Step Process for Selecting and Testing Surrogates and Indicators of Afrotemperate Forest Invertebrate Diversity

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Abstract

Background: The diversity and complexity of invertebrate communities usually result in their exclusion from conservation activities. Here we provide a step process for assessing predominantly ground-dwelling Afrotemperate forest invertebrates' (earthworms, centipedes, millipedes, ants, molluscs) potential as surrogates for conservation and indicators for monitoring. We also evaluated sampling methods (soil and litter samples, pitfall traps, active searching quadrats and tree beating) and temporal (seasonal) effects.

Methodology/Principal Findings: Lack of congruence of species richness across taxa indicated poor surrogacy potential for any of the focus taxa. Based on abundance and richness, seasonal stability, and ease of sampling, molluscs were the most appropriate taxon for use in monitoring of disturbance impacts. Mollusc richness was highest in March (Antipodal late summer wet season). The most effective and efficient methods were active searching quadrats and searching litter samples. We tested the effectiveness of molluscs as indicators for monitoring by contrasting species richness and community structure in burned relative to unburned forests. Both species richness and community structure changed significantly with burning. Some mollusc species (e.g., Macroptychia africana) showed marked negative responses to burning, and these species have potential for use as indicators.

Conclusions/Significance: Despite habitat type (i.e., Afrotemperate forest) being constant, species richness and community structure varied across forest patches. Therefore, in conservation planning, setting targets for coarse filter features (e.g., habitat type) requires fine filter features (e.g., localities for individual species). This is especially true for limited mobility taxa such as those studied here. Molluscs have high potential for indicators for monitoring, and this requires broader study.

Introduction

A systematic approach to conservation includes both the measurement of biodiversity features for prioritizing areas, and adaptive management of these, including monitoring the impacts of management or disturbance [1]. Because of limited resources and potential impact on conservation assets, both inventories and monitoring need to be as effective and efficient as possible. It is impossible to sample and identify every species, even in small areas, and this is especially true for the hyperdiverse invertebrates [2]. Invertebrates may be important in terms of their relatively high levels of endemicity [3,4], and their responsiveness to environmental change [5] makes them potential indicators for monitoring [e.g., 6,7], and for conservation planning (for overview see [8]). Invertebrates are especially poorly represented in conservation activities, largely because of their enormous abundance and diversity, and the lack of appropriate information for many taxa [9,10].

Protocols for monitoring biodiversity are not well established for terrestrial ecosystems [11], and more research is required on indicators [12]. Indicator taxa selected for monitoring must reflect environmental change and the reaction of other taxa [11,13]. Most importantly, because of differential sensitivity of taxa to environmental disturbance, empirical studies are necessary to verify the appropriateness of a particular taxon as an indicator of disturbance [14]. A stepwise, integrated, approach is necessary to properly identify biodiversity surrogates or indicators [6,15]. The following steps are recommended: (1) a survey using standardised, quantified effort [2,16]; (2) an assessment of potential taxa for use in conservation planning or monitoring [6 and references therein,15]; and (3) a test of the selected indicator taxon for monitoring a particular disturbance [6,15]). Rohr et al. [15] add in the additional dimension of testing which sampling method is most appropriate for a particular chosen taxon.

The target taxa for this study, predominantly ground-dwelling, flightless invertebrates, are potentially important for biodiversity assessment and monitoring. Invertebrates such as molluscs, earthworms, centipedes, millipedes, and onychophorans may be suitable surrogate taxa for biodiversity assessments because they have limited dispersal ability and consequently they may exhibit
high levels of endemism [4,17–22], and ants have been widely recommended as surrogates and indicators [23–26]. These taxa have relatively well known taxonomy, and are easily observed, and they may also be suitable for monitoring disturbance because they do not have complex life cycles (except for ants), adults are relatively long-lived compared to most insects [7], and because they have limited mobility they are less likely to escape and colonise other habitats after disturbance.

The aim of this study was to assess the potential for these taxa to serve as surrogates for biodiversity assessment, and as indicators for monitoring disturbance. We undertook this study in Afrotemperate forest [27] in the KwaZulu-Natal Drakensberg, South Africa, where forest patches have been small and fragmented since the last glacial maximum (18 000 y.b.p.), and have expanded and contracted prior to that [see 28 for forest history]. The objectives were: (1) to compare species richness and community structure across seasons to identify the most suitable time of year for diversity assessment and monitoring of the target taxa, and to identify taxa that do not show marked seasonal changes in diversity; (2) to determine which sampling methods used to determine species richness were most effective and efficient for use in biodiversity assessment and monitoring for the target taxa, in different months; (3) to determine which flightless invertebrate taxa were most suitable for use in biodiversity assessment (surrogates) and monitoring (indicators) using the approach and criteria of Summerville et al. [6], but including the additional component of endemism; and (4) to experimentally test these recommendations for use in monitoring, using burned and unburned Afrotemperate forest patches at a different study site.

Note that fire in Afrotemperate forests is becoming an increasing conservation concern [e.g. 29]. The study also illustrates the process that should be undertaken to evaluate the taxa selected for a large scale survey or a monitoring programme, before such activities are implemented on a large scale.

Results

We collected 4273 individual specimens representing 55 species in the four months from the three sites at Injisuthi (Table 1). The 55 species comprised 26 mollusc, four earthworm, one onychophoran, six centipede, 11 millipede and seven ant species. Because only a single onychophoran species was collected, this group was excluded from further analyses.

Seasonal Changes in Richness and Community Structure

Of the 55 species, 22 (40%) were recorded in all four seasons, 11 in three of the four seasons, seven in two of the four seasons and 11 species (20%) were collected in one season only (Table 1). Nineteen species (35%) were collected only in the two wetter (and warm) months (December and March) and no species were unique to the dry season (June and September) (Table 1), which suggests that the flightless invertebrate community in winter is merely a subset of the summer wet season community. Total species richness, mean species richness, and unique species richness for all taxa was lower in the cool, dry season (June and September) compared to the warm, wet season (March and December) (Fig. 1A). When assessing taxa separately, species richness of molluscs, centipedes, and millipedes was slightly higher in wet season months (March and December) than dry season months, but only by a couple of species (Fig. 1B). Mollusc species richness ranged across the four seasons from 14 to 18, 12 to 15, and 15 to 17 species respectively for the three different forests. Millipede richness ranged from 2 to 8, 6 to 7, and 5 to 9 respectively for the three forests across the seasons.

Community structure was significantly different between March and June (ANOSIM: R = 0.852, March and September (R = 0.815), March and December (R = 0.770), June and December (R = 0.778), and September and December (R = 0.963) (Fig. 2A). However, the community did not differ between the two dry season months (June and September) (R = 0.270). There was a significant difference in mollusc community structure between autumn (March) and winter (June) (R = 0.809), autumn (March) and spring (September) (R = 0.796), and winter (June) and summer (December) (R = 0.944), but the community differed less between autumn (March) and summer (December) (R = 0.519) (Fig. 2B). Mollusc community structure was similar between the two dry season months (June and September) (R = 0.074).

Centipedes showed distinct temporal turnover in species composition within the wet season (March and December, R = 0.899), and from spring (September to summer (December) (R = 1.000) and winter (June) to spring (September) (R = 0.556) (Fig. 2C). For millipedes (Fig. 2D), the only strong separation between seasons was between winter (September) and summer (December) (R = 0.907). Ant species composition was strongly separated between spring (September) and summer (December) (R = 0.796), but temporal turnover was also evident between autumn (March) and summer (December) (R = 0.685) (Fig. 2E).

Sampling Method Efficiency and Effectiveness

The five different sampling methods used contributed unequally to species richness (Fig. 3B). Tree beats were important for collecting live snails and ants, but did not target any other taxa. Pitfall traps and soil samples performed poorly. Active search quadrats and leaf litter samples were the sampling methods that collected the greatest number of species collected by one method only in each month (Fig. 3A), and far outperformed the other three sampling methods in terms of number of species collected. The mean efficiency (calculated as species per person hour) for soil samples was 1.0, pitfall traps was 0.2, litter samples was 2.6, active search quadrats was 2.6 and tree beats was 1.4 species per person hour respectively.

Assessment of Taxa as Biodiversity Surrogates and Indicators

When assessing the potential of the different taxa as biodiversity surrogates and indicators of disturbance, molluscs scored highest followed by millipedes (Table 2). Centipedes scored lowest, with several of the categories indicating problems with the use of this taxon (Table 2). However, none of the taxa proved to be good surrogates for the underlying diversity (all taxa excluding the target taxon) in terms of species richness, with all relationships being non-significant (Linear Regression: molluscs: F1,R = 0.983, P = 0.351; earthworms: F1,R = 0.008, P = 0.933; centipedes: F1,R = 0.521, P = 0.616; millipedes: F1,R = 0.778, P = 0.404; and ants: F1,R = 0.003, P = 0.961).

Assessment of Molluscs as Indicators of Disturbance

Since molluscs best met Summerville et al. [6]'s criteria, we tested their ability to reflect disturbance (fire history), and to act as surrogates for the responses of other taxa to this disturbance, in forest patches at Royal Natal National Park.

The species richness for molluscs was significantly lower in burned than unburned forest patches (ANOVA: F1,6 = 12.73; P = 0.012) (Fig. 4). However, species richness of all non-molluscs was not significantly different between the burn treatments (F1,6 = 0.179, P = 0.687). The species richness of millipedes was marginally non-significantly different across the treatments (F1,6 = 0.561, P = 0.055).
Table 1. Mollusc, earthworm, onychophoran, centipede, millipede and ant species collected during seasonal sampling at Injisuthi (abundance data).

| Order               | Family               | Species                                      | M   | J   | S   | D   |
|---------------------|----------------------|----------------------------------------------|-----|-----|-----|-----|
| **Class Gastropoda**|                      |                                              |     |     |     |     |
| Neritopsina         | Hydroceniidae        | Hydrocena naticola Benson, 1856              | 169 | 168 | 334 | 169 |
| Architaienioglossa  | Cyclophoridae        | *Chondrotrochus nipponensis* (Shinnyu, 1898) | 25  | 54  | 29  | 34  |
| Eupulmonata         | Pupillidae           | *Lauria adamin* (Benson, 1864)               | 12  | 9   | 5   | 7   |
| Eupulmonata         | Orculidae            | *Fauixus glauvileanus* (darglemis) (Ancey, 1888) | 23  | 36  | 88  | 50  |
| Eupulmonata         | Orculidae            | *Fauixus mcebeanianus* Melville and Ponsonby, 1901 | 15  | 17  | 25  | 56  |
| Eupulmonata         | Orculidae            | Fauixus sp.                                  |   5 | 0   | 0   |     |
| Eupulmonata         | Vertiginidae         | Pupisoma harpula (Reinhardt, 1886)           | 5   | 0   | 0   | 14  |
| Eupulmonata         | Vertiginidae         | *Lauria adamin* (Benson, 1864)               | 10  | 11  | 0   | 6   |
| Eupulmonata         | Achatinidae          | Archacatina sp.                              |     |     |     |     |
| Eupulmonata         | Charopidae           | Afrodonta novellamellaris (Burnup, 1912)     | 11  | 10  | 0   | 6   |
| Eupulmonata         | Charopidae           | *Trachycystis contabulata* Connolly, 1932    | 11  | 65  | 46  | 33  |
| Eupulmonata         | Charopidae           | *Trachycystis ceptima* (Melville and Ponsonby, 1899) | 0   | 13  | 14  | 25  |
| Eupulmonata         | Charopidae           | *Trachycystis glauvileana* (Ancey, 1893)     | 0   | 2   | 3   | 1   |
| Eupulmonata         | Charopidae           | Trachycystis ricadostata Connolly, 1923      | 94  | 91  | 51  | 31  |
| Eupulmonata         | Charopidae           | *Trachycystis subpinguis* Connolly, 1922     | 0   | 0   | 0   | 14  |
| Eupulmonata         | Charopidae           | *Trachycystis venatum* Connolly, 1932        | 23  | 0   | 0   | 0   |
| Eupulmonata         | Helicariidae         | *Kaliella euconuloides* Melville and Ponsonby, 1908 | 26  | 8   | 29  | 117 |
| Eupulmonata         | Euconulidae          | Afroconulus diaphanous (Connolly, 1922)      | 10  | 5   | 7   | 4   |
| Eupulmonata         | Achatinidae          | *Archacatina dimidia* (Smith, 1878)          | 0   | 2   | 1   | 1   |
| Eupulmonata         | Vertiginidae         | Pupisoma orcula (Benson, 1850)               | 14  | 63  | 86  | 0   |
| Eupulmonata         | Pupillidae           | *Pupilla fontana* (Krauss, 1848)             | 3   | 0   | 0   | 0   |
| Eupulmonata         | Charopidae           | Trachycystis sp.                             | 1   | 0   | 0   | 0   |
| Eupulmonata         | Chlamydephoridae     | *Chlamydephorus burnupi* Smith, 1892         | 2   | 0   | 0   | 0   |
| **Class Oligochaeta**|                      |                                              |     |     |     |     |
| Haplotaxida         | Acanthodrilidae      | Dichagaster sp.                              | 0   | 0   | 0   | 4   |
| Haplotaxida         | Acanthodrilidae      | Parachilota sp. 1                            | 0   | 0   | 0   | 29  |
| Haplotaxida         | Acanthodrilidae      | Parachilota sp. 2                            | 3   | 0   | 0   | 1   |
| Opisthodora         | Microchaetidae       | Proandricus sp.                              | 0   | 0   | 0   | 3   |
| **Class Onychophora**|                      |                                              |     |     |     |     |
| Onychophora         | Onychophora          | *Opisthatus cinctipes* Purcell, 1899         | 1   | 0   | 0   | 0   |
| **Class Chilopoda** |                      |                                              |     |     |     |     |
| Geophilomorpha      | Geophilidae          | *Rhyida afr* (afr)                           | 0   | 0   | 0   | 11  |
| Geophilomorpha      | Geophilidae          | sp. 2                                        | 0   | 0   | 0   | 36  |
| Geophilomorpha      | Geophilidae          | sp. 1                                        | 8   | 7   | 1   | 20  |
| Lithobiomorpha      | Henicopidae          | *Paralamyctes spencer*                       | 4   | 5   | 3   | 1   |
| Lithobiomorpha      | Henicopidae          | Lamyctes africana                            | 2   | 0   | 0   | 92  |
| Lithobiomorpha      | Henicopidae          | Lamyctes sp.                                 | 3   | 0   | 0   | 1   |
| **Class Diplopoda** |                      |                                              |     |     |     |     |
| Sphaerotheriida     | Sphaerotheriida      | Sphaerotherium dorsale* (Gervais, 1847)      | 2   | 3   | 63  | 23  |
| Sphaerotheriida     | Sphaerotheriida      | **Sphaerotherium mahoum Schubart, 1958       | 75  | 27  | 26  | 180 |
| Sphaerotheriida     | Sphaerotheriida      | Sphaerotherium sp.                           | 0   | 0   | 12  | 19  |
| Polydesmida         | Dalodesmida          | Gnomeskelus atmensus Verhoeff, 1939          | 1   | 1   | 0   | 23  |
| Polydesmida         | Dalodesmida          | Gnomeskelus montivagus Schubart, 1939        | 1   | 6   | 2   | 27  |
| Polydesmida         | Dalodesmida          | Gnomeskelus sp.                              | 27  | 5   | 7   | 74  |
| Polydesmida         | Gomphodesmida        | *Ulodesmus simplex* Lawrence, 1953           | 4   | 7   | 18  | 32  |
| Spirostreptida      | Odontopygida         | Spinotarsus sp. 2                            | 1   | 0   | 16  | 1   |
Therefore, in terms of species richness, molluscs were the most effective indicator of fire history.

Seven mollusc species had lower abundance in the burned forests, six species were found only in unburned forest, and two only in burned forests (Table 3). In contrast, no millipedes were found only in unburned forest, and three millipedes were found only in burned forest. A micromollusc, *Hydrocena noticola*, was the only species found in all four unburned forests, and in none of the burned forests. Of the larger species (shell length \( \geq 8 \) mm), *Macroptychia africana*, which was found in three of the four unburned forests and none of the burned forests, may be the most appropriate species for monitoring disturbance. Other potential candidates were *Nata vernicosa* and *Trachycystis subpinguis*.

The mollusc community at the unburned sites clustered out distinctly from those at the burned sites (ANOSIM: \( R = 0.719 \)) (Fig. 5B). While the non-mollusc community (earthworms, centipedes, millipedes and ants) also clustered out distinctly in terms of sites with different fire histories (\( R = 0.875 \)) (Fig. 5A), the clustering was a lot less distinct than that for molluscs (excluding site 8, which although clustering separately from the unburned sites was not closely clustered with the burned sites) (Fig. 5A). In addition, there was a lack of concordance in the communities of molluscs at burned sites relative to those at unburned sites (PROTEST: \( m^2 = 0.431, P = 0.685 \)). Therefore, mollusc community structure itself may serve as a strong indicator of disturbance, in this case fire history.

### Discussion

We provide here an additional example of the stepwise practice for selecting surrogates and indicators \([6,15]\). An important component of this process is independent testing of the indicator for monitoring, and because our test of this involved a discrete and easily identifiable disturbance event, the results should be relatively robust. Such testing follows the recommendations of Pocock and Jennings \([14]\), who highlighted that variable sensitivity of taxa to different kinds of disturbance makes them more or less effective as indicators. Note that, as in our study, even within the same higher-level taxon, species may respond differently to a particular disturbance, and each species needs to be tested \([14]\).

An important additional component that we have added to the selection process of Rohr *et al.* \([15]\), is evaluating temporal (seasonal) effects. Some taxa will be more or less abundant through the year, as they are more sensitive to temperature or moisture changes, or have different life stages that are only present in certain seasons. We highlight that, as part of the initial survey step, sampling should be conducted at different times of the year to identify (1) the best time of the year to sample, and (2) the taxa

![Figure 1](https://example.com/figure1.png)

**Figure 1.** The influence of season on species richness to determine suitability for use in biodiversity assessment and monitoring. (A) All taxa combined: total species richness (triangles), mean species richness ± one standard deviation (solid squares) and unique species (open squares); and (B) taxa separately. Data are for three forests combined.

![Figure 2](https://example.com/figure2.png)

![Figure 3](https://example.com/figure3.png)

![Figure 4](https://example.com/figure4.png)

**Table 1.** Cont.

| Order            | Family             | Species               | M | J | S | D |
|------------------|--------------------|-----------------------|---|---|---|---|
| Spirostreptida   | Odontopygidae      | *Spinotarsus* sp. 1   | 1 | 1 | 0 | 3 |
| Spirostreptida   | Spirostreptidae    | **Doratogonus montanus** Hamer, 2000 | 2 | 2 | 0 | 3 |

| Class Insecta    |                     |                       |   |   |   |   |
|------------------|----------------------|-----------------------|---|---|---|---|
| Hymenoptera      | Formicidae sp. 1     | 3                     | 0 | 1 | 0 |
| Hymenoptera      | Formicidae sp. 2     | 9                     | 0 | 2 | 520 |
| Hymenoptera      | Formicidae sp. 3     | 20                    | 0 | 18 | 18 |
| Hymenoptera      | Formicidae sp. 4     | 5                     | 0 | 0 | 6 |
| Hymenoptera      | Formicidae sp. 5     | 7                     | 12 | 17 | 16 |
| Hymenoptera      | Formicidae sp. 6     | 8                     | 1  | 14 | 0 |
| Hymenoptera      | Formicidae sp. 7     | 16                    | 1  | 4  | 0 |

\( M = \) March (autumn), \( J = \) June (winter), \( S = \) September (spring), and \( D = \) December (summer). Bold species are those that were sampled in a single season only.

\* = endemic to South Africa; ** = endemic to the Drakensberg region.

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least likely to vary with season. This is important both in carrying out biodiversity assessments or inventories, and for monitoring. Mollusc species richness was remarkably similar across seasons, although, as with other taxa, there were significant changes in community structure. The stability of the mollusc richness may be explained by the persistence and presence of dead shells, even if the live animals are not active. Any use of community assemblages should therefore control for season, and, in general, March (autumn/late summer, wet season) was the best time for sampling across taxa. Note that we only present data for a single year, and diversity may be more variable over multiple years (e.g. carabid beetles [30]). We recommend longer term survey work to assess temporal stability from year to year.

None of our taxonomic groups were good predictors of the balance of diversity, and a similar lack of congruency has been found by others [16,31–39], but see, e.g. [40]. This is despite the fact that our study focussed in one vegetation type Afrotemperate forest [27], and in a limited geographic region, where we would not expect biogeographic factors to confound relationships [see 39]. Note that local biogeographic processes such as forest area and within-valley isolation were not important drivers of our community assemblages [41].

We highlight in this work the use of community structure of a particular taxon being used as a surrogate of diversity changes in response to disturbance, rather than simply species richness [8]. Given advances in statistical analyses using ordination techniques [15], robust interpretations of assemblage changes could prove valuable. This is particularly so for the hyperdiverse invertebrates, where a single species, or species richness per se, may not be appropriate, especially for surrogates in conservation planning.

While several studies have investigated invertebrate sampling methods [e.g. 15,42–44], there are still no generally accepted, standardized sampling methods or protocols for different invertebrate taxa. The effectiveness of different sampling methods may also vary temporally depending on the activity patterns of the target taxa [45]. By focussing our monitoring on a particular indicator taxon, we can also focus our sampling effort. We consider three major aspects: (1) effectiveness at sampling the target taxon; (2) ease of implementation by managers in relatively remote areas; and (3) impact on the [protected] fauna of a reserve that is repeatedly sampled for monitoring. Considering these three aspects, we recommend for molluscs a combination of quantified litter sampling and timed active searching in restricted quadrats. This combination was effective, can be implemented in one short visit, and allows the release alive of repeatedly sampled species. Passive techniques, such as pitfall traps, should be critically assessed [see also 2] when considering monitoring programmes as they: (1) are not as effective as active searching techniques [e.g. 43]; (2) require managers to transport more equipment to remote sites, and require repeat visits to collect samples; and (3) kill (potentially large numbers of) both the target taxon and a large bycatch unnecessarily [see also 46], which can compound with repeated sampling; a problem when sampling in protected areas conserving threatened species, especially in small, patchy habitats such as the

Figure 2. Influence of season on community structure of invertebrates. (A) all taxa; (B) molluscs; (C) centipedes; (D) millipedes; and (E) ants illustrated using multidimensional scaling (MDS). Letters indicate sampling season (M = March (autumn), J = June (winter), S = September (spring) and D = December (summer), and numbers the site (forest).

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Drakensberg Afrotemperate forests. When used for monitoring purposes, it may be necessary to differentiate freshly dead from very old shells, to ensure that the current community is being sampled (we do not know the decomposition rates of shells in our habitat, or the effect of fire on old shells).

Besides the other attributes favouring molluscs as surrogates or indicators, they have high inherent conservation value, with high local species richness, and high levels of endemism, with relatively narrow distribution ranges [e.g. 47 and references therein, 48]. This would make snails particularly advantageous as fine filter features in conservation planning. Millipedes, the second most appropriate taxon, similarly have high inherent conservation value [4].

Ants have typically been used as indicators for a wide range of aspects [reviewed in 49]. In our analysis they were not as effective as surrogates or indicators relative to snails or millipedes. One reason for this may be the relatively low number of species sampled. However, they may not be suitable within this particular habitat, or within the spatial scale of the study.

Our study habitat type, i.e. Afrotemperate forest, was constant, but our results indicated wide variation in species richness across forest patches, as well as shifts in community structure (for another such example with snails see [48]). This means that setting targets and selecting habitats for management and conservation based purely on vegetation would be likely to miss potentially important species or communities. Under these circumstances, using identified species of the taxa included in this study will improve the fine-scale selection and prioritization of forest patches. We recommend that both this aspect of invertebrate conservation, and the use of molluscs as indicators of disturbance, be evaluated more broadly.

Our results also emphasise the effect of fire within these Afrotemperate patches on invertebrate communities, and careful attention needs to be paid to management of fire within these systems [see also 29].

**Methods**

The Maloti-Drakensberg Bioregion experiences summer rainfall, with 70% of the annual precipitation in the austral summer (November to March) [50]. Median rainfall values ranging across the Bioregion in the months that seasonal sampling took place are as follows: March, 100–140 mm; June, 15–50 mm; September, 20–60 mm; and December, 120–160 mm. Mean annual temperature in

Table 2. Potential of the different taxa as biodiversity surrogates and indicators of disturbance evaluated according to the criteria and scale provided by Summerville et al. [6], with endemism added.

| Taxon      | Diverse fauna (in forests)a | Well known taxonomyb | Easy to identifyc | Well known natural historyd | Readily surveyede | High ecological fidelity (forests)f | Endemismg | Total scoreh |
|------------|-----------------------------|----------------------|-------------------|-----------------------------|------------------|-----------------------------------|-----------|---------------|
| Ants       | ++                          | +++                  | +                 | +++                         | +++              | +                                 | +         | 12 (11)       |
| Onychophorans | +                          | +                    | +                 | +++                         | +                | ++                                | +         | 11 (9)        |
| Centipedes | +++                        | +                    | +                 | +                           | ++               | +                                 | +         | 9 (8)         |
| Millipedes | +++                        | +++                  | +                 | +++                         | +++              | +++                               | +++       | 17 (14)       |
| Earthworms | ++                         | +                    | +                 | +                           | +                | +++                               | +++       | 12 (9)        |
| Molluscs   | +++                        | +++                  | +++               | +++                         | +++              | +++                               | ++        | 19 (17)       |

aIn South Africa: <20 = +; 21–50 = ++; >50 = +++.

bWell known taxonomy: % of species identifiable to species level by expert: >50% = +; 50–75% = ++; >75% = +++ (based on material collected in this study).

cResources available for identification by non-expert: none = +; some but incomplete/difficult to use = ++; good = +++.

dWell known natural history: information on life history, diet, habitat available for taxon: none = +; some but incomplete = ++; good general knowledge = +++.

eReadily surveyed: require specialised sampling = +; require at least one specialised method = ++; easily sampled as part of general survey = +++.

fHigh ecological fidelity: species occur in both forest and matrix = +; most species restricted to forest = ++; all species limited to forest = +++. 

gEndemism: <10% of species regional endemics (considering entire SA fauna) = +; 10–30% regional endemics = ++; >30% regional endemics = +++.

hNumbers in parentheses indicate Summerville et al. [6] score excluding endemism.

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the Drakensberg is 16°C, with mean daily maximum temperatures ranging from 26.7°C in summer to 15.6°C in winter [50]. Temperatures can drop to below zero in winter.

We sampled three Afrotemperate forest patches (sites) at Injisuthi (Appendix S1) in March (late summer, wet season), June (winter, dry), September (spring, dry) and December (mid-summer, wet) 2004, to assess the effect of seasonal changes on invertebrate species richness and community structure. To increase our power for assessing surrogacy of molluscs or millipedes for overall diversity, we sampled an additional two forests at Injisuthi (total five), four (unburned) forests at Royal Natal and one forest at Cathedral Peak between November 2004 and January 2005 (Appendix S1).

We experimentally tested our conclusions for monitoring based on the Injisuthi data by contrasting four burned forests (in Devil’s Hock valley) with four unburned forests (in Thukela Gorge valley) at Royal Natal National Park (Appendix S1). According to Ezemvelo KZN Wildlife fire records, Devil’s Hock valley last burned in January 2003, 22 months prior to sampling, while Thukela Gorge forests had not burned during the same invasive fire and neither valley was burned again until after sampling took place at Royal Natal in November 2004.

The same sampling methods, sampling intensity and taxa were used in each forest patch at each sampling event. We collected six 0.3 L soil cores, 5 m apart in a straight line. Soil samples were kept in a cool place (refrigerator when possible) and processed within 14 days of collection. We placed soil samples in Berlese funnels for 48 h to extract invertebrates, after which we checked the soil in the funnel for large invertebrates unable to crawl through the 1 mm2 gauze.

We set six pitfall traps (plastic 0.125 L screw top jars, 75 mm deep and 40 mm diameter) per forest into the holes from the soil

![Figure 4. Effect of disturbance (fire treatment: unburned = grey line and squares; burned = black line and circles) on species richness of molluscs, millipedes, and all non-mollusc taxa combined (earthworms, centipedes, millipedes, and ants). Data are mean ±95% Confidence limits. doi:10.1371/journal.pone.0009100.g004](image)

| Table 3. The effect of disturbance (fire history) on forest mollusc species at Royal Natal National Park. |
|---|
| **Family** | **Species** | **Unburned** | **Burned** |
| Hydrocenidae | *Hydrocena noticola* Benson, 1856 | 103 | 0 |
| Pupillidae | *Lauria dadi* (Benson, 1864) | 9 | 1 |
| Orculidae | *Fauxulus glanvilleanus* (Ancey, 1888) | 46 | 6 |
| Orculidae | *Fauxulus mcbeanianus* Melville and Ponsonby, 1901 | 18 | 2 |
| Vertiginiidae | *Pupisoma harpula* (Reinhardt, 1886) | 1 | 0 |
| Vertiginiidae | *Truncatellina sykesii* (Melville and Ponsonby, 1893) | 11 | 0 |
| Clausiliidae | *Macropychia africana* (Melville and Ponsonby, 1899) | 43 | 0 |
| Streptaxidae | *Gulella justidens* (Melville and Ponsonby, 1899) | 67 | 3 |
| Streptaxidae | *Gulella mariae* (Melville and Ponsonby, 1892) | 1 | 0 |
| Streptaxidae | *Gulella sp.* | 5 | 0 |
| Rhytididae | *Nata vemicosa* (Krauss, 1848) | 8 | 1 |
| Valloniidae | *Acanthinula sp.* | 19 | 4 |
| Charopidae | *Afrodonta novemlamellaris* (Burnup, 1912) | 13 | 1 |
| Charopidae | *Trachycystis contabulata* Connolly, 1932 | 2 | 9 |
| Charopidae | *Trachycystis estima* (Melville and Ponsonby, 1899) | 46 | 61 |
| Charopidae | *Trachycystis glanvilliana* (Ancey, 1893) | 0 | 4 |
| Charopidae | *Trachycystis rudicostata* Connolly, 1923 | 41 | 14 |
| Charopidae | *Trachycystis subpunguis* Connolly, 1922 | 23 | 4 |
| Helicarionidae | *Kaliella eucnouilodes* Melville and Ponsonby, 1908 | 23 | 2 |
| Euconulidae | *Afroconulus diaphanus* (Connolly, 1922) | 0 | 5 |
| Urocyclidae | *Sheldonia truncavaelensis* (Craven, 1880) | 9 | 21 |

*Abundance scores are based on active search quadrat, litter sample and tree beat data.

Species in bold are not micromolluscs [48].

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samples. We filled traps with a glycerol-ethanol mixture and collected these pitfall traps after six days.

To specifically target micro-molluscs (adults have shells <5 mm diameter – [51]), we collected two 2 L leaf litter samples from each forest from areas that had not been disturbed by the team. The litter sample was collected from a single point covering about 0.25 m², taken at a set distance along a randomly placed transect to avoid collector bias. We collected and identified both live and dead molluscs, as well as all other target taxa. Litter samples were sorted by hand within 48 h of collection. Snails were drowned in water then preserved in 70% ethanol.

One set of five contiguous 2 x 2 m quadrats, covering an area of 20 m² on undisturbed ground was sampled in each forest. We searched all leaf litter, rocks, logs, vegetation below 0.5 m and the top 50 mm of soil, covering the entire area thoroughly for target taxa. It took one person approximately 210 min to search 20 m².

Although sampling was focussed on ground-dwelling taxa, some molluscs and ants also occur in trees. We beat ten under-storey trees per forest, selected based on their accessibility. We struck one branch of each tree five times with a large wooden stick, and collected all molluscs and ants that fell onto a white, flat, round, cotton, 0.7 m diameter collecting net.

For each 2 x 2 m quadrat and tree beat sampling, we collected representative samples in the field, recorded the number of individuals of target taxa, and released live extra specimens where we sampled a large number of individuals, and where species were readily recognisable. Earthworms were prepared as follows: each individual was rinsed in water, preserved in a weak (40%) solution of ethanol, allowed to dry for four minutes and then fixed in 4% formalin. All other invertebrates were frozen and then preserved in 70% ethanol.

Target taxa were sorted to morphospecies in the laboratory and identified to species by respective taxonomic experts as follows: molluscs, Dr Dai Herbert (Natal Museum); earthworms, Dr Danuta Plisko (Natal Museum); onychophorans, centipedes and millipedes, Prof. Michelle Hamer (UKZN/SANBI); and ants (wingless workers only, identified to morphospecies only), Dr Hylton Adie (UKZN). The reference collection is lodged in the Natal Museum, Pietermaritzburg for use in future studies.

Sampling intensity was relatively low because of the small size of the forests and the need to minimize disturbance in the forests. Sample-based species-accumulation curves [52] were plotted for each site using PRIMER [53]. We calculated effort as the approximate number of person hours taken for field sampling and processing of each replicate of each sampling method. We plotted species presence/absence against this effort using 999 permutations. Species-accumulation curves for all methods combined approached an asymptote across sites within a sampling month (Appendix S2). Observed species richness of tropical arthropods rarely reaches an asymptote, even with intensive sampling [52]. To avoid pseudoreplication [54], data from replicates taken at each site were combined into a single datum per sample method per site. These data indicate that we sampled a substantial portion of the diversity of our target taxa using our methods and effort.

**Determination of Suitable Sampling Methods for Use in Biodiversity Assessment and Monitoring**

To compare the contribution of different sampling methods to species richness counts in different seasons, we plotted species richness from each sampling method in each month for all taxa combined and separately for each target taxon. To determine which sampling method(s) were most suitable for targeting rare species and species with relatively short adult stages or short periods of surface activity, we noted the number of species collected by only one sampling method in each month for all taxa combined. We calculated efficiency of each sampling method (i.e. sampling effort) as the total number of species recorded in three forests combined, divided by the number of person hours required for sampling and processing. We calculated efficiency (species per person hour) for each sampling method in each month, and mean efficiency for each sampling method as the mean of four months.
Evaluation of Suitability of Taxa for Use as Biodiversity Surrogates and Indicators of Disturbance

Taxa were assessed according to the criteria and scale presented by Summerville et al. [6]. Data on the different taxa were obtained from experts, from the literature [17–21,55–57], or from relevant websites [58].

We assessed potential for surrogacy for overall diversity by regressing the diversity excluding a taxon against the diversity for that taxon [6]. For this analysis we used data collected in the same way in ten forests across four study sites: Royal Natal (4); Cathedral Peak (1) and Injisuthi (5). Residuals were normally distributed (Kolmogorov-Smirnov test: all P > 0.05).

Monitoring Using Our Recommendations: Assessment of Response to Environmental Disturbance

Given that we had identified molluscs as the taxon with the highest indicator potential, we focused this analysis on molluscs. We compared total and mean mollusc species richness of forests between unburned and burned valleys at Royal Natal National Park using analysis of variance (ANOVA). We compared mollusc species richness measured by quadrat, litter sample or tree beating to determine methods required to sample mollusc species richness for monitoring purposes. To determine whether mollusc species richness reflected the influence of fire on other taxa, we compared mollusc data with centipede, millipede and ant data.

We assessed the effectiveness of the mollusc community in reflecting differences associated with disturbance (fire history) by performing a Bray-Curtis similarity matrix using square-root transformed abundance data in PRIMER. We then performed an MDS plot and cluster analysis to assess differences among sites with different histories. If molluscs are a good indicator of disturbance, their communities should cluster according to fire history. In addition, we assessed whether the mollusc community reflected similar changes in the communities of other taxa by performing the same analysis for all taxa excluding molluscs. As an additional analysis, we performed a Procrustean randomisation test using PROTEST [59] which contrasts two community matrices in a more robust manner than Mantel tests [59]. Here we contrasted the community of molluscs at the four burned forests relative to the community of molluscs at the four unburned forests. In addition, we contrasted the community of all non-mollusc taxa in the burned relative to the unburned forests to assess if these were similarly (as with molluscs) different from each other.

Supporting Information

Appendix S1 Forest details with an indication of which forests were used for specific analyses. MAP = mean annual precipitation. Found at: doi:10.1371/journal.pone.0009100.s001 (0.05 MB DOC)

Appendix S2 Sampling saturation for the Injisuthi seasonal sampling. We present randomized species-accumulation curves of all target taxa combined in each month that seasonal sampling took place: (A) autumn (March), (B) winter (June), (C) spring (September), and (D) summer (December). The x-axes represent the number of person hours taken to collect and process each sampling replicate. Found at: doi:10.1371/journal.pone.0009100.s002 (0.06 MB DOC)

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Author Contributions

Conceived and designed the experiments: CU MH RS. Performed the experiments: CU. Analyzed the data: CU RS. Wrote the paper: CU MH RS.

References

1. Margules CR, Pressey RL (2000) Systematic conservation planning. Nature 405: 243–253.
2. Slotow R, Hamer M (2000) Biodiversity research in South Africa: comments on current trends and methods. South African Journal of Science 96: 222–224.
3. Ponder W (1999) Using museum collection data to assist in biodiversity assessment. In: Ponder W, Mosman D, Lunney D, eds. The Other 99%. The Conservation and Biodiversity of Invertebrates, Transactions of the Royal Zoological Society of New South Wales, Australia.
4. Hamer M, Slotow R (2002) Conservation application of existing data for South African millipedes (Diplopoda). African Entomology 10: 29–42.
5. Didham RK, Springate BD (2003) Determinants of temporal variation in community structure. Chapter 4. In: Basset Y, Novotny V, Miller SE, Kitching RL, eds. Arthropods of tropical forests, Cambridge University Press: Cambridge, UK.
6. Summerville KS, Ritter LM, Crisp TO (2004) Forest moth taxa as indicators of lepidopteran richness and habitat disturbance: a preliminary assessment. Biological Conservation 116: 9–18.
7. Lawes MJ, Kotze DJ, Bourquin SL, Morris C (2005) Epigean invertebrates as potential ecological indicators of Afromontane forest condition in South Africa. Biotropica 37: 109–118.
8. Kremen C, Colwell RK, Erwin TL, Murphy DD, Noss RF, et al. (1993) Terrestrial arthropod assemblages: their use in conservation planning. Conservation Biology 7: 796–808.
9. New TR (1999) Untangling the web: spiders and the challenges of invertebrate conservation. Journal of Insect Conservation 3: 251–256.
10. Ward DF, Lavriërve M-C (2004) Terrestrial invertebrate surveys and rapid biodiversity assessment in New Zealand: lessons from Australia. New Zealand Journal of Zoology 31: 151–159.
11. McGrosh MA (1996) The selection, testing and application of terrestrial insect as bioindicators. Biological Review 73: 181–201.
12. Lindenmayer DB, Franklin JF, Fischer J (2006) General management principles and a checklist of strategies to guide forest biodiversity conservation. Biological Conservation 131: 433–445.
13. Andersen AN (1999) My bioindicator or yours? Making the selection. Journal of Insect Conservation 3: 61–64.
14. Pocock MJO, Jennings N (2008) Testing biotic indicator taxa: the sensitivity of inverteagous mammals and their prey to the intensification of lowland agriculture. Journal of Applied Ecology 45: 151–160.
15. Rohr JR, Mahan CG, Kim KC (2007) Developing a monitoring program for invertebrates: guidelines and a case study. Conservation Biology 21: 422–433.
16. Lovell S, Hamer M, Slorow R, Herbert D (2007) Assessment of congruency across invertebrate taxa and taxonomic levels to identify potential surrogates. Biological Conservation 139: 113–125.
17. Pickford GE (1937) A monograph of the Acanthodriline earthworms of South Africa. W. Heller and Sons, Cambridge, UK.
18. Lawrence RF (1984) The centipedes and millipedes of southern Africa. A.A. Balkema, Cape Town, South Africa.
19. Hamer ML, Samways MJ, Ruhheng H (1997) A review of the Onychophora of South Africa, with discussion of their conservation. Annals of the Natal Museum 38: 283–312.
20. Ploko JD (2003) Eleven new South African earworms (Oligochaeta: Microchaetidae). African Invertebrates 44: 279–325.
21. Herbert D, Kilburn D (2004) Field guide to the land snails and slugs of eastern South Africa. Natal Museum, Pietermaritzburg, South Africa.
22. Gessner Y (2009) Patterns of distribution, diversity and endemism of terrestrial molluscs in South Africa. M.Sc. Thesis, University of KwaZulu-Natal, Durban.
23. New TR (2000) How useful are ant assemblages for monitoring habitat disturbance on grasslands in south eastern Australia? Journal of Insect Conservation 4: 153–159.
24. Parr CL, Bond WJ, Robertson HG (2002) A preliminary study of the effect of fire on ants (Formicidae) in a South African savanna. African Entomology 10: 101–111.
25. Andersen AN, Major JD (2004) Ants show the way Down Under: invertebrates as bioindicators in land management. Frontiers in Ecology and the Environment 2: 291–298.
26. Andersen AN, Fischer A, Hoffmann BD, Read JL, Richardson R (2004) Use of terrestrial invertebrates for biodiversity monitoring in Australian rangelands, with particular reference to ants. Austral Ecology 29: 87–92.
27. Mucina L, Geldenhuys CJ (2006) Afrotemperate, Subtropical and Azonal Forests. Chapter 12. In: Mucina L, Rutherford MC, eds. The vegetation of...
