Development of Models to Establish Links between Human-caused Nutrient Enrichment and Alterations to the Composition of Stream Communities

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Acronyms

AWCH_AVE – Soil Water Capacity
CHL \( a \) – Chlorophyll \( a \)
CO Plateaus – Colorado Plateaus
CV - Coefficient of Variation
DNOx – Dissolved Nitrate/Nitrite
DP – Dissolved Phosphorus
Elev_WS – Watershed Elevation
E – Ephemeroptera Richness
EPT – Ephemeroptera, Plecoptera, Trichoptera
EPTPct – Percent EPT individuals
G_NU_AVE – Mean Reactive Geology
HBI - Hilsenhoff Biotic Index
HIGH_DP – High Dissolved P Class
Lat_Dec – Latitude in Decimal Degrees
LogAnnNH4 – Log\(_{10}\)Annual Ammonium
LogAnnNO3 – Log\(_{10}\)Annual Total Nitrate
LogAnnDNO3 – Log\(_{10}\)Annual Dissolved Nitrate
LogAnnTP – Log\(_{10}\)Annual Total Phosphorus
LogAnnDP – Log\(_{10}\)Annual Dissolve Phosphorus
Log_Sq_km – Log\(_{10}\)Watershed Area
Long_Dec – Longitude in Decimal Degrees
LOWESS – Locally Weighted Scatterplot
Smoothing
N – Nitrogen
NH4 – Ammonium
NH4TOT – Total Ammonium
NOXDISS – Dissolved Nitrate/Nitrite
NOXTOT – Total Nitrate/Nitrite
O/E – Observed to Expected taxa richness
P – Phosphorus
PCA – Principal Components Analysis
PDISS – Dissolved Phosphorus
PTOT – Total Phosphorus
PRED_ALK – Predicted Alkalinity
PRED_COND – Predicted Conductivity
PRED_ALK – Predicted Alkalinity
PRED_COND – Predicted Conductivity
RFOE – O/E score
SCR – Scraper Richness
SCRCG - Scraper-Collector Gatherer Richness
S. Rockies – Southern Rockies
Taxa – Taxa Richness
TKN – Total Kjeldahl Nitrogen
TN – Total Nitrogen
TNOx – Total Nitrate/Nitrite
TP – Total Phosphorus
UDWQ - Utah Department of Water Quality
USEPA – United States Environmental Protection Agency
UT - Utah
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Background

Eutrophication of the nation’s surface water resources (lakes, reservoirs, rivers, streams and wetlands) due to excessive nutrients (nitrogen and phosphorus) is recognized as a significant water quality problem. The U.S. Environmental Protection Agency’s (USEPA) national water quality summary reports to Congress consistently identify excessive nutrients as one of the top three leading causes of impairments of the nation’s waters (along with siltation and pathogens). Natural nutrient inputs support primary producers that are essential for supporting healthy, diverse and productive ecosystems. However, excessive nutrient inputs can result in abundant growth of periphyton (diatoms and multicellular algae), macrophytes and/or phytoplankton leading to oxygen depletion, potentially toxic algal blooms, imbalances in biological community composition, an human health concerns.

As a result of the threat posed by eutrophication, USEPA published the National Strategy for the Development of Regional Nutrient Criteria (USEPA 1998) under section 304(a) of the Clean Water Act (66 Federal Register [FR] 1671). This document describes the basis and national approach for working with States in the development and adoption of nutrient criteria. Following this approach, USEPA proposed criteria for rivers, streams, lakes and reservoirs in a series of Water Quality Criteria Recommendation documents (USEPA 2000a, 2000b, 2000c, 2001a, 2001b, 2001c, 2001d, 2001e). These proposed criteria were based on nutrient ecoregions and were intended to serve as a springboard for States to develop more refined nutrient criteria, stating that “States and Tribes need to identify with greater precision the nutrient levels that protect aquatic life and recreational uses...” (USEPA 2000d, 2000e). USEPA expected States to adopt or revise the proposed nutrient ecoregion criteria into State water quality standards (WQS) by 2004 while recognizing that the time needed to develop standards and the available resources differed significantly among States. This analysis is part of Utah’s effort to develop protective and scientifically defensible nutrient criteria.

It is our understanding that Utah is considering a variety of technical approaches for criteria development. This specific summary was focused on statistical analyses with existing UT ambient monitoring data to identify stressor-response relationships of utility in developing endpoints that support the criteria development effort. This analysis focuses in particular on the available aquatic life use response variables based on macroinvertebrates that are used by the state in making aquatic life use decisions. It is valuable, at this point, to summarize the conceptual model that links nutrient enrichment to effects on macroinvertebrates (Fig. 1).
The experimental and other support for each of the relationships identified in this diagram can be found in freshwater texts (e.g., Allan 1995), summary papers (e.g., Carpenter et al. 1998), as well as an abundance of primary scientific literature (e.g., Elwood et al. 1981, Peterson et al. 1993, Rosemond et al. 1993, Feminella and Hawkins 1995, Hillebrand 2002, King and Richardson 2003, Slavik et al. 2004, Cross et al. 2006). There is, therefore, a well defined and scientific basis for establishing linkages between nutrients and responses along the causal pathway in Figure 1 up to and including aquatic life.

This report is focused on macroinvertebrate response variables since, to our knowledge, these represent the largest, available set of response data for use in developing stressor-response relationships linking nutrient enrichment to aquatic life use response in Utah. This is not a detailed report, rather it summarizes the general approaches, methods, and results of this preliminary analysis.

**Analytical Approach**

The general approach employed is consistent with EPA guidance on establishing nutrient criteria for rivers and streams (USEPA 2000d). The first step involved exploring potential classification to reduce natural variability in nutrient conditions resulting from geographic factors. After exploring classification, the second step was to explore distributional statistics of nutrients and to develop statistical stressor-response relationship models to identify candidate endpoints. We finish by reviewing analyses to strengthen support for the relationship between nutrients and aquatic life use response. Methods for each are described in order.
Classification

We worked with scientists at Utah Department of Water Quality (UDWQ) to develop a strategy for exploring potential natural classes of nutrient behavior for streams in Utah for which nutrient data were available. This process involved looking at nutrient distributions among a priori defined classifications (ecoregions) and exploring predictors of differences in nutrient concentrations within reference sites using a range of multivariate and discriminant models.

Using all the data we had in our dataset relevant to nutrients in Utah, we first examined differences in nutrient distributions by ecoregion. There was little difference among ecoregions for nitrate concentration (Fig 2) or total phosphorus concentration (Fig. 3). Non-parametric means comparison tests also indicate no significant differences among level III ecoregions.

![Box Plot (UTData.sta 164v*1510c)](image)

Figure 2 – Box and whisker plot of Log(Nitrate/Nitrite) annual values within different level III ecoregions. Units are mg/L.
We also examined reductions in coefficients of variation (CV) within level III ecoregions in comparison to all regions combined to examine whether a substantial reduction in variability could be achieved by classification. There was variation in CV across ecoregions with some higher and some lower than the combined regions CV (Table 1). In general, there was little in the CV reduction information, especially among reference sites, to recommend the use of specific classes within Utah that would not result in increased variability among some resulting classes. Any improvement would be marginal, at best, based on these data.

**Figure 3 - Box and whisker plot of Log(Total Phosphorus) annual values within different level III ecoregions. Units are mg/L.**
The next set of analyses was based on standardizations of nutrient concentrations in reference sites. Annual and seasonal nutrient mean concentrations were first identified as High P or High N if they were greater than the 75\textsuperscript{th} percentile of reference site nutrient distributions. We then looked at the difference in nutrient distributions between the two classes of reference streams – those that were high nutrients versus those that were typically low – based on a wide range of natural predictors (Table 2).

Table 1- Coefficients of variation for all sites and just reference sites (annual and summer data) for various nutrient parameters. CVs were calculated for all data combined and then split into level III ecoregions. A few sites were in unknown ecoregions because of missing latitude and longitude values. Means and standard deviations were calculated on log-transformed data.

<table>
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<th></th>
<th>Combined</th>
<th>Unknown</th>
<th>Central Basin</th>
<th>Wasatch</th>
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<th>S. Rockies</th>
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<td>All Sites</td>
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<td></td>
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<td>LogAnnNH4</td>
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<td>-44.3</td>
<td>-29.3</td>
<td>-15.8</td>
<td>-4</td>
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<tr>
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<td>-99.8</td>
<td>-520.9</td>
<td>-42.3</td>
<td>-44.7</td>
<td>-16.3</td>
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<td>-47.7</td>
<td>-21.4</td>
</tr>
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<td>-65.2</td>
<td>-21.8</td>
<td>-32</td>
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</tr>
<tr>
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<td>-36.4</td>
<td>-64.7</td>
<td>-26.3</td>
<td>-27.5</td>
<td></td>
</tr>
<tr>
<td>Reference Sites</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td>-47.5</td>
<td>-13.3</td>
<td>-2.1</td>
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<td>-20.1</td>
<td>-40.3</td>
<td></td>
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<td>LogAnnDNOx</td>
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<td>-9.4</td>
<td>-41.5</td>
<td>-39.9</td>
<td>-20.7</td>
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<td>LogAnnTP</td>
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<td>-13.6</td>
<td>-25.7</td>
<td>-10</td>
<td>-27.8</td>
<td></td>
</tr>
</tbody>
</table>

| Reference Sites     |          |         |               |         |             |            |
| LogSuNH4            | 150.2    | 141.4   | 125.6         | 223.6   |             |            |
| LogSuTNOx           | 17.6     | 24.4    | 22.5          | 8.4     | 9.8         |            |
| LogSuDNOx           | 60.3     | 141.4   | 44.7          | 88.7    |             |            |
| LogSuTP             | 45.4     | 31.3    | 0             | 44      | 56.4        |            |
| LogSuDP             | 118.8    | 0       | 72.6          | 200     |             |            |

Table 2 – Predictors generated by Utah State used in comparing nutrient distributions and in multivariate models.

<table>
<thead>
<tr>
<th>Physical</th>
<th>Temperature</th>
<th>Soils</th>
<th>Geochemistry</th>
<th>Hydrology</th>
</tr>
</thead>
<tbody>
<tr>
<td>Latitude</td>
<td>Earliest Freeze Date</td>
<td>Soil water capacity</td>
<td>Predicted conductivity</td>
<td>Min:Max flow rate</td>
</tr>
<tr>
<td>Longitude</td>
<td>Last Freeze Date</td>
<td>Soil permeability</td>
<td>Predicted alkalinity</td>
<td>Mean precipitation</td>
</tr>
<tr>
<td>Elevation</td>
<td>Mean Air Temperature of Stream Network</td>
<td>Calcium content of rocks</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Watershed Area</td>
<td>Mean Air Temperature of Watershed</td>
<td>Percent carbonate geology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Relief Ratio</td>
<td></td>
<td>Mean reactive geology</td>
<td>Percent volcanic</td>
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We examined box plots of standardized nutrient distributions between high nutrient and low nutrient reference sites for every nutrient parameter (e.g., Figs. 4 and 5). The examples shown in Figs. 4 and 5 were for dissolved phosphorus, but all the nutrient parameters showed similar results, no nutrient exhibited different distributions of potential predictors between high nutrient and low nutrient reference classes.

![Box Plot](image)

Figure 4 – Box and whisker plots of distributions of different predictor variable values between low dissolved P (N/A) and high dissolved P (Yes). This plot indicate no difference between the two nutrient regimes for latitude (Lat_Dec), longitude (Long_Dec), soil water capacity (AWCH_AVE), and predicted alkalinity (PRED_ALK).
We also ran the same analysis using reference quantiles. This was performed by coding reference sites based on the quantile of reference nutrient concentrations into which they fell and attempting to identify predictors that discriminated among the 4 quantile classes. Again, there were no natural predictors that discriminated among reference nutrient classes consistently. We then ran discriminant analysis with high and low nutrient classes and reference quantiles as groups and the list of variables in Table 2 as predictors. Once again, no significant model could be developed that adequately discriminated among reference site nutrient groups.

Figure 5 - Box and whisker plots of distributions of different predictor variable values between low dissolved P (N/A) and high dissolved P (Yes). This plot indicate no difference between the two nutrient regimes for mean reactive geology (G_NU_AVE), watershed areas (Log_Sq_Km), elevation (Elev_WS), and min:max flow rate ration (Hydr_WS).
Finally, we ran cluster analysis on the predictors in Table 2 for reference sites. We developed 2, 3, and 4 group models, but focused on the 2 group models because they showed the most promise, and this, only marginally. After defining the 2 groups with the cluster analysis, we then ran the same predictors in PCA and coded the sites by cluster group. The two clusters showed some separation with regards to the two clusters (Figure 6), and this was principally along a geologic attribute axis, reflected in predicted alkalinitities and conductivities.

Figure 6 – Plot of axes 1 and 2 from a principal components analysis of predictors in Table 2. Sites are coded according to the two cluster groups generated from a cluster analysis of the same predictors. Separation along axis 1 is evident and due to differences in geology among these two classes.
We then examined differences in nutrient distributions within reference sites using these groupings. Only dissolved phosphorus showed any indication of separation according to the cluster groupings (Fig. 7).

Given the weak relationships among natural predictors and reference site nutrient concentrations and the inability to identify a convincing basis for stream nutrient classes, we recommend against classifying streams for the basis of nutrient criteria development at this time.

**Nutrient Analysis**

**Distribution Based Numbers**

Before beginning an investigation of relationships between nutrient stressors and biological response variables, we calculated standard distributional statistics for nutrient parameters across the state. Sites were labeled according to their reference status as determined by UDWQ.
Statistics for quartiles were the focus of the analysis, after USEPA guidance (e.g., USEPA 2000d). Seasonal quartiles were estimated for reference (75\textsuperscript{th}) and all site (25\textsuperscript{th}) populations, and the median of those quartiles estimated as an annual value. Graphical distributions for total nitrate and total phosphorus were developed (Figs. 8 and 9) and a tabular summary prepared (Table 3).
Figure 9 - Box and whisker plots of seasonal total phosphorus site medians for reference and all site populations. Also shown are the 75th and 25th percentiles of these populations, respectively.

Table 3 – Seasonal 75th and 25th percentiles of nutrient variable medians for reference and all site populations, respectively. Also included are the medians of seasonal quartile values as annual values. All units in mg/L.

<table>
<thead>
<tr>
<th></th>
<th>NH4</th>
<th>TKN</th>
<th>TNOx</th>
<th>DNOx</th>
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<tr>
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<td>0.043</td>
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<td>0.048</td>
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<tr>
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<td>0.598</td>
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<tr>
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<td>0.100</td>
<td>0.026</td>
<td>0.000</td>
</tr>
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</table>

Annual reference site values for total nitrate and total phosphorus were 230 and 43 µg/L, respectively (seasonal 75th percentile ranges were 185-310 and 36-58 µg/L, respectively). Comparable values for the 25th percentile of all sites were 128 and 26 µg/L for total nitrate and total phosphorus, respectively (seasonal 25th percentile ranges were 100-160 and 24-31 µg/L, respectively).
**Stressor-Response Analysis**

Stressor-response approaches refer to a suite of analytical techniques to derive candidate endpoints by 1) identifying nutrient concentrations associated with pre-existing thresholds in response variables (interpolation) and 2) exploring and identifying thresholds in the relationships between response variables and nutrient concentrations (Figure 10).

![Figure 10](image-url)  
**Figure 10** Plot showing two stressor-response relationships, where the response is a direct measure of designated use or can be easily linked to a designated use measure. In relationship B, a preexisting threshold of the response variable exists. For example, UDWQ O/E score thresholds (e.g., 0.74) or minimum dissolved oxygen criteria (3 to 8 mg/L depending on use class) could be the response variable and the corresponding criterion value for each used as the pre-designated threshold. The hatched line from the response threshold to the regression line is extrapolated downward to estimate the nutrient concentration associated with that threshold (b). In relationship A, no pre-existing threshold in the response variable exists; however the response is non-linear and changes once the concentration (a) is reached. This changepoint becomes a candidate criterion.

Response variables for stream nutrient endpoint development for this analysis focused on aquatic life use indicators (biological metrics and O/E scores). The value of these indicators lies in their direct linkage to aquatic life use designations. They, therefore, provide a way to connect nutrient concentrations directly to designated aquatic life use support.

Response variables were first examined with general Spearman correlation analysis and visual scatterplots with smoothing functions to identify potential relationships between biological response and nutrient variables. Locally weighted scatterplot smoothing (LOWESS) was used to fit lines representing the general relationship between the response and the nutrient stressor variables. The LOWESS technique (Cleveland 1979) models nonlinear relationships where linear methods do not perform well. LOWESS fits simple models to localized subsets of the data to construct a function that describes, essentially, the central tendency of the data.
Correlations of interest (significant correlations with highest r values) were selected and used for subsequent stressor-response analyses. Highly correlated response measures were prioritized based on using those with strongest correlation with nutrient stressors and a final list of response metrics was developed (Table 4).

Table 4 – List of biological response and nutrient stressor variables that were the focus of analysis based on initial correlation screening. *Ephemeroptera richness did not show significant correlations for the summer data alone, but other response variables did.

<table>
<thead>
<tr>
<th>Biological Responses</th>
<th>Nutrient Stressors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Taxa Richness (Taxa)</td>
<td>Total Ammonium (NH4TOT)</td>
</tr>
<tr>
<td>EPT Richness (EPT)</td>
<td>Total Nitrate (NOXTOT)</td>
</tr>
<tr>
<td>Ephemeroptera Richness (E)*</td>
<td>Dissolved Nitrate (NOXDISS)</td>
</tr>
<tr>
<td>Percent EPT individuals (EPTPct)</td>
<td>Total Phosphorus (PTOT)</td>
</tr>
<tr>
<td>Scraper richness (SCR)</td>
<td>Dissolve Phosphorus (PDISS)</td>
</tr>
<tr>
<td>Scraper-Collector Gatherer Richness (SCRCG)</td>
<td></td>
</tr>
<tr>
<td>Hilsenhoff Biotic Index (HBI)</td>
<td></td>
</tr>
<tr>
<td>O/E Scores</td>
<td></td>
</tr>
</tbody>
</table>

*Interpolated endpoints based on existing thresholds*

Once correlation analyses were complete, a set of promising stressor-response relationships were identified and the stressor-response analysis proceeded. Two specific stressor-response approaches were applied: regression analysis to identify nutrient concentration associated with existing response endpoints and change-point analysis to identify thresholds in responses to increasing nutrient concentrations not having pre-existing endpoints (Figure 10).

Linear regression models were developed for each apparently linear stressor-response relationship of interest for which identifiable or pre-existing thresholds in the response variable existed. Identifiable thresholds for macroinvertebrate metrics were generated by exploring percentile values of reference site metric distributions. A standard practice in scoring individual metrics for use in bioindicator development is to use 25th percentile metric scores of reference sites for metrics that decrease with stress (75th percentile for those that increase with stress) as thresholds between reference and non-reference sites. A similar approach is used for scoring final indices (Barbour et al. 1999). We applied these criteria to develop reference site “expectations” for individual metrics that were then used as thresholds for interpolating nutrient concentrations consistent with acceptable biological conditions (Table 5). In addition to these endpoints, O/E scores of 0.54 and 0.74 were used as response thresholds, consistent with UDWQ decision rules regarding aquatic life use evaluation based on O/E scores.
Standard regression diagnostics were used to identify and evaluate appropriate models. Linear interpolation was used to identify those nutrient concentrations associated with these specific response endpoints (e.g., biocriteria levels).

An example regression and interpolation based on a nutrient stressor-response relationship is shown in Figure 11. This same analysis was repeated for each significant regression and each nutrient-metric relationship.
response pair. The summary of interpolation values is shown in Table 6 for both annual values and summer values. The figures for each of these analyses are in Appendix 1.

Table 6 – Table of interpolated nutrient values associated with biological thresholds existing (O/E) or calculated based on reference quartiles (see text). Values are given for total nitrate (NOXTOT), dissolved nitrate (NOXDISS), and total phosphorus (PTOT). All values in mg/L. Endpoints are shown based on annual values and summer values. Abbreviations for biological responses are given in Table 4.

<table>
<thead>
<tr>
<th>Biological Response</th>
<th>NOXTOT</th>
<th>NOXDISS</th>
<th>PTOT</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Annual Data</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Taxa</td>
<td>0.234</td>
<td>0.280</td>
<td>0.046</td>
</tr>
<tr>
<td>EPT</td>
<td>0.289</td>
<td>0.341</td>
<td>0.054</td>
</tr>
<tr>
<td>E</td>
<td>1.068</td>
<td>1.191</td>
<td>0.568</td>
</tr>
<tr>
<td>EPTPct</td>
<td>1.078</td>
<td>1.779</td>
<td>0.608</td>
</tr>
<tr>
<td>SCR</td>
<td>1.245</td>
<td>1.360</td>
<td>0.507</td>
</tr>
<tr>
<td>HBI</td>
<td>0.354</td>
<td>0.457</td>
<td>0.080</td>
</tr>
<tr>
<td>RFOE0.8</td>
<td>0.055</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RFOE0.74</td>
<td>0.108</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RFOE0.6</td>
<td>0.529</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RFOE0.54</td>
<td>1.040</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Summer Data</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Taxa</td>
<td>0.207</td>
<td>0.288</td>
<td></td>
</tr>
<tr>
<td>EPT</td>
<td>0.286</td>
<td>0.367</td>
<td>0.036</td>
</tr>
<tr>
<td>E</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EPTPct</td>
<td>1.513</td>
<td>1.911</td>
<td>0.731</td>
</tr>
<tr>
<td>SCR</td>
<td>0.495</td>
<td>0.545</td>
<td>0.081</td>
</tr>
<tr>
<td>SCRCG</td>
<td>0.397</td>
<td>0.455</td>
<td>0.075</td>
</tr>
<tr>
<td>HBI</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RFOE0.8</td>
<td>0.030</td>
<td>0.053</td>
<td></td>
</tr>
<tr>
<td>RFOE0.74</td>
<td>0.070</td>
<td>0.124</td>
<td></td>
</tr>
<tr>
<td>RFOE0.6</td>
<td>0.488</td>
<td>0.896</td>
<td></td>
</tr>
<tr>
<td>RFOE0.54</td>
<td>1.120</td>
<td>2.090</td>
<td></td>
</tr>
</tbody>
</table>

Endpoints based on change-point analysis

After exploring potential endpoints using interpolation of thresholds, we investigated whether apparent non-linear stressor-response relationships to nutrients also existed for the metric responses and conducted change-point analysis to identify thresholds in biological responses that may represent potential endpoints. For change-point analysis, nonparametric deviance reduction was used to identify thresholds in biological responses to nutrients (Qian et al. 2003, King and Richardson 2003). This technique is similar to regression tree models, which are used to generate predictive models of response variables for one or more predictors. The change-point, in our application, was the first significant split of a tree model with a single predictor variable (nutrient concentration). This first split is
that point along the stressor axis that splits the response data into the two groups that result in the greatest significant (based on a $\chi^2$ test) reduction in deviance (Breiman et al. 1984). Output from change-point analyses included the threshold as well as the proportion of reduction in error (PRE), analogous to the regression coefficient from general linear models.

We explored the stressor-response relationships visually using LOWESS regression for the annual nutrient chemistry variables first. Of the potential relationships, only HBI and O/E showed potential threshold responses, and those only to median nitrate concentrations (e.g., Fig. 12, additional models in Appendix 3). Additionally, three models for summer nutrient variables were explored based on apparent, potential non-linear stressor-response relationships. These were O/E and total nitrate and scraper and scraper/collector-gatherer richness and total phosphorus (Appendix 4). A summary of change-point based endpoints is provided in Table

![Graph showing HBI vs. Log$_{10}$(total nitrate) and change-point analysis results.](image)

**Figure 12 - Plot of HBI vs. Log$_{10}$(total nitrate), as well as output from a change-point analysis of HBI and Log$_{10}$(total nitrate). The change-point at Log$_{10}$(total nitrate) of -0.395 (0.403 mg/L) resulted in a 24% reduction in model error. Units in mg/L.**
Table 7 – Change-points associated with the response of various biological responses to nutrient enrichment. All values in mg/L.

<table>
<thead>
<tr>
<th>Biological Response</th>
<th>Annual NOXTOT</th>
<th>NOXDISS</th>
<th>NOXTOT</th>
<th>NOXDISS</th>
<th>PTOT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Scraper Richness</td>
<td>0.042</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Scraper Collector-Gatherer Richness</td>
<td>0.043</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HBI</td>
<td>0.403</td>
<td>0.288</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RFOE</td>
<td>0.215</td>
<td>0.23</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

An important consideration with the change-points observed in Utah that needs to inform their interpretation is the ecological phenomenon they may represent. Some change-points are identified where a lack of response to nutrients ends and nutrients begin to assert an effect on declining biological condition (e.g., Figure 10, point a). This would certainly inform the selection of an appropriately protective value. However, if there is no initial period of resistance and condition immediately responds to nutrient enrichment, then the resultant change-points may represent the point where nutrients no longer contribute to a decline in biological condition and, even worse, the point of worst possible biological condition. These change-points would be identified at the end of a response curve to a stressor (e.g., Figure 13, point c), since the statistical properties to the right of that point (flat-line) are indeed different than those to the left (declining condition).

Figure 13 – Plot showing a stressor-response relationship, where the response is a direct measure of designated use or can be easily linked to a designated use measure. In this relationship A, no pre-existing threshold in the response variable exists; however the response is non-linear and the response declines consistently with any increase in stressor level observed up to point (c).
Careful consideration of change-points derived from such relationships is warranted. Such a changepoint would arguably not protect the designated use and could be considered, perhaps, an upper limit for protection. Reference expectations for the response measure might help interpret the position of such an endpoint. The changepoints described in this section all fall into this nature of change-point response. They exist at the lower end of the response curve, rather than at the upper end.

*Causal Model Confirmation*

We conducted two additional analyses in an attempt to better refine the causal basis for nutrient effects in UT streams. The first was a set of serial regressions and the second, an attempted propensity score analysis of existing UDWQ data.

*Serial regressions*

In this approach, we attempted to build a series of models (Fig. 14) along the presumptive causal pathway linking nutrients to macroinvertebrate response (Fig. 1).

**Model 1**

![Diagram](Figure 14 – Conceptual model of serial regressions linking nutrients and potential co-variates with macroinvertebrate response.)

There was, unfortunately, limited chlorophyll data available. Moreover, true chlorophyll responses are difficult to characterize in streams, given the many factors that affect algal growth. Algae tend to integrate over far shorter time frames than invertebrates or fish, therefore identifying an algal condition linked to invertebrate response is difficult. We found only weak relationships for model 1 relating nutrients to chlorophyll and none of them were significant except for negative decline on chlorophyll with ammonium concentrations. We also found no relationship between chlorophyll and dissolved
oxygen grab samples (model 2). It should be noted that grab samples are not particularly accurate measures of oxygen conditions in streams. Lastly, we found a peculiar decrease in scraper richness with dissolved oxygen (model 3). Again, dissolved oxygen grab sampled were likely not an accurate measure of oxygen conditions and may even increase with algal growth if measured in the afternoon. We, therefore, could not use the serial modeling results to confirm the causal model.

**Propensity score analysis**

We conducted a simple propensity score analysis as an attempt to strengthen the causal basis for linking nutrients to invertebrate response (Yuan 2010). Propensity score analysis is used to investigate the strength of specific stressor-response relationships based on observational data, given the presence of covariates (Rosenbaum 2002). Nutrients co-vary with a variety of other stressors (e.g., sediment, conductivity, BOD) which make estimating the single effects of nutrients challenging, yet it is important to recognize the potential contributions of these other stressors to the observed response. The simplest way to control for effects of covariates with nutrients would be to identify sites that vary in nutrient concentration, but are similar in regards to all other stressors. In some instances this may be possible, but as the number of covariates grows, it quickly becomes difficult to do. Propensity score analysis summarizes the effects of covariates into one single variable, called the propensity score (Fig. 15). Sites with similar propensity scores are similar in regards to their covariate distribution and the effects of the factor of interest can then be explored with bins of similar propensity scores (Rosenbaum and Rubin 1983, Rosenbaum 2002, Imai and Van Dyk 2004).
We calculated propensity scores for the Utah dataset by modeling the effects of covariates on nitrate concentration. We used the following predictors in a multiple linear regression model to predict nitrate concentration: elevation, longitude, latitude, watershed area, chloride, sulfate, bicarbonate, temperature, % agriculture, % urban, and turbidity. We split the resultant scores into 6 bins and investigated the response within each bin.

The MLR model explained 60% of the variance in nitrate concentration (Figure 16) and exhibited significant negative correlations with taxa richness, EPR richness, and positive correlations with HBI. When we explored the stressor-response relationships within bins, bin 3 exhibited the most significant decline in taxa richness and EPT richness to increasing nitrate concentration (Figs. 17 and 18). In contrast, HBI scores showed increases in almost every bin to nitrate concentration, especially bins 3 and 6, but also in 2 and 4 (Fig. 19).

Figure 15 - Conceptual diagram of propensity score analysis. Covariates with nutrient concentration are used to generate a predicted nutrient concentration (propensity score). Sites with similar propensity scores are grouped together into bins representing similar covariate distributions. The nutrient-response relationship is then investigated per bin.
Nitrate and Total Taxa Richness

Figure 16 – Response of taxa richness across different propensity score bins.
Figure 17 – Scatterplots of total nitrate vs. taxa richness within each of the 6 propensity score bins. Nitrate concentration increased from bin 1 to 6.
Nitrate and EPT Richness

Figure 18 - Scatterplots of total nitrate vs. EPT richness within each of the 6 propensity score bins. Nitrate concentration increased from bin 1 to 6.
We interpret these results, especially the HBI results, to indicate support for an effect of nutrients on invertebrates in streams in Utah, independent of other stressors. Since those stressors are controlled for in the propensity scores and yet nitrate still exhibited a significant relationship to these response metrics, support for a deleterious effect of nutrients on invertebrate assemblages is strengthened.

Summary

- We explored classification options for streams in Utah for the purposes of potentially developing different nutrient criteria for different parts of the state. This analysis did not identify potential classes worth pursuing at this time, therefore, criteria would be developed for statewide application.

- We conducted a distribution-based analysis of nutrient concentrations based on reference site and all site populations. Quartiles for these populations were generated and reported as potential endpoints consistent with EPA guidance.

- The next analysis was developing stressor-response relationship models based on existing biological thresholds and interpolating associated nutrient values. This approach generated a
series of nutrient endpoints associated with biological conditions considered thresholds for adverse aquatic life use conditions.

- Change-point analysis was then performed for a few stressor-response relationships that exhibited potential non-linear responses. Change-points were identified for these relationships and reported.

- Lastly, two analyses were performed to strengthen the support for the presumptive effect of nutrients on aquatic life use endpoints: serial regression models and propensity score analysis. The propensity score analysis strengthened the basis for an effect of nutrients on aquatic life use responses.

- These analyses are part of a larger effort by UDWQ to develop nutrient criteria in streams. Hopefully the analyses performed here will inform and strengthen that effort, providing both scientifically defensible and protective values for the continued protection of aquatic life in Utah.
Citations


Appendix 1 – Interpolated endpoints based on existing thresholds figures: annual data

The following figures are regressions between nutrient variables [(\(\log_{10}\) total nitrate (*NOXTOT), \(\log_{10}\) dissolved nitrate (*NOXDISS), and \(\log_{10}\) total phosphorus (*PTOT)]. Response variable abbreviations are as in Table 4. Regression equations, significance, and coefficients of determination are shown. Lines are the mean regression (solid) and the 90% prediction intervals. Only significant models are shown.
Taxa Richness

\[ y = 16.3354 - 6.6309x; \quad r = -0.3611, p = 0.00001; r^2 = 0.1304 \]

\[ y = 14.0863 - 4.4136x; \quad r = -0.2744, p = 0.0015; r^2 = 0.0753 \]

\[ y = 15.6192 - 6.9516x; \quad r = -0.4292, p = 0.0001; r^2 = 0.1842 \]

\[ y = 16.3354 - 6.6309x; \quad r = -0.3611, p = 0.00001; r^2 = 0.1304 \]
EPT Richness

*NOXTOT:EPT:  \( y = 3.7312 - 4.2079x; \)  
\( r = -0.4966, p = 0.00001; r^2 = 0.2466 \)

*NOXDISS:EPT:  \( y = 3.9463 - 4.3979x; \)  
\( r = -0.3978, p = 0.00000; r^2 = 0.1583 \)

*PTOT:EPT:  \( y = 2.8315 - 2.4983x; \)  
\( r = -0.2731, p = 0.0016; r^2 = 0.0746 \)
E Richness

\[
y = 1.0347 - 1.2186x; \\
r = -0.4450, p = 0.00006; r^2 = 0.1980
\]

\[
y = 1.0947 - 1.2491x; \\
r = -0.3350, p = 0.00006; r^2 = 0.1122
\]

\[
y = 0.8282 - 0.6983x; \\
r = -0.2313, p = 0.0079; r^2 = 0.0535
\]
Percent EPT Individuals

- NOXTOT: $y = 13.217 - 13.3194x; r = -0.3677, p = 0.0011; r^2 = 0.1352$

- NOXDISS: $y = 15.5701 - 11.1371x; r = -0.2107, p = 0.0128; r^2 = 0.0444$

- PTOT: $y = 11.2812 - 6.9404x; r = -0.1795, p = 0.0403; r^2 = 0.0322$
Scraper Richness

- For NOXTOT:SCR, the relationship is $y = 1.1206 - 1.2657x$ with $r = -0.4342$, $p = 0.00009$, and $r^2 = 0.1885$.

- For NOXDISS:SCR, the relationship is $y = 1.1641 - 1.2276x$ with $r = -0.3324$, $p = 0.00006$, and $r^2 = 0.1105$.

- For PTOT:SCR, the relationship is $y = 0.7761 - 0.7599x$ with $r = -0.2473$, $p = 0.0044$, and $r^2 = 0.0611$. 
HBI

\[ y = 4.8956 + 1.3878x; \quad r = 0.4579, \quad p = 0.00003; \quad r^2 = 0.2097 \]

\[ y = 4.7726 + 1.463x; \quad r = 0.3746, \quad p = 0.00001; \quad r^2 = 0.1403 \]

\[ y = 5.1159 + 0.769x; \quad r = 0.2543, \quad p = 0.0034; \quad r^2 = 0.0646 \]
OE Model

*NOXTOT:RFOE:  \( y = 0.5438 - 0.2032x; \)
\( r = -0.3386, p = 0.0028; r^2 = 0.1147 \)
Appendix 2 – Interpolated endpoints based on existing thresholds figures: summer data

The following figures are regressions between summer nutrient variables [($\log_{10}$ total nitrate (Su*NOXTOT), $\log_{10}$ dissolved nitrate (Su*NOXDISS), and $\log_{10}$ total phosphorus (Su*PTOT)]. Response variable abbreviations are as in Table 4. Regression equations, significance, and coefficients of determination are shown. Lines are the mean regression (solid) and the 90% prediction intervals. Only significant models are shown.
Taxa Richness

Scatterplot of Taxa against Su*NOXTOT
UTBioChemWatershedAnnSummCleaned.sta 83v’203c
Taxa = 16.6448 - 4.9054*x; 0.9 Pred.Int.

Scatterplot of Taxa against Su*NOXDISS
UTBioChemWatershedAnnSummCleaned.sta 83v’203c
Taxa = 16.3934 - 6.679*x; 0.9 Pred.Int.
EPT Richness

Scatterplot of EPT against Su*NOXTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
EPT = 4.1354 - 3.4255*x; 0.9 Pred.Int.

Su*NOXTOT:EPT: y = 4.1354 - 3.4255*x; r = -0.4316, p = 0.0004; r^2 = 0.1863

Scatterplot of EPT against Su*NOXDISS
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
EPT = 4.2437 - 3.6231*x; 0.9 Pred.Int.

Su*NOXDISS:EPT: y = 4.2437 - 3.6231*x; r = -0.3633, p = 0.0002; r^2 = 0.1320

Scatterplot of EPT against Su*PTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
EPT = 3.8909 - 1.464*x; 0.9 Pred.Int.

Su*PTOT:EPT: y = 3.8909 - 1.464*x; r = -0.1828, p = 0.0548; r^2 = 0.0334
Scatterplot of SCR against Su*NOXTOT

UTBioChemWatershedAnnSummCleaned.sta 83v*203c

SCR = 1.2062 - 1.1472*x; 0.9 Pred.Int.

Scatterplot of SCR against Su*NOXDISS

UTBioChemWatershedAnnSummCleaned.sta 83v*203c

SCR = 1.276 - 0.9814*x; 0.9 Pred.Int.

Scatterplot of SCR against Su*PTOT

UTBioChemWatershedAnnSummCleaned.sta 83v*203c

SCR = 0.914 - 0.6324*x; 0.9 Pred.Int.

Scraper Richness
Scaper+Collector/Gatherer Richness

Scatterplot of SCRCG against Su*NOXDISS
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
SCRCG = 2.6655-1.2671*x; 0.9 Pred.Int.
Su*NOXDISS:SCRCG:   y = 2.6655 - 1.2671*x;
   r = -0.3099, p = 0.0018; r^2 = 0.0960

Scatterplot of SCRCG against Su*PTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
SCRCG = 2.2644-0.6732*x; 0.9 Pred.Int.
Su*PTOT:SCRCG:   y = 2.2644 - 0.6732*x;
   r = -0.1988, p = 0.0365; r^2 = 0.0395

Scatterplot of SCRCG against Su*NOXTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c
SCRCG = 2.5984-1.3142*x; 0.9 Pred.Int.
Su*NOXTOT:SCRCG:   y = 2.5984 - 1.3142*x;
   r = -0.3954, p = 0.0012; r^2 = 0.1564
HBI

Scatterplot of HBI against Su*NOXTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c

HBI = 4.8234+1.3667*x; 0.9 Pred.Int.

Scatterplot of HBI against Su*NOXDISS
UTBioChemWatershedAnnSummCleaned.sta 83v*203c

HBI = 4.741+1.3627*x; 0.9 Pred.Int.

Scatterplot of HBI against Su*PTOT
UTBioChemWatershedAnnSummCleaned.sta 83v*203c

HBI = 5.1419+0.7721*x; 0.9 Pred.Int.
O/E Model

Scatterplot of RFOE against Su\(^*\)NOXTOT

UTBioChemWatershedAnnSummCleaned.sta 83v*203c

RFOE = 0.5483 - 0.1657\(x\); 0.9 Pred. Int.

Su\(^*\)NOXTOT:RFOE: \(y = 0.5483 - 0.1657\(x\); \(r = -0.2958, p = 0.0176; r^2 = 0.0875\)

Scatterplot of RFOE against Su\(^*\)NOXDISS

UTBioChemWatershedAnnSummCleaned.sta 83v*203c

RFOE = 0.5922 - 0.1633\(x\); 0.9 Pred. Int.

Su\(^*\)NOXDISS:RFOE: \(y = 0.5922 - 0.1633\(x\); \(r = -0.2390, p = 0.0172; r^2 = 0.0571\)
Appendix 3 - Endpoints based on change-point analyses: annual data

The following figures are scatterplots indicating non-linear relationships between annual nutrient variables [(Log_{10} total nitrate (Su*NOXTOT), Log_{10} dissolved nitrate (Su*NOXDISS)) and biological responses. Response variable abbreviations are as in Table 4 and regression plots are as in Appendix 1. Non-linear LOWESS fits were added as curvilinear solid lines to visualize non-linearities (tension = 0.8). Also shown are results of change-point analysis indicating percent reduction in model error and a visual tree plot with change-point in nutrient concentration indicated in log units (back-transformed units).
HBI = 4.7726 + 1.463x; 0.9 Pred. Int.  
HBI = Lowess

$*NOXDISS: HBI: y = 4.7726 + 1.463x; r = 0.3746, p = 0.00001$

153 cases deleted due to missing data.
Split Variable PRE Improvement
1 VNOXDISS 0.314 0.314
Fitting Method: Least Squares
Predicted variable: HBI
Minimum split index value: 0.050
Minimum improvement in PRE: 0.050
Maximum number of nodes allowed: 4
Minimum count allowed in each node: 5
The final tree contains 2 terminal nodes
Proportional reduction in error: 0.314

Node from Count Mean SD Split Var Cut Value Fit
1 0 50 4.533 0.883 VNOXDISS -0.541 0.314
2 1 18 3.880 0.722
3 1 32 4.901 0.748

VNOXDISS < -0.541 (0.288 mg/L)
RFOE = 0.5438 - 0.2032*x; 0.9 Pred.Int.
RFOE = Lowess

*NOXTOT

153 cases deleted due to missing data.
Split Variable PRE Improvement
1 VNOXTOT 0.254 0.254
Fitting Method: Least Squares
Predicted variable: RFOE
Minimum split index value: 0.050
Minimum improvement in PRE: 0.050
Maximum number of nodes allowed: 4
Minimum count allowed in each node: 5
The final tree contains 2 terminal nodes
Proportional reduction in error: 0.254

Node from Count Mean SD Split Var Cut Value Fit
1 0 50 0.599 0.215 VNOXTOT -0.667 0.254
2 1 13 0.779 0.221
3 1 37 0.535 0.175

(0.215 mg/L)VNOXTOT<-.667
**Appendix 4 - Endpoints based on change-point analyses: summer data**

The following figures are scatterplots indicating non-linear relationships between summer nutrient variables \([\log_{10} \text{total nitrate (Su*NOXTOT)}\) and \(\log_{10} \text{total phosphorus (Su*PTOT)}\]) and biological responses. Response variable abbreviations are as in Table 4 and regression plots are as in Appendix 1. Non-linear LOWESS fits were added as curvilinear solid lines to visualize non-linearities (tension = 0.8). Also shown are results of change-point analysis indicating percent reduction in model error and a visual tree plot with change-point in nutrient concentration indicated in log units (back-transformed units).
Scatterplot of RFOE against Su*NOXTOT

UTBioChemWatershedAnnSummCleaned.sta 84v*203c

$RFOE = 0.5483 - 0.1657x; \ 0.9 \ Pred.\ Int.$

$RFOE = \text{Lowess}$

165 cases deleted due to missing data.

Split Variable PRE Improvement
1 SUNOXTOT 0.323 0.323

Fitting Method: Least Squares
Predicted variable: RFOE
Minimum split index value: 0.050
Minimum improvement in PRE: 0.050
Maximum number of nodes allowed: 22
Minimum count allowed in each node: 5
The final tree contains 2 terminal nodes
Proportional reduction in error: 0.323

Node from Count Mean SD Split Var Cut Value Fit
1 0 38 0.601 0.222 SUNOXTOT -0.638 0.323
2 1 8 0.843 0.257
3 1 30 0.537 0.164

SUNOXTOT<0.638 (0.230 mg/L)
Scatterplot of SCR against Su*PTOT
UTBioChemWatershedAnnSummCleaned.sta 84v*203c
SCR = 0.914-0.6324*x; 0.9 Pred.Int.
SCR = Lowess

-1.8 -1.6 -1.4 -1.2 -1.0 -0.8 -0.6 -0.4 -0.2 0.0 0.2 0.4
Su*PTOT
-1 0 1 2 3 4 5 6 7
SCR

167 cases deleted due to missing data.
Split Variable PRE Improvement
1 SUPTOT 0.201 0.201
Fitting Method: Least Squares
Predicted variable: SCR
Minimum split index value: 0.050
Minimum improvement in PRE: 0.050
Maximum number of nodes allowed: 4
Minimum count allowed in each node: 5
The final tree contains 2 terminal nodes
Proportional reduction in error: 0.201
Node from Count Mean SD Split Var Cut Value Fit
1 0 36 1.526 0.893 SUPTOT -1.381 0.201
2 1 5 2.507 0.583
3 1 31 1.367 0.836

SUPTOT< -1.381 (0.042 mg/L)
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Scatterplot of SCRCG against Su*PTOT

UTBioChemWatershedAnnSummCleaned.sta 84v*203c

SCRCG = 2.2644-0.6732*x; 0.9 Pred.Int.

SCRCG = Lowess

167 cases deleted due to missing data.

Split Variable PRE Improvement
1 SUPTOT 0.103 0.103

Fitting Method: Least Squares

Predicted variable: SCRCG

Minimum split index value: 0.050
Minimum improvement in PRE: 0.050
Maximum number of nodes allowed: 4
Minimum count allowed in each node: 5

The final tree contains 2 terminal nodes

Proportional reduction in error: 0.103

Node from Count Mean SD Split Var Cut Value Fit
1 0 36 3.021 1.143 SUPTOT -1.381 0.103
2 1 5 3.921 0.942
3 1 31 2.876 1.118