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# Evaluating AUSRIVAS predictive model performance for detecting simulated eutrophication effects on invertebrate assemblages

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Abstract: Confidence in any bioassessment method is related to its ability to detect ecological improvement or impairment. We evaluated Australian River Assessment (AUSRIVAS)-style predictive models built using referencesite data sets from the Australian Capital Territory (ACT), the Yukon Territory (YT; Canada), and the Laurentian Great Lakes (GL; North America) area. We evaluated model performance as ability to correctly assign reference condition with independent reference-site data. Evaluating model ability to detect human disturbance is generally more problematic because the actual condition of test sites is usually unknown. Independent reference-site data underwent simulated impairment by varying the proportions of sensitive, intermediate, and tolerant taxa to simulate degrees of eutrophication. Model performance was related to differences in data sets, such as number and distribution of invertebrate taxa. Sensitive taxa tended to have lower expected probabilities of occurrence than more-tolerant taxa, but the distribution of taxa grouped by tolerance categories also differed by data set. Thus, the models differed in ability to detect the simulated impairment. The ACT model performed best with respect to Type 1 error rates (0%) and the GL model the worst (38%). The YT model performed best (10% error) for detecting moderate impairment, and the ACT model detected all severely impaired sites. AUSRIVAS did not assign most mildly impaired sites to below-reference condition, but a reduction in observed/expected values for some of the mildly impaired sites was observed. Models did not detect mild impairment that simply changed taxon abundances because presence-absence data were used for models. However, in comparison with other models described in this special issue (that did use abundance data), the AUSRIVAS model performance was comparable or better for detecting the simulated moderate and severe impairments.

**Key words:** reference condition approach, AUSRIVAS, simulated impairment, bioassessment, predictive modeling, macroinvertebrates

The Australian River Assessment System (AUSRIVAS) has been Australia's national standard method for biological assessment of river health for over a decade (Davies 2000, Simpson and Norris 2000, eWater CRC 2012). AUSRIVAS consists of a standardized invertebrate sampling method, predictive models, and software for assessing river health (Simpson and Norris 2000) that uses the reference-condition approach (Reynoldson et al. 1997). Adoption of AUSRIVAS bioassessment by water and environment agencies was rapid with implementation into state policy and regulatory frameworks and a variety of environmental management settings, by government, community, and industry (Davies 2000). AUSRIVAS has been used for targeted impact assessment (e.g., Marchant and Hehir 2002, Sloane and Norris 2003, Nichols et al. 2006, Growns et al. 2009, White

et al. 2012), state/regional assessments of river condition (e.g., Turak et al. 1999, ACT Government 2006, Rose et al. 2008, Norris and Nichols 2011), community-based river assessment programs (e.g. WaterWatch; Davies 2007), and very broad-scale assessment at multijurisdictional and national levels (Turak et al. 1999, Norris et al. 2001a, b, EPA 2004, Davies et al. 2010, Harrison et al. 2011). A major strength of national systems like AUSRIVAS, River Invertebrate Prediction and Assessment System (RIVPACS) and Canadian Aquatic Biomonitoring Network (CABIN) is the broad-scale bioassessment and biomonitoring opportunities such programs allow (Rosenberg et al. 2000, Wright et al. 2000, Norris et al. 2001a, 2007). For example, AUSRIVAS data were the only data with national coverage used to report in-stream biological condition for Australia's 2011 State

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of the Environment report (Harrison et al. 2011). Thus, AUSRIVAS has national significance for monitoring and assessing river condition in Australia.

In a review of alternatives to the River Invertebrate Prediction and Classification system (RIVPACS)-style predictive models, Johnson (2000) concluded that it was a robust approach for predicting assemblage structure and found no compelling reason justifying a change to other techniques. The AUSRIVAS method has produced models that work well in many of Australia's varied environments, and they have proved useful for river assessment (for further examples see Marchant and Hehir 2002, Hose et al. 2004, Metzeling et al. 2006, Nichols et al. 2010). However, since Johnson's review, other modeling methods have been used more extensively (Linke et al. 2005, Van Sickle et al. 2006, Chessman 2009, Webb and King 2009, Aroviita et al. 2010, Feio and Poquet 2011), and investigators have identified some limitations of the AUSRIVAS approach. For example, model performance is poor where reference sites are problematic or lacking (Chessman et al. 2010).

Implementation of national-scale water reforms and statutory water planning (Tomlinson and Davis 2010, Connell 2011, EU 2012) will necessitate evaluation of interventions designed to improve river conditions. Renewed pertinence of adequate assessment tools and continued advances in river assessment methods have prompted interest in development of new and improved tools for assessing ecological effects of human activities. Given the large initial investment in the AUSRIVAS approach, the utility of the method, and almost 20 y of experience since its inception, an appraisal seems timely and was one motivation prompting this special series of papers.

User confidence in any bioassessment modeling method is related to the method's ability to detect ecological improvement or impairment. Updating of predictive models that are in widespread use or introduction of alternative modeling options should involve careful evaluation and comparison of their performance. Evaluations of model performance generally are based on how well models predict group membership of reference sites and how well models predict the taxa found at new reference sites (Coysh et al. 2000, Hawkins et al. 2000). Such validation usually involves a data set of reference sites that are independent of those used to create the predictive model. However, evaluating the ability of models to detect human disturbance is more problematic than validating with reference sites because the biological condition of test sites is usually unknown. One approach is to use simulated impairments to determine the sensitivity of a method for detecting impairment (Cao and Hawkins 2005, Bailey et al. 2012). Evaluating both Type 1 and Type 2 error rates provides a better indication of model performance.

We used independent reference sites and simulated impairment (Bailey et al. 2014) to evaluate AUSRIVAS-style models built from reference-site data collected in Australia, the Yukon Territory (Canada), and the Laurentian Great Lakes (GL) area of North America to compare model performance for 3 very different environments. Independent reference-site data were artificially impaired to simulate 3 degrees of eutrophication. Evaluating model performance in this way allowed us to test the ability of models to detect known impairment. The results provided by the standard AUSRIVAS method used in our study form the basis for comparison with other modeling methods presented in this special series.

## **METHODS** Data sets

Authors of all papers in this special series analyzed the same data sets (described in full by Bailey et al. 2014). The reference-site data (invertebrate and environmental data) were collected from wadeable streams in the Australian Capital Territory (ACT) region (i.e., the upper Murrumbidgee River catchment), the Yukon Territory (YT), and from near-shore sites in the Laurentian Great Lakes (GL; North America). Each region had 2 reference-site data sets, 1 for model training and another independent data set consisting of 20 sites for model validation (D0). The invertebrate data from the validation sites were artificially impaired to simulate the effects of 3 degrees (D1 = mild, D2 = moderate, and D3 = severe) of eutrophication by varying the proportions of sensitive, intermediate, and tolerant taxa (Bailey et al. 2014). Impairment was simulated at each site for each level by altering the abundance of taxa or by removing some taxa. The simulated impairment was applied to randomly selected taxa within tolerance categories (e.g., sensitive, intermediate, tolerant) that were based on regionspecific tolerance scores (Barbour et al. 1999) for the YT data, Hilsenhoff tolerance values (Hilsenhoff 1988) for the GL data, and Stream Invertebrate Grade Number (SIGNAL) values (Chessman 2003) for the ACT data.

# **AUSRIVAS** modeling methods

We developed a standard AUSRIVAS model (Smith et al. 1999, Simpson and Norris 2000) using the referencesite training data for each data set. AUSRIVAS developers adapted the modeling approach originally described by the authors of the RIVPACS models (Wright et al. 1984, Moss et al. 1987, Wright 1995). In accordance with AUSRIVAS methods, we excluded sites with <6 taxa and taxa that occurred at <10% of sites in the training data. For each model, we grouped reference sites based on the similarity of their invertebrate assemblages using Unweighted Pair Group Method with Arithmetic Mean (UPGMA) cluster analysis on presence-absence data (Belbin 1993). We then selected a subset of the environmental variables (predictor variables) by using stepwise discriminant function analysis to determine which environmental variables best discriminated among reference-site groups (Smith et al. 1999, Simpson and Norris 2000). We used the discrimination cross-validation procedure as an internal model check regarding error rates for assigning training sites to correct groups (Smith et al. 1999). AUSRIVAS models predict the taxa expected at a site by summing the individual probabilities of occurrence for all the taxa predicted to have ≥50% probability of occurrence (Simpson and Norris 2000), resulting in a site-specific expected taxa list.

#### **AUSRIVAS** bands of biological condition

How much the observed invertebrate assemblage (O) deviates from that expected (E) is a measure of the severity of environmental impairment. AUSRIVAS assigns O/E scores to quality bands that represent different levels of biological condition (Coysh et al. 2000). Sites with O/E scores in band A are similar to reference condition, whereas sites with O/E values in band B or lower are considered impaired (Table 1). The distributions of the training reference-site O/E scores were used to determine the width of the quality bands. Thus, band widths are specific to each model (Table 1).

#### Model validation and performance evaluation

**Outside model experience** We used standard AUSRIVAS methods to assess whether validation sites were within the environmental scope of the reference data set for each model. We calculated the Mahalanobis distance of each site to each canonical variate (as per Clarke et al. 1996). We then used a  $\chi^2$  test to determine whether each site was within the 99% confidence interval of the centroid of  $\geq 1$ 

reference-site group. If a site's environmental characteristics differed significantly from the training data set (which could indicate underrepresentation of that site type in the training data set) then that site would have no appropriate reference group for comparison. At that stage, the site may be identified as 'outside the experience of the model', and the model predictions and site assessments treated as suspect (Coysh et al. 2000).

**Model validation** We assessed the ability of each model to correctly assign a new reference-site O/E to band A (Table 1) with the validation data set. Assuming the validation sites were truly in reference condition, <10% of the sites should mistakenly fall below AUSRIVAS band A (Coysh et al. 2000). A failure rate >10% would indicate that the model had a greater than expected Type 1 error rate (sites failed that should have passed). Thus, we based the Type 1 error rate on % validation sites with O/E values <10<sup>th</sup> percentile of the training-data O/E distribution.

**Model performance for detecting impairment** We used the simulated impairment validation data sets to assess Type 2 error rates (sites passed that should have failed). We tested the ability of models to detect the 3 degrees of simulated impairment as the percentage of sites assessed as below band A. Thus, we based the Type 2 error rate on % simulated impairment sites with O/E values >10<sup>th</sup> percentile of the training-data O/E distribution. We used the other AUSRIVAS quality bands to assess the ability of the models to detect a disturbance gradient.

Table 1. Australian River Assessment System (AUSRIVAS) bands of biological condition for the Australian Capital Territory (ACT; upper Murrumbidgee River catchment in ACT region of Australia), the Yukon Territory (YT), and the Laurentian Great Lakes (GL; North America) models, showing observed/expected (O/E) taxa range, band descriptions, and interpretations (Coysh et al. 2000).

| Band | O/E value                                   | Band description  | O/E interpretations  |
|------|---|---|--|
| X    | ACT > 1.15<br>YT > 1.28<br>GL > 1.18        | More biologically diverse than reference; O/E > 90 <sup>th</sup> percentile of reference sites used to create the model | More taxa found than expected; site is potentially rich in biodiversity or may have mild organic enrichment that initially could increase the number of taxa because of increased food resources resulting from the increase in nutrients; test site requires further consideration before drawing conclusions |
| A    | ACT 0.85-1.15<br>YT 72-1.28<br>GL 0.82-1.18 | Similar to reference; O/E within range of central 80% of reference sites used to create the model                       | Most/all of the expected taxa found; water quality or habitat condition similar to reference sites   |
| В    | ACT 53-84<br>YT 0.15-71<br>GL 0.43-0.81     | Significantly impaired; O/E < 10th percentile of reference sites used to create the model; band B width = band A width  | Fewer families than expected; water quality or habitat quality possibly impaired, resulting in loss of expected taxa   |
| С    | ACT 0.21-52<br>YT 0-0.14<br>GL -0.08-0.44   | Severely impaired; O/E value < band B; band C width = band A width  | Many fewer families than expected; poor water quality or<br>habitat quality resulting in loss of expected invertebrate<br>richness   |
| D    | ACT 0-0.20<br>YT No band D<br>GL 0-0.07     | Extremely impaired; O/E value < band C to 0   | Few expected families and only hardy and pollution-<br>tolerant taxa remain; extremely poor water quality or<br>habitat quality resulting in severe impairment   |

Low-probability taxa E for a site is the sum of the sitespecific expected probabilities of the individual taxa with ≥0.50 probability. We tested whether sensitive taxa (as defined by Bailey et al. 2014), on average, had lower expected probabilities of occurrence than more-tolerant taxa. If so, excluding sensitive taxa with low probability of occurrence might obscure the simulated impairment that removed selected sensitive taxa. For each model, we evaluated the effect of the tolerance category of low-probability taxa on the model and the model's ability to detect the 3 levels of simulated impairment. Taxa with 0 (to 3 decimal places) expected probabilities were excluded from the analyses because some taxa are naturally restricted to particular stream types and, therefore, are naturally excluded from a proportion of the sites (as per Clarke and Murphy 2006). Including many 0 values would have distorted the frequency distributions. We calculated the expected probability of occurrence of each taxon across all the validation sites and compared the distributions of the probability values within each tolerance category (box plots).

# **RESULTS**

### Model summary

We created 1 model from each of the 3 data sets (Tables 2, 3). The number of taxa removed from the data set because they were considered rare and, therefore, excluded from the models was high (ACT: 46%, YT: 52%, GL: 59%). The ACT model used all available training sites, but sites were removed from the YT (n = 17) and GL (n = 17)40) data sets because they had too few taxa for modeling purposes (≤5). The cross-validation error for the YT model (44%) was greater than usually desired for an AUSRIVAS model (Table 2). The YT model also produced the widest range of O/E values for the training sites and had the widest quality bands of all models (Table 1). A wide band A equates to a wide range of accepted reference condition.

The data sets differed in the total number of taxa and total taxa used for modeling (Table 2). Taxon richness was ≤6 at 50% of the GL and 17% of the YT sites, where-

as no ACT sites had <10 taxa and 50% had >18 taxa. The YT and GL data sets had 8 and 14 fewer taxa, respectively, than the ACT data set. The models developed with these data sets also varied in the number of taxa expected to occur at sites (using the standard 0.5 probability cutoff) (Table 2). The ACT model predicted 8 to 10 more taxa than the low estimates predicted by the YT and GL models (Table 2), a greater percentage than the difference in total taxa. This result indicates that factors other than the difference in total number of taxa present in the data sets are required to explain differences among models regarding predicted taxa.

# Model validation and performance for detecting simulated impairment

Some validation sites appeared dissimilar to training sites used for model development based on ordination of the biological data (Fig. 1A-C), particularly in the YT and GL models where some validation sites and the removed low-richness training sites shared a similar ordination space (Fig. 1B, C). However, for all models, no validation sites were outside the model experience regarding their environmental character.

Type 1 errors The ACT model correctly assigned all validation sites to band A and had the lowest Type 1 error rate (Table 4). The Type 1 error rates for the YT and GL models were >10%. The GL model had the highest Type 1 error rate (Table 4).

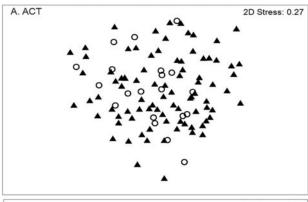
Type 2 errors The ACT model detected all severely impaired (D3) sites, which were allocated to AUSRIVAS band C (severely impaired) or near the boundary of bands C and B (Table 4, Fig. 2A). Most (80%) of ACT sites with moderate (D2) levels of impairment were allocated to AUSRIVAS band B (significantly impaired) (Table 4, Fig. 2A). Except for 2 sites, the mildly impaired (D1) ACT sites did not fall below band A (Fig. 2A). However, some D1 sites had lower O/E values within band A (Fig. 2A) than did the original unimpaired validation sites (D0). The ACT

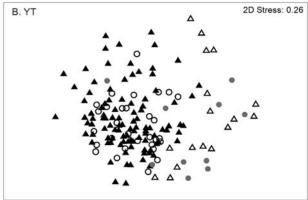
Table 2. Summary details for Australian River Assessment (AUSRIVAS) predictive models developed for the Australian Capital Territory (ACT; upper Murrumbidgee River catchment in ACT region of Australia), the Yukon Territory (YT), and the Laurentian Great Lakes (GL; North America). O/E = observed/expected taxa.

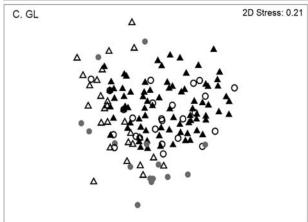
|       | Initial Model<br>data set data set |      |       |      | Model details                |                            |               |                                   |                      |                     |  |
|-------|------------------------------------|------|-------|------|------------------------------|----------------------------|---------------|-----------------------------------|----------------------|---------------------|--|
| Model | Sites                              | Taxa | Sites | Taxa | Groups (and sites per group) | Cross-<br>validation error | Band<br>width | Upper value of impaired O/E range | Predicted taxa range | Expected taxa range |  |
| ACT   | 87                                 | 67   | 87    | 36   | 3 (45, 25, 17)               | 0.28                       | 0.31          | 0.84                              | 13-19                | 10.3-13.9           |  |
| YT    | 118                                | 59   | 101   | 28   | 4 (27, 25, 31, 18)           | 0.44                       | 0.56          | 0.71                              | 5-12                 | 3.6-9.0             |  |
| GL    | 124                                | 54   | 84    | 22   | 3 (21, 24, 39)               | 0.31                       | 0.36          | 0.81                              | 5–9                  | 4.3 - 7.2           |  |

Table 3. Mean (SD) predictor variables and number of taxa per group for Australian River Assessment (AUSRIVAS) predictive models developed for the Australian Capital Territory (ACT; upper Murrumbidgee River catchment in ACT region of Australia), the Yukon Territory (YT), and the Laurentian Great Lakes (GL; North America). EC = ecological condition, UCA = upstream catchment area, EC = electrical conductivity, nonprod = nonproductive, snow = total annual snowfall, rif = riffle, boul = boulders.

| ACT   Activated (m)   UCA (km²)   EC (µS/cm)   Riffle depth (m)   % rif boal   Velocity: depth   Chock (µs)   18 (117)   147 (3.0)   boulder-sized sub-boulder sized size   | Groups  | Statistic | Taxa      |                  |               | Predicto        | Predictor variables |               |                  | Summary of group characteristics  |
|---|---------|-----------|-----------|------------------|---------------|-----------------|---------------------|---------------|------------------|---|
| Mean   18 (2.7)   740 (1893)   499 (1178.5)   96 (82.8)   18 (8.4)   18 (11.7)   147 (3.0)   8     Range   11–24   380–1095   12–5709   24–459   4-40   0–50   10–20     Range   11–24   380–1095   12–5709   22–28   4-40   0–50   10–20     Range   15–25   620–1360   33–572   17–245   3–36   0–40   10–10   1     Range   15–25   620–1360   35–6589   417–509   6–39   0–30   7–19   1     Range   10–20   380–910   55–6589   417–509   6–39   0–30   7–19   1     Range   6–20   0–100   0–100   0–100   0–93.4   0–26.9   37 (32.6)   1     Mean   (1.58)   15 (6.24)   66 (44.2)   18 (25.9)   37 (32.6)   14 (3.3)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)   14 (3.4)  | ACT     |           |           | Altitude (m)     | $UCA (km^2)$  | EC (µS/cm)      | Riffle depth (cm)   | % rif boul    | Velocity: depth  |   |
| Mean   19 (2.5)   982 (207.4)   80 (142.5)   52.58 (45.87)   18.28 (8.78)   11.4 (10.56)   13.4 (2.6)   Hone     Mean   19 (2.5)   982 (207.4)   80 (142.5)   52.58 (45.87)   18.28 (8.78)   11.4 (10.56)   13.4 (2.6)   H     Mean   14 (2.7)   665 (153.7)   1539 (2405.8)   22.22 (162.8)   19.94 (9)   11.76 (11.45)   14.13   1.0-18     Mean   14 (2.7)   665 (153.7)   1539 (2405.8)   22.22 (162.8)   19.94 (9)   11.76 (11.45)   14.13   1.0-18     Mean   10 (3.3)   15 (36.2)   55 -6589   41.7-509   6-39   0-30   7-19   1.0-18     Mean   10 (3.2)   50 (48.3)   15 (35.2)   50 (48.3)   15 (35.4)   14 (13.4)   15 (4.3)   7.19   1.1     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   14 (2.3)   8.1   1.25   1.25   98-167   1.3-11   1.25   1.25   98-167   1.3-11   1.25   1.3-11   1.25   1.25 <td>Group 1</td> <td>Mean</td> <td>18 (2.7)</td> <td>740 (189.9)</td> <td>499 (1178.5)</td> <td>96 (82.8)</td> <td>18 (8.4)</td> <td>18 (11.7)</td> <td>14.7 (3.0)</td> <td>Streams with high proportion of houlder-sized substrate mid-range</td>  | Group 1 | Mean      | 18 (2.7)  | 740 (189.9)      | 499 (1178.5)  | 96 (82.8)       | 18 (8.4)            | 18 (11.7)     | 14.7 (3.0)       | Streams with high proportion of houlder-sized substrate mid-range               |
| Mean   19 (2.5)   982 (2074)   80 (142.5)   52.58 (45.87)   18.28 (8.78)   114 (10.56)   134 (2.6)   H     Range   15-25   620-1360   39-572   17-245   3-36   0-40   10-18   H     Mean   14 (2.7)   665 (153.7)   1539 (240.88)   2222 (162.8)   1994 (9)   11.76 (11.45)   14 (3.4)   La     Range   10-20   380-910   5.5-6889   41,7-509   6-39   0-30   7-19   La     Mean   10 (3.3)   15 (36.2)   50 (48.3)   15 (25.3)   20 (24.2)   133 (21.6)   7-19   La     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   1.03-167   7-19   La     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   1.26 (18.5)   4 (2.3)   S     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   1.46 (18.8)   7.3 (3.7)   C     Range   6-12   0-100   <  |         | Kange     | 11-24     | 380-1095         | 17-5/09       | 74-429          | 4-40                | 0-20          | 10-20            | EC values   |
| Range   15–25   620–1360   3.9–572   17–245   3–36   0–40   10–18     Range   10–20   380–910   5.5–6589   41.7–509   6–39   0–40   11.6 (1145)   14 (3.4)   1.8     Range   10–20   380–910   5.5–6589   41.7–509   6–39   0–30   7–19   1.3     Mean   10–20   380–910   5.6–6589   41.7–509   6–39   0–30   7–19   1.2     Mean   10–30   0–100   0–99   0–99   0–80   1.3   1.0   1.2   <   | Group 2 | Mean      | 19 (2.5)  | 982 (207.4)      | 80 (142.5)    | 52.58 (45.87)   | 18.28 (8.78)        | 11.4 (10.56)  | 13.4 (2.6)       | High altitude sites with small  |
| Range   14 (2.7)   665 (153.7)   1539 (2405.8)   2222 (162.8)   19.94 (9)   11.76 (1145)   14 (34)   Lakange     Range   10-20   380-910   5.5-6589   41.7-509   6-39   0-30   7-19     Mean   10 (33)   15 (36.2)   56 (48.3)   15 (25.3)   20 (24.2)   133 (21.6)   7.3 (6)     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   103-167   1-25     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   103-167   1-25     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   103-167   1-25     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   103-167   1.3-11     Mean   10     |         | Range     | 15–25     | 620–1360         | 3.9–572       | 17–245          | 3–36                | 0-40          | 10–18            | catchment area, low EC values,<br>greater taxon richness                        |
| Range   10-20   380-910   55-6589   41.7-509   6-39   0-30   7-19     Mean   10 (33)   metamorphic   % sedimentary   % alpine   % nonprod forest   Snow (mm)   Wetted width (m)     Mean   6-13   15 (36.2)   50 (48.3)   15 (55.3)   20 (24.2)   133 (21.6)   7.3 (6)   M     Range   6-20   0-100   0-997   0-88.5   103-167   1-25   37 (35.6)   123 (15.3)   41.33     Mean   8 (15.8)   16 (37.4)   66 (44.2)   18 (15.9)   37 (35.6)   124 (13.3)   41.33     Mean   1.3   0 (0)   0 -100   0 -93.4   0 -95.6   148 (19.8)   7.3 (3.7)   C2     Mean   1.0   0 (0)   94 (13)   35 (40.3)   14 (21.4)   15 (14.4)   60.4 (6.3)   H     Range   6-14   0 -0   51-100   0 -39   0 -61   10-16   0 -13   12 (14.4)   15 (14.4)   60.4 (6.3)   H     Range   6-14   0 -0 <td< td=""><td>Group 3</td><td>Mean</td><td>14 (2.7)</td><td>665 (153.7)</td><td>1539 (2405.8)</td><td>222.2 (162.8)</td><td>19.94 (9)</td><td>11.76 (11.45)</td><td>14 (3.4)</td><td>Larger rivers, greater EC values,</td></td<>  | Group 3 | Mean      | 14 (2.7)  | 665 (153.7)      | 1539 (2405.8) | 222.2 (162.8)   | 19.94 (9)           | 11.76 (11.45) | 14 (3.4)         | Larger rivers, greater EC values,   |
| Mean   10 (3.3)   % metamorphic metam |         | Range     | 10-20     | 380-910          | 5.5–6589      | 41.7–509        | 6–39                | 0-30          | 7–19             | substrate of smaller particle size  |
| Mean   10 (3.3)   15 (36.2)   50 (48.3)   15 (25.3)   20 (24.2)   133 (21.6)   7.3 (6)   M     Range   6-20   0-100   0-100   0-99.7   0-88.5   103-167   1-25   M     Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   122 (18.5)   4 (2.3)   Sr     Range   6-12   0-100   0-100   0-93.4   0-95.6   98-167   1.3-11   Sr     Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   7.3 (3.7)   CG     Range   8-20   0 -0   0 -100   0 -73.7   0 -67.9   106-167   2-14.6   H     Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     Range   6-14   0 -0   51-100   0 -99   0 -61   106-167   1.3-30   N     Mean   7 (1.4)   46.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)  | YT      |           |           | % metamorphic    | % sedimentary | % alpine        | % nonprod forest    | Snow (mm)     | Wetted width (m) |   |
| Range   6-20   0-100   0-997   0-88.5   103-167   1-25     sup 2   Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   122 (18.5)   4 (2.3)   Sr     sup 2   Range   6-12   0-100   0-100   0-93.4   0-95.6   98-167   1.3-11   Sr     sup 3   Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   7.3 (3.7)   CG     sup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     sup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     sup 5   G-14   0 -0   51-100   0 -99   0 -61   106-167   1.3-30   1.3-30     sup 6   G-14   45.5 (1.8)   % clay   Water depth (m)   % sediment MgO   1.3-30   1.3-30     sup 7   Mean   7 (14)   45.5 (1.8)  | Group 1 | Mean      | 10 (3.3)  | 15 (36.2)        | 50 (48.3)     | 15 (25.3)       | 20 (24.2)           | 133 (21.6)    | 7.3 (6)          | Metamorphic/sedimentary geology,  |
| oup 2   Mean   8 (1.58)   16 (37.4)   66 (44.2)   18 (25.9)   37 (32.6)   122 (18.5)   4 (2.3)   Sr     oup 3   Range   6-12   0-100   0-93.4   0-95.6   98-167   1.3-11   Sr     oup 3   Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   7.3 (3.7)   C3     oup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     sange   8-20   0 -0   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     kange   6-14   0 -0   51-100   0 -99   0 -61   106-167   2-14.6   H     oup 5   Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   27 (1.4)   M     sange   6-10   42-49   6.7-77   8-51   1-8   0.6-18   1.3-30     sange   6-14   43.8-48.9   0-83.8 <td< td=""><td></td><td>Range</td><td>6-20</td><td>0 - 100</td><td>0 - 100</td><td>0-99.7</td><td>0-88.5</td><td>103–167</td><td>1-25</td><td>higher taxon richness</td></td<>  |         | Range     | 6-20      | 0 - 100          | 0 - 100       | 0-99.7          | 0-88.5              | 103–167       | 1-25             | higher taxon richness   |
| up 3   Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   7.3 (3.7)   CG     uup 4   Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   7.3 (3.7)   CG     uup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   105 (14.4)   6.04 (6.3)   Hi     uup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   Hi     uup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   Hi     uup 5   Mean   6-14   0 -0   51-100   0 -99   0 -61   106-167   1.3-30   M     uup 5   Mean   7 (1.4)   45.5 (18)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     Range   6-10   46.8 (1.9)   206 (22.1)   36 (22.9)   3.2 (2.1)   M     Range   6-14   44.3 (1.4)  | Group 2 | Mean      | 8 (1.58)  | 16 (37.4)        | 66 (44.2)     | 18 (25.9)       | 37 (32.6)           | 122 (18.5)    | 4 (2.3)          | Small streams, metamorphic/   |
| oup 3   Mean   13   0 (0)   75 (41.6)   30 (19.9)   37 (25.6)   148 (19.8)   73 (3.7)   C3     sup 4   (3.23)   0-0   0-100   0-73.7   0-67.9   106-167   2-14.6   2-14.6     oup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     Range   6-14   0-0   51-100   0-99   0-61   106-167   1.3-30   H     oup 1   Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     sange   6-10   42-49   6.7-77   8-51   1-8   M     sange   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   M     sup 2   Mean   11(2.3)   44.3 (1.4)   20 (18.7)   7 (6.6)   2.1 (1.1)   0.95-7.3   O     sup 3   Mean   11(2.3)   42.3-48.9   0-76.1   0.55-27.5   0.955-7.3   0.95-7.3   O  |         | Range     | 6-12      | 0-100            | 0-100         | 0-93.4          | 0-95.6              | 98–167        | 1.3–11           | sedimentary geology, less snow, wide sediment size range, lowest taxon richness |
| Range   8-20   0-0   0-100   0-73.7   0-67.9   106-167   2-14.6     oup 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   H     Range   6-14   0 - 0   51-100   0-99   0-61   106-167   1.3-30     up 1   Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     up 2   Mean   8 (2.6)   46.8 (1.9)   20.6 (22.1)   36 (22.9)   3.2 (2.1)   U     sange   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   U     sange   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   U     sange   6-17   42.3-48.9   0-76.1   0.5-27.5   0.95-7.3   O  | Group 3 | Mean      | 13 (3.23) | (0) 0            | 75 (41.6)     | 30 (19.9)       | 37 (25.6)           | 148 (19.8)    | 7.3 (3.7)        | Catchments with high % alpine and % nonproductive forest, greatest              |
| up 4   Mean   10   0 (0)   94 (13)   35 (40.3)   14 (21.4)   125 (14.4)   6.04 (6.3)   Hi     (2.17)   (2.17)   0-0   51-100   0-99   0-61   106-167   1.3-30     up 1   Mean   7 (1.4)   45.5 (1.8)   % clay   Water depth (m)   % sediment MgO   1.3-30     up 1   Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     sange   6-10   42-49   6.7-77   8-51   1-8   M     nup 2   Mean   8 (2.6)   46.8 (1.9)   20.6 (22.1)   36 (22.9)   3.2 (2.1)   Up     Range   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   0.6-8.4     Range   6-17   42.3-48.9   0-76.1   0.5-27.5   0.95-7.3   O   |         | Range     | 8-20      | 0-0              | 0 - 100       | 0-73.7          | 6.79-0              | 106 - 167     | 2 - 14.6         | snowfall, greatest taxon richness   |
| Range   6-14   0-0   51-100   0-99   0-61   106-167   1.3-30     up 1   Mean   7 (1.4)   45.5 (1.8)   % clay   Water depth (m)   % sediment MgO   M     up 2   Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     sup 2   Mean   7 (1.4)   45.5 (1.8)   6.7-77   8-51   1-8   M     sup 2   Mean   8 (2.6)   46.8 (1.9)   20.6 (22.1)   36 (22.9)   3.2 (2.1)   Uj     sange   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   Uj     sup 3   Mean   11(2.3)   44.3 (1.4)   20 (18.7)   7 (6.6)   2.1 (1.1)   0     sange   6-17   42.3-48.9   0-76.1   0.5-27.5   0.95-7.3   0   | Group 4 | Mean      | 10 (2.17) | 0 (0)            | 94 (13)       | 35 (40.3)       | 14 (21.4)           | 125 (14.4)    | 6.04 (6.3)       | High % alpine and low % nonproductive forest, sedimentary                       |
| up 1 Mean 7 (1.4) 45.5 (1.8) % clay Water depth (m) % sediment MgO   up 1 Mean 7 (1.4) 45.5 (1.8) 46.3 (20.9) 20 (10.2) 2.7 (1.4) M   up 2 Range 6-10 42-49 6.7-77 8-51 1-8 M   up 2 Mean 8 (2.6) 46.8 (1.9) 20.6 (22.1) 36 (22.9) 3.2 (2.1) UJ   Range 6-14 43.8-48.9 0-83.8 6.8-87 0.6-8.4   up 3 Mean 11(2.3) 44.3 (1.4) 20 (18.7) 7 (6.6) 2.1 (1.1) O   Range 6-17 42.3-48.9 0-76.1 0.5-27.5 0.95-7.3 O   |         | Range     | 6–14      | 0-0              | 51-100        | 66-0            | 0–61                | 106–167       | 1.3–30           | bedrock geology   |
| Mean   7 (1.4)   45.5 (1.8)   46.3 (20.9)   20 (10.2)   2.7 (1.4)   M     Range   6-10   42-49   6.7-77   8-51   1-8   M     Mean   8 (2.6)   46.8 (1.9)   20.6 (22.1)   36 (22.9)   3.2 (2.1)   Uj     Range   6-14   43.8-48.9   0-83.8   6.8-87   0.6-8.4   Uj     Mean   11(2.3)   44.3 (1.4)   20 (18.7)   7 (6.6)   2.1 (1.1)   Oj     Range   6-17   42.3-48.9   0-76.1   0.5-27.5   0.95-7.3   Oj   | Э       |           |           | Latitude (dec °) | % clay        | Water depth (m) | % sediment MgO      |               |                  |   |
| Range 6-10 42-49 6.7-77 8-51 1-8   Mean 8 (2.6) 46.8 (1.9) 20.6 (22.1) 36 (22.9) 3.2 (2.1) UJ   Range 6-14 43.8-48.9 0-83.8 6.8-87 0.6-8.4   Mean 11(2.3) 44.3 (1.4) 20 (18.7) 7 (6.6) 2.1 (1.1)   Range 6-17 42.3-48.9 0-76.1 0.5-27.5 0.95-7.3  | Group 1 | Mean      | 7 (1.4)   | 45.5 (1.8)       | 46.3 (20.9)   | 20 (10.2)       | 2.7 (1.4)           |               |                  | Most lakes in this group, large %   |
| Mean 8 (2.6) 46.8 (1.9) 20.6 (22.1) 36 (22.9) 3.2 (2.1) Up   Range 6-14 43.8-48.9 0-83.8 6.8-87 0.6-8.4   Mean 11(2.3) 44.3 (1.4) 20 (18.7) 7 (6.6) 2.1 (1.1)   Range 6-17 42.3-48.9 0-76.1 0.5-27.5 0.95-7.3   |         | Range     | 6-10      | 42-49            | 6.7-77        | 8–51            | 1–8                 |               |                  | clay substrates, low taxon richness   |
| Mean 11(2.3) 44.3 (1.4) 20 (18.7) 7 (6.6) 2.1 (1.1) Or  | Group 2 | Mean      | 8 (2.6)   | 46.8 (1.9)       | 20.6 (22.1)   | 36 (22.9)       | 3.2 (2.1)           |               |                  | Upper oligotrophic lakes (Huron, Superior), sediment with high %                |
| Mean 11(2.3) 44.3 (1.4) 20 (18.7) 7 (6.6) 2.1 (1.1) Or<br>Range 6-17 42.3-48.9 0-76.1 0.5-27.5 0.95-7.3   |         | Nalige    | 0-14      | 40.0-40.0        | 0-03.0        | 0.0-0,0         | 0.010.4             |               |                  | MgO, deeper water   |
| 6–17 42.3–48.9 0–76.1 0.5–27.5 0.95–7.3   | Group 3 | Mean      | 11(2.3)   | 44.3 (1.4)       | 20 (18.7)     | 7 (6.6)         | 2.1 (1.1)           |               |                  | Only group with lower lake (Erie)   |
| 200 25 20 25  |         | Range     | 6–17      | 42.3–48.9        | 0-76.1        | 0.5–27.5        | 0.95–7.3            |               |                  | sites (more mesotrophic),<br>shallow water depth, high taxon                    |







- ▲= training sites used for model development
- △= low diversity training sites removed from dataset
- O = validation reference sites
- = failed validation reference sites

Figure 1. Two-dimensional ordinations of the invertebrate presence-absence data for Australian Capital Territory (ACT) (A), Yukon Territory (YT) (B), and Laurentian Great Lakes (GL) (C).

model produced O/E values that distinguished best between D2 and D3 sites (Fig. 2A).

O/E values produced by all models were distributed along a gradient, but the gradients produced by the YT and GL models did not always correspond to the simulated impairment levels (Fig. 2A-C). The YT and GL models generally did not distinguish D1 sites from the original D0 sites (Fig. 2B, C) because the data did not differ. Between 46 and 59% of the total taxa in the original data sets were not used for model creation because they occurred at <10% of reference sites, and many of those excluded taxa were also involved in the simulated impairment process that was applied to data sets prior to model development (ACT: 45%, YT: 42%, GL: 53%). Thus, this situation contributed to nondetection of mild simulated impairment.

**Low-probability taxa** For ACT, the median probability of occurrence for predicted taxa in the sensitive category was 0.34, which is lower than the medians for taxa in the tolerant (0.65) and intermediate (0.46) categories. Compared with the other models, the ACT model had the most taxa above the 0.5 probability cut-off value (Fig. 3A, Table 5). For YT, taxa in the intermediate category had the greatest median value (0.47), and most probability values >0.5 were for taxa in the intermediate category (Table 5, Fig. 3B). For GL, most taxa with probability values >0.5 were in the tolerant category, but the median values for probabilities in all tolerance categories were <0.5 indicating the presence of many low-probability taxa (Fig. 3C, Table 5). This difference between models in taxon probabilities in tolerance categories (combined with fewer taxa overall) contributed to differential ability to detect simulated impairment among models and explains the low number of expected taxa for the YT and GL models (Table 2).

# **DISCUSSION**

We built an AUSRIVAS-style predictive model for reference-site data sets from very different environments (ACT, YT, GL; Tables 2, 3) and used the simulated impairment data sets to evaluate the ability of each model to detect impairment. The data sets differed in total number

Table 4. Type 1 and Type 2 error rates for the Australian Capital Territory (ACT), Yukon Territory (YT), and Laurentian Great Lakes (GL) Australian River Assessment System (AUSRIVAS) predictive models. Each category had 20 sites artificially impaired to simulate degrees of eutrophication impact (D0 = validation sites with no impairment, D1 = mild, D2 = moderate, and D3 = severe).

|       | Type 1 error rate<br>(% validation sites<br>assessed as impaired) | Type 2 error rate<br>(% simulated impairment<br>sites assessed as equal<br>to reference (%)) |      |      |  |
|-------|---|--|------|------|--|
| Model | D0  | D1   | D2   | D3   |  |
| ACT   | 0   | 90   | 20   | 0    |  |
| YT    | 25  | 75   | 10   | 12.5 |  |
| GL    | 37.5  | 57.5   | 32.5 | 12.5 |  |

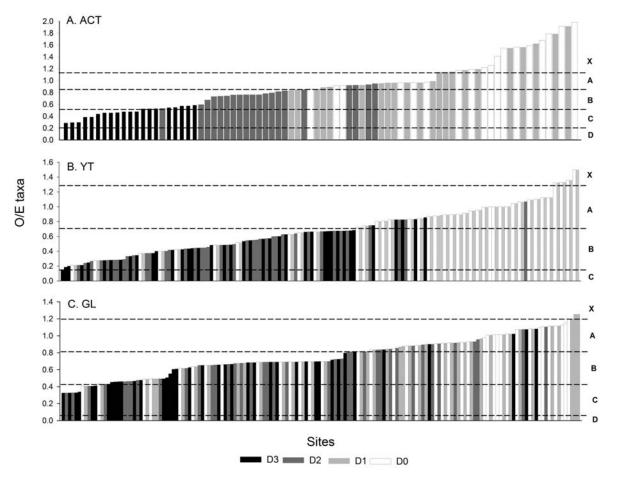


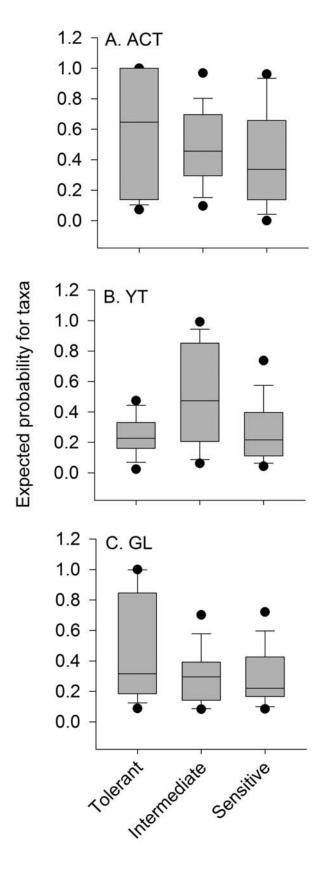
Figure 2. Distribution of observed/expected (O/E) taxa values for the validation sites (D0) and the 3 degrees of simulated impairment (D1 = mild, D2 = moderate, D3 = severe) for the Australian Capital Territory (ACT) (A), Yukon Territory (YT) (B), and Laurentian Great Lakes (GL) (C) data sets. The horizontal dotted lines represent the Australian River Assessment System (AUSRIVAS) bands of biological condition. Band A is equivalent to reference condition, and lower bands represent increasingly impaired condition. Band X is more diverse than reference.

of taxa (Table 2) and in the distribution of taxa (Bailey et al. 2014). These major differences and inherent characteristics of each data set influenced Type 1 and Type 2 error rates of the models.

The Type 1 error for the ACT model was 0%, but the YT and GL models had greater-than-expected Type 1 error rates. The YT and GL data sets contained sites that were similar in terms of measured environmental variables but that differed in their invertebrate assemblages, a combination that makes modeling difficult. The failed sites were within the environmental scope of the models (i.e., not outside model experience based on the predictor variables used) but biologically, many were dissimilar to the training sites used in the models (Fig. 1B, C). The invertebrate assemblages of these failed validation sites were similar to those of the unused, low-richness sites that were considered to have too few taxa for modeling. These low-richness sites may constitute a particular site type that was consequently not represented in the model.

If such sites could be characterized (e.g., if they were all sites from harsh glacial environments in the Yukon Territory or oligotrophic systems in the Laurentian Great Lakes region), the model limitations could be characterized and subsequent model users could be advised that assessment of these types of sites will underestimate the O/E value. Knowledge of the model's limitations would enable users to identify particular site types that a model will not adequately match to reference sites. Users could then select an alternative assessment method or biological group more suitable for assessing those sites. Knowledge that test sites were being compared with an appropriate set of reference sites would provide users with greater confidence in the site assessments provided by the predictive models.

Models with wide biological-quality bands may have lower probabilities of misbanding than models with narrow bands (Barmuta et al. 2003). However, wide bands mean wide ranges of acceptable reference condition and, possibly, less sensitivity to impairment because the im-



Tolerance categories

paired condition is more likely to fall within the range of acceptable condition. Regardless of the cause of wide bands (an inadequate set of reference sites or a naturally wide range of reference condition), such a model may have low power to detect impairment. However, the GL model was least able to detect impairment even though the YT model had the greatest band widths.

Other potential sources of error in estimates of the expected taxa and reference condition include an inadequate set of reference sites or insufficient environmental predictor variables to distinguish among reference-site groups (Ostermiller and Hawkins 2004, Clarke and Hering 2006, Bailey et al. 2012). New spatial tools are becoming increasingly available, particularly geographic information system (GIS) tools and an array of catchment-scale map layers describing attributes, such as geology, landuse, vegetation type, and climate (Frazier et al. 2012). GIS layers and remotely sensed data offer alternative approaches to defining reference sites (Yates and Bailey 2010) and are sources of potential predictor variables (Armanini et al. 2012). The predictor variables used in our study (Table 3) were a selected subset of the available data set, but variables that more completely characterize the factors controlling invertebrate distribution might improve the models (Ostermiller and Hawkins 2004).

Simulating mild impairment involved decreasing the abundance of sensitive taxa, increasing the abundance of tolerant taxa, and removing 2 randomly selected sensitive taxa (Bailey et al. 2014). With a few exceptions for the ACT model, the AUSRIVAS models did not perform well in detecting such mild impairment. Three factors contributed to the nondetection of the mild level of simulated impairment. First, the AUSRIVAS observed taxa list will not change if the taxa selected for simulated removal were not used for model development and, thus, were not included in the list of expected taxa. The standard AUSRIVAS modeling procedure is to remove (rare) taxa that occur at <10% of reference sites in the training data set. For all models, the number of taxa removed before model creation was high (Table 2). Second, where the artificially impacted taxa had <0.5 probability of occurrence at the site, they would not contribute to the O/E score. Third, we developed the models using presence-absence data, and thus, the models will not detect impacts within the data sets that are manifested only by a change in abundance. Thus, the

Figure 3. Distribution of the expected probability of occurrence for taxa across all validation sites for the Australian Capital Territory (ACT) (A), Yukon Territory (YT) (B), and Laurentian Great Lakes (GL) models. Taxa are grouped by their tolerance category (see Bailey et al. 2014 for assignments of taxa to categories). Lines in boxes are medians, box ends are quartiles, whiskers are 90th and 10th percentiles, and circles are 5th and 95<sup>th</sup> percentiles.

Table 5. Number of taxa probability values > 0.5 by taxon tolerance category for the Australian Capital Territory (ACT), Yukon Territory (YT), and Laurentian Great Lakes (GL) Australian River Assessment System (AUSRIVAS) predictive models (number of sites used for each model in brackets).

| Model             | Tolerant | Intermediate | Sensitive | Total |
|-------------------|----------|--------------|-----------|-------|
| ACT (sites = 87)  | 40       | 177          | 96        | 313   |
| Taxa per category | 5        | 18           | 13        | 36    |
| YT (sites = 101)  | 7        | 187          | 64        | 258   |
| Taxa per category | 6        | 10           | 12        | 28    |
| GL (sites = 84)   | 199      | 29           | 20        | 248   |
| Taxa per category | 14       | 5            | 3         | 22    |

taxon richness (the basis for the O/E score) for most of the mildly disturbed sites was similar to that of the original validation data set and the taxa observed (O) (i.e., the taxa captured from the list of predicted taxa) differed little, or not at all, between the validation sites and those that were mildly impaired for all models (Table 4, Fig. 2A-C).

Consequently, the models had large Type 2 error rates regarding mildly disturbed sites (particularly the ACT model; Table 4). The Type 2 errors for the D1 sites were largely an inverse reflection of the Type 1 error rates. The difference among models regarding the Type 2 errors for mildly impaired sites is related to the random nature of taxa removed to simulate the impairment and to the differential effects that missing taxa have in relation to the number of taxa expected (which differed by model; Table 2). For example, removing 2 taxa from a YT site at which 3.6 taxa (56% of expected taxa) are expected will have greater effect on the O/E value than removing 2 taxa from an ACT site at which 13.9 taxa (14.4% of expected taxa) are expected.

If the simulated impairment data sets accurately represented eutrophication disturbance, then the AUSRIVAS model better detected such disturbance in the ACT region than did the models developed for the GL or YT regions. The ACT model most accurately displayed the gradient from moderate to severe impairment (Fig. 2A). The other 2 models displayed a gradient of O/E values, but the impaired and validation sites were more randomly distributed along that gradient. Often the different levels of impairment at specific sites in the GL and YT data sets were not distinguishable (Fig. 2B, C). The ability to detect the simulated impairment depended on whether the simulated disturbance was severe enough to remove taxa and whether those same taxa were used for modeling. The ACT model was created with a data set that had more taxa per site and more uniformly distributed taxa than in the other data sets, so a greater proportion of the biological data were used for model development. Thus, the taxa that underwent simulated impairment had a greater chance of being used in the ACT model than in the YT and GL models, which increased the probability of detecting the ACT impairment.

Regardless of whether abundance or presence-absence data (as for AUSRIVAS) are used for modeling, detection of eutrophication or any other disturbance in the real world will depend on the invertebrate sampling and processing methods. As sites become increasingly stressed, more of the sensitive taxa will disappear from the sites and the samples (Cao and Hawkins 2005). The sampling and subsampling methods will influence the proportion of locally rare taxa observed in a sample (Clarke and Murphy 2006). For example, sampling methods that collect the maximum number of different taxa regardless of their abundance may cause the model to have trouble detecting a mild disturbance that simply changes the relative abundances of taxa (Nichols and Norris 2006), whereas a sampling method that collects taxa relative to their abundance at the site (Nichols et al. 2000, Nichols and Norris 2006, Environment Canada 2012) could enable the model to detect a change in abundance before the impact removes taxa from the site, even when relying on taxon richness measures for assessment. Data sets collected with different sampling methods at the same stressed site can give the appearance of different responses to the disturbance simply because of the sampling or subsampling method (Ostermiller and Hawkins 2004). When simulated impairments were applied to an existing data set, the assumption made was that the sampling methods had not influenced the assemblage structure of the data set. Clearly, this assumption was not correct. Nonetheless, a simulated impairment data set is the only way to evaluate method performance regarding Type 2 errors. The accuracy of the representation of the impairment is less important than knowing the level to which the data set was impaired. Moreover, we evaluated only the O/E taxa modeling method, which is only 1 component of the AUSRIVAS bioassessment protocol, which includes standardized sampling methods and other outputs and indices to aid interpretation of the O/E result.

AUSRIVAS models from the different regions varied in the number of taxa predicted and, thus, expected (Table 2). In models with low numbers of expected taxa, the O/E score is vulnerable to the chance omission of observed taxa at a site (Barmuta et al. 2003). Such chance omissions could result in misbanding a site and failing it when it should have passed (Type 1 error). Marchant (2002) suggested that O/E scores calculated from <20 expected taxa may be too variable and unreliable to use. In reality, the argument regarding the chance omission of observed taxa should be viewed relative to the probability of missing or misidentifying taxa at a site (Barmuta et al. 2003). If taxa at the low-richness sites also have a low probability of being misidentified or missed during sampling then the problem may not be great (Barmuta et al. 2003). Replicated sampling at naturally low-richness sites used for modeling (such as those from harsh environments, e.g., some YT sites) may help to ensure that the reference condition is not underestimated for initial model development (Barmuta et al. 2003). Such replicated sampling could reduce the chance of Type 2 errors (sites passing that should fail) by providing a more reliable estimate of reference condition at lowrichness sites.

We used the standard AUSRIVAS method in our study so that we could compare our results with those produced by other methods presented in this special series. Thus, we used the standard (although somewhat arbitrary) AUSRIVAS probability cut-off of 0.5, which excludes lowprobability taxa from the expected taxa list. AUSRIVAS uses a 0.5 probability cut-off because taxa with ≤0.5 probability of occurrence have an equal or greater probability of not being observed at a site. Decreasing the cut-off value to <0.5 may increase the number of expected taxa. However, any new taxa will add increasingly slowly to the count of expected taxa and will not necessarily strengthen confidence in the O/E scores (see Marchant 2002, Clarke and Murphy 2006). Excluding taxa with a low probability of occurrence will result in less variable O/E estimates (Clarke and Murphy 2006), but the optimal cut-off value does not have universal consensus (Hawkins et al. 2000, Marchant 2002, Ostermiller and Hawkins 2004, Clarke and Murphy 2006, Van Sickle et al. 2007). Clark and Murphy (2006) found the marginally best cut-off value to be 0.2, but the power of detecting impacts was similar up to 0.5. Van Sickle et al. (2007) found that excluding taxa with < 0.5 probability increased ability to detect impairment. Including low-probability taxa in the O/E calculations assumes they are reliable and not simply absent by chance from new assessment sites. Marchant (2002) concluded that low-probability taxa play no useful role in predictive models, such as AUSRIVAS. The results of such studies caused AUSRIVAS developers to use the 0.5 cut-off. Moreover, O/E cut-off thresholds must be standardized when comparing site assessments in multijurisdictional bioassessment programs, or they should be treated as different indices (Clarke and Murphy 2006). Nevertheless, by comparing the performance differences among models from the 3 regions, we found that the distribution of lowprobability taxa contributed to whether a particular data set could produce an adequate predictive model for the detection of impairment. Compared with the other 2 data sets, the GL data set had the most severely skewed invertebrate frequency distribution (i.e., more of the lowprobability taxa) (Bailey et al. 2014), and therefore, was more vulnerable to the effects of excluding taxa with <0.5 probability of occurrence.

Taxa assigned to the sensitive category tended to have lower expected probabilities of occurrence than moretolerant taxa (Fig. 3A-C). Other investigators found similar patterns in the expected probabilities of sensitive taxa for RIVPACS-style predictions (Clarke and Murphy 2006).

The sensitive taxa tended to be less widespread among reference sites particularly for the YT and GL data sets and, thus, had considerably lower average expected probabilities (Fig. 3B, C). Thus, use of the 0.5 probability threshold excluded more sensitive taxa than taxa in other categories and contributed to nondetection of impairment by the YT and GL models.

AUSRIVAS and most other RIVPACS-style predictive models use discriminant function analysis (DFA), which requires identification of reference-site groups (Van Sickle et al. 2006). However, in most reference data sets with many sites, invertebrate data are not characterized by discrete community assemblages (Hawkins and Vinson 2000). Rather, the data structure displays sites along a continuum of ≥1 taxonomic gradients. Each taxon's array of environmental requirements and habitat preferences determine the gradients evident in the invertebrate data sets (Resh et al. 1994, Menezes et al. 2010). The spatial scale of sampling also influences the underlying structure revealed by analysis, such as classification and ordination (Marchant et al. 1999), and gradients may become more obvious as the size of the reference data set (or the density and spatial scale of reference-site coverage) increases (Turak et al. 1999). The ACT data were collected from a relatively small area (12,000 km<sup>2</sup>) compared with both the YT (840,000 km<sup>2</sup>) and GL (244,160 km<sup>2</sup>) data sets (Bailey et al. 2014). Thus, the density of ACT reference-site coverage also was greater. Our results indicated that the ACT model performed best, and the spatial ordination and density of reference sites may have contributed to this outcome.

Classifying discrete groups of sites is a requirement of DFA rather than a representation of the reality of the invertebrate assemblages. Other modeling approaches may explicitly acknowledge the continuum in taxon distributions and avoid the use of classification groups by using the ordination space of reference sites as the basis for predicting site-specific invertebrate assemblages (Linke et al. 2005). However, AUSRIVAS does not base the probability of taxon occurrence on just 1 classification group that is most similar to a site, unlike some other methods, e.g., Benthic Assessment of Sediment (BEAST; Reynoldson et al. 2001). Rather, AUSRIVAS uses the weighted probabilities of the site membership to all of the groups, in a sense accounting for the assemblage continuum. The use of weighted probabilities of the site membership to all classification groups may moderate the effects of misclassification errors associated with large cross-validation errors, as for the YT model (Table 2).

#### Conclusion

The ability of our models to detect the simulated impairment depended on whether the simulated disturbance was severe enough to remove taxa from the data set and on whether the removed taxa had been used for modeling. Rare taxa, which have a patchy distribution in the data sets, were removed before developing the AUSRIVAS models. To further improve predictive performance, we removed sites with naturally low richness (from the YT and GL data sets). Thus, this low-richness site type was not represented in the models, a limitation for the particular model. Other methods or biota may be better for assessing the condition of sites with naturally low invertebrate richness. Thus, effectiveness and performance of the models was related to differences in the total number of invertebrate taxa and to the distribution of taxa in the data sets. In short, data sets with highly skewed taxon distributions are difficult to model.

Careful evaluation of model performance should consider both Type 1 and Type 2 errors because confidence in the assessment is related to the model's ability to detect impairment. Use of a simulated impairment data set is the only way to evaluate Type 2 errors in model performance. The YT model was the best for detecting moderate impairment (10% error), and the ACT model detected all severely impaired sites. All models detected a gradient in O/E scores, but the ACT model best distinguished between moderate and severe simulated impairment. Moreover, AUSRIVAS was able to assign site O/E scores to a band of biological quality, thereby indicating the level of impairment. AUSRIVAS did not assign most mildly impaired sites to below reference condition, but a reduction in O/E values within band A was observed for some mildly impaired sites. AUSRIVAS did not detect simulated mild impairment that simply changed taxon abundance in a data set because presence-absence data were used for model development. Nevertheless, in comparison with other models described in this special issue (that did use abundance data), the AUSRIVAS model performance was comparable or better for detecting the simulated moderate and severe impairments.

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