-sample). We collated smolt migration data from Norwegian Atlantic salmon rivers to generate a model of the spatial and environmental features predicting the timing of smolt migration for years between 1984 and 2018. We used regression modelling to test the relationships and validate the predictive power of the model, generate out-of-sample predictions and compare the predicted timings to estimates used by national management councils.

2 | METHODS

2.1 | Data collection

2.1.1 | Smolt migration data

The goal of our study was to collate available smolt migration data from Norway, in an attempt to make a predictive model of the timing of smolt migration in this area. Data were extracted from three sources: (a) published scientific articles, (b) Norwegian reports, and (c) unpublished data available from the authors' research institutions. Data were updated from previous compilations using the same methodology (Ugedal et al., 2014). The database does not contain daily counts but summarizes the timing of smolt emigration by percentiles, recording the dates of 25% passage, 50% passage and 75% passage (Table 1).

Smolt migration was monitored in 348 river years, comprising 47 rivers between 58.02 and 70.50 degrees latitude from 1984 to 2018 (Figure 1). Monitoring was conducted using different methods of observation: traps ($N = 252$ river years), video counting ($N = 84$ river years) and tagging ($N = 12$ river years). Note Eio and Vigda are the only two rivers that had multiple counting methods, so they are counted multiple times. The placement of these monitoring tools was not consistent among rivers nor was the timing of deployment standardized.

2.1.2 | River morphology data

Morphological data from the river catchments were downloaded from Nevina (<http://nevina.nve.no/>). This includes elevation data from the catchment, land composition (e.g. per cent of catchment covered by agriculture, forest, lake and urban areas) and air temperature throughout the year (summer, winter, July, August temperatures). In addition, modelled average discharge, average rainfall and average air temperature were extracted from each of the catchments from the same database.

2.1.3 | Annual environmental data

Seasonal water temperature measurement data were not available for the majority of rivers. However, air temperature data were

TABLE 1 Summary of modelled variables including those in the full and reduced models. Rivers in Norway are characterized by ID numbers assigned by the national resource authority, which are included for reference. Observed dates of 25% outmigration ($\pm SD$) from the data are included along with the number of years of data, the counting method, river coordinates, elevation and grade, whether the river has lakes, and the mean air temperatures and river flows extracted for modelling (see below for details)

River	Norway ID	25% Migration	Lon	Lat	Method	Grade	Lakes	Elevation	Years	Mean temp	Avg. first julian day of 10% flow
Mandalselva	022.Z	128 \pm 9	7.5	58.0	Trap	8.6	0	1,160	16	1.5	95.4
Tovdalselva	020.Z	132 \pm 8	8.1	58.2	Trap	6	0	1,146	6	1.9	93.7
Kvina	025.Z	124	6.9	58.3	Trap	9.3	0	1,430	1	0.6	96.0
Nidelva	019.Z	130 \pm 7	8.7	58.4	Trap	5.8	1	1519	6	1.8	95.8
Storelva	018.Z	127 \pm 6	9.1	58.7	Trap	6.4	1	506	12	1.4	101.3
Imsa	029.2Z	122 \pm 4	6.0	58.9	Trap	9.3	0	487	19	1.7	91.8
Numedalslågen	015.Z	120 \pm 9	10.1	59.0	Trap	3.5	0	1537	3	1.4	117.7
Suldalslågen	036.Z	117 \pm 4	6.3	59.5	Trap	13.2	1	1686	16	0.9	100.2
Etneelva	041.Z	128 \pm 6	5.9	59.7	Trap	30.5	1	1,294	3	1.3	91.0
Uskedalselva	045.2Z	120	5.9	59.9	Video	40.5	0	1,245	1	2.3	91.0
Guddalselva	045.32Z	124 \pm 8	6.0	60.0	Trap	2.7	0	1,430	18	1.1	92.8
Oselva	055.7Z	127	5.5	60.2	Tag	29.5	1	955	1	0.1	95.0
Kinso	050.1Z	145	6.7	60.4	Video	30	0	1686	1	-0.5	115.0
Arna/Storelva	061.2Z	119	5.5	60.4	Video	43.9	1	983	1	0.5	91.0
Eio	050.Z	130	7.1	60.5	Tag	0.4	1	1853	1	-0.9	130.0
Eio	050.Z	146 \pm 11	7.1	60.5	Trap	0.4	1	1853	2	-1.3	130.0
Tveito	050.Z	156 \pm 12	7.1	60.5	Video	0.4	1	1853	3	-1.2	125.0
Lone	060.4Z	128	5.5	60.5	Video	43	0	671	1	0.5	91.0
Granvin	052.1Z	126	6.7	60.5	Tag	43.7	1	1573	1	-0.6	100.0
Daleelva	061.Z	136 \pm 8	5.8	60.6	Trap	22	0	1,194	12	1.3	92.3
Vosso	062.Z	134 \pm 6	6.0	60.6	Trap	15.2	1	1602	17	1.0	97.9
Flåm	072.2Z	136 \pm 9	7.1	60.9	Trap	31.3	0	1761	5	0.1	132.0
Aurland	072.Z	136 \pm 8	7.2	60.9	Trap	30.3	0	1806	6	-0.3	132.3
Vikja	070.Z	110 \pm 12	6.6	61.1	Trap	52	0	1647	10	0.7	102.4
Lærdalselva	073.Z	129 \pm 9	7.5	61.1	Tag	18.4	0	1917	3	0.4	126.7
Osenelva	085.Z	144	5.4	61.5	Video	15	1	1,117	1	1.8	91.0
Stryn	088.Z	127 \pm 1	6.7	61.9	Tag	27.2	1	1933	2	0.9	101.5
Driva	109.Z	130 \pm 11	8.5	62.7	Trap	12.1	0	2,283	5	1.0	115.6
Eira	104.Z	133 \pm 5	8.1	62.7	Trap	16.5	1	1961	18	0.8	109.3
Hustadelva	107.6Z	135 \pm 10	7.1	63.0	Video	19.8	1	905	9	1.2	91.8
Surna	112.Z	143 \pm 7	8.7	63.0	Trap	8.3	0	1668	6	0.1	97.0
Vigda	122.2Z	128 \pm 4	10.2	63.3	Tag	12.9	0	533	2	-0.3	119.5
Orkla	121.Z	129 \pm 5	9.8	63.3	Trap	8	0	1,640	26	-0.4	114.4
Stjørdalselva	124.Z	136 \pm 6	10.9	63.4	Trap	10.1	0	1,245	6	-0.5	99.5
Moelva/ Salsvassdrag	140.Z	159 \pm 8	11.4	64.7	Video	6.2	1	763	11	-0.6	94.1
Urvollelva	144.5Z	147 \pm 9	12.6	65.1	Video	39.2	1	1,048	8	-0.5	99.8
Sila	157.52Z	162	13.1	66.3	Video	78.9	1	1,020	1	-2.9	110.0
Flostrand	157.42Z	158	13.4	66.3	Video	38.1	1	1,152	1	-3.1	110.0
Saltdalselva	163.Z	159 \pm 8	15.4	67.1	Trap	15.2	0	1741	4	-1.6	104.3
Skjoma	173.Z	151 \pm 7	17.4	68.2	Video	1.5	0	1,880	12	-2.5	119.7
Roksdalvass	186.2Z	153 \pm 7	15.9	69.1	Video	15.7	1	584	20	-1.7	95.6

(Continues)

TABLE 1 (Continued)

River	Norway ID	25% Migration	Lon	Lat	Method	Grade	Lakes	Elevation	Years	Mean temp	Avg. first julian day of 10% flow
Lakselva, Senja	194.Z	172 ± 6	17.8	69.2	Video	8.9	1	894	7	-2.7	100.4
Altaelva	212.Z	174 ± 9	23.4	70.0	Trap	2.2	0	1,085	10	-4.0	144.2
Halselva	212.2Z	162 ± 7	23.0	70.0	Trap	31.7	0	1,146	25	-3.7	139.2
Repparfjordelva	213.Z	171	24.3	70.4	Tag	9.1	0	712	1	-5.8	136.0
Utsjok, Tana	234.Z	176 ± 6	28.3	70.5	Video	1.4	0	1,064	7	-4.5	130.3
Kvalsundelva	213.6Z	171	24.0	70.5	Tag	21.5	0	707	1	-5.4	136.0

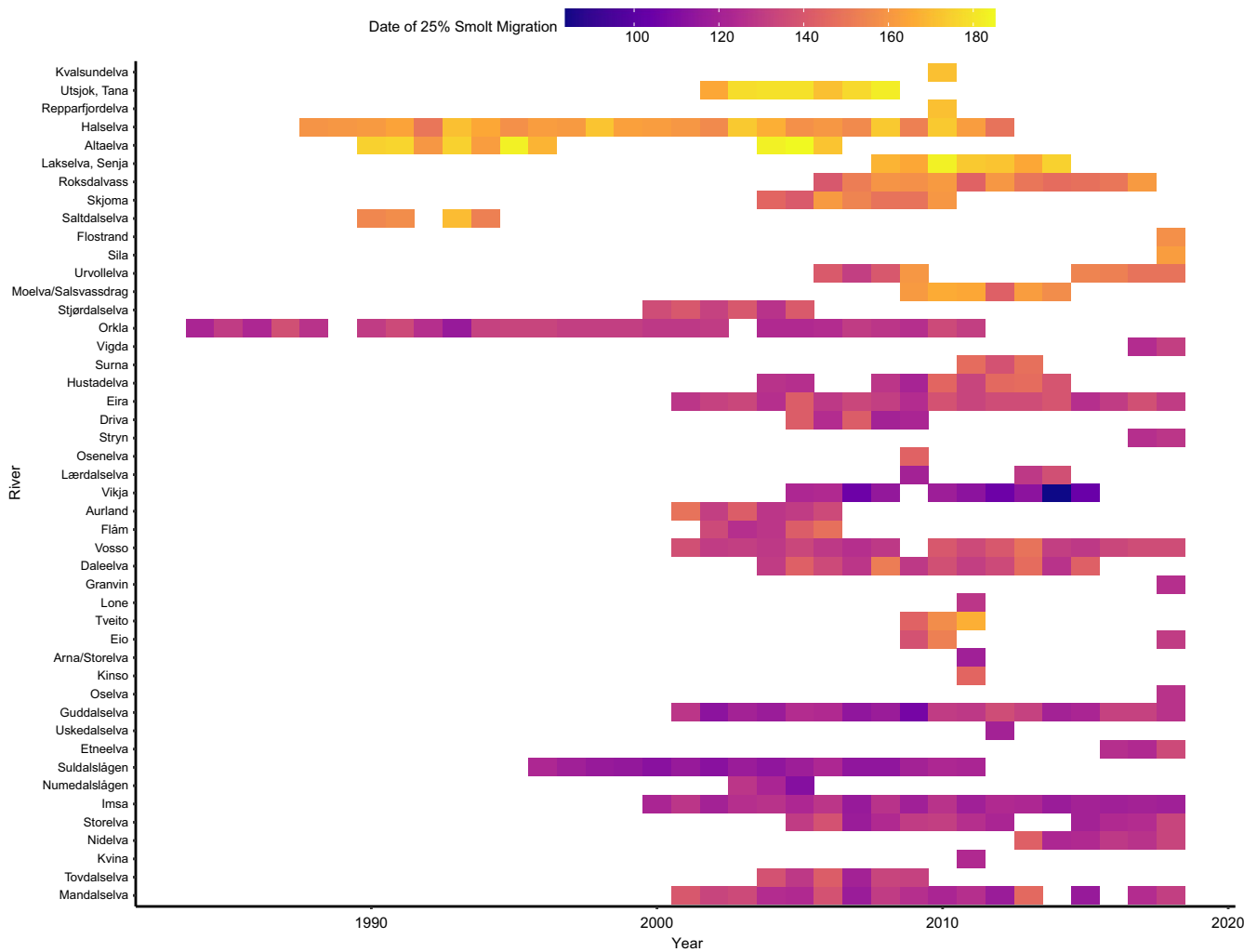


FIGURE 1 Tile plot of raw data for rivers represented in the dataset. Rivers are sorted from north to south. Filled in areas representing river years in which data were available for modelling with colour fill representing the Julian date of 25% smolt migration

collected by monitoring stations throughout Norway, for which historical data are freely available through an API using the *esd* package in R (Benestad et al., 2019). We downloaded temperature records from all available stations. A generalized additive model was fit to each individual year to explain the air temperature recorded by the longitude, latitude and altitude of the station using the *gam* function

in the *mgcv* package (Wood, 2017). The gam models were then carried forward to predict the air temperatures for each river in each year using coordinates of the river mouth, and average grade of the river using the *predict.gam* function. Air temperature for each river in each year was summarized as the average between 1 January and 31 March.

Modelled water discharge data were available for each river from the NorKyst800 model (Albretsen et al., 2011). The Norwegian river discharges were modelled by the NVE (Norwegian Water Resources and Energy Directorate) using a distributed version of the HBV model with 1 km horizontal resolution (Beldring et al., 2013; Huang et al., 2019). We summarized water discharge for each river in each year by extracting the first day of the year when the flow first hit 10 and 25% of the maximum flow from 1 March to 18 July, which was considered to be the maximal likely window for onset of smolt migration. We used different temporal windows for temperature and flow because temperature should control physiological readiness (proximate cause) and flow should drive the exact timing of migration (ultimate cause). Both flow extractions gave very similar model outputs.

2.2 | Data analysis

We started with a large dataset including physical river characteristics, land use, geographic and climate variables. We initially considered a principal component analysis (PCA) to summarize the data but opted to manually select variables of interest because (a) many were highly correlated, and (b) we wanted to understand the relationship between smolt run timing and specific variables and not a hybrid variable produced by PCA. Therefore, we discarded land use variables and several temperature variables that were highly correlated with the average air temperature variable (1 January to 31 March) that we retained.

We constructed generalized additive models using the *gam* function in the R package *mgcv* (Wood, 2017) to model the influence of longitude and latitude at the river mouth, mean air temperature in the first quarter of the year and first increase in spring discharge (25% as explained above). The independent variable of interest was the timing of 25% outmigration from the river, which was selected because it was more complete than the 50%, 75% and 100% estimates. However, these were all highly correlated so this selection should not influence the interpretations from the model, except that had we modelled 50% emigration the estimate would be shifted later in the year. We constructed two models, one including more variables and one with simpler, more accessible data. The first model included spatial variables latitude and longitude as a combined smoother (Pedersen et al., 2019) and linear effects for mean air temperature, date of flow being 10% of the annual maximum for the first time, the river height and gradient, whether the river included lakes in the anadromous section and the sampling method (video, PIT tagging, acoustic tagging, smolt screw trap, wolf trap), in addition to a random effects of river and year to account for measurements coming from the same river and in the same year (specified as a smoother using the argument *bs="re"*). The second model was simpler, with only the smooth terms for latitude and longitude together, linear effects of temperature and flow, counting method and a random intercept for each river and year. Predictions were generated using the *predict.gam* function (Wood, 2017).

Predictions were generated in sample and out of sample to estimate the run timing in rivers where we did not have estimates of the date of 25% smolt migration. For the out-of-sample predictions, we set the random effects of year and river to zero. A mixed-effects model of the in-sample predictions against the known values for the 25% smolt migration date was run with river as a random effect with the *lme* function in the *nlme* package (Pinheiro et al., 2019). Predictions were extracted for 2018 and compared with the national estimate of 25% smolt outmigration using a linear model with the *lm* function. Model performance was assessed by k-fold cross-validation by splitting the dataset into ten groups, training the simplified GAM model on nine subsets and using the tenth subset as a test set to compare predictions to the true values, rotating through all ten combinations of models and testing the correlation of predictions generated from the 10 models against the known values.

2.3 | Data visualization

Data were plotted using *ggplot2* (Wickham, 2016). The map was accessed from the *cshapes* R package with the *cshp* function (Weidmann & Gleditsch, 2016).

3 | RESULTS

We used 348 observations of the date of 25% smolt emigration from 47 rivers from 1984 to 2018 to derive our model (Figure 1). Observations ranged from 24 March (Vikja River, 2014) to 4 July (Alta River, 2005). The full and the simplified model fit similarly, but the simpler model was better ($\Delta AIC = 2.7$). We therefore proceeded with the simplified GAM model that included the random effect for river, and smoothers for latitude and longitude together, mean air temperature in the first quarter of the year and the first date of 25% flow. The model had a strong fit to the data ($R^2 = .86$; deviance explained = 88%). The smoother on longitude and latitude was highly significant ($F = 6.64, p < .01$). Air temperature in the first quarter was significant ($F = -4.81, p < .01$), but flow was not significant ($F = -0.93, p = .35$). The counting method was also significant, with estimates from tagging studies yielding earlier results than video counting ($t = 2.74, p = .01$) but not trap catches ($t = 1.66, p = .10$). Predictions were generated in-sample to determine how well the predictions from the model fit the known data, showing a significant correlation with the observations ($t = 15.37, p < .01$; Figure 2).

Out-of-sample predictions were then generated to estimate outmigration timing in river years throughout Norway ($N = 1,753$) based on the longitude, latitude, air temperature and flow in a given year; the capture method was set to the factor of trap (Figure 3). There was a strong relationship between GAM-predicted timing and the national estimate of outmigration timing for 2018 ($t = 46.45, p < .01, R^2 = 84\%$); the model fit (intercept 39.73, slope 0.78) suggested that

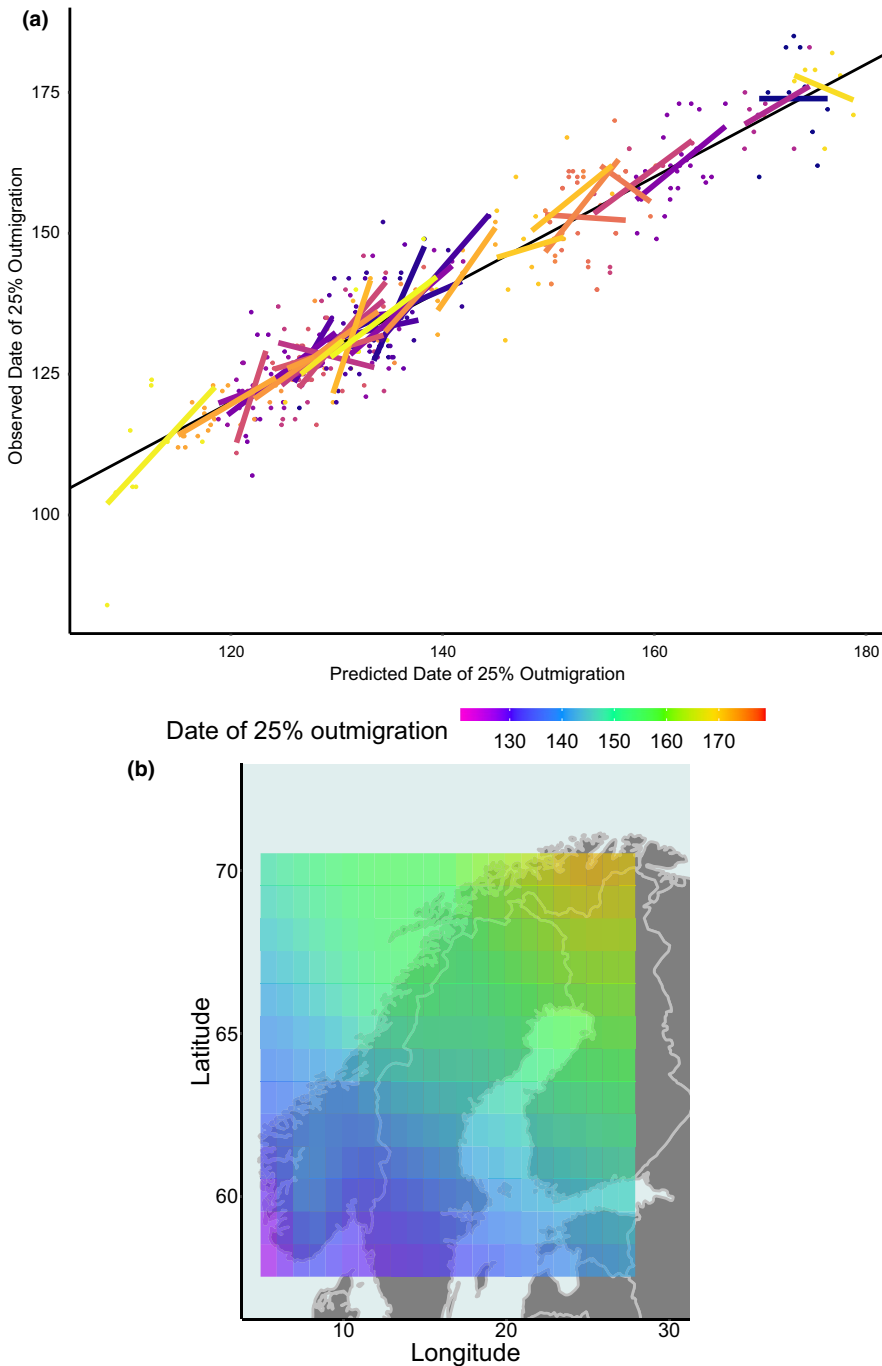


FIGURE 2 (a) Observed and predicted dates of 25% smolt emigration for each river, coded by colour. Coloured lines represent the fit for individual rivers between observed and predicted values. The solid black line is the line of identity. (b) Grid of predictions for date of 25% outmigration superimposed over Norway. Date is julian date

the national estimates were generally earlier than the GAM predictions in southerly latitudes, but was later in the north (Figure 4). Ten-fold cross-validation of the model predictions against observations revealed an R^2 of 83%.

4 | DISCUSSION

We constructed a model that effectively described the variance associated with the timing of 25% smolt migration out of Norwegian rivers. The model was robust, showing good predictive accuracy, based on the spatial and environmental model inputs. We derived

air temperature from predictions of a separate generalized additive model that interpolated temperatures based on weather stations throughout Norway, which was a strong predictor of smolt migration timing. These results are highly relevant to understanding Atlantic salmon ecology and managing this culturally and ecologically important migration that is threatened by human development and climate change (Otero et al., 2014).

Estimated air temperature was a significant predictor of the timing of smolt migration from the rivers that we modelled in Norway. Spatial effects were very strong predictors of the outmigration timing, and given that temperature is correlated with latitude, this may have influenced the interpretation of the effect of temperature in

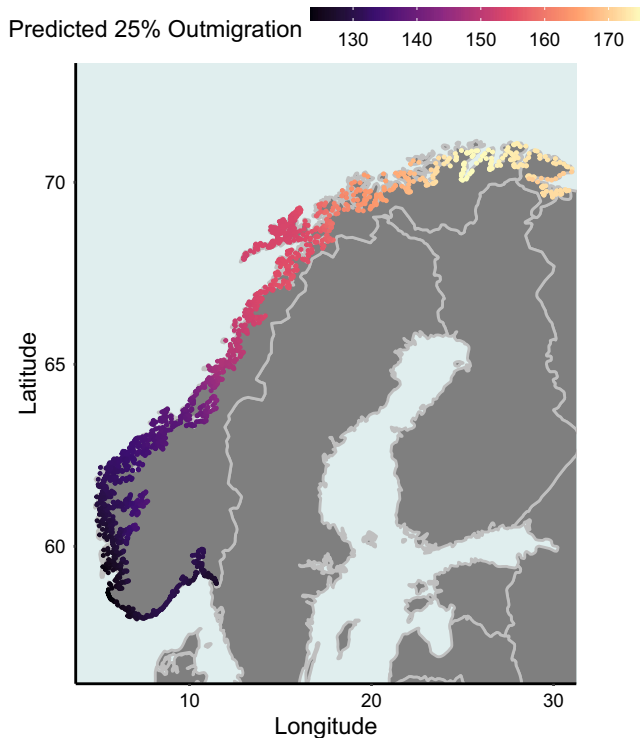


FIGURE 3 Out-of-sample (i.e. predictions made for rivers not included in the model) predictions for the mean timing of outmigration in 401 Norwegian rivers. The generalized additive model from which the predictions were derived included mean air temperature in the first quarter of the year, water flow, longitude and latitude, and monitoring method (set to the factor level “trap”). Predictions are shown for 2018 temperature/flow values. The random intercepts of year and river were set to zero to avoid adjustment for these effects

the model. The smoother on longitude and latitude was included to account for potential spatial effects. Moreover, latitude provides information about photoperiods that would otherwise be unavailable to the model. The spatial dynamics here are likely complex and we considered using more specialized statistical tools for this purpose by generating a mesh and stochastic partial differential equations in the INLA (integrated nested Laplace approximation) framework (Bakka et al., 2018), which did not improve the model, so we continued with the GAM framework. Likely the interacting factors of temperature, photoperiod and other effects related to latitude and experimental studies are needed to more effectively separate these effects.

Temperature controls physiological rates and development of animals, particularly ectothermic species such as most fishes (Brett, 1971; Fry, 1971). In Atlantic salmon, juvenile growth in rivers is controlled by water temperature (Elliott & Hurley, 1997). There is also a strong relationship between the smolt age, that is the time it takes for a juvenile salmon to develop to the stage at which it initiates seaward migration, with photoperiod and water temperature (Metcalf & Thorpe, 1990). Temperature is believed to have a role in the timing of smolt migrations (Zydlewski et al., 2005). Jonsson and Ruud-Hansen (1985) found that water temperature was a significant predictor of smolt run timing in the River Imsa, measuring temperature from 9 April to 16 May but suggesting that the models were relatively insensitive to the period of time-temperature data were collected. We used air temperature rather than water temperature because it was more accessible from historic records. In addition, air temperature can easily be accessed for any river in Norway or estimated by interpolation using the same modelling approach as we implemented for our study making it a versatile tool (Benestad et al., 2019).

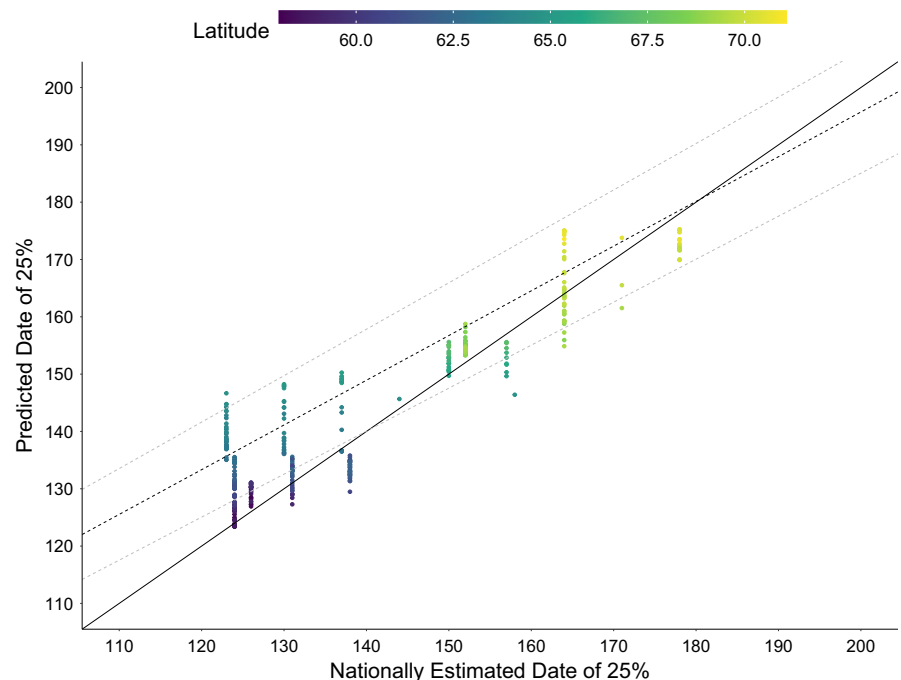


FIGURE 4 Relationship between the estimated migration date for Atlantic salmon smolts and the mean out-of-sample model predictions across years for rivers in Norway. The broken line is the regression line from the linear model (intercept = 39.73, slope = 0.78) and the solid line is the line of identity, suggesting that the national model yields earlier dates of 25% migration relative to the GAM model predictions. The grey dashed lines are the 2.5% and 97.5% interval around the regression line. Date is Julian date

We had access to data from many rivers collected across multiple years that used different counting methods to monitor the smolt run. Standardizing these methods would be ideal to construct a robust model, for example, using the same monitoring methods with the same installation dates across years in different rivers. The data were not structured as such, however, and we capitalized on what was available to construct our model. There was evidence in the model that there were differences among the enumeration methods up to 11 d (e.g. video recordings on average having the latest outmigration dates). Although monitoring efforts should ideally be standardized across rivers to make the most comparable models, barriers to this exist. Smolt traps are effective in rivers where the trap can be placed in areas where they will not be damaged by flooding and can capture a consistent proportion of the run each year. Methods of remote observation such as PIT or video counting may be best in areas where personnel for operating traps or smolt screws is limited, but these require access to power sources. Acoustic telemetry can work in large rivers that cannot be adequately covered by a trap, PIT antenna or video counter and has the benefit of internal power from batteries so that it does not require an outside power source (as PIT systems do). However, PIT and acoustic methods require the fish to be captured, handled and tagged, which may affect the timing of their migration and representativeness of these individuals as sentinels for the total smolt run (Vollset et al., 2018). Tagging also then applies to a relatively small proportion of the total population, which can bias results. Tagged fish may also exit differently than untagged fish, potentially biasing PIT and acoustic methods (Hulbak et al., in review). High flows can flood nets and traps and make them ineffective, which can also affect estimates. Placement of a trap, weir or antenna also affects the timing estimation; all are most effective at narrow points to increase detection probability but should also be as close to sea as possible for a representative estimate of timing.

We expected that water discharge would influence the outmigration timing expressed by the Atlantic salmon smolts; however, this did not have a significant effect on outmigration in our model. Whereas water temperature affects salmon development by acting directly upon the physiology of the fish such that accumulated thermal units across time presumably influences the preparedness of a fish to migrate, water discharge is more likely a threshold cue such that peak flows will provide a stimulus to the fish to migrate downriver (Urke et al., 2013). Snow melts later in cold, northern rivers, and therefore, the peak floods are expected to be later on average than in the south; yet, we observed no correlation between the average temperature and the peak discharge variables that we included in our model. This might be surprising given that many studies have shown a very clear response to either discharge or increase in discharge when studying outmigration of salmon smolts (Hvidsten et al., 1995). However, it is important to note that 25% outmigration may not change much even though the timing of first migration may change a lot. This may be because not all of the population migrates during the first increase in discharge. The proportion not migrating is likely controlled by the developmental rate, which is influenced by temperature during the winter season preceding migration; this

temperature development probably transpires in a more nonlinear fashion than we could model and will require some experimental approaches to derive empirically in the future. Consequently, the median or 25% outmigration time may not be as sensitive to these changes although individual fish may respond very clearly to discharge triggers.

Predictive models are important for both improving the fundamental understanding of ecological processes and making effective evidence-based management decisions. Perhaps the best example of application of a predictive model is within the management of effects of salmon lice (*Lepeophtheirus salmonis* Krøyer 1837) on the survival of out-migrating salmon smolts. Models that attempt to estimate the salmon lice-induced mortality on out-migrating post-smolts have all identified the timing of outmigration of the modelled smolts as one of the most sensitive parameters (Johnsen et al., 2021; Kristoffersen et al., 2018); this is because sea lice infestation pressure is strongly temperature-dependent and increases dramatically throughout the season in areas with high fish farming activity. This is partially due to the density dependence in the sea lice population on fish farms (Jansen et al., 2012) and because the development rates in critical life stages of salmon lice increase nonlinearly with temperature (Stien et al., 2005). Consequently, modelled parasite-induced mortality can vary from negligible to above 30% if smolts migrate two weeks later. In Norway, these models have become even more critical for management and the salmon farming industry as a new management system that relates allowable biomass in fish farms to estimated parasite-induced mortality has been implemented (Vollset et al., 2018). Timing of outmigration, which is such an essential part of this equation, has heretofore been based on a subjective evaluation on outmigration time based on data from nearby sentinel rivers. Our model is a step forward in making an objective evaluation of the outmigration time that can be used in models that are used in management. It is important to note, however, that the differences between the new predictive model and the national evaluation were not very large; the difference was greatest in the southern rivers, but our model can be parameterized with temperatures and flow in a given year to yield more accurate estimates.

5 | CONCLUSIONS

Managing Atlantic salmon populations is a challenge for many nations, regions and municipalities. In Norway, there are hundreds of salmon-producing rivers that cannot all be monitored effectively. Yet, monitoring is increasingly important given that a progressively destabilizing climate means that changes to temperature regimes and animal populations can be expected to occur rapidly. Models using local geographic and environmental information can be used to better understand macroecological processes such as migration and manage threats to salmon populations more proactively. Salmon provides a great case study for other migratory species for which migration timing is critical to management; generalized additive models accounting for the spatial variation seem to be strong tools for generating

predictions that can inform management. Our results show the potential for using accessible parameters to estimate the timing of smolt migrations in rivers, a tool that has the capacity to make contributions to the management of industry particularly through estimation of sea lice burdens on different populations. Our model is refinable with additional data and standardization of collection methods, and more data would allow for improved model calibration using training and test sets. Indeed, the model could be updated with additional data from other countries to expand our understanding of smolt run timing. Nonetheless, this model with available data represents an improvement to the methodologies used by fisheries management agencies to estimate run timing of Atlantic salmon smolts.

ACKNOWLEDGEMENTS

The authors thank Gavin Simpson for the helpful stack exchange post detailing how to set random effects to zero for predictions with GAM models (<https://stats.stackexchange.com/questions/131106/predicting-with-random-effects-in-mgcv-gam>). This work was carried out with financing and support from the Ministry of Trade, Industry and Fisheries.

CONFLICT OF INTEREST

The authors wish for readers to be aware that none of their interests are in conflict with the content of the manuscript.

PEER REVIEW

The peer review history for this article is available at <https://publons.com/publon/10.1111/ddi.13285>.

DATA AVAILABILITY STATEMENT

Data are available in the Dryad Repository: <https://doi.org/10.5061/dryad.p2ngf1vq9>. The file is a post-processed (i.e. cleaned and joined with relevant metadata) spreadsheet. The spreadsheet includes river-specific smolt migration estimates joined with river metadata, temperature and flow as used in the models.

ORCID

Knut Wiik Vollset  <https://orcid.org/0000-0003-0210-4316>

Robert J. Lennox  <https://orcid.org/0000-0003-1010-0577>

REFERENCES

- Albretsen, J., Sperrevik, A. K., Staalstrøm, A., Sandvik, A. D., Vikebø, F., & Asplin, L. (2011). NorKyst-800 report no. 1: User manual and technical descriptions. IMR Res Rep Ser Fiske og Havet2/2011. Institute of Marine Research, Bergen.
- Bakka, H., Rue, H., Fuglstad, G. A., Riebler, A., Bolin, D., Illian, J., Simpson, D., & Lindgren, F. (2018). Spatial modeling with R-INLA: A review. *Wiley Interdisciplinary Reviews: Computational Statistics*, 10(6), e1443.
- Beldring, S., Engeland, K., Roland, L. A., Sælthun, N. R., & Voksø, A. (2013). Estimation of parameters in a distributed precipitation-runoff model for Norway. *Hydrology and Earth System Sciences Discussions*, 7(3), 304–316. <https://doi.org/10.5194/hess-7-304-2003>
- Benestad, R. E., Mezghani, A., Parding, K. M., Erlandsen, H. B., & Tunheim, K. (2019). *esd: Climate analysis and empirical-statistical downscaling (ESD) package for monthly and daily data. R package version 1.9.12*. Retrieved from <http://github.com/metno/esd>
- Brett, J. R. (1971). Energetic responses of salmon to temperature. A study of some thermal relations in the physiology and freshwater ecology of sockeye salmon (*Oncorhynchus nerka*). *American Zoologist*, 11(1), 99–113.
- Duston, J., & Saunders, R. L. (1990). The entrainment role of photoperiod on hypoosmoregulatory and growth-related aspects of smolting in Atlantic salmon (*Salmo salar*). *Canadian Journal of Zoology*, 68(4), 707–715.
- Elliott, J. M., & Hurley, M. A. (1997). A functional model for maximum growth of Atlantic salmon parr, *Salmo salar*, from two populations in northwest England. *Functional Ecology*, 11(5), 592–603. <https://doi.org/10.1046/j.1365-2435.1997.00130.x>
- Fry, F. E. J. (1971). The effect of environmental factors on the physiology of fish. In W. S. Hoar, & D. J. Randall (Eds.), *Fish physiology* (Vol. 6, pp. 1–98). Academic Press.
- Hansen, L. P., & Jonsson, B. (1989). Salmon ranching experiments in the River Imsa: Effect of timing of Atlantic salmon (*Salmo salar*) smolt migration on survival to adults. *Aquaculture*, 82(1–4), 367–373. [https://doi.org/10.1016/0044-8486\(89\)90422-5](https://doi.org/10.1016/0044-8486(89)90422-5)
- Huang, S., Eisner, S., Magnusson, J. O., Lussana, C., Yang, X., & Beldring, S. (2019). Improvements of the spatially distributed hydrological modelling using the HBV model at 1 km resolution for Norway. *Journal of Hydrology*, 577, 123585.
- Hvidsten, N. A., Heggberget, T. G., & Jensen, A. J. (1998). Sea water temperatures at Atlantic salmon smolt entrance. *Nordic Journal of Freshwater Research*, 74, 79–86.
- Hvidsten, N. A., Jensen, A. J., Vivås, H., Bakke, Ø., & Heggberget, T. G. (1995). Downstream migration of Atlantic salmon smolts in relation to water flow, water temperature, moon phase and social interaction. *Nordic Journal of Freshwater Research*, 70, 38–48.
- Jansen, P. A., Kristoffersen, A. B., Viljugrein, H., Jimenez, D., Aldrin, M., & Stien, A. (2012). Sea lice as a density-dependent constraint to salmonid farming. *Proceedings of the Royal Society B: Biological Sciences*, 279(1737), 2330–2338.
- Johnsen, I. A., Harvey, A., Sævik, P. N., Sandvik, A. D., Ugedal, O., Ådlandsvik, B., ... Karlsen, Ø. (2021). Salmon lice-induced mortality of Atlantic salmon during post-smolt migration in Norway. *ICES Journal of Marine Science*, 78(1), 142–154. <https://doi.org/10.1093/icesjms/fsaa202>
- Jonsson, B., & Ruud-Hansen, J. (1985). Water temperature as the primary influence on timing of seaward migrations of Atlantic salmon (*Salmo salar*) smolts. *Canadian Journal of Fisheries and Aquatic Sciences*, 42(3), 593–595.
- Klemetsen, A., Amundsen, P. A., Dempson, J. B., Jonsson, B., Jonsson, N., O'Connell, M. F., & Mortensen, E. (2003). Atlantic salmon *Salmo salar* L., brown trout *Salmo trutta* L. and Arctic charr *Salvelinus alpinus* (L.): A review of aspects of their life histories. *Ecology of Freshwater Fish*, 12(1), 1–59.
- Kristoffersen, A. B., Qviller, L., Helgesen, K. O., Vollset, K. W., Viljugrein, H., & Jansen, P. A. (2018). Quantitative risk assessment of salmon louse-induced mortality of seaward-migrating post-smolt Atlantic salmon. *Epidemics*, 23, 19–33. <https://doi.org/10.1016/j.epidem.2017.11.001>
- Lennox, R. J., Chapman, J. M., Souliere, C. M., Tudorache, C., Wikelski, M., Metcalfe, J. D., & Cooke, S. J. (2016). Conservation physiology of animal migration. *Conservation Physiology*, 4(1). <https://doi.org/10.1093/conphys/cov072>
- Lothian, A. J., Newton, M., Barry, J., Walters, M., Miller, R. C., & Adams, C. E. (2018). Migration pathways, speed and mortality of Atlantic salmon (*Salmo salar*) smolts in a Scottish river and the near-shore coastal marine environment. *Ecology of Freshwater Fish*, 27(2), 549–558.

- McCormick, S. D., Hansen, L. P., Quinn, T. P., & Saunders, R. L. (1998). Movement, migration, and smolting of Atlantic salmon (*Salmo salar*). *Canadian Journal of Fisheries and Aquatic Sciences*, 55(Suppl. 1), 77–92.
- Metcalfe, N. B., & Thorpe, J. E. (1990). Determinants of geographical variation in the age of seaward-migrating salmon, *Salmo salar*. *The Journal of Animal Ecology*, 135–145. <https://doi.org/10.2307/5163>
- Otero, J., L'Abée-Lund, J. H., Castro-Santos, T., Leonardsson, K., Storvik, G. O., Jonsson, B., ... Dionne, M. (2014). Basin-scale phenology and effects of climate variability on global timing of initial seaward migration of Atlantic salmon (*Salmo salar*). *Global Change Biology*, 20(1), 61–75.
- Pedersen, E. J., Miller, D. L., Simpson, G. L., & Ross, N. (2019). Hierarchical generalized additive models in ecology: An introduction with mgcv. *PeerJ*, 7, e6876. <https://doi.org/10.7717/peerj.6876>
- Pinheiro, J., Bates, D., DebRoy, S., Sarkar, D., & R Core Team. (2019). *nlme: Linear and Nonlinear Mixed Effects Models*. R package version 3.1-141. Retrieved from <https://CRAN.R-project.org/package=nlme>
- Scheuerell, M. D., Zabel, R. W., & Sandford, B. P. (2009). Relating juvenile migration timing and survival to adulthood in two species of threatened Pacific salmon (*Oncorhynchus* spp.). *Journal of Applied Ecology*, 46(5), 983–990.
- Stich, D. S., Zydlewski, G. B., Kocik, J. F., & Zydlewski, J. D. (2015). Linking behavior, physiology, and survival of Atlantic salmon smolts during estuary migration. *Marine and Coastal Fisheries*, 7(1), 68–86. <https://doi.org/10.1080/19425120.2015.1007185>
- Stien, A., Bjørn, P. A., Heuch, P. A., & Elston, D. A. (2005). Population dynamics of salmon lice *Lepeophtheirus salmonis* on Atlantic salmon and sea trout. *Marine Ecology Progress Series*, 290, 263–275. <https://doi.org/10.3354/meps290263>
- Thorstad, E. B., Whoriskey, F., Uglem, I., Moore, A., Rikardsen, A. H., & Finstad, B. (2012). A critical life stage of the Atlantic salmon *Salmo salar*: Behaviour and survival during the smolt and initial post-smolt migration. *Journal of Fish Biology*, 81(2), 500–542. <https://doi.org/10.1111/j.1095-8649.2012.03370.x>
- Ugedal, O., Kroglund, F., Barlaup, B., & Lamberg, A. (2014). *Smolt-en kunnskapsoppsummering*. Miljødirektoratet M136–2014.
- Urke, H. A., Kristensen, T., Ulvund, J. B., & Alfredsen, J. A. (2013). Riverine and fjord migration of wild and hatchery-reared Atlantic salmon smolts. *Fisheries Management and Ecology*, 20(6), 544–552. <https://doi.org/10.1111/fme.12042>
- Vollset, K. W., Dohoo, I., Karlsen, Ø., Halttunen, E., Kvamme, B. O., Finstad, B., Wennevik, V., Diserud, O. H., Bateman, A., Friedland, K. D., Mahlum, S., Jørgensen, C., Qviller, L., Krkošek, M., Åtland, Å., & Barlaup, B. T. (2018). Disentangling the role of sea lice on the marine survival of Atlantic salmon. *ICES Journal of Marine Science*, 75(1), 50–60. <https://doi.org/10.1093/icesjms/fsx104>
- Weidmann, N. B., & Gleditsch, K. S. (2016). *cshapes: The CShapes dataset and utilities*. R Package version 0.6. Retrieved from <https://CRAN.R-project.org/package=cshapes>
- Whalen, K. G., Parrish, D. L., & McCormick, S. D. (1999). Migration timing of Atlantic salmon smolts relative to environmental and physiological factors. *Transactions of the American Fisheries Society*, 128(2), 289–301.
- Wickham, H. (2016). *ggplot2: Elegant graphics for data analysis*. Springer-Verlag.
- Wood, S. N. (2017). *Generalized additive models: An introduction with R* (2nd ed.). Chapman and Hall/CRC.
- Zydlewski, G. B., Haro, A., & McCormick, S. D. (2005). Evidence for cumulative temperature as an initiating and terminating factor in downstream migratory behavior of Atlantic salmon (*Salmo salar*) smolts. *Canadian Journal of Fisheries and Aquatic Sciences*, 62(1), 68–78.

BIOSKETCH

Knut Wiik Vollset is a senior researcher at NORCE LFI in Bergen, Norway. Knut's research explores the pressures placed on wild fish populations from human activities and aims to provide scientific tools that inform management and restoration efforts. Atlantic salmon are highly valued in Norway, and recent declines necessitate tools such as these to help managers better address population-specific conservation challenges.

How to cite this article: Vollset KW, Lennox RJ, Lamberg A, et al. Predicting the nationwide outmigration timing of Atlantic salmon (*Salmo salar*) smolts along 12 degrees of latitude in Norway. *Divers Distrib*. 2021;27:1383–1392. <https://doi.org/10.1111/ddi.13285>