**Methods**

Study Area

Summers (May-September) are relatively dry, with seasonal drought and temperatures ranging from 15°C-25°C, and winters (October-April) are wet with temperatures ranging from -5°C-11°C. The average precipitation for the area is 720 mm (Northwest River Forecast Center, NOAA).

Because of the distinct seasonal patterns (described in the paragraph you will), eastern Cascadesfgpplp-

A major defoliator of the coniferous forests of Central Washington, as well as western North America in general (Senf et al. 2016), is the western spruce budworm (WSB) (*Choristoneura freemani)*—a native lepidopteran that ranges from Southern British Columbia to Arizona and New Mexico (Fellin and Dewey, 1982). These insects emerge during budburst around mid-May to feed on the new growth of short needle conifers, specifically Douglas fir (*Pseudotsuga menziesii*) and grand fir (*Abies grandis*) in my study area, until late June or early July. They then pupate and emerge as adults, taking flight around mid to late July for oviposition. Larvae then emerge the following year in mid-May to repeat their life cycle. In a more natural fire regime that maintained an open forest structure, WSB outbreaks would occur about once every decade. In recent years, thicker forests from fire suppression and increased drought stress from climate change has created conditions that encourage more frequent and further spreading WSB outbreaks (Willis et al, 2008; Lovett et al, 2006). This shift in forest structure and herbivore behavior has the potential to change forest ecosystem dynamics with implications for forest-stream connectivity. Furthermore, the cold weather that would have normally killed off pests in the past is occurring less often,. This allows these pests to stay out longer, causing more damage to plants more often than they otherwise would (Griffin and Turner, 2012). It has also been suggested that pest outbreaks can lead to increased fires due to the dead and dying trees they leave behind (Schlesinger et al, 2015), but new research has shownthat this may not be the case, and in fact may have the opposite effect. These insects are defoliators as opposed to wood burrowers and therefore potentially have a different effects on ecosystem dynamics.

The low budworm sites for this study were located in the Teanaway Community Forest in Washington State, approximately 40 miles X direction from Central Washington University on public land. These sites were located near creeks called: Stand Up Creek (903 m above sea level), Jungle Creek (824 meters above sea level), Jack Creek (963 meters above sea level), and Moonbeam Creek (973 meters above sea level). The high budworm sites were located in the Swauk drainage in the Okanogan-Wenatchee National Forest in Washington State approximately 45 miles X direction from Central Washington University and east of the low budworm sites, also on public land. The creeks near these sites were called: Cougar Creek (984 meters above sea level), Hurley Creek (978 meters above sea level), Hovey Creek (1050 meters above sea level), and Blue Creek (1055 meters above sea level). Although each individual site varied based on microclimatic factors, sites were exposed to fairly similar temperature and precipitation patterns.

I used a nested study design with repeated sampling through time to investigate how budworm herbivory influenced throughfall composition, litter decomposition, and soil nutrient dynamics. Within each budworm herbivory level (low versus high), I established study sites along 4 different streams (n=8). At each stream I established three replicate plots approximately 15 m from each other from upstream to downstream. At each replicate plot, I measured throughfall water chemistry, litter decomposition rates, and soil chemistry 8 times between early September 2015 and early November 2016. Frass collectors were also deployed at each site to measure budworm activity over the course of the study.

Throughfall

Throughfall collectors were installed under the tree canopy close to each decomposition bag line. Two rainfall collectors were set up in an area with no canopy cover in the low budworm activity sites and two in the high impacted sites as controls. During rain events, water came through the canopy into a funnel ( \_ mm) with polywool filtering out any litter that happened to fall into the aparatus. The funnel was attached to a hose with nylon mesh ( \_ μm) between the two and was held in place with parafilm. Water traveled through the tubing into an acid washed 4 L jug. Water was then transferred to an acid washed nalgene bottle and was taken to the lab, filtered through a 1.0 μm fiberglass filter using vacuum filtration. The volume of each collector was also measured in order to calculate mg of nutrient per L of water. Water samples were frozen until analysis of solutes at a later date. Throughfall collectors were taken down just before snowpack and redeployed just after snowmelt to prevent damage to the apparatus.

Frass and Litter Measurements

Frass and litter fall were also collected at each site to quantify and differentiate organic matter inputs to the forest floor. Funnels ( 0.25 m2 diameter) made of tarp and garden hose connected to a one liter Nalgene bottle were placed on the ground under trees affected by budworms. and litter, and samples were collected approximately monthly until snowpack. Upon return the laboratory, the sample was by frass versus litter Weights were then converted to a daily litter or frassfall rate by XXXXX. . Frass collectors were taken down during the winter months to prevent damage, and reinstalled on XXXX for sampling during 2016. Unfortunately, due to frequent rains in the spring months of 2016, samples decomposed before they could be collected and measured, so no data are available for the second half of the study.

Decomposition

At each replicate plot I deployed twenty 20x20cm mesh litter bags (García-Palacios et al. 2016) for a total of 480 bags across all sites. Ten bags at each site contained a mixed conifer needle sample of Douglas fir (*Pseudotsuga menziesii)*, grand fir (*Abies grandis*), and ponderosa pine (*Pinus ponderosa*), to represent the most abundant species in the study area. To make this mixed sample, I harvested needle material in an approximate ratio of 1 part ponderosa pine, two parts Douglas fir, and one part grand fir from the forest and air dried the needles until they reached constant weight. The other ten bags at each replicate plot contained sugar maple (*Acer saccharum*) leaves which are non-native to the area but are commonly used in decomposition studies for comparison across biomes.

Within each litter bag, I placed ~3-5 grams of air dried needles (Benfield, 1996) after recording the needle mass, and I added an aluminum tag with a unique ID. The bottom side of the mesh bags were made of a smaller sieve size ( 0.5 mm) (Schweitzer et al, 2005) than the top ( 2 mm) (Genung et al, 2013) to reduce the loss of its contents while still allowing small detritivores to enter the bags. Bags were assembled by stapling the two sieve sizes together and by reinforcing them with super glue at the corners. The bags stayed intact throughout the 9 month deployment. Mesh bags with leaves were subsequently placed into red peanut bags (mesh size XX Clay will get) to further protect them during deployment and to simplify sample collection, and each individual bag was placed into a Ziploc for transport to the field.

In the field, bags were strung together on an approximately 6 m line held in place by a 2 ft piece of rebar driven into the ground on either side. The rebar anchors prevented bags from being moved by the wind or displaced by hillslope runoff. A coin flip determined which bags (conifers or deciduous maple) were placed upstream and downstream at each site. Handling loss was applied to the mass of the material by deploying twenty bags, ten deciduous and ten coniferous and extracting them immediately to see if there was any mass loss from deployment and extraction. This was done separately for conifer and deciduous leaves.

Bags were collected 7 times beginning XXX and ending XXX in approximately 1-2 month intervals with a 5 month break during winter snowpack from XXX to XXX when the sites were inaccessible. On each retrieval from the field, one conifer bag and one maple bag were collected from each plot for a total of 48 bags per sampling time. Bags were randomly chosen for pickup and returned to the lab in a Ziploc bag to prevent additional leaf mass loss. On the final collection day, all remaining bags were collected from the sites (n=? per plot and leaf type).. Upon return to the lab, decomposition bags were air dried to constant mass (Schweitzer, 2005) in paper bags (Genung et al. 2013) hung on a clothesline. Once the bags were air dried, each bag was sorted to remove any noticeable debris that had become incorporated in the sample (Chapman et al. 2013). Because of natural loss of conifer needles from the canopy, it was difficult to determine what was originally in the bag and what had fallen into it, so the mass of conifer needles accumulated in the maple decomposition bags was sorted and used as a correction factor for the mass of conifer needles that entered the conifer bags.

Paragraph to describe how you took mass remaining to calculate a decomposition rate K. Cite appropriate sources.

*Soil Analyses*

Upon each collection of decomposition bags, I also used a thermocouple to measure temperature at three soil depths: 2 cm, 10 cm, 20 cm. These corresponded approximately to the O horizon, the top of the A horizon, and within the A horizon respectively. A soil core of ~10 cm depth was also collected from each replicate plot at each stream site each time I collected litter bags. Soil cores were stored on ice for return to the laboratory whereupon each core was homogenized in a Ziploc bag. Soils were immediately analyzed for moisture content and percent organic matter, and soils were frozen for later analysis of ammonia, nitrate, inorganic P, and using methods detailed below.

*Moisture Content and Percent Organic Matter:*

Soil was sieved at 2 mm and a subsample was placed into an ashed aluminum pan and weighed immediately for field mass. Pans were then placed in a drying oven at 60ºC until constant mass, cooled to room temperature, and weighed to obtain dry mass (DM). The difference between field mass and dry mass was used to calculate percent moisture. Then dried soil samples were placed in a muffle furnace at 500ºC for 48 h. to combust all organic compounds. After ashing, samples were cooled to room temperature, rehydrated with Milli-Q water to rehydrate clays and colloids containing water molecules, and then placed again into a drying oven until constant mass. Pans were cooled to room temperature and reweighed to obtain ash-free dry mass, with the difference between dry mass and ash-free dry mass used to calculate percent organic matter.

*Phosphorous Analysis*

The Bray P1 method was used to extract phosphorus from each soil sample (Hamilton, 1997; Patton and Kryskalla, 2003). One gram of air dried soil was added to 10 mLs of the Bray P1 extractant solution (30 mL 1 N NH4F to 50 mL 0.5 HCl) and shaken on a shaking table at 100 rpm for 15 minutes. The sample was then filtered with a syringe through a 1.0 µm glass fiber filter and stored in the freezer until analysis. Samples were analyzed for inorganic phosphorous using the XXX method on a Seal AQ1 Discrete Analyzer (Seal AQ1, Seal Analytical; Mequon, Wisconsin, USA) with EPA equivalent methods.

*Nitrogen Analyses*

A 2M KCl extraction method was used to extract inorganic nitrogen from each soil sample. Ten grams of air dried soil were added to 70 mLs of 2M KCl and shaken at 100 rpm for 2 hours on a shaker table. The sample was then filtered with a syringe through a 1.0 µm fiberglass filter and stored in the freezer until analysis. Samples were analyzed for NO3-+NO2- (hereafter referred to as NO3-)using the cadmium reduction and NH4+ using the phenate method on a Seal AQ1 Discrete Analyzer ((Seal AQ1, Seal Analytical; Mequon, Wisconsin, USA) using EPA equivalent methods.

Each site also contained a resin bag made of bleached nylons (to prevent color leaching that may affect results) filled with 30 g of ion exchange beads to measure changes in the DIN pool in soils throughout the deployment. and was also extracted using 2M KCl and were analyzed for NO3- and NH4+ to look at net nitrification on a Seal AQ1 Discrete Analyzer using EPA equivalent methods. These bags were deployed 10 cms deep in the soil during while initial soil samples were taken to examine the change in soil chemistry just before snowpack and just after snow melt.

**Statistical Analysis**

I used a two-sample t-test to compare the two treatments; coniferous litter vs deciduous. This was done for both the rate of decomposition and for net nitrification. I used generalized least squares (GLS) models and linear mixed effects (LME) models (Senf et al. 2016) to compare the variances within each of the 8 sites and each of the 3 replications within to compare percent soil moisture, percent organic matter, NO3-, NH4+, SRP, N:P ratio, total inorganic N, and net nitrification/mineralization. Data was analyzed using R.