

REVIEW

# Minimising exposure of amphibians to pathogens during field studies

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**ABSTRACT:** Many of the recent global amphibian mass mortalities, declines and extinctions have been attributed to the emerging infectious disease chytridiomycosis. There have been mass mortalities due to ranaviral disease but no major declines or extinctions. Controlling the transmission and spread of disease is of utmost importance, especially where there is the potential for human involvement. We have reviewed current hygiene guidelines for working with wild frogs, identified potential flaws and recommended those most suitable and effective for the field environment. Our within-site hygiene measures aim to reduce the risk of transmission among individuals. These measures encompass the capture, handling and holding of amphibians, skin disinfection before and after invasive procedures, marking frogs, sealing open wounds and treatment of accessory equipment. Our between-site hygiene measures aim to mitigate the risk of pathogen spread among populations. We have designed a risk calculator to help simplify and standardise the decision-making process for determining the level of risk and appropriate risk mitigation strategies to reduce the risk of increasing pathogen spread above background levels. Calculation of an overall risk score for pathogen spread takes into account the prior activity of field workers, the proposed activity, remoteness of the site, presence of known pathogens and the consequences of increased pathogen spread for amphibians in a given area.

**KEY WORDS:** Chytridiomycosis · Hygiene · *Batrachochytrium dendrobatidis* · Ranaviral disease · Ranavirus · Field study

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## INTRODUCTION

The past 30 yr has seen alarming declines and extinctions of amphibians world wide. The recently emerged disease, chytridiomycosis, has caused mass mortalities, major declines and extinctions, and ranaviral disease has caused mass mortalities (Cunningham et al. 1996, Berger et al. 1998, Daszak et al. 1999, Long-

core et al. 1999, Green et al. 2002, Muths et al. 2003, Lips et al. 2006, Schloegel et al. 2006, Picco et al. 2007, Skerratt et al. 2007). The consequences of introducing these or other serious pathogens into naïve amphibian populations are likely to be severe (Berger et al. 1999, Pearman et al. 2004, Weldon et al. 2004, Cunningham et al. 2005, Jancovich et al. 2005, Lips et al. 2006, 2008, Fisher & Garner 2007, Skerratt et al. 2007).

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Unregulated trade in amphibians, as well as unintentional shipment, is suspected to have been a major contributor to the spread of these diseases (Weldon et al. 2004, Fisher & Garner 2007, Skerratt et al. 2007). To curb their spread, the World Organisation for Animal Health (OIE) has made these diseases notifiable (OIE 2008). Preventing the spread of these amphibian diseases across international borders is of great importance, but it is also important to control disease spread within national borders (i.e. domestically).

The potential for human involvement in transmission (passing of disease from an infected to an uninfected individual) and spread (movement of disease geographically) of these diseases, within and among amphibian populations, appears to be significant (Mutschmann et al. 2000, Raverty & Reynolds 2001, Parker et al. 2002, Mazzoni et al. 2003, Hanselmann et al. 2004, Weldon et al. 2004, Jancovich et al. 2005, Oevermann et al. 2005, Daszak et al. 2006, Picco et al. 2007, Picco & Collins 2009). Of particular concern is the potential for field biologists to spread disease amongst wild populations.

While evidence directly implicating field biologists in the spread of amphibian diseases is currently lacking, the hazard of disease transmission and spread by biologists must be considered. For researchers working with amphibians or within areas of amphibian habitat, the risk of disease transmission within these habitats and the spread of disease among populations may be increased due to (1) frequent handling of amphibians, (2) frequent or rapid movement between isolated areas of habitat and (3) rapid or frequent movement between captive husbandry and laboratory facilities and the field. It is therefore important that biologists working with amphibians or within amphibian habitats take care to minimise disease transmission and spread both within and among amphibian populations. This is especially important where rare, geographically restricted or threatened amphibian species are concerned and when the spread of disease can have serious consequences for species survival.

Field researchers should, therefore, evaluate their activities to determine their relative risk of pathogen transmission and spread compared with background levels (i.e. the risk posed by other mechanisms of disease transmission or pathogen dispersal) and implement appropriate strategies to minimise this risk during field studies. It must be recognised that the overall risk of transmission and spread of pathogens cannot be reduced below the background level by risk mitigation during research or monitoring activities. Therefore, the aim of risk mitigation is to prevent risk increasing above background levels (i.e. the risk posed by other mechanisms of disease transmission or pathogen dispersal). The risk of transmission and

spread should be evaluated by researchers, animal ethics committees and government agencies issuing permits. However, we do not advocate the cessation of field-based research or monitoring as a risk management strategy. The value of the research outcome needs to be balanced with the disease risk posed to individuals and, more importantly, to populations of amphibians. In addition, the presence of infectious disease (caused by pathogens) should not be confused with that of non-infectious diseases such as most cancers, which cannot be transmitted between animals.

Current hygiene guidelines for handling wild frogs (Daszak et al. 2001, NSW NPWS 2001, NWHC 2001, HACCC 2004, Speare et al. 2004, Aguirre & Lampo 2006, CCADC 2008) seek to minimise transmission of infectious diseases; however, some guidelines are contradictory and potentially harmful to amphibians (e.g. the use of iodine compounds for surgical disinfection). Here we review the suggested practices for minimising the spread of infectious diseases within sites and make recommendations for those most suitable and effective in the field environment. In addition, existing hygiene protocols identify the importance of between-site measures to reduce pathogen spread but lack a framework for determining the stringency and scale at which between-site measures are applied. To help inform such decisions we present a risk calculator that enables field biologists to assess the relative risk (above background levels) of their activities and the hazard they pose to amphibian populations. This approach is similar to that currently employed by Biosecurity Australia and other member countries of the OIE during import risk assessments (AGDAFF 2009). Our recommendations are aimed at amphibian pathogens known to result in widespread mortality, namely *Batrachochytrium dendrobatidis* and ranaviruses, but will also be effective against other pathogens.

## WITHIN-SITE HYGIENE MEASURES

### Capture, handling and holding wild amphibians

Capture, handling and housing of wild amphibians should be minimised or avoided where possible, as stressed animals are at greater risk of infection (Carey et al. 1999, Daszak et al. 2003). Direct microbial transfer during capture and handling of successive adult amphibians can be reduced by wearing single-use gloves (latex, nitrile or vinyl) or capturing and handling frogs in single use lightweight plastic bags. Some tadpoles suffer lethal effects when exposed to latex, nitrile and, to a lesser extent, vinyl gloves, although some of these effects can be mitigated by washing the gloves before handling (Gutleb et al. 2001, Cashins et

al. 2008). Larvae should, therefore, be handled with gloves that have been proven or rendered safe for that species. Data suggest the causative pathogen of chytridiomycosis, *Batrachochytrium dendrobatidis*, does not survive on bare human hands for long; however, this killing effect is reduced with repeated hand washing in water (Mendez et al. 2008). It is unknown whether ranaviruses remain viable following contact with human skin. We therefore recommend that single-use gloves be worn during all capture and handling events involving high risk species (critically endangered or narrowly endemic) or locations (e.g. isolated areas with endemic species, disjunct populations or high species richness). In situations where gloves are not available or suitable, hand washing with 70% ethanol (then allowing hands to air dry) between handling individual frogs is an alternative. If ethanol contact is maintained for 60 s, most bacteria, fungi and viruses will be killed (see Table 1); however, repeated use on human skin is not recommended. If 70% ethanol is not available or suitable, the minimum recommended treatment is hand-washing in the water body to which the amphibian is naturally exposed. While rinsing in source water will not kill pathogens on hands, it may dislodge them and so exposure will not be increased greatly above background environmental levels. This treatment, however, needs to be tested further and should only be used for low risk species or locations.

When amphibians must be held temporarily, individuals should be housed in single-use containers (e.g. plastic bags) or in containers disinfected between each animal (Johnson et al. 2003, Webb et al. 2007, see Table 1). Adults should never be held in groups. Tadpoles from the same pond or stream section may be housed for short periods in a common container, though care must be taken to avoid overcrowding as this can increase transmission rates and cause physiological stress (Denver & Crespi 2006). Longer holding times (>60 min) will require changes to water and the provision of appropriate food (when holding time is >24 h). Tadpoles of some species, particularly pond-breeding species with high fins, are easily damaged on capture and with movement of water within the holding container (E. Meyer pers. obs.). Rough handling of tadpoles of these species may, therefore, increase their susceptibility to trauma and infection. The requirements of tadpoles of each species should be considered when designing the capture and holding components of a field study. If animals are removed from the field and then returned, it should always be to the same site.

Dead amphibians or live animals showing clinical signs of disease must be regarded as having a high infection risk to healthy animals and rigorous hygiene measures are needed. No effective field treatment for

chytridiomycosis has been demonstrated; however, infected frogs in captivity can be treated with elevated environmental temperatures (Woodhams et al. 2003, Retallick & Miera 2007), itraconazole (Aguirre & Lampo 2006, Pessier 2007) or chloramphenicol (Poulter et al. 2008) to clear *Batrachochytrium dendrobatidis*. These treatments require further validation in a range of species and some could be tested in the field in certain situations such as when recapture probabilities are high. No treatment regimes for ranaviral infection of frogs have been described; however, Johnson & Wellehan (2005) suggest Zovirax<sup>®</sup> may be successful in controlling the disease. If successful these treatments could be used to lower the burden of disease in populations and reduce the risk of transmission and spread.

Sick and dead frogs should be collected for disease diagnosis. Disease surveillance may detect disease introduction and enable emergency responses. It is also useful to assess the risk of pathogen transmission to other individuals or spread to populations.

#### **Skin disinfection before and after invasive procedures**

Identifying individual amphibians during mark-recapture studies is essential for determining survival, reproduction, home range, habitat preferences and incidence of infection. However, the method used for recognising animals must not significantly affect survival or behaviour. Therefore, strict hygiene standards must be maintained during amphibian marking procedures including implanting internal radio transmitters, passive integrated transponder (PIT) tags, visible implant alphanumeric (VIA) tags, visible implant elastomer (VIE) tags and toe tipping or clipping.

Amphibian skin is more permeable than skin of other vertebrates and requires special disinfectants. Alcohol, phenol and iodine based disinfectants will destroy mucus and wax that prevent dehydration and microbial infection of amphibian skin. These disinfectants are also potentially toxic and may be absorbed systemically (Wright 2001). The only suitable commercially available preparation for disinfecting wounds is Bactine<sup>®</sup> spray (active ingredient 0.14% w/w benzalkonium chloride and 2.6% w/w lidocaine hydrochloride in a non-alcohol base). Bactine<sup>®</sup> has been successfully used to treat amphibian wounds (Martin & Hong 1991). Chlorhexidine (0.75% diluted from 2% Nolvasan<sup>®</sup>) is also suitable for surgical disinfection (Wright 2001).

Betadine<sup>®</sup> disinfectant (active ingredient 1% povidone iodine) has been recommended for use on frogs before surgical procedures such as toe clipping (NSW NPWS 2001). However, iodine has caused toxic reac-

tions in some amphibians (Wright 2001). Sensitivity, however, appears to be species-specific. Treatment with povidone iodine did not have adverse effects on the green tree frog *Hyla cinerea* (Brown 1995), but was implicated in rapid mortality of the harlequin poison frog *Dendrobates histrionicus* (Stoskopf et al. 1985). Therefore, we do not recommend Betadine® or other povidone-iodine products be used as disinfectants for an amphibian species until toxicity has been determined.

Application of a disinfectant before and after invasive procedures helps minimise pathogen entry from the skin to the wound. However, the protection provided is probably lost when the treated animal is released at the site of capture into a water body. The water-soluble antiseptics suitable for amphibians will dissolve upon contact with water. Therefore, for these treatments to be effective an extended holding time (e.g. 1 h) is necessary before animals are released into or near water.

### Marking frogs

Toe tipping (removal of most distal phalange) or toe clipping (amputation of a greater proportion of the digit) should occur through the interphalangeal joint. Johnson et al. (2003) recommend 70% ethanol as the most suitable disinfectant for use in field sterilisation of scissors to minimise the risk of transmitting *Batrachochytrium dendrobatidis* or ranaviruses. This treatment should be effective against most bacterial, viral and fungal pathogens.

PIT, VIE and VIA tags should be inserted with a sterile, single-use applicator. There is no clinical documentation of infection from any marking or tagging method; however, further controlled studies are required to conclusively demonstrate a limited effect on animal health and mobility (Phillott et al. 2007).

### Sealing open wounds

The use of a cryanoacrylate compound such as Vetbond® (active ingredient n-butyl cryanoacrylate) as a tissue adhesive after toe tipping or clipping is possible in larger amphibians, but becomes increasingly difficult in smaller animals because it is difficult to isolate toes for application. Vetbond® can also be used to seal incisions made during subdermal injection of VIA, VIE and PIT tags. Moisture can interfere with setting times and adhesion (see Boothe 2003) so care must be taken to ensure setting has occurred before release. Problems may be experienced in their application to stream- or pond-dwelling amphibians, but can be

avoided by using a small piece of sterile absorbent dressing to draw surplus water from the wound before application of the adhesive (H. B. Hines pers. obs.). Less expensive industrial adhesives ('superglues') should not be used as a replacement for surgical tissue glues as they have resulted in tissue death in other animals (see Wright 2001). Adhesives, such as Vetbond®, may also trap microbes in the wound and promote infection (see Boothe 2003). To alleviate this problem a disinfectant such as Bactine® should be applied before the adhesive.

### Treatment of accessory equipment

Appropriate disinfection strategies for accessory equipment such as nets, carry bags, clothing and footwear are listed in Table 1, which describes the minimum exposure time and disinfectant concentration shown to be effective on clean surfaces. No disinfectants should be applied where runoff could wash these into nearby water bodies. All equipment should be dry and cleaned of mud and organic debris before use, as this may reduce disinfection efficacy, then treated before use between sites (see next section).

### ENTRY, EXIT AND BETWEEN-SITE HYGIENE MEASURES

Field biologists exposed to ill animals or their pathogens may pose an increased risk (above background levels) of spreading wildlife diseases. This risk is exacerbated by the ease and speed with which biologists can now move between sites and, more importantly, across natural barriers to disease spread (e.g. expanses of habitat unsuitable to amphibians or their pathogens such as deserts, mountain ranges and seas). It is therefore important that field biologists take steps to avoid increased risk of transmission of wildlife diseases between sites and populations.

Potential measures for reducing the likelihood of disease spread between sites are listed in Table 2. These measures range in rigour from simple rinsing and drying of equipment through to the use of site-dedicated equipment and disinfection of vehicles. The purpose of these measures is not to eliminate pathogen spread entirely (as this may not be possible), but rather to avoid increased risk of spread above background levels. The stringency of hygiene measures to be implemented should, therefore, reflect the risk of field biologists spreading disease relative to background risk levels. This will depend on a range of factors including (1) activities undertaken before field work, (2) the nature of the proposed field work, (3) the remoteness

Table 1. Disinfection strategies suitable for killing *Batrachochytrium dendrobatidis* and ranaviruses in field studies. Information for *Batrachochytrium dendrobatidis* is adapted from Berger (2001), Johnson et al. (2003) and Webb et al. (2007), and for ranaviruses from Langdon (1989), Miocevic et al. (1993) and Bryan et al. (2009)

Application	Disinfectant	Strength	Time	Target pathogen	
Disinfecting surgical equipment and other instruments (e.g. scales, callipers)	Benzalkonium chloride	2 mg ml <sup>-1</sup>	1 min	<i>B. dendrobatidis</i>	
	Ethanol	70 %	1 min	<i>B. dendrobatidis</i> , Ranaviruses	
Disinfecting collection equipment and containers	Sodium hypochlorite (bleach contains 4 % sodium hypochlorite)	1 %	1 min	<i>B. dendrobatidis</i>	
		3 %	1 min	Ranaviruses	
Trigene	Path X or quaternary ammonium compound 128	1 in 500 dilution	0.5 min	<i>B. dendrobatidis</i>	
	1 in 5000 dilution	1 min	<i>B. dendrobatidis</i>		
	F10	1 in 1500 dilution	1 min	<i>B. dendrobatidis</i>	
	Virkon	2 mg ml <sup>-1</sup>	1 min	<i>B. dendrobatidis</i>	
			1 min	Ranaviruses	
	Nolvasan	0.75 %	1 min	Ranaviruses	
	Potassium permanganate	1 %	10 min	<i>B. dendrobatidis</i>	
	Complete drying		>3 h	<i>B. dendrobatidis</i>	
	Heat	60°C	30 min	<i>B. dendrobatidis</i> , Ranaviruses	
	Heat	37°C	8 h	<i>B. dendrobatidis</i>	
Sterilising UV light		1 min	Ranaviruses only		
Disinfecting footwear	Sodium hypochlorite (bleach contains 4 % sodium hypochlorite)	1 %	1 min	<i>B. dendrobatidis</i>	
		3 %	1 min	Ranaviruses	
	Path X or quaternary ammonium compound 128	1 in 500 dilution	0.5 min	<i>B. dendrobatidis</i>	
	Trigene	1 in 5000 dilution	1 min	<i>B. dendrobatidis</i>	
	F10	1 in 1500 dilution	1 min	<i>B. dendrobatidis</i>	
	Complete drying		>3 h	<i>B. dendrobatidis</i>	
	Disinfecting cloth (e.g. carry bags, clothes)	Hot wash	60°C or greater	30 min	<i>B. dendrobatidis</i> , Ranaviruses

and frequency of human visitation of sites where work is being undertaken and (4) the presence or absence of known pathogens at a site.

In deciding on the rigour of between-site hygiene measures, consideration must be given not only to the aforementioned risk factors, but also the severity of the consequences of disease introduction for rare (geographically restricted) and threatened amphibian species. Since the implications of pathogen spread between sites for these species is probably more serious (potentially resulting in extinction of species or populations), more rigorous hygiene protocols may be appropriate, even where the risk posed by biologists is only slightly above background levels.

Given the above, deciding which measures are necessary and appropriate can be difficult, as can deciding on the geographical scale at which to apply be-

Table 2. Hygiene measures to be employed before visiting a study site grouped according to stringency

Stringency	Hygiene measures
Low	Clean and dry equipment and clothing
Moderate	Equipment that has been in contact with amphibians or their habitat cleaned and disinfected
	Vehicle that has been at a site with amphibians is cleaned
High	Site- or area-dedicated or disposable field equipment or thorough cleaning and disinfection of all clothing and field equipment that has been in contact with amphibians or their habitat
	Vehicle that has been at a site with amphibians is washed and disinfected



tween-site hygiene measures. Current hygiene guidelines provide limited assistance in defining the scale at which between-site hygiene protocols are applied. For example, NSW NPWS (2001) state that defining the boundary of a site may be problematic, but provide no criteria for making such assessments. We suggest that different levels of hygiene measures are appropriate between catchments of various sizes depending on the level of risk. For example highly stringent hygiene protocols are applied at the scale of small subcatchments when risk is high; less stringent protocols are required for larger catchments when relative risk is low.

To help simplify and standardise this decision-making process, we have developed a risk calculator, incorporating the risk factors and consequences identified above. Using this approach, field biologists can assess their risk of increased pathogen spread above background levels. An overall risk estimate is calculated that provides a quantitative measure that can assist field workers in determining the scale at which to apply between-site hygiene measures and the level of stringency required to reduce their risk (see Table 5). Given the diversity of environments encountered or activities undertaken during field studies, this risk calculator is best viewed as a framework that can be modified to suit particular circumstances. Its use, however, will ensure that there is greater consistency among field biologists in their application of hygiene protocols.

To assess the risk of increased pathogen spread, each of the 4 risk factors identified previously is scored against criteria described below and listed in Table 3.

### Risk 1: prior activity

Activities in the days (~2 to 3 d) before field work begins can greatly increase risk of pathogen spread particularly where these involve work with exotic (nonlocal) captive amphibians, pathogen cultures or infected amphibians. This is because pathogen survival on fomites or mechanical vectors generally decreases exponentially with time. Involvement in such activities necessitates the implementation of more stringent hygiene measures, particularly entry protocols. For this reason we have given activities associated with working with pathogens or known infected or captive amphibians an order of magnitude greater score (100 points) versus the maximum of 10 points for other risk factors. This high score can be reduced to 10 if working under biosecure conditions such as in a level 2 biosecurity (PC2) facility. Working on apparently healthy amphibians and aquatic environments in areas harbouring known pathogens or diseases receives a score of 10 unless those amphibians have disease, in which case the score is 100. Working on amphibians and aquatic environments in uninfected areas receives a score of 7 because pathogens of amphibians are poorly known and it is

Table 3. Risk factors, criteria and scores used to determine the total risk score for pathogen spread to a study site. The overall risk score is obtained by summing the scores for each of the 4 risk factors

Risk factor	Criteria	Score
1. Prior activity (~2–3 d)	(a) Worked with exotic (nonlocal) captive amphibians or amphibians with significant infectious or potentially significant infectious diseases or in a laboratory with significant amphibian pathogen cultures	100
	(b) Targeted apparently healthy amphibians or aquatic environments in areas where significant amphibian pathogens are known to be present or status is unknown	10
	(c) Targeted apparently healthy amphibians or aquatic environments in areas where significant amphibian pathogens are known to be absent	7
	(d) Did not target amphibians or aquatic environments	1
2. Proposed activity at the study site	(a) Directed towards amphibians or aquatic environments	10
	(b) Not directed towards amphibians or aquatic environments	1
3. Remoteness of the study site	(a) Isolated or difficult to access area; human visitation rare (e.g. <100 visitors yr <sup>-1</sup> )	10
	(b) Visitation low (e.g. 100–999 visitors yr <sup>-1</sup> )	7
	(c) Visitation moderate (e.g. 1000–9999 visitors yr <sup>-1</sup> )	3
	(d) Visitation high; area highly disturbed and/or populated	1
4. Presence of pathogen at the study site	(a) Significant amphibian pathogens are known to be absent or status is unknown	10
	(b) Significant amphibian pathogens are known to be present	5

possible that an undiscovered pathogen occurs there. When field workers did not target amphibians or aquatic environments this receives a score of 1, as encountering amphibian pathogens is much less likely to occur (Table 3).

### Risk 2: proposed activity

The nature of the proposed field work influences the risk of spread. Activities directly targeting amphibians or aquatic environments, such as mark–recapture studies, are likely to increase the risk of pathogen spread between sites and are given a score of 10. Other field activities not directly targeting amphibians but which take place within known amphibian habitat, such as a mammal survey, may also increase the risk of pathogen spread, although at a much lower likelihood, and are given a score of 1 (Table 3).

### Risk 3: remoteness of site

For accessible areas subject to high levels of human visitation, the value in field workers applying rigorous between-site hygiene measures may be reduced, as the background risk of pathogen spread by human activities is high. Where field biologists are accessing remote rarely visited areas, however, the potential for introducing new pathogens is relatively much higher compared with background levels, necessitating the adoption of more stringent hygiene measures. Consequently, we have scaled the relative risk by setting the criterion scores based on the level of total human visitation to the area (Table 3).

### Risk 4: presence of known pathogens

If the area to be visited is known to be free of significant amphibian pathogens, then the risk posed by disease spread is likely to be higher, necessitating more stringent hygiene measures. Where the disease status of sites is unknown or poorly known, we recommend a precautionary approach whereby a site is assumed to be pathogen-free and is also given a score of 10. Sites with known pathogens are given a score of 5 as pathogens of amphibians are poorly known and it is possible that an undiscovered significant pathogen could be introduced (Table 3). The status of amphibian disease at a site is particularly important for exit protocols, i.e. if a disease capable of causing mass mortality (e.g. chytridiomycosis, ranaviral disease) is known from an area then it is important to thoroughly disinfect everything before leaving that area. Transferring

water, substrates, plant material or animals between sites can spread pathogens and should be strictly avoided unless they can be guaranteed to be pathogen-free.

### Calculating and applying the overall risk estimate

The scores for each of the above factors are summed to produce a total risk score. This total risk score is multiplied by the consequence score, a value reflecting the International Union for Conservation of Nature (IUCN) conservation status (IUCN 2001) of amphibian species known or likely to occur within the area of the proposed activity (Table 4). The resulting overall risk score is then used to decide what hygiene measures are required to ensure pathogen transmission is not increased above background levels, as well as the scale at which these measures should be applied between sites (i.e. between major or minor subcatchments) (see Table 5).

An overall risk score of less than 150 indicates that the relative risk of a field biologist spreading pathogens is low and that between-site hygiene measures required are simple and applied at the broad scale of major catchments. An overall risk score in excess of 300 indicates that the relative risk posed by the field biologist is high and that between-site hygiene measures required to reduce this risk are stringent and applied at the fine scale of minor subcatchments (i.e. stream orders 1 and 2; Strahler 1952) within the area to be visited (Table 5). While the scores calculated by our system are relative and not absolute, they are useful in ensuring that key aspects of the risk of pathogen spread are considered when deciding which between-site hygiene measures to use.

Five examples of the application of the risk calculator are provided in Table 6. The highest overall risk score is for a graduate student conducting research on endemic frog species in remote areas where the status

Table 4. Criteria and consequence score for pathogen spread to the major catchment around the study site

Criteria	Score
The major catchment where the proposed activity is to be undertaken supports:	
(a) at least 1 critically endangered species or has endemic species	10
(b) at least 1 endangered species	8
(c) at least 1 vulnerable species	6
(d) at least 1 near-threatened species	4
(e) at least 1 amphibian (native or exotic) species	3
(f) no amphibians (known or likely)	1

Table 5. Overall relative risk score for pathogen transmission between sites and subsequent hygiene requirements. The overall risk score is obtained by summing the scores for risk factors in Table 3 (the total risk score) and multiplying by the consequence score from Table 4. Stream order is as defined by Strahler (1952)

	High	Medium	Low
Overall risk score	>300	150–300	<150
Entry and exit protocols	Stringent	Moderate	Low
Scale at which hygiene is required	Between minor subcatchments (e.g. stream order 1–2)	Between major subcatchments (e.g. stream order 3–4)	Between major catchments (e.g. stream order ≥5)

of amphibian diseases is not known. As this student is targeting frogs, their probability of spreading a new disease is relatively high and the consequence of this is also high due to the presence of locally endemic species. Therefore, highly stringent hygiene protocols applied at a small scale are recommended.

An amphibian keeper from a zoo that holds exotic frogs and who wishes to undertake monitoring of frogs in a local urban area, also has a relatively high risk of spreading a new disease, due to the keeper's potential exposure to exotic pathogens. These captive frogs have been sourced from a number of localities, including international ones, so they have the poten-

tial to carry subclinical diseases and parasites not native to the local area. The zookeeper, therefore, must apply stringent hygiene measures before commencing monitoring. In contrast a volunteer from the local community helping the zookeeper with monitoring poses a much smaller relative risk as they have not been in contact with exotic captive amphibians and the area targeted for monitoring is highly disturbed. We recommend the use of low stringency hygiene measures by this volunteer (i.e. use of clean and dry personal equipment and clothing) due to this low relative risk.

The remaining 2 examples of the risk calculator in Table 6 compare 2 cases where a government officer is undertaking amphibian survey and monitoring activities. If human visitation is high and the targeted area is only likely to support frogs with a relatively low risk of extinction, then the relative risk is low and only a low stringency of hygiene measures is required. However, where the officer is moving from site to site, including areas with chytridiomycosis present, and targeting endangered frogs, the relative risk is higher and so the hygiene measures need to be more stringent. The measures will become even more stringent if any of the monitored frogs have chytridiomycosis.

Table 6. Examples of the application of the risk calculator using 5 scenarios. RF1–4 refers to risk factors 1 to 4 and their criteria scores as shown in Table 3. The total risk score is the sum of the scores for risk factors 1 to 4. The overall risk score is calculated by multiplying the total risk score by the consequence score (Table 4)

Scenario	RF1	RF2	RF3	RF4	Total risk score	Consequence score	Overall risk score
1. A graduate student is studying frog speciation and endemism on remote mountain tops in Borneo where disease status is unknown	10	10	10	10	40	10	400
2. An amphibian keeper from a zoo that holds exotic frogs is planning to undertake monitoring of nonthreatened frogs in an urban area where disease outbreaks have not been observed	100	10	1	10	121	3	363
3. A community volunteer who does not hold a collection of exotic amphibians is assisting in the above survey of urban frogs	1	10	1	10	22	3	66
4. A government officer is undertaking a one-time survey of amphibian species at a popular swimming spot in a national park where a species considered vulnerable is likely to occur	1	10	3	10	24	6	144
5. A government officer is regularly monitoring populations of apparently healthy endangered frogs in national parks with moderate levels of visitation and where chytridiomycosis is known to occur	10 <sup>a</sup>	10	3	5	28	8	224

<sup>a</sup>This risk factor score would rise to 100 if any of the monitored frogs had chytridiomycosis



### Determining the order of visitation of multiple field sites

When a field trip encompasses several field sites, or a number of locations are being visited in succession within 2 to 3 d of each other, the order of visitation should be determined according to the presence of known pathogens and diseases. While unknown pathogens obviously cannot be accounted for, areas known to be absent of disease should be visited first, followed by areas of unknown status and then known infected areas.

### Return of captive animals to the wild

If wild amphibians are housed for any period of time in a captive situation (e.g. laboratory, zoo or captive breeding facility), we do not recommend that they be returned to their natural population unless they have been kept in isolation, their captive history is free of undiagnosed morbidity or mortality and they have had rigorous pathogen screening before release. This is usually beyond the means of most studies. Thus, when assessing permits for such an activity, wildlife conservation agencies and ethics committees should view animals taken from the wild as a permanent take.

### CONCLUSIONS

In the absence of detailed studies quantifying the effect of research and monitoring procedures on transmission and spread of pathogens, these guidelines provide researchers with the most appropriate and effective methods for minimising exposure of amphibians to pathogens within and among field sites. Prevention or failure to prevent transmission and spread of infection despite best hygiene practice should be documented to improve guidelines.

In addition to minimising exposure of amphibians to pathogens during field studies, our between-site hygiene recommendations may also be useful in preventing the spread of nonamphibian pathogens, such as the plant fungal pathogen *Phytophthora cinnamomi*, and weeds. Our hygiene principles may be used as a basis when developing pathogen transmission and spread-risk minimisation guidelines for other people such as recreational fishers or for pathogens of other taxa.

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*Submitted: March 16, 2009; Accepted: October 4, 2009  
Proofs received from author(s): January 19, 2010*