



# Lymphocyte Inhibition by the Salamander-Killing Chytrid Fungus, *Batrachochytrium salamandivorans*

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**ABSTRACT** Amphibian populations have been declining around the world for more than five decades, and the losses continue. Although causes are complex, major contributors to these declines are two chytrid fungi, *Batrachochytrium dendrobatidis* and *Batrachochytrium salamandivorans*, which both cause the disease termed chytridiomycosis. Previously, we showed that *B. dendrobatidis* impedes amphibian defenses by directly inhibiting lymphocytes *in vitro* and *in vivo* by release of soluble metabolites, including kynurenone (KYN), methylthioadenosine (MTA), and spermidine (SPD). Here, we show that *B. salamandivorans* cells and cell-free supernatants also inhibit amphibian lymphocytes as well as a human T cell line. As we have shown for *B. dendrobatidis*, high-performance liquid chromatography (HPLC) and mass spectrometry revealed that KYN, MTA, and SPD are key metabolites found in the *B. salamandivorans* supernatants. Production of inhibitory factors by *B. salamandivorans* is limited to mature zoosporangia and can occur over a range of temperatures between 16°C and 26°C. Taken together, these results suggest that both pathogenic *Batrachochytrium* fungi have evolved similar mechanisms to inhibit lymphocytes in order to evade clearance by the amphibian immune system.

**KEYWORDS** amphibian declines, *Batrachochytrium dendrobatidis*, *Batrachochytrium salamandivorans*, chytrid, immunomodulation, kynurenone, lymphocyte, methylthioadenosine, polyamine, spermidine

Chytridiomycosis is a skin disease caused by two pathogenic chytrid fungi, *Batrachochytrium dendrobatidis* and *B. salamandivorans*. *Batrachochytrium dendrobatidis* was first described in 1998 (1–3) and is a globally distributed pathogen associated with declines of hundreds of amphibian species (4). *Batrachochytrium salamandivorans* was first described in 2013 and is thought to be native to parts of Asia but was lethal when accidentally introduced into populations of fire salamanders (*Salamandra salamandra*) in northern Europe (5–7). Although published information about the number of amphibian species tested for susceptibility to *B. salamandivorans* is limited, this pathogen appears to cause greater disease in urodeles (salamanders and newts) than in anurans (frogs and toads) (6, 8, 9). Previous studies showed that *B. dendrobatidis* releases soluble factors that can inhibit both amphibian and mammalian lymphocyte proliferation, induce apoptosis of lymphocytes and epithelial cells (10, 11), induce apoptosis in frog skin (12), and inhibit a delayed type of hypersensitivity (DTH) response in frog skin (13). The supernatants were shown to contain at least three small immunomodulatory metabolites, kynurenone (KYN), methylthioadenosine (MTA), and spermidine (SPD) (14, 15). Studies designed to examine an immune response to *B. salamandivorans* in salamanders showed that prior exposure to this pathogen followed by heat-induced clearance

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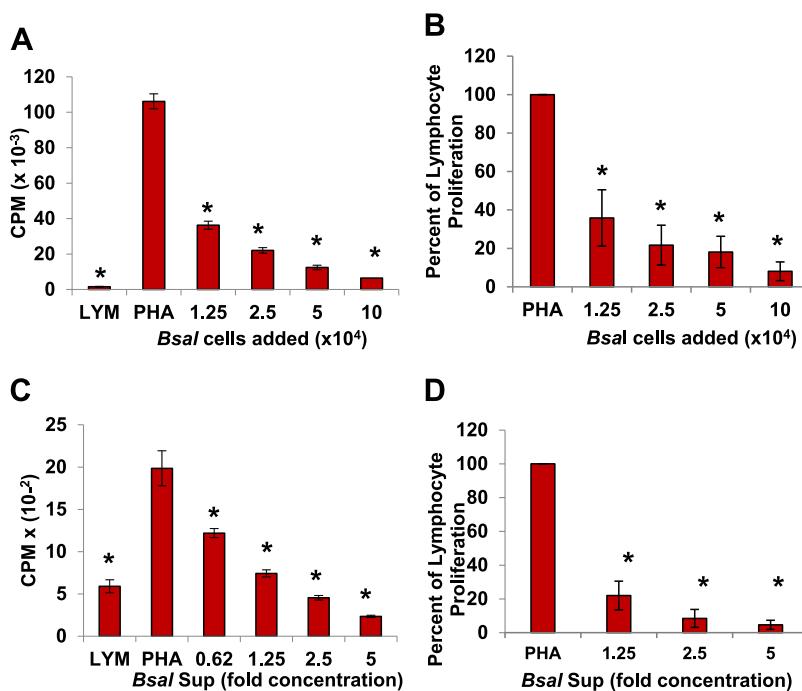
did not appear to protect from subsequent exposures, suggesting that adaptive immunity is slow to develop or absent (7, 16). This suggests that *B. salamandivorans*, like its most closely related sister species, *B. dendrobatidis*, has evolved mechanisms to evade immune recognition and clearance. Here, we investigated whether *B. salamandivorans* also directly inhibits amphibian lymphocyte functions and produces a similar suite of inhibitory metabolites as those produced by *B. dendrobatidis*.

## RESULTS

**Effects of *B. salamandivorans* cells and supernatants on proliferation of amphibian lymphocytes.** To examine the effects of *B. salamandivorans* zoosporangia on activated amphibian lymphocytes, increasing numbers of zoosporangia were cocultured with the lymphocyte-enriched fraction of splenocytes (LYM) from *Xenopus laevis* that were induced to proliferate with the plant lectin, phytohemagglutinin (PHA). Frogs do not have lymph nodes, and thus, the greatest source of lymphocytes is the spleen. PHA specifically induces proliferation of the T lymphocyte subset (17). When  $1.25 \times 10^4$  zoosporangia were added to  $10^5$  lymphocytes (a ratio of about 1 zoosporangium to 8 lymphocytes), proliferation of the lymphocytes was consistently and significantly reduced by approximately 75% (\*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test). As the number of added zoosporangia was increased, proliferation was reduced in a dose-dependent fashion and was almost completely inhibited when zoosporangia and lymphocytes were cultured in equal numbers (Fig. 1A and B; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test). To determine whether the inhibition of lymphocyte proliferation was due to direct cell-to-cell signaling, we prepared and tested cell-free supernatants from the *B. salamandivorans* cultures (*B. salamandivorans* Sups). The supernatant factors were also significantly inhibitory to proliferating lymphocytes. At a 5-fold concentration of the supernatants, proliferation of lymphocytes was almost completely prevented. Even at a 1.25-fold concentration of the supernatants, proliferation of the lymphocytes was reduced by greater than 75% (see Materials and Methods for supernatant preparation) (Fig. 1C and D; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test.). Although live fungal cells were very effective inhibitors of activated T lymphocytes, dead zoosporangia also inhibited proliferating lymphocytes. However, it required at least one fungal cell for each two lymphocytes (Fig. 2A and B; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test). This suggested that the heat-killed fungal cells are still capable of delivering an inhibitory signal.

*Batrachochytrium salamandivorans* is transmitted by a swimming flagellated zoospore that differentiates into the mature zoosporangium within the skin (5, reviewed in reference 18). The zoospores have been shown to express protease genes (16), and thus, it was important to test whether zoospores, as well as mature zoosporangia, could inhibit *Xenopus* lymphocyte proliferation. In contrast to the mature cell and cell-free supernatants of zoosporangia, purified zoospores did not inhibit lymphocyte proliferation in coculture assays (Fig. 3A and B).

**Effects of temperature on capacity of *B. salamandivorans* zoosporangia to inhibit lymphocyte proliferation.** *Xenopus laevis* lymphocyte proliferation is optimal at about 26°C, and thus, the initial coculture assays with live *B. salamandivorans* were conducted at this temperature (Fig. 1A and B). However, 26°C is likely to be a very stressful temperature for this cold-loving pathogen (5). Therefore, the capacity of the zoosporangia to inhibit lymphocyte proliferation was also tested at both 16°C and 21°C. In a series of coculture experiments, we tested the capacity of *B. salamandivorans* zoosporangia to inhibit proliferation at all three temperatures, 16°C, 21°C, and 26°C. In a representative coculture experiment with lymphocytes from one large frog, significant inhibition was observed at all three temperatures in a similar dose-dependent fashion (\*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test). It should be noted that the lymphocyte proliferation is highly temperature dependent, and thus, the maximum proliferation was very different at each temperature (Fig. 4A to C). Although results were variable, multiple-replicate coculture experiments at the three different temperatures suggested that the warmer temperatures of 21°C and 26°C resulted in a greater level of inhibition (Fig. 4D).

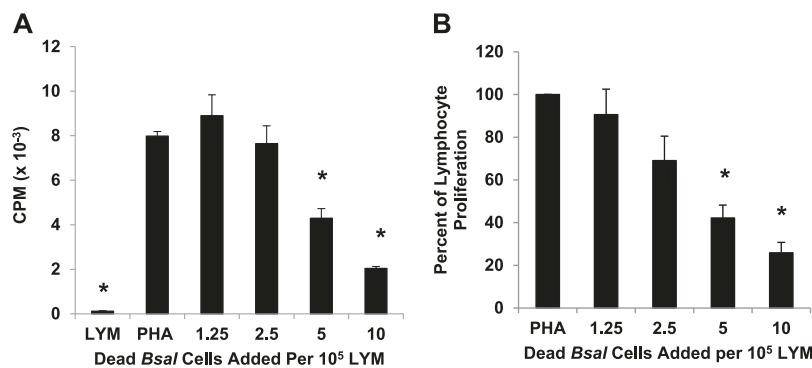


**FIG 1** *B. salamandrovorans* (*Bsal*) zoosporangia and supernatants inhibit proliferation of amphibian lymphocytes. (A) Lymphocytes (LYM) ( $10^5$ /well) from *X. laevis* were cultured alone or with phytohemagglutinin (PHA). PHA-stimulated lymphocytes were cultured alone or with increasing numbers of *B. salamandrovorans* zoosporangia. One representative experiment of six. \*, significantly reduced [ $^3$ H]-thymidine uptake detected as counts per minute (CPM) compared to control PHA-stimulated lymphocytes. (B) Summary of six experiments reported as percentage of proliferation of lymphocytes with added *B. salamandrovorans* zoosporangia in comparison with [ $^3$ H]-thymidine uptake by PHA-stimulated control lymphocytes. (C) LYM ( $10^5$ /well) from *X. laevis* were cultured alone or with PHA. PHA-stimulated lymphocytes were cultured alone or with increasing concentrations of *B. salamandrovorans* supernatants (*Bsal* Sup). One representative experiment of four. \*, significantly reduced [ $^3$ H]-thymidine uptake detected as CPM compared to control PHA-stimulated lymphocytes. (D) Summary of four experiments reported as percent proliferation of lymphocytes with added *Bsal* Sup in comparison with [ $^3$ H]-thymidine uptake by PHA-stimulated control lymphocytes. For all panels, comparisons were made to examine differences from PHA-stimulated lymphocytes. \*, significant differences with  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test. Lymphocytes and zoosporangia were cultured at 26°C.

**Effects of *B. salamandrovorans* supernatants on proliferation of a human T cell line.** To determine whether the inhibition of lymphocytes was limited to amphibian cells, the capacity of *B. salamandrovorans* Sups to affect the survival and viability of a human T cell line, Jurkat T cells were also tested. Similar to the effects of the supernatants on amphibian lymphocytes, *B. salamandrovorans* Sups also significantly inhibited the survival of Jurkat T cells at concentrations greater than 1.25-fold (\*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test). There was greater than 95% reduction of survival at the highest concentration tested (5-fold) (Fig. 5A and B).

**Comparison of *B. salamandrovorans* and *B. dendrobatidis* inhibition.** Because we hypothesized that both pathogens may use similar immune evasion strategies, the zoosporangia from each pathogen were tested in equal numbers for inhibition of the same sets of *Xenopus* lymphocytes incubated at 21°C, a temperature that is within the normal range of both pathogens (5, 19). In three experiments, inhibition by *B. dendrobatidis* zoosporangia was significantly greater than that of *B. salamandrovorans* on a per-cell basis for several concentrations of added cells (Fig. 6A; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey *post hoc* test, and Fig. 6B, significantly different by multiple comparison of means with Tukey *post hoc* test; \*,  $P < 0.05$ ; #,  $P = 0.0668$ ).

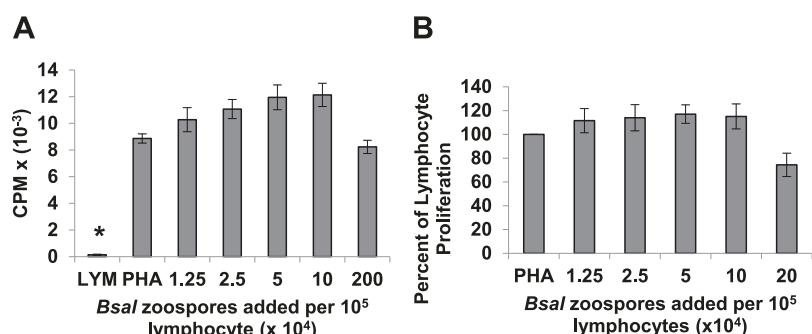
**Detection of immunomodulatory metabolites in *B. salamandrovorans* cell-free supernatants.** Since *B. salamandrovorans* Sups showed lymphocyte inhibition similar to that observed previously with *B. dendrobatidis* supernatants (*B. dendrobatidis* Sups), the concentrated *B. salamandrovorans* Sups were analyzed by high-performance liquid



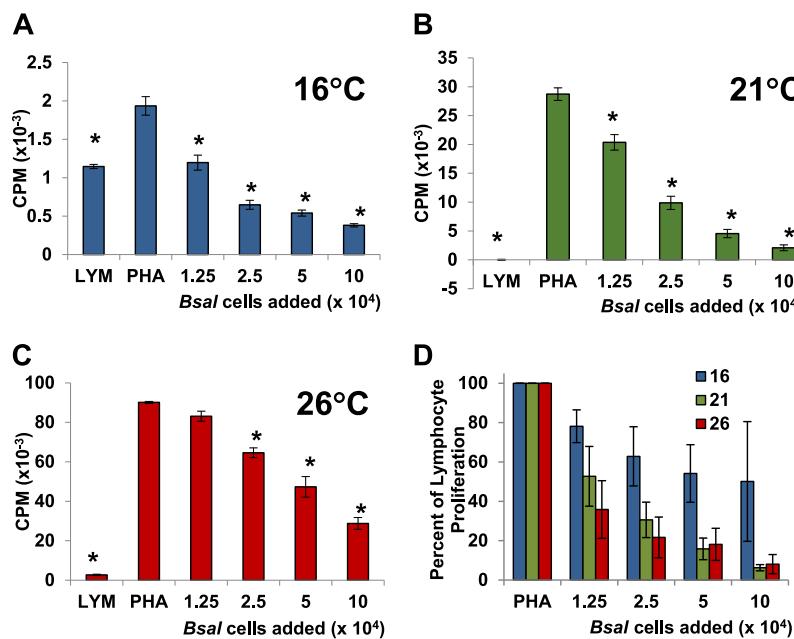
chromatography (HPLC), UV-visible spectroscopy (UV-Vis), and mass spectrometry (MS) using multiple-reaction monitoring (MRM) (see Materials and Methods). Three metabolites previously found in *B. dendrobatidis* Sups were also detected in *B. salamandrovorans* Sups generated at  $16^\circ\text{C}$  or  $21^\circ\text{C}$ . Cultures maintained at  $26^\circ\text{C}$  could not be maintained long enough to develop the high numbers of cells needed to produce *B. salamandrovorans* Sups at this temperature. The metabolites found in *B. salamandrovorans* Sups included high levels of the amino acid tryptophan, methylthioadenosine (MTA), and kynurenone (KYN) (Fig. 7). A third metabolite previously found in *B. dendrobatidis* Sups is the polyamine spermidine (SPD). SPD cannot be visualized with UV-Vis spectroscopy unless it is conjugated with a light-absorbing chromophore such as a dansyl group. Therefore, the *B. salamandrovorans* Sups were treated with dansyl chloride and then analyzed (see Materials and Methods). Both supernatants generated at  $16^\circ\text{C}$  and  $21^\circ\text{C}$  also contained spermidine (Fig. 8).

## DISCUSSION

**Lymphocyte inhibition by *B. salamandrovorans* cells and cell-free metabolites.** Our results showed clearly that *B. salamandrovorans*, like its closest fungal relative, *B.*

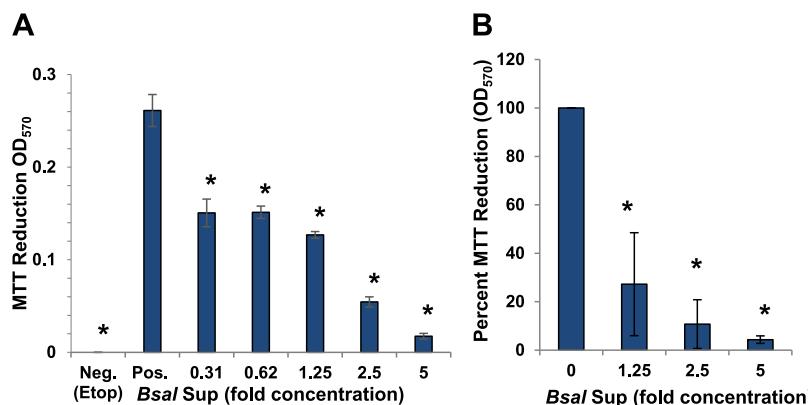


**FIG 3** In coculture, purified *B. salamandrovorans* (*Bsal*) zoospores do not inhibit proliferation of PHA-stimulated amphibian lymphocytes. (A) Lymphocytes (LYM) ( $10^5$ /well) from *X. laevis* were cultured alone or with phytohemagglutinin (PHA). PHA-stimulated lymphocytes were cultured alone or with increasing numbers of purified *B. salamandrovorans* zoospores. One representative experiment of five. Only unstimulated control lymphocytes showed less proliferation than PHA stimulated lymphocytes; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey post hoc test. (B) Summary of five experiments reported as percentage of lymphocyte proliferation with added *B. salamandrovorans* zoospores in comparison with [ $^3\text{H}$ ]-thymidine uptake by PHA-stimulated control lymphocytes. Lymphocytes and zoospores were cultured at  $26^\circ\text{C}$ .

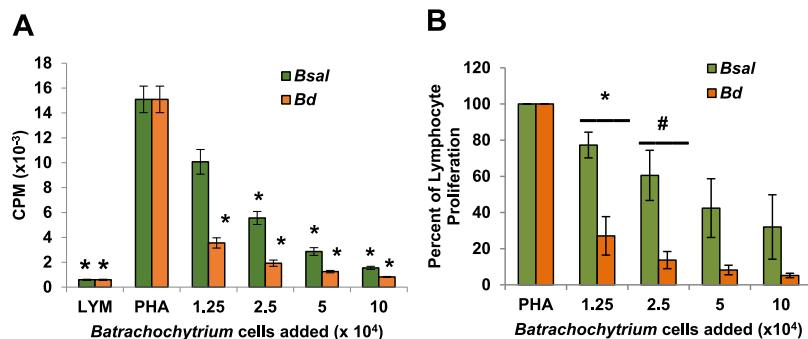


**FIG 4** *B. salamandrovorans* (*Bsal*) zoosporangia cocultured with PHA-stimulated amphibian lymphocytes inhibit proliferation of lymphocytes at a wide range of temperatures. (A to C) Lymphocytes (LYM) ( $10^5$ /well) from one *X. laevis* were cultured alone or with phytohemagglutinin (PHA). PHA-stimulated lymphocytes were cultured alone or with increasing numbers of *B. salamandrovorans* zoosporangia at 16°C (A), 21°C (B), and 26°C (C). \*, significantly reduced [ $^3$ H]-thymidine uptake detected as counts per minute (CPM) compared to control PHA-stimulated lymphocytes; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey post hoc test. (D) Summary of three experiments using individual frogs at 16°C, five experiments at 21°C, and six experiments at 26°C reported as percent proliferation of lymphocytes with added *B. salamandrovorans* zoosporangia in comparison with [ $^3$ H]-thymidine uptake by PHA-stimulated control lymphocytes.

*dendrobatis*, inhibited amphibian lymphocytes in coculture, and the factors produced by this fungus inhibited both amphibian and mammalian lymphocytes. The inhibitory molecules included kynurenone (KYN), methylthioadenosine (MTA), and spermidine (SPD). It should be noted that MTA is a by-product of spermidine synthesis, and the two can act together to enhance amphibian lymphocyte inhibition (15).

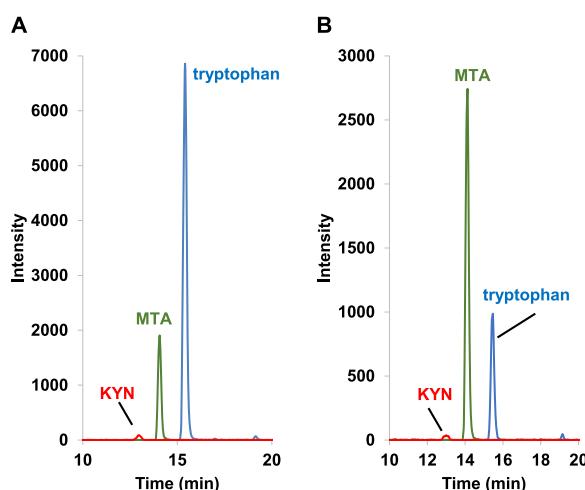


**FIG 5** *B. salamandrovorans* supernatants (*Bsal* Sup) inhibit proliferation of Jurkat T cells. (A) Jurkat T cells ( $2 \times 10^5$ /mL) were cultured alone (Pos.) as a positive control for growth or with increasing concentrations of *B. salamandrovorans* Sup as shown or with 25  $\mu$ g/mL etoposide (Etop) as a negative control for growth. Survival was measured as 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide (MTT) reduction after 3 days. (A) One representative of two similar experiments (Sup generated by cells cultured at  $10^7$ /mL). \*, proliferation was significantly reduced determined by one-way analysis of variance, with Tukey post hoc test;  $P < 0.01$ . (B) Average of two similar experiments as percentage of positive growth in control wells (Sup generated by cells cultured at  $10^7$ /mL or  $10^8$ /mL). \*, differences were significant by one-way analysis of variance with Tukey post hoc test;  $P < 0.05$ .

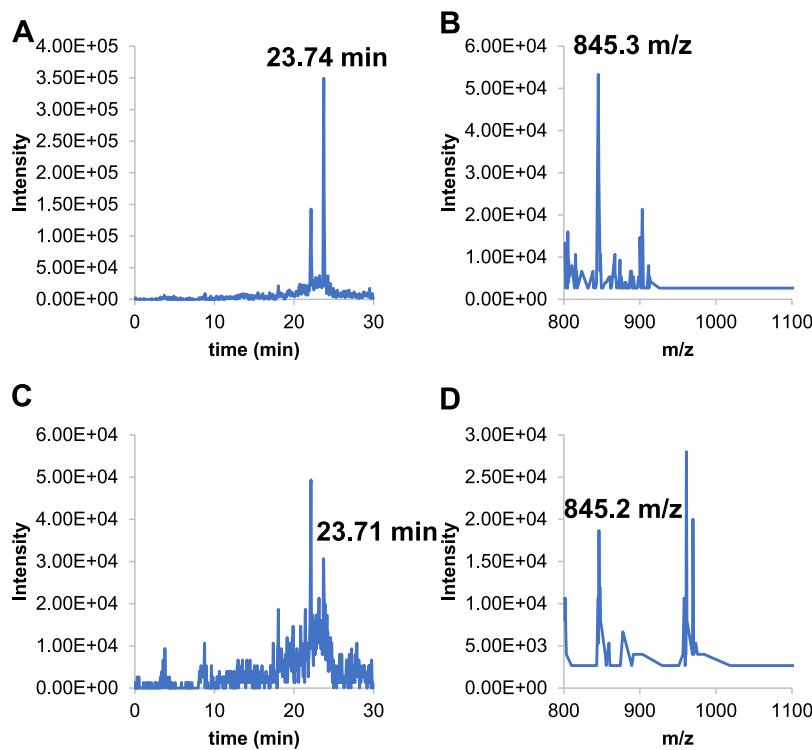


**FIG 6** *B. salamandivorans* (*Bsal*) zoosporangia inhibit less well than *B. dendrobatidis* (*Bd*) zoosporangia at a common temperature. (A) Lymphocytes (LYM) ( $10^5$ /well) from *X. laevis* were cultured alone or with phytohemagglutinin (PHA). PHA-stimulated lymphocytes were cultured alone or with increasing numbers of *B. salamandivorans* or *B. dendrobatidis* zoosporangia. One representative experiment of three. \*, significantly reduced [<sup>3</sup>H]-thymidine uptake detected as counts per minute (CPM) compared to control PHA-stimulated lymphocytes; \*,  $P \leq 0.01$  by one-way analysis of variance with Tukey post hoc test. Lymphocytes and zoosporangia were cultured at 21°C. (B) Summary of three independent experiments reported as percentage of proliferation of lymphocytes with added *B. salamandivorans* or *B. dendrobatidis* zoosporangia in comparison with [<sup>3</sup>H]-thymidine uptake by PHA-stimulated control lymphocytes. Significantly different by multiple comparison of means with Tukey post hoc test; \*,  $P < 0.05$ ; #,  $P = 0.0668$ .

An effective immune response to a fungal pathogen in the skin would likely involve detection by resident antigen-presenting cells such as macrophages or Langerhans cells. Damaged skin cells would trigger activation of macrophages and neutrophils. The antigen-presenting cells would also recruit effector T lymphocytes. The T lymphocytes would proliferate and release cytokines to recruit additional phagocytes to clear the pathogen. Thus, it is thought that T cells are critical for effective antifungal responses (reviewed in reference 20). However, many fungal pathogens have evolved molecular defenses to evade immune responses (reviewed in reference 21). Here, we showed that *B. salamandivorans* released at least three small metabolites that have previously been shown to inhibit T lymphocytes from amphibians and mammals (14, 15). These factors target fundamental processes that are shared by both amphibian and human lymphocytes. Previously, we and others showed that *B. dendrobatidis* induced apoptosis of frog lymphocytes and frog epithelial and skin cells (10–12). Thus, it is likely that the factors released by *B. salamandivorans* may also induce apoptosis of lymphocytes and epithelial cells in the local environment of the skin. Cells undergoing apoptosis release a pool of small molecules, including



**FIG 7** Mass spectrometry with multiple-reaction monitoring (MRM) of *B. salamandivorans* supernatants to detect immunomodulatory compounds. Mass spectrometric detection of protonated kynurenine (red; 209→146 m/z), MTA (green; 298→136 m/z), and tryptophan (blue; 205→188 m/z) in *B. salamandivorans* supernatants cultured at 16°C (A) and 21°C (B).



**FIG 8** Positive-ion electrospray ionization-mass spectrometry (ESI-MS) detection of tridansylated spermidine (expected retention time, 23.7 min; expected mass to charge [ $m/z$ ], 845.3) in *B. salamandrovorans* supernatant samples from chytrids grown at 16°C (A and B) and at 21°C (C and D). (A) Extracted-ion chromatogram (XIC) of a 16°C sample showing detection of a compound eluting at 23.74 min. (B) Mass spectrogram of the compound eluting at 23.74 min in the 16°C sample with  $m/z$  of 845.3. (C) XIC of 21°C sample showing detection of a compound eluting at 23.71 min. (D) Mass spectrogram of the compound eluting at 23.71 min in the 21°C sample with  $m/z$  of 845.2.

spermidine that induces anti-inflammatory responses, including production of the immunosuppressive cytokine interleukin 10 (22). Thus, the metabolites released by *B. salamandrovorans*, as well as metabolites released by cells undergoing apoptosis, may create a local immunosuppressive environment that permits the fungus to survive.

**Immune responses of other amphibians to *B. salamandrovorans*.** Because *B. salamandrovorans* was detected and described more recently than *B. dendrobatidis*, less is known about the immune responses of amphibians against this pathogen than those against its sister species; however, initial data are not encouraging in terms of a protective immune response. One small study, designed to demonstrate protection of fire salamanders (*Salamandra salamandra*) after multiple rounds of infection and clearance with heat, showed no significant difference in probability of infection or infection burden detected by quantitative PCR (qPCR) (7). Accordingly, fire salamanders surviving in the wild in areas where the pathogen is still present remain very susceptible (7). A further study of the transcriptome of the skin of another salamander species, the Wenxian knobby newt (*Tylopolypus wenxianensis*), showed very little induction of immune response genes after *B. salamandrovorans* infection, whereas *T. wenxianensis* did respond by activation of immune genes against *B. dendrobatidis* (16). Other recent studies of the responses of eastern newts (*Notophthalmus viridescens*) to *B. salamandrovorans* demonstrated that this species is highly susceptible to lethal disease following experimental exposures (6, 23). Although *N. viridescens* appears to have some defensive factors and inhibitory bacteria in the skin secretions, these defenses did not sufficiently protect from lethal infections (6, 23). Studies of the skin transcriptome of *N. viridescens* infected with *B. salamandrovorans* showed that some immune genes were upregulated relative to control skin (e.g., genes involved in interferon signaling and antigen presentation), while other

genes involved in T cell differentiation and signaling were downregulated (24). This demonstrated that the adaptive immune system was engaged, but the response was not effective. Thus, some studies suggested that there was little or no immune system activation following *B. salamandrivorans* infection in two salamander species, while the eastern newt study suggested immune activation that may have been dysregulated. Failure to respond and clear chytrid infections could be due to the stealthy infection mode by germ tube invasion demonstrated for *B. dendrobatidis* (25, 26) and presumed to occur in *B. salamandrivorans* (5). By injecting the contents of the encysted zoospore directly into the cytoplasm of the infected skin cells, the fungus may evade detection by patrolling antigen-presenting cells. Dying skin cells undergoing apoptosis (11, 12) may release some small metabolites, including spermidine that is immunosuppressive (22). However, our current data support the hypothesis that *B. salamandrivorans* appears to release inhibitory metabolites that have the potential to locally impair immunity in the skin. Thus, *B. salamandrivorans*, like *B. dendrobatidis*, has counterdefensive capabilities.

**Possible effects of infection by both *Batrachochytrium* pathogens.** Because *B. dendrobatidis* is widespread globally, it is likely that accidental introduction of *B. salamandrivorans* into amphibian communities will result in a dual infection with both chytrid pathogens in the same hosts. Here, we compared the capacity of *B. dendrobatidis* and *B. salamandrivorans* to inhibit lymphocytes from the same frog in several replicate experiments. In our studies, the *B. dendrobatidis* zoosporangia were more suppressive at comparable ratios of chytrid cells to lymphocytes, suggesting that production or release of the inhibitory factors may differ in the two pathogens. Alternatively, because *B. salamandrivorans* prefers somewhat cooler temperatures (15°C is optimal) (5), 21°C may have been more favorable for production and release of inhibitory factors by *B. dendrobatidis*.

In animal studies, experimental coinfection of eastern newts with both *B. dendrobatidis* and *B. salamandrivorans* demonstrated greater mortality in the coinfecting hosts than those infected by a single pathogen (27), raising the possibility that dual infection is more harmful. Related transcriptomic analysis of the skin and spleen showed differential gene expression patterns in the skin and spleen of coinfecting newts in comparison with singly infected or control newts. Immune suppression was suggested in the coinfecting newts (24). Another recent study of urodele species suggested that prior infection with nonvirulent isolates of global panzootic lineages of *B. dendrobatidis* (*B. dendrobatidis* GPL) had variable effects on susceptibility to *B. salamandrivorans*, depending on the species. That is, marbled newts (*Triturus marmoratus*) appeared to be protected, but fire salamanders were not (28). Much more research is needed to further understand the immune evasion capabilities of each *Batrachochytrium* pathogen. If coinfections occur, it is likely that environmental temperature will affect growth of each pathogen, immune defenses of the host, and counterdefenses of the chytrids.

## MATERIALS AND METHODS

**Culture of *Batrachochytrium* pathogens and enrichment of zoospores.** *Batrachochytrium dendrobatidis* isolate JEL 197 was cultured in 1% tryptone broth (T-broth) at 21°C and subcultured twice weekly as previously described (10, 29). *Batrachochytrium salamandrivorans* isolate AMFP13/1 was a generous gift of An Martel and Frank Pasmans, Ghent University, and was cultured and maintained in tryptone-gelatin hydrolysate-lactose (TGhL) broth (16 g tryptone, 4 g gelatin hydrolysate, and 2 g lactose in 1 L distilled water). The culture was incubated at 16°C and passaged once or twice weekly, depending on the observed rate of growth of cells. Zoospores were purified as previously described by washing the agar surface of 5- to 7-day-old cultures of *B. dendrobatidis* growing on 1% tryptone agar at 21°C or *B. salamandrivorans* growing on TGhL agar plates at 16°C three times using 3 to 5 mL of sterile broth (10, 29). The combined broth containing zoospores was passed over sterile nylon spectra/mesh filters (BioDesign Inc. of New York, Carmel, NY, USA) of 20-μm mesh opening size to remove mature zoosporangia.

**Preparation of fungal supernatants.** Concentrated cell-free supernatants were prepared as previously described (10, 15). Briefly, a large volume of cells was cultured in TGhL broth at 16°C to ensure there would be sufficient material. These cells were grown for about 7 to 10 days, counted, and resuspended in sterile distilled water at 10<sup>7</sup> cells/mL (unless indicated in figure legends), counting only mature zoosporangia. After a 24-h incubation at 16°C, the cells were centrifuged, and supernatants were passed through 0.2-μm filters to remove any remaining cells. Supernatants were frozen and lyophilized. Lyophilized supernatants were resuspended at 1/10 of the original volume and diluted in culture to

achieve 0.625-fold to 5-fold concentration above the original concentration. For addition of supernatant factors to cultures of lymphocytes, the lyophilized supernatants were resuspended in complete Leibovitz-15 (L-15) medium, which had been adjusted to amphibian osmolarity and supplemented with 100 IU/mL penicillin, 100  $\mu$ g/mL streptomycin, 12.5 mM sodium bicarbonate, 50  $\mu$ M 2-mercaptoethanol, 2 mM L-glutamine, and 1% heat-inactivated fetal calf serum.

**Frog lymphocyte culture.** Splenic lymphocytes from *Xenopus laevis* were obtained and enriched over a Ficoll gradient as previously described (10). Briefly, the lymphocytes were cultured in complete L-15 medium at a density of  $5 \times 10^5$ /mL ( $10^5$  per well, replicates of five or six per parameter tested), and T lymphocytes were induced to proliferate by addition of phytohemagglutinin (PHA) at a final concentration of 2  $\mu$ g/mL. Cells were incubated at temperatures ranging from 16°C to 26°C in the presence of 5% CO<sub>2</sub>-95% air for 3 days and were pulsed with 0.5  $\mu$ Ci [<sup>3</sup>H]-thymidine (5  $\mu$ Ci/mL; specific activity, 2 Ci/mmol) (Perkin Elmer, Waltham, MA, USA) during the last 24 h prior to harvesting. Proliferation was measured by uptake of [<sup>3</sup>H]-thymidine in the last 24 h and recorded as counts per minute (CPM). Use of animals was approved by the Institutional Animal Care and Use Committee (IACUC) of Vanderbilt University School of Medicine.

**Coculture of lymphocytes with *B. salamandrovorans* or *B. dendrobatidis* zoosporangia or supernatant factors.** To determine the effects of coculture of cells from *B. salamandrovorans* or *B. dendrobatidis* or cell-free supernatants of *B. salamandrovorans* (*B. salamandrovorans* Sup) on frog lymphocyte proliferation, lymphocytes were cultured at a density of  $5 \times 10^5$ /mL ( $10^5$  per well, five or six replicates) alone or stimulated with PHA (2  $\mu$ g/mL). PHA-stimulated cells received the addition of various numbers of maturing zoosporangia or zoospores ranging from  $1.25 \times 10^4$  to  $1 \times 10^5$  per well or with *B. salamandrovorans* supernatant at a 0.31- to 5-fold concentration of the supernatant. For each set of cells or supernatants tested, there were five or six replicate cultures. Control wells contained lymphocytes only with no addition of PHA (negative) or lymphocytes with PHA but no added cells (positive). For experiments using dead *B. salamandrovorans* zoosporangia, the cells were killed by heat treatment at 60°C for 10 min (10). Proliferation was measured by uptake of [<sup>3</sup>H]-thymidine in the last 24 h of a 3-day culture and recorded as CPM.

**Detection of the presence of inhibitory metabolites in cell-free supernatants of *B. salamandrovorans*.** Cell-free supernatants prepared in the Vanderbilt laboratory were lyophilized and sent to the Villanova laboratory for chemical identification of potential immunomodulatory components as previously described (14, 15). Briefly, a lyophilized cell-free supernatant (CFS) was reconstituted in 1.0 mL of HPLC-grade methanol and analyzed by liquid chromatography-mass spectrometry (LCMS) (Shimadzu LC-20 liquid chromatograph equipped with an ACE C<sub>18</sub> column [3  $\mu$ m, 150 by 4.6 mm], a Shimadzu SPD-M20A diode array detector, and an Applied Biosystems Sciex API 2000 triple quadrupole mass spectrometer). After elution (ramping from acidified H<sub>2</sub>O to acidified acetonitrile as per Umile et al. [30]), the chromatogram was compared to that of lyophilized water and commercial standards of KYN, MTA, and phenylalanine.

The mass spectrometric detection of MTA, kynurenone, and tryptophan was accomplished by building a multiple-reaction monitoring (MRM) method to isolate both the molecular ion [M+H]<sup>+</sup> and a significant fragment ion. The following terms apply to the Sciex mass spectrometry source optimization and output. DP refers to declustering potential; FP refers to focusing potential; EP refers to entrance potential; CE refers to collision energy; CXP refers to collision cell exit potential. Tryptophan (205→188 m/z) shows the loss of an amine (DP = 80; FP = 400; EP = 10, CE = 25; CXP = 3). MTA (298→136 m/z) demonstrates a probable loss of the methythiopentose ring (DP = 80; FP = 400; EP = 10, CE = 25; CXP = 3). Kynurenone (209→146 m/z) has a loss of the amine followed by the loss of a carboxylic acid (DP = 60; FP = 400; EP = 10, CE = 35; CXP = 3).

SPD present in *B. salamandrovorans* cell-free supernatant was derivatized as previously reported (15); in a scintillation vial, 100  $\mu$ L of supernatant was mixed with 200  $\mu$ L saturated sodium carbonate and 400  $\mu$ L of a dansyl chloride solution (7.5 mg/mL; 28 mM) in acetone. Reactions were stirred at 40°C in the dark for 1 to 2 h for a complete reaction. After the reaction was completed, 400  $\mu$ L of 28 mM aqueous proline was added to quench any unreacted dansyl chloride. Dansylated products were extracted with dichloromethane, the dichloromethane was evaporated under a gentle air stream, and the residue was reconstituted in 1 mL of methanol for LCMS analysis as above.

**Statistical analysis.** Differences in lymphocyte proliferation were assessed by one-way analysis of variance (ANOVA) with Tukey *post hoc* tests using the VassarStats online program (<http://vassarstats.net/anova1u.html>). For most experiments, control lymphocytes were cultured in sets of five or six replicate wells, and each experimental parameter (*B. salamandrovorans* cells or *B. salamandrovorans* supernatant concentrations added) was tested in sets of five or six replicate wells. A *P* value of  $\leq 0.05$  was considered to be statistically significant.

**Data availability.** All data needed to understand these results are summarized in the figures and figure legends. Further details about the data are available upon request.

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