BC Caribou Research Program - Mortality Site Investigation Form

	LOCATION	
Date of site visit (DD/MM/YYYY):	Date mort. signal received:	Date of death
General location:	Personnel:	
Waypoint: UTM Zone: E:	N: or	Lat:Long:
	ANIMAL IDENTIFICATION	
WLH ID: OTHER ID:	Herd:	
Found dead Euthanized Euth. method:	If Euth. collect b	lood 2 x gold top
State of decomposition (Circle): Fresh / Bloated / Active	e decay (maggots) / Desiccated /	Bones Collar only
Ear tag(s) (number and colour): Left:	_ Right:	
Collar recovered: Yes / No Freq.:	_ Ser. No.:	Circle: Functional / Damaged / Destroyed / N/A
	ORTALITY SITE DESCRIPTION	
Describe mortality site and document with pho Photos should include wide angle, medium and	•	•
	·	
Circle: No snow / Snow / Fluffy Snow depth (cm):	Ice crust □ Circle: Light/Mod	
		Pictures □ Video □
		video 🗆

Select all that	appl	y. Describe any	abno	ormal find			AL EXAM Other' in con	nme	nts s	ection on	page 3	3.			
Carcass Location	on			Body Condition			Body/Skin/Hair			Eyes			Ears/Nose		
In open		Fresh#		Excellent			Hair loss			Clear			Ear cru		
In water		Frozen		Good			Ticks/parasite	es		Swollen			Ulcers		
In cover		Decomposed		Fair			Lumps/warts	5		Cloudy			Discha	rge *	
Buried		Intact #		Poor			Wounds			Discharge	*		Other		
On roadside		Disarticulated		Emaciated	d		Besnoitia cys	ts		Other					
Collar only		Scattered		Unknown	l		eyes/legs/fac	ce		One / L / R	/ Both				
Other		Scavenged					Other								
Oral Cavity		Teeth		Bones a	and Join	nts	Hooves/	/Feet		Fe	ces		Re	productiv	ve
Ulcers/sores		Teeth worn		Fracture(s	s)		Excessive wea	ar		No feces			Lactati	ng	
Rumen content		Teeth irregular		Joints swo	ollen		Abnormal we	ear		Diarrhea			Udder	abnormal	
Blood		Teeth broken		Joint fluid	I		Overgrown			Fecal stain	ing		Vagina	ıl	
Other		Feed impacted		clear/pus/	/blood		Infection			mild/mod/	extr.		discha	rge *	
		Other		Antler def	form.		Other			Blood			Aborti	on	
				Retained	velvet					Rectal prol	apse		Testes	abnorm.	
				Other						Other			Penis		
													discha	rge *	
													Other		
* Discharge Clea	r / Clo	oudy / Purulent (Pเ	ıs) / B	lood Colle	ect sam	ples/	swabs 🗆								
# Consider sling	ing o	ut/removing inta	ct and	d fresh car	rcasses	for n	ecropsy by a	proje	ect ve	terinarian	– For h	nigh p	riority	herds	
Winter tick coun															
Hair Loss: None Before samplii intact gastroin	ng, ta	ake pictures of o	ppen	ed chest (en). Take	INT (showi	ERN ng h	AL EXAM eart and lun	igs) a	and a						
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COMMENTS FOR EXTERNAL EXAM, INTERNAL EXAM, SUSPECTED PROXIMATE vs. ULTIMATE CAUSE OF DEATH					
If animal is found alive, describe symptoms (e.g. lying down, circling, vocalizing, aggressive, dull, etc.)					
Questions in the field or the lah? Co.	ntaci	t Dr. Helen Schwantje 250-751-3234 or cell 250-361-7619			
Questions in the field of the lab.	iitac	1 Dr. Helen 301 Wange 230 731 3234 01 cen 230 301 7013			
CARIROLL TICCUE CANA	DI EC	TO COLLECT IN THE FIELD (AC AVAILABLE)			
		TO COLLECT IN THE FIELD (AS AVAILABLE)			
•	offic	ce. Post-field sub-sampling described on the following processing sheet	-		
Head (or obex and RPLN in field for CWD)		Spleen (palm size piece)			
Pictures of jaw and incisors#		Lymph nodes (if abnormal)			
Mandible left and right with incisors		Intestine (if abnormal + fresh) - Open and assess a few sections of large			
Ear tips x 2		and small intestines and abomasum. Collect parasites if found.			
Hairx100 (pluck from between shoulders)		Rumen contents (palm full)			
Intact long bone #1 (femur or humerus)		Feces (10-20 pellets from colon)			
Intact metatarsus (hind foot left or right)		Fetus and placenta			
Skeletal muscle (from leg, palm size piece)		Uterus and ovaries (collect only if abnormal)			
Lung front lobe (palm size piece, right)		Calf (if newborn and dead)			
Lung middle lobe (palm size piece, left and right)		Cysts and tumors (if unknown cause, include adjacent normal tissue)			
Lung back lobe (palm size piece, left and right)		Winter ticks (10+ all life stages, engorged and not engorged)			
Heart (full cross section of atria and both ventricles)		PREDATION SAMPLES			
Blood (heart/jugular in 2 x gold top)		* Fill out and attach predator ID data form if swabs collected*			
Whole left kidney + fat		DNA (hide/collar punctures/bite/rake wounds; Swab in field is best)			
Whole right kidney (keep separate from left kidney)		Predator hair			
Liver (palm size piece x 3 in separate bags)		Predator scat			
	<u> </u>	Other			

Take three photos of teeth and jaw – one from each side and one from the front showing the incisors

ALL SAMPLES IN SEPARATE WHIRL PAK BAGS, EACH LABELLED WITH: WLH ID, SPECIES, HERD, SEX, SAMPLE TYPE, DATE

Caribou Mortality Sample Processing and Storage

SAMPLE	PROCESSING	STORAGE
Intact head	Double heavy garbage bag/seal well	Frozen ^a
Obex and retropharyngeal	Collect and subsample as per CWD	a)Whole obex, ½ of each RPLN: Fixed
lymph nodes (RPLNs)	sampling protocols	b
lymph hodes (Kr ENS)	Sampling protocols	h) 1/ of cook DDI N. France
NA or dibloc	Disco in hone has/s//seel well	b) ½ of each RPLN: Frozen
Mandibles	Place in bone bag(s)/seal well	Frozen
2 x Ear tips	Place in 2 separate non-manila	Room temperature ^d
	envelopes (air dry) ^c	
Hair x 100	Separate non-manila envelope per body	Room temperature
	region collected (air dry)	
Long bones	Place in bone bag(s)/seal well	Frozen
Metatarsus	Place in bone bag(s)/seal well	Frozen
Skeletal muscle	Place in whirl-pak /seal well	Frozen
Lung front lobe	Subsample at office e, f	a) 2, 1 cm thick sections (1 from each
Right	- Fixed portions in 10% formalin	lobe, if abnormal take up to 4 per
	- Frozen in separate whirl-pak/seal well	lobe): Fixed
	·	b) Remaining tissue: Frozen
Lung middle lobe	Subsample at office	a) 2, 1 cm thick sections (1 from each
Left and Right	- Fixed portions in 10% formalin	lobe, if abnormal take up to 4 per
	- Frozen in separate whirl-pak/seal well	lobe): Fixed
		b) Remaining tissue: Frozen
Lung back lobes	Subsample at office	a) 2, 1 cm thick sections (1 from each
Left and Right	- Fixed portions in 10% formalin	lobe, if abnormal take up to 4 per
	- Frozen in separate whirl-pak/seal well	lobe): Fixed
		b) Remaining tissue: Frozen
Heart	Subsample at office	2, 1 cm thick sections: Fixed
	- Fixed portions in 10% formalin	
Heart blood	Place blood tubes in whirl-pak/seal well	Frozen
Left kidney + fat	Place in whirl-pak/seal well	Frozen
Right kidney	Subsample at office	a) 1-2, 1 cm thick cross sections:
	- Fixed portions in 10% formalin	Fixed
	- Frozen in separate whirl-pak/seal well	b) Remaining tissue, divided into <u>two</u>
		separate whirl-paks: Frozen
Liver		
	Subsample at office	a) 1-2, 1 cm thick cross sections:
	- Fixed portions in 10% formalin	Fixed
	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well	Fixed b) Remaining tissue, divided into three separate whirl-paks: Frozen
Spleen	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office	Fixed b) Remaining tissue, divided into
Spleen	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed
Spleen	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two
	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen
Spleen Lymph nodes	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections:
	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed
Lymph nodes	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well	Fixed b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen
	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but
Lymph nodes	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but only if fresh! Fixed
Lymph nodes Various intestine	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but
Lymph nodes	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but only if fresh! Fixed
Lymph nodes Various intestine	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but only if fresh! Fixed b) Remaining tissue: Frozen
Lymph nodes Various intestine GI parasites	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well 70% ETOH in well-sealed container	b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but only if fresh! Fixed b) Remaining tissue: Frozen Room temperature
Lymph nodes Various intestine GI parasites Rumen contents	- Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well Subsample at office - Fixed portions in 10% formalin - Frozen in separate whirl-pak/seal well 70% ETOH in well-sealed container Place in whirl-pak/seal well	Fixed b) Remaining tissue, divided into three separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue, divided into two separate whirl-paks: Frozen a) 1-2, 1 cm thick cross sections: Fixed b) Remaining tissue: Frozen a) 1-2, 1 cm thick cross sections but only if fresh! Fixed b) Remaining tissue: Frozen Room temperature Frozen

SAMPLE	PROCESSING	STORAGE
Uterus	Subsample at office	a) If abnormal, 1-3, 1 cm thick cross
	- Fixed portions in 10% formalin	sections: Fixed
	- Frozen in separate whirl-pak/seal well	b) Palm size part: Frozen
Calf	Double heavy garbage bag/seal well	Frozen
Abscesses, cysts and	Subsample at office	a) 1-2, 1 cm thick cross sections:
tumors	- Fixed portions in 10% formalin	Fixed
	- Frozen in separate whirl-pak/seal well	b) Remaining tissue: Frozen
Winter ticks ^g	Subsample at office	a) Room temperature for sample
	- Half in 70% ETOH in well-sealed	with ETOH
	container (i.e. cryovial or similar)	b) Frozen for sample without ETOH
	- Half frozen in separate cryovial without	
	ETOH	
Predator scat h	Place in whirl-pak/seal well	Frozen
Predator DNA	Collect as per double swab protocol ⁱ	Room temperature
Predator hair	Non-manila paper envelope (air dry)	Room temperature

NOTES ON HANDLING, STORING, SHIPPING SAMPLES and TISSUES

Note: Most supplies are provided by the Wildlife Health Program. Contact us before you run out.

- a) Frozen tissue samples must be stored and shipped at minimum -20°C. For long-term storage, only freeze tissue samples in whirl-paks (or similar). Do not use Ziplocs. Avoid freeze/thaw.
- b) Fixed tissue samples in 10% Neutral Buffered Formalin. Fixed tissue must be stored at room temperature in a leak proof, puncture-proof container with a 10:1 formalin: tissue ratio. Fixed tissue must not be frozen. In addition, fixed tissue must not be shipped in the same box/cooler as frozen samples, as formalin fumes can kill live pathogens which limit the efficacy of tissue culture and other diagnostics.
- c) Air dry samples at room temperature in an area protected from excessive heat (i.e. not near a stove, heater, or on a truck dashboard), light, and moisture. If samples or envelopes are wet when initially collected in the field, transfer to a fresh, dry envelope immediately on return to the lab and before leaving to air dry. Be sure to label the new envelope.
- **d)** Hair and tissue samples stored at room temperature must always be protected from heat, light, and moisture. Envelopes can be stored in a cardboard box and sent to the WLH program lab. Please do NOT stockpile dry samples.
- e) Subsampling usually requires collection of both fixed (in 10% formalin) and frozen samples.
- f) Collecting fixed tissue in 10% formalin:
- Tissues must be fixed as soon as possible after collection to preserve for microscopic exams.
- To ensure proper penetration of formalin, tissue samples must be ≤ 1 cm thick.
- To reduce artefact, always trim tissues to size using a sharp knife or scalpel on a plastic cutting board. Handle tissues carefully. Use forceps and do not crush or squeeze. Handle from the edge.
- If lesions are found, collect and fix several sections of the abnormal area. Include the edge of where abnormal meets normal tissue.
- All tissue to be fixed must be placed into a leak-proof, puncture-proof, container(s) with a 10:1 ratio of formalin: tissue.

- With the exception of intestines and CWD samples (obex and RPLNs), which should be placed in their own containers, different tissues can be fixed in the same container.
- Please record the types of fixed tissues on the container's label and also in a separate Excel file.
- REMEMBER: FORMALIN IS TOXIC. DO NOT BREATHE IT AND USE ONLY WHERE THERE IS GOOD VENTILATION. ALWAYS WEAR GLOVES AND EYE PROTECTION.

g) Winter tick - tick associated hair loss scoring in caribou

HAIR LOSS CATEGORY	PATTERN
None (No Picture)	No hair loss or breakage
Mild (Picture 1)	Few small to medium sized patches of broken hair or hair loss
Moderate (Picture 2)	Several or large patches broken hair or hair loss - NO EXPOSED SKIN
Severe (Picture 3)	Several or large patches broken hair or hair loss with 1-2 small areas exposed skin
Extreme (Picture 4)	Several or large patches broken hair or hair loss with large or > 2 areas of exposed skin

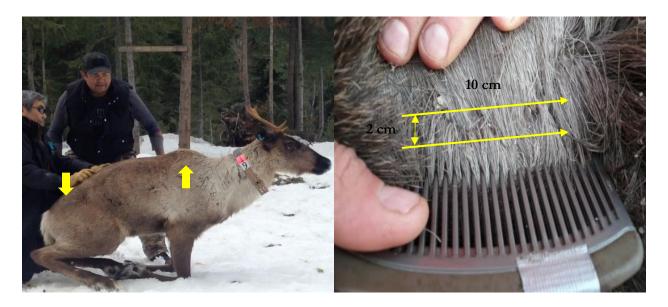
^{*}Note degree of tick associated hair loss observed in caribou is not always correlated with infestation burden.



* Photos and hair loss classification score: D. Culling, Diversified Environmental Services Inc., Fort St. John, BC.

Tick burden assessment

- Part the hair along the upper edge of the shoulder blade with a comb or ruler.
- Count the number of ticks observed along a single 10 cm x 2 cm transect.
- Part the hair along the rump.
- Count the number of ticks observed along a single 10 cm x 2 cm transect.
- If there is significant hair loss on the shoulder perform the assessment only on the rump.



- Collect a representative sample (e.g. various life stages, engorged, not engorged) of ticks.
- Record WLH ID, date of collection, management unit, and host (i.e. caribou)

For Identification Purposes:

- Store half of the ticks in a cryovial with alcohol (ethanol, 70-95%), with collection information on an external label written in pencil (or ink that is not affected by alcohol (indelible). Be sure to use enough ethanol in the cryovial, especially for engorged ticks, which will dilute the alcohol content (minimum 10:1 ratio; ethanol: ticks).
- Ensure the cryovials are well-sealed to prevent evaporation.
- Keep at room temperature, protected from heat and light.

For DNA Isolation and/or Pathogen Detection:

- Store **the other half of the ticks** frozen (-20C), <u>without</u> **ethanol** in a cryovial with collection data recorded on the outside of the cryovial as above.
- Keep in -20C freezer and ship to Wildlife Health Lab in Nanaimo, ensuring they stay frozen.

Frozen ticks and ticks in alcohol must be stored and shipped separately.

h) CAUTION: THERE IS A ZOONOTIC DISEASE RISK FROM PREDATOR SCAT - *Echinococcus* spp. tapeworms from wolf, coyote, and fox feces.

- Always wear gloves and coveralls when doing necropsies and if collecting scat.
- Collect carnivore feces with a stick or disposable utensil.
- Do not contaminate clothing, field or laboratory equipment, helicopters, trucks etc.

• Predator DNA is best obtained from the outside of scat samples. To maintain accuracy, do not crush scat (i.e. try to maintain the sample's original shape) and collect in whirl-pak(s) significantly larger than the sample itself.

i) PREDATOR DNA SWAB PROTOCOL

This protocol can be used when the predator species is unclear (i.e. predator hair and/or scat were not available to collect for DNA analysis).

Equipment needed for double swab protocol:

- Sharp knife and scalpel (with multiple, disposable blades) or disposable scalpels
- Nitrile gloves
- Sterile swabs successful identification may be decreased depending on type of swab used. Prefer individually packaged, fine tipped, cotton or poly swabs with plastic handle.
- Paper envelopes
- Whirl-paks
- Silica desiccant
- Sharpie
- Stapler and staples
- Small ethanol tubes for swab collection
- Large ethanol tube for sterilizing knife
- Kleenex/paper towel
- Lighter
- ** 95% ETOH is recommended as the best wetting agent for collecting swabs. Other wetting agents
 (i.e. denatured alcohol, isopropyl alcohol) or sterile water can be used if 95% ETOH is not available
 however, there is potential for decreased success.

If carcass present:

- Put on new nitrile gloves.
- Carefully examine the carcass for killing wounds as identified by the presence of haemorrhage.
- Take pictures of the wound (wide angle and close-up perspectives) without disturbing the wound site.
- Dip swab in the **SMALL** ethanol tube.
- Swab wound (~10 seconds).
- Place swab in non-Manila envelope and snap off shaft.
- Staple envelope shut and label envelope with Wildlife Health ID and swab number using a sharpie (e.g. xx-xxxx 1A).
- Place envelope in a whirl-pak with silica desiccant.
- Fill out Predator Identification Data Form. Attach a copy of this form to the Caribou Mortality Site Data Form
- Repeat process on the same wound with a second swab (Identified as xx-xxxx 1B)
- Change gloves.
- Identify other killing wounds or feeding wounds defined below (Identified as xx-xxxx 2A, xx-xxxx 2B, etc.) and repeat the double swab process.

If only bones and/or collar present:

• Perform the double swab procedure on any remains (including the collar) that appear to have been chewed on by predators.

Considerations: Avoid cross-contaminating wounds

- Small amounts of DNA can easily be transferred between wounds.
- Do not touch multiple wounds with the same gloves during wound identification.

Identification of wounds

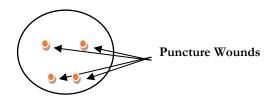
- Carefully skin the animal trying to preserve puncture marks and determining areas of hemorrhage.
- Sterilize knife after examining each wound wipe knife with new Kleenex or paper towel dip knife blade in the **LARGE** ethanol filled tube, carefully, burn ethanol off blade using lighter.

Bite description

- Killing wound is a wound that caused death or that occurred before death as indicated by hemorrhage.
- **Feeding wound** is any other wound on a carcass such as bites or chewing with no hemorrhaging or an area of the carcass that has been consumed.

Swabbing Technique

- AVOID DRENCHING THE SWAB IN BLOOD.
- For killing wound swab entire area around puncture wounds.
- For feeding wound concentrate swabbing on areas that appear to have been bitten or chewed



Predator Identification Data Form *Attach Copy to Caribou Mortality Site Data Form*					
Date (DD/MM/YYYY):					
Summary: Predator species 1: Unknown □	species 2:	species 3:			
WLH ID (for the mortality):					
Swab No. 1		Wound Location and Description			
Check boxes as swabs are completed.	Wound type (Circle) Killing Feeding				
	Other				
Swell No. 2		Pictures Pictures			
Check boxes as swabs are completed.	Wound type (Circle) Kill Feed Other	Wound Location and Description			
		Pictures □			
Swab No.3Check boxes as swabs are completed.AB	Wound type (Circle) Kill Feed Other	Wound Location and Description			
		Pictures □			
Swab No. 4 Check boxes as swabs are completed.	Wound type (Circle) Kill Feed Other	Wound Location and Description			
		Pictures			

ALL SAMPLES MUST BE RETURNED TO:

Wildlife Health Program
Attention: Dr. H. Schwantje
Ministry of Forests, Lands, Natural Resource Operations and Rural Development
2080 Labieux Road
Nanaimo BC, V9T 6J9

Phone Numbers:

Helen: (250) 751-3234 Lab: (250) 751-7246 Cait: (250) 751-3219

Shipping:

- Frozen samples MUST remain frozen during transport or their use is compromised.
- Appropriate insulated shipping containers and icepacks and can be purchased at low cost from ULINE.ca or contact the Wildlife Health Program.
- Formalin/fixed samples must be shipped separately from frozen samples. If tissues are appropriately trimmed and have been fixed for > 36 hours, excess formalin can be drained off prior to shipping (leave samples covered by a piece of paper towel wetted with a small amount of formalin).
- When shipping tissue samples in formalin ensure they are in leak-proof, puncture-proof containers and double bagged with ample absorbent material (paper towel etc.) in case of leaks.
- Please notify the Wildlife Health Lab in Nanaimo BEFORE samples are shipped

Shari Willmott, (250) 751-7246 Shari.Willmott@gov.bc.ca
Maeve Winchester, (250) 751-7246 Maeve.Winchester@gov.bc.ca
Cait Nelson, (250) 751-3219 Cait.Nelson@gov.bc.ca
or Helen Schwantje, (250) 751-3234 Helen.Schwantje@gov.bc.ca

Try to ship samples on Monday or Tuesday, never past Wednesday.

For health-related questions that arise during caribou necropsies

Or

Any questions related to caribou sample collection, handling, processing and storage

Contact:

Dr. Helen Schwantje 250-751-3234 or cell 250-361-7619