



Comments on the

Custer Gallatin National Forest

Draft Environmental Impact Statement for the

Draft Revised Forest Plan

Submitted by:

North American Packgoat Association

June 6, 2019

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VIA ELECTRONIC SUBMITTAL

RE: Comments on the Custer Gallatin National Forest Draft Environmental Impact Statement for the Draft Revised Forest Plan

To: Custer Gallatin National Forest
Attn: Forest Plan Revision
10 E Babcock, P.O. Box 130
Bozeman, MT 59771
Electronic Submittal: <https://cara.ecosystem-management.org/Public/CommentInput?Project=50185>

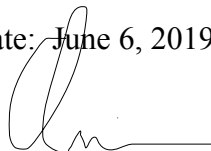
Responsible
Official: Mary C. Erickson, Forest Supervisor
Custer Gallatin National Forest
10 E Babcock, P.O. Box 130
Bozeman, MT 59771

From: Andrew A. Irvine
of Andrew A. Irvine, P.C.
P.O. Box 3221
Jackson, WY 83001
Phone: (307) 690-8383
Email: andy@andrewirvinelaw.com

On behalf of: North American Packgoat Association
Curtis King, President
P.O. Box 170166
Boise, ID 83717
Phone: (509) 539-0982
Email: curtis.king66@yahoo.com

On behalf of the North American Packgoat Association, I hereby timely submit these Comments on the Custer Gallatin National Forest ("Gallatin NF") Draft Environmental Impact Statement ("DEIS") for the Draft Revised Forest Plan ("Forest Plan"). If you have any questions concerning these comments or need further information, you may contact NAPgA or Andrew Irvine at the emails and phone numbers indicated above.

Date: June 6, 2019



Andrew A. Irvine
of Andrew A. Irvine, P.C.

I. Introduction to Comments

The North American Packgoat Association (“NAPgA”) timely submits comments on the Custer Gallatin National Forest (“Gallatin NF”) Draft Environmental Impact Statement (“DEIS”) for the Draft Revised Forest Plan (“Forest Plan”). *See* 84 Fed. Reg. 8524 (Mar. 8, 2019) (Notice of Availability). Comments on the DEIS and Forest Plan were requested by the Gallatin NF as required by 40 C.F.R. §§ 1502.9, 1503.1. *See id.*; *see also* Letter from Mary C. Erickson, Forest Supervisor, Gallatin NF, to Interested Parties, dated March 1, 2019, available at https://www.fs.usda.gov/Internet/FSE_DOCUMENTS/fseprd611353.pdf (requesting comments). The comment period ends on June 6, 2019. *See* 84 Fed. Reg. 8524.

The North American Packgoat Association, Inc. is an organization established specifically for promoting packing with pack goats. The organization was incorporated in March 2001 as a 501(c)(3) non-profit organization. NAPgA seeks to further the pursuit of goatpacking by sharing the knowledge, ideas and experiences of its members, by promoting the use of pack goats to the public as a means of low impact wilderness transportation and recreation, by serving as an advisory group on local and national land use issues, and by engaging in other activities related to educating the public about goatpacking.

NAPgA appreciates this opportunity to comment on the Custer Gallatin National Forest DEIS for the Forest Plan. NAPgA and its numerous goatpacking-members will be affected by the management direction proposed in the draft goals and standards. The proposed management direction would result in closure of one of the premier goatpacking areas in the nation, and set a bad precedent for other forests to follow in managing goatpacking as a recreational use. These comments will better inform the DEIS and Forest Plan and further develop the efficacy of the management direction as defined by the draft goals and standards.

II. Legal Background for the Comments

A. NEPA Prohibits Uninformed Agency Action

In passing NEPA, Congress “recogniz[ed] the profound impact of man’s activity on the interrelations of all components of the natural environment” and set out “to create and maintain conditions under which man and nature can exist in productive harmony.” 42 U.S.C. § 4331(a). To bring federal action in line with Congress’ goals and to foster environmentally informed decision-making by federal agencies, NEPA “establishes ‘action-forcing’ procedures that require agencies to take a ‘hard look’ at environmental consequences.” *W. Watersheds Project v. Kraayenbrink*, 632 F.3d 472, 486 (9th Cir. 2011) (citing *Metcalf v. Daley*, 214 F.3d 1135, 1141 (9th Cir. 2000)). Foremost among those procedures is the preparation of an environmental impact statement (“EIS”). *Id.*

Agencies considering “major Federal actions significantly affecting the quality of the human environment” are required to prepare an EIS. 42 U.S.C. § 4332(C). The EIS “shall provide full and fair discussion of [the] significant environmental impacts” of the proposed action. 40 C.F.R. § 1502.1. That discussion serves two purposes:

First, it ensures that the agency, in reaching its decision, will have available, and will carefully consider, detailed information

concerning significant environmental impacts. Second, it guarantees that the relevant information will be made available to the larger audience that may also play a role in both the decisionmaking process and the implementation of that decision.

W. Watersheds Project, 632 F.3d at 487 (quoting *Dep't of Transp. v. Pub. Citizen*, 541 U.S. 752, 768 (2004)). This process does not mandate particular substantive results, but “NEPA . . . prohibits uninformed . . . agency action.” *Robertson v. Methow Valley Citizens Council*, 490 U.S. 332, 351 (1989). By focusing agency and public attention on the environmental effects of proposed action, “NEPA ensures that the agency will not act on incomplete information, only to regret its decision after it is too late to correct.” *Marsh v. ONRC*, 490 U.S. 360, 371 (1989).

B. Review Under the APA

The Administrative Procedure Act (“APA”), 5 U.S.C. §§ 701-706, provides for judicial review of agency actions, such as those at issue here.¹ Under the APA, a reviewing court shall “hold unlawful and set aside agency action, findings, and conclusions found to be . . . arbitrary, capricious, an abuse of discretion, or otherwise not in accordance with law; . . . [or] without observance of procedures required by law.” 5 U.S.C. § 706(2)(A), (D). Although the arbitrary and capricious standard is a “narrow one,” the court is required to “engage in a substantial inquiry” and a “thorough, probing, in-depth review.” *Native Ecosystems Council v. U.S. Forest Serv.*, 418 F.3d 953, 960 (9th Cir. 2005) (quoting *Citizens to Preserve Overton Park, Inc. v. Volpe*, 401 U.S. 402, 415-16 (1971)).

Under this standard, an agency decision is to be reversed as arbitrary and capricious if the agency has “. . . entirely failed to consider an important aspect of the problem, [or] offered an explanation that runs counter to the evidence before the agency. . . .” *Motor Vehicle Mfrs. Ass’n v. State Farm Mutual Auto. Ins. Co.*, 463 U.S. 29, 43 (1983). “The reviewing court should not attempt itself to make up for such deficiencies.” *Id.* (citation omitted). Most fundamentally, the agency must “examine the relevant data and articulate a satisfactory explanation for its action including a ‘rational connection between the facts found and the choice made.’” *Motor Vehicle*, 463 U.S. at 53 (quotation omitted).

Where, as here, there has been a change in policy from allowing goatpacking on the Gallatin NF to eliminating goatpacking on the Forest, judicial review starts with the presumption that the change in policy is *not* justified by the administrative record. *Motor Vehicle*, 463 U.S. at 42. Additionally, the traditional presumption of agency expertise “‘may be rebutted if the decisions, even though based on scientific expertise, are not reasoned.’” *W. Watersheds Project v. Ashe*, No. 11-462, 2013 WL 2433370 at *5 (D. Idaho June 4, 2013) (citations omitted).

¹ NEPA claims are subject to judicial review under the APA, 5 U.S.C. § 706(2)(A). See *Dep't of Transp. v. Pub. Citizen*, 541 U.S. at 763; *Marsh*, 490 U.S. at 375–76; *League of Wilderness Defenders-Blue Mtns. Biodiversity Project v. U.S.*, 549 F.3d 1211, 1215 (9th Cir. 2008) (the APA provides authority for the court’s review of decisions under NEPA); *W. Watersheds Project v. U.S. Forest Serv.*, 2006 WL 292010, *2 (D. Idaho) (same).

In addition to the requirements of the NEPA and the APA, Forest Service regulations require that “best available science” be taken into account in forest planning. 36 C.F.R. § 219.3. In taking “best available science” into account, the Forest Service must “document how the best available science information was used to inform the assessment, the plan decision, and the monitoring program” and such documentation must “[i]dentify what information was determined to be the best available scientific information, explain the basis for that determination, and explain how the information was applied to the issues considered.” *Id.*

III. Background on the Forest Plan

The Forest Plan makes radical changes to the Gallatin NF’s existing management of goatpacking on the Forest. At General Recreation, Section 2.4.15, and with regard to Suitability 01, the Forest Plan states that under Alternatives B and C “[r]ecreational use of pack goats is not suitable in the Madison, Henry’s Lake, and Gallatin Mountains; Absaroka –Beartooth; or Pryor Mountain Geographic Areas. Under Alternative D, “[r]ecreational use of pack goats is not suitable forestwide,” and under Alternative E, “[r]ecreational use of pack goats is suitable forestwide.” Thus, three of the four proposed alternatives would render recreational use of pack goats unsuitable on the Forest.

In following, at Section 2.4.25 Recreational Opportunities—Outfitter Guides (RECOG), and with regard to Standards (FW-STD-RECOG) 01, the Forest Plan states:

Alternatives B and C: Use of pack goats under new special use permits shall not be permitted in the Madison, Henrys Lake, and Gallatin Mountains; Absaroka-Beartooth; or Pryor Mountain Geographic Areas. Use of pack goats under new special use permits may be permitted in the Bridger, Bangtail, and Crazy Mountains; Ashland; and Sioux Geographic Areas only if a risk assessment indicates that spatial or temporal separation, or other mitigation can effectively minimize risk of disease transmission between livestock and bighorn sheep.

Alternative D: Use of pack goats under new special use permits shall not be permitted.

Alternative E: Use of pack goats under new special use permits shall be permitted only if a risk assessment indicates that spatial or temporal separation, or other mitigation can effectively minimize risk of disease transmission between livestock and bighorn sheep.

See also DEIS at 434-35 (repeating management direction). As a result, pack goats are banned from most of the Forest under Alternatives B and C, and from the whole Forest under Alternative D. Moreover, even where pack goats are not banned, they are subject to an undefined “risk assessment” under Alternatives B, C and E that must indicate, “spatial or temporal separation, or other mitigation can effectively minimize risk of disease transmission between livestock and bighorn sheep.”

The Forest Plan adds that special use permits must provide the following:

02 Written instructions shall be included in the permit to address management, retrieval and disposition of stray pack goats.

03 Notification procedures shall be included in the permit for situations when wandering bighorn sheep may come into contact with pack goats, prompt notification of interaction shall be required by permittees.

04 The Forest Service shall require permittees to take appropriate measures to prevent use of sick or diseased pack goats

Finally, in the Glossary at page 215, the Forest Plan defines “effective separation” as “[t]he spatial or temporal separation between wild sheep and domestic sheep or goats to minimize the potential for association and the probability of transmission of diseases between species (Wild Sheep Working Group 2012).”

IV. Comments on the DEIS and Forest Plan

To assist the Gallatin NF, NAPgA’s comments generally refer to specific pages of the DEIS and Forest Plan that form the basis for each comment; however, some comments may apply more broadly. Comments are intended to apply to all listed pages, or generally, and should be addressed in the context of each of the listed pages or in general.

NAPgA looks forward to the Gallatin NF’s responses to its comments. In addition to its general obligation to respond to public comments under 40 C.F.R. § 1503.4(a), the Gallatin NF must specifically “discuss at appropriate points in the final [EIS] any responsible opposing view which was not adequately discussed in the draft [EIS] and . . . indicate the agency’s response to the issues raised.” *Ctr. for Biological Diversity v. U.S. Forest Serv.*, 349 F.3d 1157, 1167 (9th Cir. 2003) (quoting 40 C.F.R. § 1502.9(b)). A failure to do so is itself a NEPA violation. *Id.* at 1168. The Gallatin NF must also “insure the professional integrity, including scientific integrity, of the discussions and analyses” included in its DEIS. 40 C.F.R. § 1502.24.

1. The No Action Alternative is Mischaracterized in the DEIS and Must be Revised so as NOT to Include Policy which has Not been Subject to NEPA Review and Public Comment

Under Alternative A – No Action (the Current Plans), the DEIS states, “[w]hile no specific management direction is stated related to disease transmission to bighorn sheep from domestic sheep and goats, the Forest Service would follow current policy to only allow this use if a risk assessment indicates risk of disease transmission to bighorn sheep can be minimized.” DEIS at 18.

As the Gallatin NF recognizes, the current plans do not provide specific management direction related to disease transmission from pack goats. As a result, Alternative A – No Action in the DEIS should represent an alternative where no restrictions are placed on pack goat use on the Forest. The Gallatin NF mischaracterizes Alternative A – No Action by adding a “policy” that has not been approved in a Forest Plan and has not been subject to NEPA. The policy does not represent the No Action alternative and cannot be incorporated as part of Alternative A – No

Action. This attempt by the Gallatin NF to include such policy as part of the existing Forest Plan in order to avoid NEPA review and public comment of the policy is improper.

Further, this “policy,” although mentioned in the DEIS, is not specifically named, discussed or presented in the document, so the public is uninformed about the policy. This policy must be named, discussed and presented in the DEIS, so that the public can review the policy and comment on its inclusion as part of the DEIS.

2. The “Risk Assessment” Referenced in the DEIS Must be Presented to the Public and Discussed in the DEIS

At 2.5.4 Alternative B and throughout the DEIS, the DEIS references a “risk assessment.” For example, the DEIS states, “[e]lsewhere on the national forest, . . . permitted recreational goat packing would be allowed only if a risk assessment indicated risk of disease transmission to bighorn sheep can be minimized.” DEIS at 19. What is this “risk assessment?” This “risk assessment” must be presented to the public and subject to public comment as part of the DEIS.

3. NAPGA Generally Supports Alternative A and Alternative E to the Extent the Alternatives Allow for Continued Goatpacking on the Gallatin NF

Although the Gallatin NF fails to provide or discuss the “policy” made part of Alternative A and likewise fails to provide or discuss the “risk assessment” made part of Alternative E, both of these alternatives would appear to allow goatpacking to continue on the Gallatin NF. As a result, NAPgA urges the responsible official to choose Alternative A or Alternative E as the preferred alternative. Public recreational goatpacking is definitely a suitable use and should be allowed on the Forest with or without a risk assessment, as there is little to no risk of disease transmission to bighorn sheep posed by the use of pack goats on the Gallatin NF.

4. The DEIS Misrepresents the Science on Disease Transmission from Domestic Goats, Especially Pack Goats. To Ensure the Scientific Integrity of the DEIS and Forest Plan, the Gallatin NF Must Correct and/or Remove False or Unsupported Statements Concerning Pack Goats from the DEIS and Forest Plan

In evaluating the environmental impacts of a proposed action, NEPA requires federal agencies to ensure the scientific integrity of an EIS by considering appropriate studies and data. 40 C.F.R. § 1502.24. The Gallatin NF must “insure the professional integrity, including scientific integrity, of the discussions and analyses” included in its DEIS. *Id.* An agency may not rely on conclusory statements unsupported by data, authorities, or explanatory information. *Seattle Audubon Soc’y v. Moseley*, 798 F. Supp. 1473, 1480-83 (W.D. Wash. 1992), *aff’d*, 998 F.2d 699 (9th Cir. 1993). NEPA requires that an agency candidly disclose in its EIS the risks and effects of its proposed actions, and that it respond to adverse opinions held by respected scientists. *Seattle Audubon*, 798 F. Supp. at 1482 (*citing Friends of the Earth v. Hall*, 693 F. Supp. 904, 937 (W.D. Wash. 1988)). Further, under NEPA, courts have held that agency actions based on unexplained assumptions are arbitrary and capricious. *Ctr. for Biological Diversity v. U.S. Dep’t of the Interior*, 623 F.3d 633, 650 (9th Cir. 2010); *see also Dow*

Agrosciences LLC v. Nat'l Marine Fisheries Serv., 707 F.3d 462, 470 (4th Cir. 2013) (agency must explain why lab tests reflect nature).

The Gallatin NF has failed to ensure the professional integrity, including scientific integrity, of the discussions and analyses in the DEIS as required under NEPA. The Gallatin NF appears to be operating on incomplete information concerning disease transmission from domestic goats, including packgoats, to bighorn sheep, and also appears to be ignoring important aspects of the problem of disease transmission as well as offering explanations in the DEIS that run counter to the evidence before the Gallatin NF. Much of the analysis and discussion in the DEIS lacks factual or scientific support.

At Section 3.10.4 General Wildlife, the DEIS cites Wild Sheep Working Group 2012 for the statement that “[a]n extensive review of scientific literature and available data on bighorn sheep populations in the western United States concluded that contact with domestic sheep and goats was the source of most of the disease resulting in major die-offs of bighorn sheep.” This cite is to a collection of “Recommendations for Domestic Sheep and Goat Management in Wild Sheep Habitat,” not a scientific research paper. To the extent there is any scientific evidence reference in the Wild Sheep Working Group 2012 to support this statement as it pertains to goats, particularly pack goats, such science should be directly cited and the public should be allowed to review and comment on such science. Otherwise, the Gallatin NF should remove the reference, as it is not a scientific research paper providing any evidence concerning disease transmission between pack goats and bighorn sheep.

Further, the DEIS indicates that “[t]he presence of disease-carrying domestic sheep and goats in close proximity to bighorn sheep is a key stressor for bighorn sheep.” DEIS at 432. What is the basis for this statement as it applies to goats, particularly pack goats? There is absolutely no science indicating that “[t]he presence of disease-carrying domestic . . . [pack goats] in close proximity to bighorn sheep” is a “key stressor” for bighorn sheep. This statement must be revised to exclude pack goats.

Finally, the DEIS states, “[c]onsequently, comingling of bighorns with domestic sheep and goats continues to be a major concern today,” citing Garrott et al. 2015. DEIS at 432. Where in Garrott et al. 2015 is there a concern raised about comingling of bighorns with domestic goats, particularly pack goats? Garrott et al. 2015 does not mention any risk of disease transmission from pack goats. As a result, this statement must be revised to exclude pack goats.

5. Statements in the DEIS Concerning Effects of the Current Plans Must be Revised to Reflect Current Science on Disease Transmission from Pack Goats

In the DEIS under Effects of the Current Plans, the DEIS states that domestic goats “may carry some of the same strains of disease, and can transmit disease to bighorn sheep in the wild.” DEIS at 433. No cite is provided for this statement. When has a domestic goat, particularly a pack goat, ever transmitted disease to bighorn sheep in the wild? That has never happened. This sentence must be revised to exclude pack goats.

The DEIS also references a “primary threat of disease transmission from domestic sheep and goats to bighorn sheep.” DEIS at 433. What is this “threat of disease transmission” from pack goats to bighorn sheep? No such threat has been established. As a result, this statement should be revised to exclude pack goats. Furthermore, because pack goats do not pose a threat of disease transmission, as discussed further below, this section should be revised to indicate that existing plans are more than sufficient to minimize disease transmission from pack goats, as pack goats do not pose a threat of disease transmission to bighorn sheep.

6. The Gallatin NF Must Consider Dr. Margaret Highland’s Research Concerning the Limited Prevalence of *Mycoplasma ovipneumoniae* in Pack Goats

The Gallatin NF has failed to consider recent scientific research indicating that pack goats do not commonly carry *Mycoplasma ovipneumoniae*. This research by Dr. Margaret Highland, Research Veterinarian with the Animal Disease Research Unit-ARS-USDA is presented in Exhibit B. Dr. Highland’s research indicates that pack goats do not commonly carry the disease-causing organisms associated with bighorn sheep die-offs. The results of the testing performed for Dr. Highland’s research are also included in Exhibit B, so that the Gallatin NF can consider the results and verify the legitimacy and scientific method in the research. Dr. Highland’s research is in the process of being published, but has already been presented, *see, e.g.*, <https://pdfs.semanticscholar.org/presentation/4bb7/616fa740f42ceda2c55d275f0a8032fc6ca8.pdf>, and has been considered by the Forest Service on numerous other occasions (except on the Gallatin NF).

Under the APA and NEPA, the Gallatin NF is required to consider the fundamental aspect of the problem of disease transmission, namely, whether pack goats can actually carry and transmit *M. ovi* to bighorn sheep in the wild. *See Motor Vehicle*, 463 U.S. at 43. The Gallatin NF is also required to examine relevant data, consider opposing viewpoints, ensure the scientific integrity of its discussions, and articulate a satisfactory explanation for its action. *See id.* at 42-43, 53; *Ctr. for Biological Diversity v. U.S. Forest Serv.*, 349 F.3d at 1167 (quoting 40 C.F.R. § 1502.9(b)).

Moreover, and in addition to the requirements of the APA and NEPA, Forest Service regulations require that “best available science” be taken into account in forest planning. 36 C.F.R. § 219.3. In taking “best available science” into account, the Forest Service must “document how the best available science information was used to inform the assessment, the plan decision, and the monitoring program” and such documentation must “[i]dentify what information was determined to be the best available scientific information, explain the basis for that determination, and explain how the information was applied to the issues considered.” *Id.* The Forest Service Land Management Planning Handbook, FSH 1909.12, directs the Gallatin NF’s use of the best available scientific information and provides that where research is relevant, accurate and reliable, the Forest Service should include it as the best available scientific information. *See* FSH 1909.12, 42.13.

As a result, this science presented by Dr. Highland must be considered in the DEIS under the APA and NEPA, as well as the implications of pack goats not being carriers of *M. ovi*. If

pack goats are not carriers of disease-causing pathogens, then they do not pose a risk of disease transmission to bighorn sheep on the Gallatin NF.

In sum, the Gallatin NF must review and consider Dr. Highland's research in the DEIS. Such consideration is required by the APA, NEPA and the Forest Service's own planning regulations. Dr. Highland's research indicates that pack goats are rarely carriers of *M. ovi*. As a result, pack goats do not pose a significant risk of disease transmission to bighorn sheep on the Gallatin NF. Pack goats cannot transmit disease they do not have. These points must be considered in the DEIS.

7. Cooperation and Collaboration in Decision-Making is Required Before, Not After, the Gallatin NF Makes a Decision to Ban Pack Goat Use on the Forest

The DEIS indicates that a goal of the Forest Plan is "cooperation and collaboration with . . . livestock permittees, and other interested parties to develop livestock management protocols and habitat management strategies to minimize risk of disease transmission between domestic livestock and bighorn sheep." DEIS at 434. How is the Gallatin NF achieving this goal? It would seem important, as well as required under NEPA, to cooperate and collaborate with NAPgA prior to banning pack goat use from the Forest. In order to avoid uniformed agency decision-making, the Gallatin NF must consult with NAPgA before, not after, deciding to ban pack goat use from the Forest.

8. The DEIS Must Specifically Identify and Discuss the "Threat of Disease Transmission" from Pack Goats to Bighorn Sheep

As touched on above, the DEIS refers to a "threat of disease transmission from domestic livestock to bighorn sheep." DEIS at 434. The DEIS further indicates that plan alternatives include components to address this "threat" and that where pack goats are not completely banned from the Forest outright, they will be banned according to an undefined "risk assessment." *Id.* at 434-35. While the Gallatin NF is quick to ban pack goats use because of the "risk" or "threat" of disease transmission, it does not define what this "risk" or "threat" actually is?

The Gallatin NF should explain in the DEIS what it means by risk of disease transmission between pack goats and bighorn sheep. Notably, contact between pack goats and bighorn sheep has never occurred before on the Forest, so risk does not mean that contact is more likely that not, otherwise such contact would have already occurred. There is no scale of risk to inform the reader about the actual likelihood of contact. The Gallatin NF should explain what they mean by "risk," including the various scales of risk from high to low. Also, the Gallatin NF should explain how contact between pack goats and bighorn sheep on the Forest would actually occur. What does the Gallatin NF mean by "contact?" Would a bighorn sheep approach a pack goat on a trail, in the presence of the pack goat's human owner and make "contact?" Would a bighorn sheep enter into a camp in a forested area where there is a pack goat, again in the presence of its human owner, and make contact there? Is this nose-to-nose or sexual contact? When the Gallatin NF refers to "risk" and "contact" in the DEIS it is unclear what the Gallatin NF is talking about and how such "contact" would occur. These things should be explained. Likewise, the Gallatin NF should discuss the likelihood of contact in understandable terms and present how

such contact would occur based on the behavior of bighorn sheep and use and training of pack goats.

At page 431, the DEIS indicates that “[a] few individuals have used domestic pack goats for personal (not outfitting or guiding) recreational purposes, but to date, such use has been very limited on the Custer Gallatin National Forest.” Considering this very limited use, the Gallatin NF is basically saying there is a strong likelihood that on one of the two or three goatpacking trips taken on the Gallatin NF each year, a bighorn sheep would (1) leave its herd and its summer habitat in the high country, (2) find a human and pack goat camp, (3) sneak into that camp without causing any disturbance in the pack goats and without being detected by the humans, (4) ask the pack goats to not be alarmed, to remain still and to muffle their bells and collars, (5) find a tethered goat that is infected by and shedding strains of *M. ovi*, (6) make physical contact with that goat sufficient for disease transmission, and (7) sneak back out of camp and return to its herd and infect other bighorn sheep. It is a far-fetched scenario that has never happened before.

In reality, there is almost no overlap in time or space between pack goats and bighorn sheep on the Gallatin NF; bighorn sheep are not prone to leave their herd/habitat and wander into human and pack goat camps; pack goats react noisily when they are alarmed by other wildlife, including bighorn sheep; the vast majority of pack goats do not carry and shed strains of *M. ovi*; and it is unknown whether bighorn sheep can even be infected with strains of *M. ovi* from pack goats resulting in fatal respiratory disease. The facts do not support the Gallatin NF’s assumption that there is a likelihood of disease transmission from pack goats to bighorn sheep on the Forest.

Before undertaking management action concerning the risk of contact and disease transmission between pack goats and bighorn sheep on the Gallatin NF, the Forest should provide an analysis of the current risk posed by pack goats. This could be done with a quantitative risk assessment. Regardless, the Gallatin NF has not presented any scientific information indicating that pack goats pose a significant risk. Rather, pack goats rarely use the Gallatin NF, rarely carry disease and are very unlikely to contact a bighorn sheep, particularly when handled according to established guidelines, so pack goats would appear to pose negligible risk. Why then are they being prohibited from the Gallatin NF? The Gallatin NF must answer this threshold question. The Gallatin NF’s explanation for prohibiting pack goat use runs counter to the evidence before the agency. Without establishing significant risk, the Gallatin NF’s prohibition on pack goat use is unjustified.

9. The Gallatin NF Arbitrarily and Capriciously Treats Potential Disease Transmission from Pack Goats Different than that From Llamas and Alpacas on the Forest

Curiously, with regard to llamas and alpacas, the Gallatin NF provides, “[u]ntil more definitive science verifies disease transmission from llamas and alpacas to bighorn sheep in the wild, the Custer Gallatin would track this issue related to the forestwide desired condition for low or no disease transmission between domestic livestock and wildlife, under all revised plan alternatives.” DEIS at 436. Although pack goats are likewise seldomly used on the Gallatin NF, with no known or suspected disease transmission to wild sheep or goats, the Gallatin NF has taken an approach opposite of that taken on llamas and alpacas—the Gallatin NF has banned

pack goats from the Forest. This decision is arbitrary and capricious. There is no science and certainly no known or suspected disease transmission from pack goats to wild sheep or goats. As a result, and similar to treatment of llamas and alpacas, the Gallatin NF should track the issue of disease transmission as it related to pack goats rather than institute a ban on pack goat use.

10. The Gallatin NF Mischaracterizes the Results of Besser's Research and Must Correct Statements in the DEIS Concerning Disease Transmission from Pack Goats to Bighorn Sheep

The DEIS states that “[d]isease transmission from recreational use of domestic pack goats is a potential threat to bighorn sheep.” DEIS at 440. As discussed above, there is no scientific support for this statement. The Gallatin NF adds that “Besser and associates (2017) found that while domestic goats carry disease that can be transmitted to bighorn sheep, the severity of disease impacts on wild sheep populations was milder than impacts from disease transmitted from domestic sheep.” *Id.* This is a gross misstatement of the research by Besser and associates (2017). The domestic goats in from Besser's research did not “carry disease.” Rather, they were infected by disease by Besser during his research. Pack goats have not been infected by disease by Besser and thus are very different than the domestic goats used for Besser's research. Pack goats, in fact, rarely carry *M. ovi*, the primary disease of concern for disease transmission to bighorn sheep. The DEIS grossly mischaracterizes the research by Besser. If anything, Besser's research showed that domestic goats do not post a threat of disease transmission resulting in mortality in bighorn sheep. During Besser's research, not a single bighorn sheep died as a result of disease transmission from a domestic goat. As a result, the Gallatin NF must correct the misstatements and provide an accurate description of Besser's research, including the information discussed below.

Further, the Gallatin NF is cautioned about relying on Besser and associates (2017) as the research article is filled with inaccuracies and exaggerations and lacks objectivity. *See* <http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0178707>. Indeed, the publisher *PLOS ONE* issued a correction to the article to correct some of the inaccuracies and exaggerations. *See* <http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0192006>.

Importantly, based on the data and findings in Besser and associates (2017), and as stated above, not a single bighorn sheep died from exposure to domestic goats in any context throughout Besser's experiments. Indeed, as discussed on pages 5 through 7 of 13 of the article, to the extent bighorn sheep exhibited signs of respiratory problems when initially commingled with domestic goats, all bighorn sheep exhibited fewer signs of respiratory problems over time, indicating recovery from such problems prior to being euthanized. In short, Besser and associates (2017) shows that even when domestic goats are purposefully infected with *Mycoplasma ovipneumoniae*, comingling of such goats with bighorn sheep does not result in fatal respiratory disease in bighorn sheep. In other words, exposure of bighorn sheep to domestic goats colonized with *M. ovi* does not induce fatal pneumonia.

To the extent the Gallatin NF continues to rely on Besser and associates (2017) in the DEIS and Forest Plan, the Gallatin NF should update the reference to the article to the recently corrected version. The Gallatin NF should also recognize and discuss that comingling of domestic goats, even those purposefully infected with *M. ovi*, does not lead to fatal respiratory

disease in bighorn sheep. When domestic goats are not infected with *M. ovi*, as is the common case with pack goats, there is no risk of transmission of *M. ovi* leading to fatal respiratory disease in bighorn sheep.

11. The Gallatin NF Should Focus on Herd Density Issues in Managing Bighorn Sheep Populations and Should Further Acknowledge that Pack Goats Do Not and Have Never Posed a Threat of Disease Transmission to Wild Bighorn Sheep

The DEIS indicates that the “primary issue driving bighorn sheep populations on the Custer Gallatin National Forest and surrounding areas is major die-offs associated with disease spread among and possible between herds. Although wild sheep can carry disease and transmit to others, many of the same diseases can be carried by domestic sheep and goats, and can be transmitted to wild sheep.” DEIS at 441. To start, the conclusion that the primary issues is disease transmission from domestic livestock is not necessarily true.

Recent science indicates that herds at high density are at a much, much greater risk of die-offs than those at low density (Sells et al. 2015). Sells et al. (2015) found that “[r]isk of a pneumonia epizootic increased >5-fold when herds were at a medium density and nearly 15-fold when herds were at a high density compared to when they were at a low density.” Further, Sells et al. (2015) indicated, “[d]ensity is a component of risk that has previously received little attention because the positive association between risk of pneumonia and higher densities had not been quantified. The association between higher herd density and risk may appear to contradict the idea that herds of larger population size should be less threatened by extirpation than smaller herds [].” (citations omitted). So, according to Sells et al. (2015), the most important consideration, by a long shot, in managing to avoid pneumonia epizootics is control of bighorn sheep herd density and, in particular, ensuring that bighorn sheep herd density does not get too high. The Gallatin NF should consider this best available science.

Second, there are no facts or science indicating disease transmission between pack goats and bighorn sheep, in the wild. That has never happened. There simply is no credible threat of disease transmission from pack goats to bighorn sheep in the wild. As a result, the Gallatin NF’s conclusion about disease transmission must be revised to give a true description of the facts, science and “risk” about disease transmission from pack goats to bighorn sheep in the wild.

12. The Gallatin NF Must Consult the Agricultural Research Service, within the United States Department of Agriculture, Before Preparing the Final EIS and Record of Decision

NEPA imposes on federal agencies conducting environmental review a duty to consult with certain other agencies.” Prior to making any detailed statement, the responsible Federal official shall consult with and obtain the comments of any Federal agency which has jurisdiction by law or special expertise with respect to any environmental impact involved [in the proposed action].” 42 U.S.C. § 4332(2)(C). Further, to promote NEPA’s policies of public participation and informed decisionmaking, copies of the EIS and comments thereon from other agencies “shall accompany the proposal through the existing agency review processes.” *Id.*

The regulations implementing these provisions state that “[a]fter preparing a draft environmental impact statement and before preparing a final environmental impact statement the agency shall . . . [o]btain the comments of any Federal agency which has jurisdiction by law or special expertise with respect to any environmental impact involved” 40 C.F.R. § 1503.1(a)(1); *see also id.* § 1500.1(b) (“Accurate scientific analysis, expert agency comments, and public scrutiny are essential to implementing NEPA.” (emphasis added)). “Special expertise” is defined as “statutory responsibility, agency mission, or related program experience.” *Id.* § 1508.26. Under the statute and its implementing regulations, the Gallatin NF has a duty to consult with the Agriculture Research Service (“ARS”) before issuing the Final EIS. *See Idaho Wool Growers Ass’n v. Vilsack*, 816 F.3d 1095, 1103 (9th Cir. 2016).

ARS has “special expertise” concerning significant aspects of the proposed decision, including the mechanics of pathogen transmission in domestic sheep and goats. For example, 7 C.F.R. § 2.65 delegates to ARS, among other matters, the authority to “[c]onduct research concerning domestic animals and poultry, their protection and use, [and] the causes of contagious, infectious, and communicable diseases.” Also, ARS’s mission statement proclaims: “ARS conducts research to develop and transfer solutions to agricultural problems of high national priority and provide information access and dissemination to . . . enhance the natural resource base and the environment” U.S. Department of Agriculture, Agricultural Research Service, ARS: About US, <http://www.ars.usda.gov/aboutus/aboutus.htm>.

Thus, considering the language establishing NEPA’s consultation requirement is expansive, NEPA mandates consultation with any federal agency that has “special expertise with respect to any environmental impact involved.” 42 U.S.C. § 4332(2)(C) (emphasis added); *see also* 40 C.F.R. § 1503.1(a)(1) (“[T]he agency shall . . . [o]btain the comments of any Federal agency which has jurisdiction by law or special expertise with respect to any environmental impact involved. . . .” (emphasis added)). And, further considering that *Warm Springs Dam Task Force v. Gribble* suggests that for the consultation requirement to apply, the particular expertise of an agency does not have to encompass the proposed project as a whole or the issue the proposed project was designed to address. Rather, the expertise need relate only to one of the project’s anticipated environmental effects. *See* 621 F.2d 1017, 1020-21 (9th Cir. 1980) (per curiam); *see also Idaho Wool Growers Ass’n*, 816 F.3d at 1103. It is a clear requirement that the Gallatin NF MUST consult with ARS on issues of disease transmission, such as those presented in the DEIS and Forest Plan, prior to issuing a Final EIS. As a result, the Gallatin NF MUST consult with ARS and should detail such consultation in the Final EIS.

13. The Gallatin NF Fails to Account for the Important Differences Between Pack Goats and Herd Domestic Goats and Domestic Sheep

The Gallatin NF fails to acknowledge the important differences between pack goats and herd domestic sheep and goats. These differences must be considered in the DEIS and Forest Plan. NEPA prohibits this type of uninformed agency action. *See Robertson*, 490 U.S. at 352 (“NEPA . . . prohibits uninformed . . . agency action.”); *Marsh*, 490 U.S. at 371 (“NEPA ensures that the agency will not act on incomplete information, only to regret its decision after it is too late to correct.”). These differences are critical to the Gallatin NF’s analysis of disease transmission from pack goats to bighorn sheep and must be considered by the Forest under NEPA.

Pack goats are very different from other domestic goats (and domestic sheep), both by breed and by use. These differences result in far less risk to bighorn sheep than the risk posed by domestic goats (or domestic sheep) on grazing allotments. The Gallatin NF DEIS must account for these differences. To consider pack goats the same as other domestic goats (or domestic sheep) for purposes of analyzing the risk of disease transmission to bighorn sheep on the Gallatin NF would be a critical error.

Pack goat owners go to great lengths and expense to find and train particular goats that will not stray from the security of a finite string of pack goats and their owner. Pack goats are inextricably bonded to their owners, which represent the “alpha goat” in their small herd. This is achieved through the processes of imprinting and socialization of pack goats from birth. As a result, pack goats are not prone to straying and remain in very close proximity to the “alpha goat.” Other domestic goats (and domestic sheep), while often included in herds that number in the hundreds or thousands (compared to a string of pack goats ranging from two to ten goats), are not individually trained and, thus, there may be some risk of individual domestic herd goats (or domestic sheep) straying from the herd. The risk associated with domestic sheep or domestic goats transmitting disease to bighorn sheep requires “physical contact” between the domestic animal and the bighorn sheep, therefore, a pack goat that is less likely to stray and thereby come into contact with a bighorn sheep poses a much lower risk of transmission than any number of herd domestic sheep or goats which can wander and stray.

Domestic goat and sheep herds typical to grazing allotments on public land represent larger populations of animals that are more difficult to maintain, and which may not be in immediate proximity of their caretaker at all times. Pack goats, on the other hand, require their owner or “alpha goat” to be present to monitor the herd at all times, and are always in their owner’s immediate presence and control. The small size of a pack goat string and perpetual control of the owner allows pack goats to be tied in unison while on trails, and tethered or high-lined at night (among other best management practices that can be easily implemented) to reduce the risk of contact between a pack goat and a bighorn sheep. Furthermore, if ever in sight of a bighorn sheep, there is always a human present in close proximity to the pack goats, making it extremely unlikely that a bighorn sheep would approach the string. In the presence of wild animals, such as bighorn sheep, pack goats are also on heightened alert and retreat to a position near the “alpha goat,” i.e., their human caretaker. This and the other defining traits of pack goats, and the nature of their use and training, make pack goats far less of a risk of coming into contact with a bighorn sheep than herd domestic goats and/or domestic sheep.

Further, the lifestyle and care of a pack goat differs greatly from that of a typical herd domestic goat or domestic sheep. This difference in care means that pack goats are healthier and less likely to be the carrier of a disease. Pack goats are seen by their owners as a significant investment in time and resources. A pack goat is not viable for packing purposes until at least the age of three or four, and often pack goats do not reach their packing prime until the age of five or six. Thus, a goatpacker will have had to invest a number of years into a pack goat before it is ready to hit the trail. During this time, and throughout a pack goat’s life, pack goats see personalized veterinary care in order to keep the goat healthy and prolong their useful life, a luxury that other free ranging herd domestic goats or domestic sheep do not enjoy.

Because of their overall health and stamina, a trained pack goat can bring a sale price of over \$450. This means that a pack goat owner has a large financial interest in each of his or her pack goats. This high financial interest means that the owner of pack goats is likely to see to their care and protection whether that is protection from disease at home, or from contact with other wildlife when on public lands.

Further, typical herd domestic goats and domestic sheep may be sold and intermixed with goats from other herds. In contrast, pack goats—which are treated more like household pets than livestock—are not likely to change owners. The higher frequency that typical herd domestic goats and domestic sheep may be exposed to other domestic stock, would increase the opportunity for disease to spread between individual animals. On the other hand, pack goats are infrequently transferred between owners because of the nature of their function and required bonding. This greatly reduces the risk of exposure of pack goats to various diseases as compared to herd domestic goats and domestic sheep.

Perhaps most critical to the Gallatin NF’s analysis of disease transmission from pack goats to bighorn sheep is the fact that the overwhelming majority of pack goats are not known to carry *M. ovi*. If a pack goat did not carry *M. ovi* it would be impossible for that goat to transmit disease to a bighorn sheep. Thus, the risk of disease transmission from that pack goat to a bighorn sheep would be zero. Further, even if a pack goat were to carry *M. ovi* and directly contact a bighorn sheep, there is no science indicating that the pack goat would transmit this pathogen to the bighorn sheep and that the bighorn would succumb to pneumonia as a result. The Gallatin NF did not consider these important factors in its analysis.

Finally, goatpackers limit their visits to the Gallatin NF, as well as their time on the Forest when they do visit. With only a few pack goats per goatpacker and only a few visits by goatpackers per year, for a limited amount of time, the chance that a pack goat would come into contact with a bighorn sheep is extremely unlikely. This factor was not considered by the Gallatin NF.

Here, the Gallatin NF’s analysis in the DEIS is completely silent on the differences between pack goats and herd domestic goats and how those differences affect the risk of disease transmission between pack goats and bighorn sheep. These differences are critical and must be considered by the Gallatin NF. An agency decision is to be reversed as arbitrary and capricious if the agency has “entirely failed to consider an important aspect of the problem.” *Motor Vehicle Mfrs. Ass’n v. State Farm Mutual Auto. Ins. Co.*, 463 U.S. 29, 43 (1983). The Gallatin NF’s silence on the issue will not suffice. The agency’s path must be reasonably discerned. *Id.* A court “cannot infer an agency’s reasoning from mere silence or where the agency failed to address significant objections and alternative proposals.” *Beno v. Shalala*, 30 F.3d 1057, 1073 (9th Cir. 1994) (citing *Motor Vehicle*, 463 U.S. at 57); *see also, e.g., SEC v. Chenery Corp.*, 332 U.S. 194, 196-97 (1947) (“[i]t will not do for a court to be compelled to guess at the theory underlying the agency’s action.”).

In conclusion, pack goats are very different than other herd domestic goats or domestic sheep that are grazed on or near the Gallatin NF, and the use of pack goats on the Gallatin NF is very different than the use of other herd domestic goats and domestic sheep. The Gallatin NF DEIS and Forest Plan fail to account for these differences in the analysis of disease transmission

from domestic sheep and domestic goats to bighorn sheep on the Gallatin NF. As a result, the DEIS must be revised to consider (1) pack goats separate from other herd domestic goats and domestic sheep and (2) the unlikelihood that pack goats carry disease and (3) the unlikelihood that pack goats would ever come in close contact with bighorn sheep on the Gallatin NF. Further, the Gallatin NF must consider that the nature and use of pack goats on the Forest already achieves the spatial and/or temporal separation recommended by the Gallatin NF to minimize potential disease transmission. Thus, there is no justification and no need for the prohibition of pack goats on the Gallatin NF.

14. The Gallatin NF Fails to Consider Implementation of Mitigation Measures to Ensure the Separation of Pack Goats and Bighorn Sheep

Rather than consider implementation of minimization and mitigation measures to prevent contact and possible disease transmission between pack goats and bighorn sheep on the Gallatin NF, the Gallatin NF has simply closed a large portion of the Forest to pack goats. Under NEPA, however, the Gallatin NF must consider and discuss mitigation measures that would allow the use of pack goats on the Forest. NAPgA has attached a proposed suite of best management practices (“BMPs”) and other minimization and mitigation measures at Exhibit A to prevent contact and possible disease transmission between pack goats and bighorn sheep on the Gallatin NF. These, as well as other available practices and measures must be considered by the Gallatin NF in the DEIS.

For example, the Gallatin NF DEIS fails to consider that separation between pack goats and bighorn sheep is maintained by the presence of a human with pack goats, by nighttime tethering or high-lining of pack goats, and by the nature and training of pack goats. The DEIS also failed to consider the use of GPS tracking collars on pack goats, pathogen testing, permitting for pack goat trips, designation of corridors for pack goats, and a host of other measures. Certainly, if pack goats do not carry disease and do not come into contact with bighorn sheep, there is zero risk of disease transmission from pack goats to bighorn sheep. Neither of these scenarios were considered in the DEIS. Instead of considering any of these measures, in violation of NEPA, the Gallatin NF fails to provide any consideration of these best management practices to maintain separation between pack goats and bighorn sheep on the Gallatin NF.

BMPs are mitigation measures that can be employed by goatpackers to prevent contact between pack goats and bighorn sheep. 40 C.F.R. § 1508.20 (defining “mitigation measures” to include “[a]voiding the impact” and “[m]inimizing impacts by limiting the degree or magnitude of the action and its implementation”). For a reasonable range of alternatives, the Gallatin NF DEIS must consider implementation of BMPs and mitigation measures, rather than simply concluding that goatpacking on the Gallatin NF must be prohibited. 40 C.F.R. § 1502.14.

An EIS must discuss “mitigation . . . in sufficient detail to ensure that environmental consequences have been fairly evaluated.” *Robertson*, 490 U.S. at 352. An agency is required to “discuss possible mitigation measures in defining the scope of the EIS, 40 CFR § 1508.25(b), in discussing alternatives to the proposed action, § 1502.14(f), and consequences of that action, § 1502.16(h), and in explaining its ultimate decision, § 1505.2(c).” *Id.*; see also *Okanogan Highlands Alliance v. Williams*, 236 F.3d 468, 473 (9th Cir. 2000) (An EIS must contain a “reasonably complete discussion of possible mitigation measures.” (quoting *Robertson*, 490

U.S. at 352)). To be sure, an agency's final decision must "[s]tate whether all practicable means to avoid or minimize environmental harm from the alternative selected have been adopted, and if not, why they were not." 40 C.F.R. § 1505.2(c).

Further, NEPA mandates that federal agencies "provide legitimate consideration to alternatives that fall between the obvious extremes." *Colorado Envtl. Coalition v. Dombeck*, 185 F.3d 1162, 1175 (10th Cir. 1998). More specifically, NEPA is violated when an agency dismisses the consideration of an alternative "in a conclusory and perfunctory manner that [does] not support a conclusion that it was unreasonable to consider them as viable alternatives." *Davis v. Mineta*, 302 F.3d 1104, 1122 (10th Cir. 2002). "The existence of reasonable but unexamined alternatives renders an EIS inadequate." *Ilio 'ulaokalani Coalition v. Rumsfeld*, 464 F.3d 1083, 1095, 1101 (9th Cir. 2006).

Without an alternative that describes and analyzes the implementation of mitigation measures to prevent contact between pack goats and bighorn sheep, instead of simply eliminating pack goats from the Gallatin NF, the DEIS contains an inadequate range of alternatives. Alternatives considering BMPs and mitigation measures are both reasonable and feasible under the circumstances, and must be analyzed in the DEIS.

In conclusion, the Gallatin NF has violated NEPA by failing to discuss and consider mitigation measures that would allow use of pack goats on the Forest while preventing the risk of disease transmission between pack goats and bighorn sheep. As a result, the Gallatin NF must revise the DEIS and Forest Plan to discuss and consider appropriate mitigation measures to prevent the risk of disease transmission between pack goats and bighorn sheep. Proper consideration of such measures should include consideration and adoption of an alternative to allow the use of pack goats on the Gallatin NF. This alternative should consider maintenance of the separation of pack goats and bighorn sheep on the Forest and, thus, achieve avoidance of any potential for disease transmission between pack goats and bighorn sheep.

15. The Gallatin NF Must Evaluate Alternatives that Consider Strengthening Bighorn Sheep Immunity to Disease

Established epidemiology shows that disease occurs in bighorn sheep populations in the absence of contact with domestic sheep and other animals, including pack goats. These data indicate that infectious agents and other contributing factors involved in the disease process are present within bighorn sheep populations. It appears that most bighorns are getting pneumonia from other bighorns because most of the herds that have outbreaks of pneumonia, are not in contact with domestic sheep or domestic goats. This indicates that the major problem is the lack of a good immune system in the bighorns. As discussed below, there are inherent risks in choosing a management strategy that attempts to isolate bighorn sheep populations from all perceived transmission risks (when complete isolation is not possible); instead the focus should be on managing population immunity.

The critical component of managing infectious diseases is population immunity. A decision to isolate a given population of bighorn sheep from contact with potential sources of infection assumes the ability for that population to maintain isolation. The wisdom of this management scheme (maintaining immunological naivety) in animal populations within the

United States, when sources of infection are present in nature, is questionable at best. Two methods which provide population immunity are vaccination and/or exposure of populations through natural exposure (transmission). This latter situation is also referred to as premonition (resistance to a disease due to the existence of its causative agent in a state of physiological equilibrium in the host and/or by immunity to a particular infection due to previous presence of the causative agent).

A primary risk associated with incomplete immunologic isolation of animal populations is cycles of disease when isolation is broken as opposed to a continuum of managed population immunity through vaccines and/or natural exposure and premonition. When multiple sources of a given pathogen or group of pathogens exist, the prudent long-term health management dictates that population immunity be the primary tool. As an example of population immunity being the most effective management tool, the Lostine River herd of bighorns experienced a die-off in the 1980s, but is now considered the most viable herd in the Hells Canyon area due to successful population immunity. Since bighorn sheep are infecting each other, building up their immune systems could have a beneficial effect on survival from many forms of disease.

Likewise, bighorn sheep face the risk of infection from domestic sheep and other animals on and off the Gallatin NF. Consequently, the elimination of pack goats on the Gallatin NF, even if there was evidence that pack goats carried and transmitted disease, would not eliminate the risk of disease transmission to bighorns. This fact is not adequately considered in the DEIS. It will be impossible for the Gallatin NF to eliminate the risk of disease transmission to bighorns because of the numerous variables besides pack goats (which are not even a known carrier or transmitter of disease) on the Gallatin NF. As a result, the Gallatin NF must analyze alternative solutions to maintaining bighorn sheep viability.

The Gallatin NF must also analyze the possibility that without interaction between bighorn sheep and other animals, bighorn sheep tolerance to disease may become worse, leading to more widespread die-offs, instead of fewer die-offs. Instead of considering this likelihood, the DEIS only considers one course of action: total separation. Based on the analysis in the DEIS, the most prudent and most logical management action would be to encourage development of immunity in bighorns because total separation is impossible. This action must be considered by the Gallatin NF in the DEIS.

16. Epidemiological Modeling is Needed to Understand How a Range of Factors Affect the Dynamics of Disease Spread Under Various Management Alternatives

The very limited disease review in the DEIS is generally based on geographic characteristics of the disease in the context of interaction between domestic sheep and bighorn sheep. While this is a useful component of much needed research, it is not in itself enough to make well-informed recommendations on policy alternatives. There remains limited knowledge of transmission dynamics. Clinical studies have shown bighorn sheep susceptibility to disease from contact with domestic sheep. However, epidemiologic modeling is needed to understand how contacts with domestic sheep, bighorn sheep, and other disease carriers (elk, deer, wild goats, birds, etc.), forage and climatic conditions, and other factors affect the dynamics of the

disease spread under various management alternatives. The Gallatin NF does not appear to apply any sort of modeling for the risk of disease transmission on the Forest.

NEPA's procedures require the presentation of "complete and accurate information to decision makers and to the public to allow an informed comparison of the alternatives considered in the EIS." *NRDC v. U.S. Forest Service*, 421 F.3d at 813. Here, further modeling and additional study is needed to determine the added probability of disease transmission among bighorns and from other animals. The probability that healthy "carrier" bighorns are infecting "non-carrier" bighorns is likely high, since a large number of the bighorns on the Gallatin NF may be disease-carriers. Additionally, more information and study should be undertaken to determine the exact mechanism for developing pneumonia in bighorn sheep following association with domestic sheep or other animals. Further, the Gallatin NF must study the development of immunity to disease in bighorn sheep. All of this information should be considered and addressed by the Gallatin NF in the DEIS.

17. The Gallatin NF Fails to Consider the Most Important Aspects of the Problem in the DEIS

Under the APA, agency decisions under NEPA and NFMA will be set aside if they are "arbitrary, capricious, an abuse of discretion, or otherwise not in accordance with law." 5 U.S.C. § 706(2)(A). Under this standard, judicial review of agency action seeks to determine whether an agency "has relied on factors which Congress has not intended it to consider, entirely failed to consider an important aspect of the problem, offered an explanation for its decision that runs counter to the evidence before the agency, or is so implausible that it could not be ascribed to a difference in view or the product of agency expertise." *See Motor Vehicle Mfrs. Ass'n v. State Farm Mut. Auto. Ins. Co.*, 463 U.S. 29, 43 (1983); *Utah Environmental Congress v. Bosworth*, 443 F.3d 732, 739 (10th Cir. 2006).

In its DEIS, the Gallatin NF has failed to consider and acknowledge that the proposed alternative is unlikely to control disease transmission and is implausible. Disease could still be a factor for bighorn sheep populations on the Gallatin NF, regardless of the closure of the Forest to pack goats. The DEIS fails to give importance to the fact that bighorn sheep themselves on the Gallatin NF in fact already carry the pathogens that lead to disease. Thus, bighorn sheep are at risk of contacting other bighorn sheep that carry the pathogens that can lead to disease.

Because bighorn sheep are carriers of the pathogens that can lead to disease, contact with other bighorn sheep not only puts bighorn sheep populations at risk, but renders irrelevant pack goats as the vector for transmission of the pathogens (assuming that pack goats on the Gallatin NF are carriers of the pathogens). This can mislead readers to believe that eliminating risk of contact on the Gallatin NF between pack goats and bighorn sheep will eliminate the threat of disease transmission. Under this misleading premise, the DEIS appears to be designed to depict pack goats as a significant cause of disease transmission or even risk of disease transmission, which is not accurate. The alternatives and the discussion in the DEIS must acknowledge more fully the potential futility of alternatives and explain the need for more comprehensive solutions to the problem of disease transmission, such as the development of a vaccine, or the improvement of bighorn sheep immunity, or the improvement of bighorn sheep habitat.

The DEIS also fails to consider that other animals on the Gallatin NF, like elk, deer, birds, etc., may carry the pathogens that can lead to diseases. Thus, contact between cattle and other animals, besides pack goats, and bighorn sheep may lead to disease transmission on the Gallatin NF. The DEIS does not discuss this possibility. In addition, the DEIS fails to acknowledge that bighorn sheep are at risk of contact with domestic sheep and other animals off the areas controlled by the Gallatin NF, and which risk is not mitigated by the alternatives or the ban on pack goat use.

Because the DEIS wholly fails to consider the risks of disease transmission from other bighorns, the risks of disease transmission off the Forest, and risks of disease transmission from other sources, the DEIS is inadequate under NEPA. As a result, the DEIS must be revised to consider risks of disease transfer from other bighorns, off of the forest and from other sources.

18. The DEIS Does Not Properly Address the Relevance of Unavailable or Incomplete Scientific Information

The Gallatin NF acknowledges in the DEIS that it lacks complete information to assess the potential effects of disease transmission between domestic sheep and domestic goats and bighorn sheep, let alone pack goats and bighorn sheep. The DEIS does little to address the lack of information with its subsequent conclusions.

In situations such as this, where the relevant information for assessing impacts is incomplete or unavailable, the agency preparing the EIS must take the following steps: first, if the incomplete information relevant to reasonably foreseeable adverse effects is essential to a reasoned choice among alternatives and the overall costs of obtaining the information is not exorbitant, the agency must include that information in the EIS. Next, if the relevant information cannot be obtained because the overall costs are exorbitant or the means of obtaining the information are not known, then an agency must include in an EIS:

- (1) a statement that such information is incomplete or unavailable;
- (2) a statement of the relevance of the incomplete or unavailable information to evaluating reasonably foreseeable significant impacts on the human environment; (3) a summary of existing credible scientific evidence which is relevant to evaluating the reasonably foreseeable significant adverse impacts on the human environment; and (4) the agency's evaluation of such impacts based upon theoretical approaches or research methods generally accepted in the scientific community.

40 C.F.R. § 1502.22(b).

Here, the Forest Service fails to take these required steps to address the incomplete or unavailable information relevant to ascertaining the possibility and consequences of disease transmission between domestic sheep/domestic goats and bighorns, and further fails to do so pertaining to the lesser risks of disease transmission from domestic goats in comparison to domestic sheep. The evidence provided in the DEIS suggests the likelihood or risk of disease transfer is largely specific to domestic sheep and not to domestic goats. The DEIS fails here to

include contrasting scientific points of view that have studied the differences in disease transfer risk between domestic sheep and bighorns, and domestic goats and bighorns. Here also, the DEIS fails to distinguish relevant information pertaining to disease transfer between other free ranging animals as comparable to easily managed and controlled animals like pack goats. Likewise, the DEIS fails to contain a clear and direct statement that the required information is incomplete or unavailable. The DEIS also fails to discuss the relevance of incomplete or unavailable information in light of evaluation of a reasonably foreseeable environmental impact. Lastly, the DEIS fails to contain the Forest Service's own evaluation of such impacts "based upon theoretical approaches or research methods generally accepted in the scientific community." *Id.*

Instead of honestly evaluating the range of potential scientific opinion applicable to disease transmission between pack goats and bighorns, the Forest Service impermissibly fails to comply with the requirements of the CEQ regulations to address incomplete or unavailable scientific information. Based on this fundamental flaw in the evaluation of environmental consequences in the DEIS, the DEIS should be revised to provide further analysis.

19. The Gallatin NF must Obtain Additional Information for the DEIS

When particular information "relevant to reasonably foreseeable significant adverse impacts is essential to a reasoned choice among alternatives," the agency must obtain that information and include it in the EIS, unless the cost is "exorbitant or the means to obtain it are not known." 40 C.F.R. § 1502.22. If obtaining the information is too costly or infeasible, the agency can forego its collection, providing full explanation in the EIS. *Id.* § 1502.22(b). "In that case the agency must include in the EIS: (1) A statement that the information is incomplete or unavailable; (2) a statement of the relevance of the incomplete or unavailable information; (3) a summary of relevant "existing credible scientific evidence;" and (4) the agency's evaluation of impacts based on "theoretical approaches or research methods generally accepted in the scientific community." *Id.*

The Gallatin NF has not included the following relevant information in the DEIS:

- Information indicating the differences between pack goats and other domestic goats;
- Information indicating that pack goats carry disease that can be transmitted to bighorn sheep;
- Information indicating that pack goats may come into contact or have come into contact with bighorn sheep on the Gallatin NF;
- Information indicating that BMPs and/or mitigation measures are not effective to ensure separation between pack goats and bighorn sheep on the Gallatin NF;
- Information indicating that pack goats may transmit or have transmitted disease to bighorn sheep on the Gallatin NF;

- Information indicating that bighorn sheep have contracted disease transmitted by pack goats on the Gallatin NF;
- Information indicating that bighorn sheep that have contracted disease transmitted by pack goats on the Gallatin NF have returned to their herds and infected other bighorn sheep;
- Information indicating that bighorn sheep that have contracted disease transmitted by pack goats on the Gallatin NF have returned to their herds and infected other bighorn sheep, which has led to a die-off;
- Information indicating that there is a risk of disease transmission from pack goats to bighorn sheep on the Gallatin NF;
- Information indicating the risk of disease transmission from other animals on and off of the Gallatin NF to bighorn sheep;
- Information indicating the impacts of wolves, mountain goats, and hunting on bighorn sheep populations on the Gallatin NF; and
- Information indicating the recreational, social and economic impacts on goatpackers of a closure of all or part of the Gallatin NF to pack goats.

EXHIBIT A

NAPgA Best Management Practices (BMP'S)

The BMP document is a living document which is open to editing and updating as needed.

NAPgA created the BMP's to establish responsible common sense guidelines for goatpacking. They are not intended to be overly restrictive or to discourage packgoat use in any way or in any location.

NAPgA will use best available science as a guide in which to measure and develop the BMP's to address wildlife and other resource concerns.

BMP#1: Individually Identify Your Packgoats

Each packgoat shall be individually identified. Each goat shall have a collar with a tag attached to it containing, at a minimum, the current owner's name and phone number.

Packgoats may be identified with a tattoo or microchip which is specific to each individual goat in conjunction with a collar.

Tattoos containing the individual packgoat's Scrapie Herd Number & ID or an official Scrapie ear tag may be used in conjunction with a collar.

BMP#2: Control

All packgoats shall be under direct human supervision at all times. They shall be on leads or have leads attached to their collar/halter.

In camp all packgoats shall be in direct sight or tethered in some fashion (picketing, high lining, etc.).

All packgoats shall be tethered at night within 30 feet of humans and bells will be attached to their collars.

BMP#3: Separation

Goatpackers shall minimize packgoat contact with wildlife.

BMP#4: Lost Packgoat

If a packgoat becomes lost every effort will be exhausted to locate and recover it.

If the owner is unable to locate and recover the lost packgoat the following agencies shall be contacted by telephone as soon as possible.

Information given should include a detailed description of the packgoat (size; color; ears erect, hanging or none, horned or not), any equipment they are carrying and the last known location. A photograph of the packgoat, if possible.

The local County Sheriff's office. Call 911 or the non-emergency line to dispatch of that county. Most hikers, hunters, land owners or citizens will call the sheriff's office first if they find a lost pack stock animal.

The state's Department of Fish and Game or Fish.

The local land management agency responsible for the area where the packgoat was lost. (Forest Service/BLM/DNR).

Post information, including photos if available, at convenience stores, trail heads and camp grounds with owners contact information, goat and gear descriptions.

Contact the North American Pack Goat Association (NAPgA) to report the loss. NAPgA will maintain a documentation file on all lost pack goats. NAPgA will request an initial report as well as an after-action report from the packgoat's owner/user. The information will be used for documentation as well as continued training and educational awareness training for pack goat users.

Contact the North American Pack Goat Association (NAPgA) to report the loss. NAPgA will maintain a documentation file on all lost pack goats. NAPgA will request an initial report as well as an after-action report from the packgoat's owner/user. The information will be used for documentation as well as continued training and educational awareness training for pack goat users.

BMP#5: Leave No Trace

Leave No Trace principles are strongly encouraged.

Leave No Trace principles are found on this website: <https://lnt.org/learn/7-principles>

EXHIBIT B

Packgoats and *Mycoplasma ovipneumoniae* Prevalence Study 2016 **North American Packgoat Association Summary of Understanding**

Mycoplasma ovipneumoniae, often referred to by the nickname “Movi” (or some variation of that) is the pathogen currently believed to be the most likely primary cause of outbreaks of bighorn sheep pneumonia that have threatened recovery of that species. On November 10, 2015 information was presented at The Technical Packgoat Meeting to NAPgA and the Blue Mountain Forest Plan Revision team in Pendleton, Oregon that goats had a 90% prevalence rate of *M. ovipneumoniae*. In clarifying this information Dr. Tom Besser noted in an email Dec 15, 2015 that this information was obtained from a “report of a large US survey of sheep operations tested for MOVI”. Domestic goats are different than domestic sheep and most certainly packgoats are very different from domestic sheep on public lands grazing allotments.

To consider packgoats the same as sheep for purposes of analyzing the risk of disease (pathogen) transmission to bighorn sheep is in error. Packgoat owners train packgoat prospects from a young age. Packgoats are inextricably bonded to their owner, which represents the “alpha goat” in their small herd. The lifestyle and care of a packgoat in herds of 2 to 10 differs greatly from that of a typical herd of domestic sheep or goats which can range in size of hundreds to thousands. Packgoats are seen by their owners as a significant investment in time and resources for 3 or 4 years before they are viable for packing purposes. Throughout a packgoat’s life, the packgoat receives routine veterinary care in order to keep the goat healthy and prolong their useful life.

Available literature at the time of this 2015 meeting quoted decades-old science in its discussion of evidence for “disease transmission” from domestic goats to BHS. There was no, and to date remains no, scientific support to implicate packgoats in BHS die-offs. Goats and sheep are different species and the scientific data from captive commingling experiments concerning pathogen (*M. ovipneumoniae* or other historically examined pathogens, such as members of the Pasteurellaceae family of bacteria) transmission to bighorn sheep and subsequent disease is vastly different. The types of *M. ovipneumoniae* carried by domestic sheep differ genetically from those carried by domestic goats (Maksimovic, Cassirer, unpublished data). Goat types or “strains” of *M. ovipneumoniae* have resulted in relatively mild (non-fatal) respiratory illness, dramatically different than the nearly 100% fatality reported from captive commingling with domestic sheep. To group sheep and goats together, and even packgoats and other types of domestic goats, in the discussion of pathogen or disease transmission falsely implicates packgoats in BHS die-off’s.

In more recent research by Besser *et al.* (2016), not a single domestic goat or bighorn sheep succumbed to any sort of pneumonia before or after being infected with a “goat type” of *M. ovipneumoniae* and not a single animal died as a result of disease during the study. Domestic goats were not shown to cause deaths of bighorn sheep as a result of pathogen (“disease”) transmission, even when the 3 study goats, were inoculated/infected with a “goat type” of *M. ovipneumoniae* and forced to commingle with bighorn sheep for 100 days. All animals in the study, both the domestic goats and bighorn sheep began showing symptoms of respiratory illness, and all of them recovered prior to being euthanized by the researchers. While the publication would imply that “sub-lethal pneumonia” was

induced in the bighorn sheep in this study, this is not consistent with the histopathology reports from lung tissue that was submitted to the Washington Animal Disease Diagnostic Laboratory in Pullman, WA. Those reports indicated that there were minimal to mild changes that are typically seen in small ruminants that are infected with *M. ovipneumoniae* (bronchiolar associated lymphoid tissue (BALT) hyperplasia and hyperplasia of the bronchial/bronchiolar epithelium); but no diagnosis of pneumonia was reported.

NAPgA is the leading organization in making recommendations on how to safely recreate with packgoats around BHS habitat. The complete lack of relevant research regarding *M. ovipneumoniae* prevalence in packgoats lead NAPgA to contact the USDA - Agricultural Research Unit - Animal Disease Research Unit (ARDU) in December of 2015. ARDU and APHIS (Animal and Plant Health Inspection Service) developed a packgoat *M. ovipneumoniae* surveillance research project.

In the spring of 2016 NAPgA recruited packgoat owners to participate in this research project. Consent was obtained from each packgoat owner. The majority of samples were collected by APHIS personnel and the remainder by Margaret Highland, DVM, PhD, Dipl. ACVP. Duplicate swabs were collected by both APHIS personnel and Dr. Highland. One swab was tested in the ARDU-ARS-USDA laboratory and the other was tested in the Washington Animal Disease Laboratory (except for kids <6 months of age and some of the non-packers that were also tested, which were tested only in the USDA-ARS-ARDU laboratory, as a means to save on research funds, since these animals are not used for packing).

A **packgoat owner survey** was completed. Information obtained was as follows:

- Goat information: Age, Sex, Breed
- Number of goats on premises (packers, non-packers)
- Illness(es) within the last year, including pinkeye/respiratory disease
- Any recent (last month) use of antibiotics
- Vaccination and antiparasitic regimen
- Use of packgoats on public lands? Proximity to bighorn sheep?

Samples collected (spring-fall 2016)

Packgoats

- 3 sets of duplicate nasal swabs collected at 4 week minimum intervals (few premises had only 1 or 2 sample collections)
- 1 blood sample for serum
- Other goats (milkers/breeders/etc) on premises were also tested
- At a minimum, 1 or 2 nasal swabs collected, at 1 to 3 time points
- Not all premises had “non-packer” goats on premises sampled
- All samples processed within 72 hours of collection

Sample Testing

- Nasal Swab samples tested by PCR and/or qPCR; positive samples confirmed by DNA sequencing
- PCR = polymerase chain reaction = technique that amplifies a segment of the bacteria’s genome to determine if it is present
- Duplicate nasal swabs from the first sample collection submitted to the Washington Animal Disease Diagnostic Laboratory (qPCR analysis)
- Serum samples are currently banked frozen

Distribution

State	#premises	#packgoats	# other goats	Total
AZ	3	16	23	39
CA	6	16	42	58
CO	8	29	12	41
ID	25	101	35	136
KS	1	13	51	64
MT	5	21	6	27
NM	1	2	0	2
NV	2	8	0	8
OR	9	32	3	35
UT	5	34	2	36
WA	14	65	17	82
WY	4	40	3	43
Total	83	377	194	571

“Other goats” = milkers, bucks, kids under 4 months of age which would not be out packing or on long hikes

WADDL Test Results

# Goats Tested	Detected	Indeterminate *	Not Detected
485 (83premises)	18 (5 premises)	20 (9 premises, 3 overlapped with the detected premises)	474 (72 premises)
	3.7% (6.0%premises)	4.1% (10.8%premises)	92.2% (86.7%premises)

* Indeterminate indicates that either there was an extremely low number of *M. ovipneumoniae* present in the sample OR the sample is truly negative, and the low detection is a false positive

WADDL Laboratory Test Results

NAPgA believes the large number of samples tested by the AAVLD accredited state diagnostic laboratory (WADDL) provide sufficient and valid evidence as to the very low prevalence of *M. ovipneumoniae* in packgoats.

ADRU-ARS-USDA Laboratory Results

8.2%, or 47, of all goats tested (n=571) had at least 1 sample in which *M. ovipneumoniae* was detected. Twenty-six of the positive animals were ≤4 months old, 35 were ≤12 months, and when considering only the “packers”, 3.3% overall had *M. ovipneumoniae* detected on at least 1 sample collection. 10 of the 14 premises with at least 1 positive detection were premises reported to house kids or were a premises in which the packgoat(s) were in recent contact with a positive packgoat or kids from a positive premises. These results have not yet been published in a peer-reviewed venue. Overall NAPgA will provide the complete report after peer-reviewed publication.

This is a living document and will be updated as new scientific evidence-based information is available.

From: Highland, Margaret
Sent: Friday, May 05, 2017 9:59 AM
To: 'Steve Kilpatrick' <skilpatrick@wyomingwildsheep.org>; 'Ron Smith' <rsagebrushsmith@aol.com>; canyonshadows@wyoming.com; johnmionne@gmail.com; packgoat@icloud.com; ctrulock@fs.fed.us; sschacht@fs.fed.us; brandonjhouck@fs.fed.us; rvandervoet@blm.gov; Lander_WYMail@blm.gov; daryl.lutz@wyo.gov; pat_hnilicka@fws.gov; sara@bighorn.org
Cc: 'Knowles, Don' (dknowles@vetmed.wsu.edu)' <dknowles@vetmed.wsu.edu>
Subject: RE: Pack Goat Meeting rescheduled

Since this may not occur before a final decision is made on the Shoshone NF, I would like to share with this group the data from the large scale pack goat study that was performed in 2016. While the ocular swabs are now and finally being tested after developing and validating PCR assays for detecting the 4 most common bacterial agents of pink eye (this process was much slower than anticipated by me), the *Mycoplasma ovipneumoniae* results are completed. The following, in quotes, is an email that I shared with Jim Wilder on 12/16/17. Since then we have retested all of the pack goat nasal swabs a 3 time with a more sensitive standard PCR method, the update on the findings from this follow the email correspondence.

“Over the last year we (ADRU-ARS-USDA), in collaboration with APHIS, were able to complete a fairly large scale surveillance study testing nasal shedding/presence of *Mycoplasma ovipneumoniae* in pack goats. We also tested goats that were housed with or on the same premises as domestic goats that were reported by the owner to be used specifically for packing. We also collected ocular swabs from participating goats to test for the presence of the common agents of small ruminant pink eye (*Chlamydophila* sp and *Mycoplasma conjunctivae*, *Moraxella ovis*, and *Acheloplasma oculi*); the ocular swabs are still being analyzed, with hopes of completing analysis this month. Upon analysis completion of the ocular swabs, the plan is to report the results by publishing in a peer-reviewed scientific journal by the end of winter/early spring.

I would like to share with you the following results from the nasal swab samples that were collected:

Nasal swabs were collected 3 times, at 1 month minimum intervals, from participating goats (aside from the handful of animals that were sold, removed from the study as per the owners discretion, or entered into the study late so had fewer sample time points). A couple of the premises had 4 or 5 samples collected. Duplicate nasal swabs were collected at each time point. 1 swab was tested in our USDA laboratory and samples that tested negative were then submitted to an independent laboratory for confirmation of the results (WADDL in Pullman, WA was the independent laboratory).

We tested a total of 576 domestic goats from 84 premises which included the following states (# of premises in parentheses after each): AZ (3), CA (6), CO (7), ID (26), KS (1), MT (5), NM (1), NV (2), OR (9), UT (5), WA (14), WY (4), VT (1). (I believe I had reported that there were 88 premises in earlier info that I shared with Mark P.....I forgot to deduct the 4 premises scattered in 4 eastern states that we didn't get tested).

Of all of the premises tested, we confirmed *M. ovipneumoniae* to be present in nasal

secretions from goats on 2 premises, limited to kids ≤ 2 weeks of age at only one test time. We collected additional swabs from 1 of these premises for 5 times total sample collections and the last 3 collection points had no detected *M. ovipneumoniae* and interestingly, all of the adult goats (9 of them) never had *M. ovipneumoniae* detected....the kids (there were 15 of them total) had 3 positives at time point 1, and 2 different kids positive at time point 2 (1st 3 positive were negative at this 2nd time point) and all goats on the premises were negative the last 3 sample collections.

As for the other premises that had a handful of positive kids: I repeat swabbed several of them 1 or 2 more times and they too were subsequently negative on the repeat samplings. This “kid phenomenon” is interesting.....I’ll leave it at that as to save typing time in this already lengthy email, but am happy to discuss further some time if you are interested. One additional premises that had *M. ovipneumoniae* detected 2 of the 3 sample times had a small group of yearling pack goats that were being housed at fence line with an ‘open’ breeding herd of registered Boer goats that were used for shows and sent out to farms for sire purposes. I instructed that owner to move his packers as soon as possible away from the large group of traveling Boer goats.....I suspect that his pack goats may clear (not shed) *M. ovipneumoniae* without the constant potential exposure, as all of his goats were negative on the 3 sample collection (I’d be happy to discuss why I suspect this may be possible with you too, if you’re interested).

The other 81 premises had no confirmed *M. ovipneumoniae* present on any of the nasal swabs collected. Of interest to your local and nearby area, none of the WY, UT, CO, MT herds had confirmed *M. ovipneumoniae* detection at any of the time points. 1 of the places with “kid detected *M. ovipneumoniae*” was in ID, but these kids are the ones that have sense been negative and the adults never positive.

While nothing is ever 100% risk free in life, I think this data strongly supports that there is a very low prevalence of *M. ovipneumoniae* in goats, at least those raised and kept in closed and typically small groups (however, a few of the premises that I tested had 20+ goats though and still negative....even the premises that tested their milk goats).

I would also like to take the time here to give warning that unless researchers and/or diagnosticians are looking beyond the common published techniques for identifying *M. ovipneumoniae*, there is a chance that false positive results will occur...particularly in goats. For example, we know that the published PCR primers, referred to as “LM primers” and qPCR techniques that have been developed in the past based on these primers can (and do) result in false positive results. By “looking beyond” I mean perform standard PCR to amplify a minimum of 2 regions of the bacterial genome and sequence the products/amplicons.....and making sure that the products/amplicons match well-characterized strains of *M. ovipneumoniae* (ie. strains that are characterized by reputable groups such as ATCC). Mycoplasmas are tricky, to say the least. Again, I’m happy to discuss more should you be interested.

Please feel free to let me know, either by email or phone (listed in signature line), if you have questions, comments, or concerns about the information provided herein or if you have anything that you would like to further discuss with me regarding the bighorn pneumonia phenomenon.”

Update following repeated testing using a more sensitive method of detection:
Five of the 83 premises tested (6%) had *M. ovipneumoniae* identified during the repeat nasal sample collections. Premises that had *M. ovipneumoniae* detected in any the goats

had at least 7 goats housed on the premises. *M. ovipneumoniae* was confirmed to be present on the nasal swabs collected from 30 of the 576 total goats tested, meaning that 94.8% of the goats tested had no *M. ovipneumoniae* detected at any of the sample collection time points. Of the 30 total *M. ovipneumoniae* positive goats, 27 (or 90%) of the were ≤ 1 year of age, and 23 of them were < 5 months of age.

During the 2016 North American Pack goat annual gathering (“the Rendy”) held in Oregon, I sampled in total 27 adults and 2 kid goats whose owners brought them to the sample collection site that I set up. Most of these goats were already part of the large pack goat/domestic goat surveillance study and I asked owners if they minded me taking an extra nasal swab from their animals with the thought that perhaps the stressor of travelling or bringing a large group of goats together may result in shedding of *M. ovipneumoniae* from animals that it hadn’t been detected on during the first round of sample collections and it also gave the opportunity to add a couple more premises to the study. *M. ovipneumoniae* **was not detected** on any of the swab samples collected at the Rendy.

It’s unfortunate how long research takes, particularly with something as time sensitive as this seems to be, as I had truly hoped that this entire study would be out in published in a peer-reviewed form at this point (April was my goal). Hoping now for June with fingers crossed that all of the ocular swab testing goes smoothly....and more importantly accurately with good specificity and sensitivity.

Thank you and I look forward to participating in the Pack Goat meeting whenever the final date is decided upon.

Maggie

Margaret A. Highland, DVM, PhD, Dipl. ACVP
Animal Disease Research Unit-ARS-USDA (VMO Researcher)
Washington Animal Disease Diagnostic Laboratory (Adjunct Pathologist)
School for Global Animal Health (Adjunct Faculty)
Washington State University
Pullman, WA 99164

Office phone: 509-335-6327
Cell phone: 608-213-3025
Fax: 509-335-8328

ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University

Web Site: <http://waddl.vetmed.wsu.edu>

US Postal Service mailing address:
 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
 Bustad Hall, Rm.155-N
 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

2016-6030
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routed: and

05/10/16
 form 2 pages

Please type or use black ink and print clearly.

Veterinarian or Last Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF-WSU		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99164	
Phone: 509-335-6327	Fax: 509-335-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☒ Owner ☐ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.
Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted:		Date Collected: April 2016	
nasal swabs		Date Shipped:	
(Please use WADDL Animal ID Sheet for multiple animals.)			
Tests Requested: <input type="checkbox"/> Necropsy <input type="checkbox"/> Virology <input type="checkbox"/> Bacteriology <input type="checkbox"/> IHC <input type="checkbox"/> Histopathology <input type="checkbox"/> Serology <input type="checkbox"/> Mycoplasma culture <input checked="" type="checkbox"/> PCR <input type="checkbox"/> Toxicology <input type="checkbox"/> Fungal Culture <input type="checkbox"/> Parasitology <input type="checkbox"/> Other:			
Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.			
Animal ID (name/tag#)	Species	Breed	Age
see multiple animal form	goat	multiple	1mo-12yrs
Sex	Animal Weight		
Location of Lesion	No. in group	No. Dead	No. Sick
N/A			
* Was animal euthanized? If so, what method? N/A			

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

Please save any remaining DNA isolations and call Maggie for pick up.

Bill to ADRU-ARS-USDA acct #RSA 2540-1080

Samples were maintained ~~on ice~~ on ice then frozen w/in 2 days of collection + kept at -20°C since.

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's or Owner's Signature:	Condition(s) Suspected:
---	-------------------------

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: Shipping address:
 P.O. Box 647034 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 FAX: (509) 335-7424
 E-Mail: waddl@vetmed.wsu.edu
 Web Site: http://waddl.vetmed.wsu.edu

2016-6030
 Ref Vet: Highland, Margaret
 Owner: USDA-ARS-ADRU
 Breed: Domestic Goat
 Routing: ,md

Owner: ADRU-ARS-USDA

Veterinarian: Maggie Highland

TEST(S) REQUESTED: Mycoplasma ovipneumoniae qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal
1	3_A	26	5_F	51	3_D	76	
2	3_B	27	5_G	52	7_A	77	
3	3_C	28	5_H	53	7_B	78	
4	11_A	29	6_A	54	7_C	79	
5	11_B	30	6_B	55	7_D	80	
6	11_C	31	6_C	56	7_E	81	
7	11_D	32	6_D	57	11_A	82	
8	16_A	33	6_E	58	11_B	83	
9	16_B	34	6_F	59	11_C	84	
10	4_A	35	6_G	60	11_D	85	
11	4_B	36	8_A	61	12_A	86	
12	4_C	37	8_B	62	12_B	87	
13	4_D	38	8_C	63	12_C	88	
14	4_E	39	8_D	64	12_D	89	
15	4_F	40	9_A	65	12_E	90	
16	4_G	41	9_B	66	12_F	91	
17	10_A	42	9_C	67	12_G	92	
18	10_B	43	2_A	68	12_H	93	
19	10_C	44	2_B	69	12_I	94	
20	10_D	45	2_C	70	12_J	95	
21	5_A	46	2_D	71	12_K	96	
22	5_B	47	2_E	72	12_L	97	
23	5_C	48	3_A	73		98	
24	5_D	49	3_B	74		99	
25	5_E	50	3_C	75		100 *	

05/10/16

* For over 100 samples, please copy this form and continue numbering.

Washington Animal Disease Diagnostic Lab

P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424

Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF

Case#: 2016-6030
Report Date: 05/16/16

Pullman, WA 99164-6630

Submittal Date: 05/10/16
Owner: USDA-ARS-ADRU

Species: Domestic Goat

Age:
Sex:

Final Report:

Molecular Diagnostics- Reported on 05/16/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
3_A	Nasal swab	Not detected
3_B	Nasal swab	Not detected
3_C	Nasal swab	Not detected
11_A	Nasal swab	Not detected
11_B	Nasal swab	Not detected
11_C	Nasal swab	Not detected
11_D	Nasal swab	Not detected
16_A	Nasal swab	Not detected
16_B	Nasal swab	Not detected
4_A	Nasal swab	Detected
4_B	Nasal swab	Detected
4_C	Nasal swab	Detected
4_D	Nasal swab	Detected
4_E	Nasal swab	Detected
4_F	Nasal swab	Detected
4_G	Nasal swab	Detected
10_A	Nasal swab	Indeterminate
10_B	Nasal swab	Not detected
10_C	Nasal swab	Not detected
10_D	Nasal swab	Not detected
5_A	Nasal swab	Not detected
5_B	Nasal swab	Not detected
5_C	Nasal swab	Not detected
5_D	Nasal swab	Not detected
5_E	Nasal swab	Not detected
5_F	Nasal swab	Not detected
5_G	Nasal swab	Not detected
5_H	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
V1.6.A	Nasal swab	Not detected
V1.6.B	Nasal swab	Not detected
V1.6.C	Nasal swab	Not detected
V1.6.D	Nasal swab	Not detected
V1.6.E	Nasal swab	Not detected
V1.6.F	Nasal swab	Not detected
V1.6.G	Nasal swab	Not detected
V1.8.A	Nasal swab	Not detected
V1.8.B	Nasal swab	Not detected
V1.8.C	Nasal swab	Not detected
V1.8.D	Nasal swab	Not detected
V1.9.A	Nasal swab	Not detected
V1.9.B	Nasal swab	Not detected
V1.9.C	Nasal swab	Not detected
V1.2.A	Nasal swab	Not detected
V1.2.B	Nasal swab	Not detected
V1.2.C	Nasal swab	Not detected
V1.2.D	Nasal swab	Not detected
V1.2.E	Nasal swab	Not detected
V1.3.A	Nasal swab	Not detected
V1.3.B	Nasal swab	Not detected
V1.3.C	Nasal swab	Not detected
V1.3.D	Nasal swab	Not detected
V1.7.A	Nasal swab	Not detected
V1.7.B	Nasal swab	Not detected
V1.7.C	Nasal swab	Not detected
V1.7.D	Nasal swab	Not detected
V1.7.E	Nasal swab	Not detected
V1.11.A	Nasal swab	Not detected
V1.11.B	Nasal swab	Not detected
V1.11.C	Nasal swab	Not detected
V1.11.D	Nasal swab	Not detected
V1.12.A	Nasal swab	Not detected
V1.12.B	Nasal swab	Not detected
V1.12.C	Nasal swab	Not detected
V1.12.D	Nasal swab	Not detected
V1.12.E	Nasal swab	Not detected
V1.12.F	Nasal swab	Not detected
V1.12.G	Nasal swab	Not detected
V1.12.H	Nasal swab	Not detected
V1.12.I	Nasal swab	Not detected
V1.12.J	Nasal swab	Not detected
V1.12.K	Nasal swab	Not detected
V1.12.L	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

72 nasal swabs

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms: <input type="checkbox"/> Yes <input type="checkbox"/> No Explain below:	Opened by:
Samples Received Via:	<input type="checkbox"/> US Mail	<input type="checkbox"/> FedEx	<input checked="" type="checkbox"/> Drop off			
	<input type="checkbox"/> UPS	<input type="checkbox"/> FedEx-R	<input type="checkbox"/> Other:			

Comments for Case Tracking:

by Margaret Highland

2016-6030

Ref Vet: Highland, Margaret
Owner: USDA-ARS-ADRU
Breed: Domestic Goat
Routed: ,md05/10/16
Page 1 of 1

Sample Label

[Signature]

ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Web Site: <http://waddl.vetmed.wsu.edu>

US Postal Service mailing address:
 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

2016-6160
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routed: and

05/12/16
 Item: 3 pages

Please type or use black ink and print clearly.

Veterinarian or Last Case Coordinator Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF-WSU		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99164	
Phone: 509-335-6327	Fax: 509-335-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☒ Owner ☐ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.
Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted:		Date Collected: 4/16-5/16	
(Please use WADDL Animal ID Sheet for multiple animals.)		Date Shipped:	
nasal swabs			
Tests Requested:	<input type="checkbox"/> Necropsy <input type="checkbox"/> Virology <input type="checkbox"/> Bacteriology <input type="checkbox"/> IHC		
	<input type="checkbox"/> Histopathology <input type="checkbox"/> Serology <input type="checkbox"/> Mycoplasma culture <input checked="" type="checkbox"/> PCR		
	<input type="checkbox"/> Toxicology <input type="checkbox"/> Fungal Culture <input type="checkbox"/> Parasitology <input type="checkbox"/> Other:		
<i>Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.</i>			
Animal ID (name/tag#)	Species	Breed	Age
see multiple animal form	domestic goats	multiple	1mo-12yrs
Sex	Animal Weight	No. in group	No. Dead
N/A	N/A	No. Sick	No. on Premises
Duration of Problem		N/A	

* Was animal euthanized? If so, what method? **N/A**

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

Nasal swabs for *M. ovipneumoniae* qPCR

Please save any remaining DNA isolations and call Maggie for pick up.

Bill to ADRU-ARS-USDA acct #RSA 2540-1080

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's or Owner's Signature: <i>Maggie Highland</i>	Condition(s) Suspected: None/Healthy Animals
--	---

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: Shipping address:
 P.O. Box 647034 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 FAX: (509) 335-7424
 E-Mail: waddl@vetmed.wsu.edu
 Web Site: http://waddl.vetmed.wsu.edu

2016-6160
 Ref Vet: Highland, Margaret
 Owner: USDA-ARS-ADRU
 Breed: Domestic Goat
 Routing: md

05/12/16

Owner: ADRU-ARS-USDA

Veterinarian: Maggie Highland

TEST(S) REQUESTED: M. ovipneumoniae qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
1	1_A	26	4_N	51	5_S	76	14_C
2	1_B	27	4_O	52	5_T	77	17_A
3	1_C	28	4_P	53	5_U	78	17_B
4	1_D	29	4_Q	54	5_V	79	17_C
5	1_E	30	4_R	55	5_W	80	17_D
6	1_F	31	4_S	56	5_X	81	17_E
7	1_G	32	4_T	57	5_Y	82	17_F
8	7_A	33	5_A	58	5_Z	83	17_G
9	7_B	34	5_B	59	8_A	84	17_H
10	7_C	35	5_C	60	8_B	85	17_I
11	7_D	36	5_D	61	8_C	86	22_A
12	7_E	37	5_E	62	9_A	87	22_B
13	4_A	38	5_F	63	9_B	88	22_C
14	4_B	39	5_G	64	9_C	89	23_A
15	4_C	40	5_H	65	9_D	90	23_B
16	4_D	41	5_I	66	9_E	91	23_C
17	4_E	42	5_J	67	19_A	92	23_D
18	4_F	43	5_K	68	19_B	93	23_E
19	4_G	44	5_L	69	10_A	94	23_F
20	4_H	45	5_M	70	10_B	95	23_G
21	4_I	46	5_N	71	6_A	96	2_A
22	4_J	47	5_O	72	6_B	97	2_B
23	4_K	48	5_P	73	6_C	98	12_A
24	4_L	49	5_Q	74	14_A	99	12_B
25	4_M	50	5_R	75	14_B	100 *	12_C

* For over 100 samples, please copy this form and continue numbering.

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: Shipping address:
 P.O. Box 647034 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 FAX: (509) 335-7424
 E-Mail: waddl@vetmed.wsu.edu
 Web Site: <http://waddl.vetmed.wsu.edu>

2016-6160
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routing: .md

Owner: ADRU-ARS-USDA

Veterinarian: Maggie Highland

TEST(S) REQUESTED: M. ovipneumoniae qPCR

05/12/16

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Anim
1	12_D	26		51		76	
2	20_A	27		52		77	
3	20_B	28		53		78	
4		29		54		79	
5		30		55		80	
6		31		56		81	
7		32		57		82	
8		33		58		83	
9		34		59		84	
10		35		60		85	
11		36		61		86	
12		37		62		87	
13		38		63		88	
14		39		64		89	
15		40		65		90	
16		41		66		91	
17		42		67		92	
18		43		68		93	
19		44		69		94	
20		45		70		95	
21		46		71		96	
22		47		72		97	
23		48		73		98	
24		49		74		99	
25		50		75		100 *	

* For over 100 samples, please copy this form and continue numbering

Washington Animal Disease Diagnostic Lab

**P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424**

**Case#: 2016-6160
Report Date: 05/16/16**

**Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF**

Pullman, WA 99164-6630

Submittal Date: 05/12/16
Owner: USDA-ARS-ADRU

Species: Domestic Goat

Age:
Sex:

Final Report:

Molecular Diagnostics- Reported on 05/16/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
1.A	Nasal swab	Not detected
1.B	Nasal swab	Not detected
1.C	Nasal swab	Not detected
1.D	Nasal swab	Not detected
1.E	Nasal swab	Not detected
1.F	Nasal swab	Not detected
1.G	Nasal swab	Not detected
7.A	Nasal swab	Not detected
7.B	Nasal swab	Not detected
7.C	Nasal swab	Not detected
7.D	Nasal swab	Not detected
7.E	Nasal swab	Not detected
4.A	Nasal swab	Not detected
4.B	Nasal swab	Not detected
4.C	Nasal swab	Not detected
4.D	Nasal swab	Not detected
4.E	Nasal swab	Not detected
4.F	Nasal swab	Not detected
4.G	Nasal swab	Not detected
4.H	Nasal swab	Not detected
4.I	Nasal swab	Detected
4.J	Nasal swab	Indeterminate
4.K	Nasal swab	Not detected
4.L	Nasal swab	Not detected
4.M	Nasal swab	Not detected
4.N	Nasal swab	Not detected
4.O	Nasal swab	Indeterminate
4.P	Nasal swab	Indeterminate

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
4-Q	Nasal swab	Indeterminate
4-R	Nasal swab	Not detected
4-S	Nasal swab	Detected
4-T	Nasal swab	Detected
5-A	Nasal swab	Not detected
5-B	Nasal swab	Not detected
5-C	Nasal swab	Not detected
5-D	Nasal swab	Not detected
5-E	Nasal swab	Not detected
5-F	Nasal swab	Not detected
5-G	Nasal swab	Not detected
5-H	Nasal swab	Not detected
5-I	Nasal swab	Not detected
5-J	Nasal swab	Not detected
5-K	Nasal swab	Not detected
5-L	Nasal swab	Not detected
5-M	Nasal swab	Not detected
5-N	Nasal swab	Not detected
5-O	Nasal swab	Not detected
5-P	Nasal swab	Not detected
5-Q	Nasal swab	Not detected
5-R	Nasal swab	Not detected
5-S	Nasal swab	Not detected
5-T	Nasal swab	Not detected
5-U	Nasal swab	Not detected
5-V	Nasal swab	Not detected
5-W	Nasal swab	Not detected
5-X	Nasal swab	Not detected
5-Y	Nasal swab	Not detected
5-Z	Nasal swab	Not detected
8-A	Nasal swab	Not detected
8-B	Nasal swab	Not detected
8-C	Nasal swab	Not detected
9-A	Nasal swab	Not detected
9-B	Nasal swab	Not detected
9-C	Nasal swab	Not detected
9-D	Nasal swab	Not detected
9-E	Nasal swab	Not detected
19-A	Nasal swab	Not detected
19-B	Nasal swab	Not detected
10-A	Nasal swab	Not detected
10-B	Nasal swab	Not detected
6-A	Nasal swab	Not detected
6-B	Nasal swab	Not detected
6-C	Nasal swab	Not detected
14-A	Nasal swab	Not detected
14-B	Nasal swab	Not detected
14-C	Nasal swab	Not detected
17-A	Nasal swab	Not detected
17-B	Nasal swab	Not detected
17-C	Nasal swab	Not detected
17-D	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
17.E	Nasal swab	Not detected
17.F	Nasal swab	Not detected
17.G	Nasal swab	Not detected
17.H	Nasal swab	Not detected
17.I	Nasal swab	Not detected
22.A	Nasal swab	Not detected
22.B	Nasal swab	Not detected
22.C	Nasal swab	Not detected
23.A	Nasal swab	Not detected
23.B	Nasal swab	Not detected
23.C	Nasal swab	Not detected
23.D	Nasal swab	Not detected
23.E	Nasal swab	Not detected
23.F	Nasal swab	Not detected
23.G	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
12.A	Nasal swab	Not detected
12.B	Nasal swab	Not detected
12.C	Nasal swab	Not detected
12.D	Nasal swab	Not detected
20.A	Nasal swab	Not detected
20.B	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

103 nasal swabs

- Dropped off by M. Highland

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No Explain below:	Opened by: <i>WTH</i>
	Samples Received Via: <input type="checkbox"/> US Mail <input type="checkbox"/> FedEx <input checked="" type="checkbox"/> Drop off <input type="checkbox"/> UPS <input type="checkbox"/> FedEx-R <input type="checkbox"/> Other:					

Comments for Case Tracking:

2016-6160
Ref Vet: Highland, Margaret
Owner: USDA - ARS - ADRI
Breed: Domestic Goat
Routed: .md05/12/16
Index: 1 page

Sample Label ✓

WTH

ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University

Web Site: <http://waddl.vetmed.wsu.edu>

US Postal Service mailing address:
 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
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 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

2016 - 7117
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routed: md

06/02/16
 form 2 pages

Please type or use black ink and print clearly.

Veterinarian or Case Coordinator: Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF 3033		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99163	
Phone: 5-6327	Fax: 5-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☐ Owner ☒ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.
Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted:		Date Collected: May 2016	
nasal swabs		Date Shipped:	
(Please use WADDL Animal ID Sheet for multiple animals.)			
Tests Requested:	<input type="checkbox"/> Necropsy <input type="checkbox"/> Histopathology <input type="checkbox"/> Toxicology	<input type="checkbox"/> Virology <input type="checkbox"/> Serology <input type="checkbox"/> Fungal Culture	<input type="checkbox"/> Bacteriology <input type="checkbox"/> Mycoplasma culture <input type="checkbox"/> Parasitology
		<input type="checkbox"/> IHC <input checked="" type="checkbox"/> PCR <input type="checkbox"/> Other: Mycoplasma ovipneumoniae qPCR	
<i>Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.</i>			
Animal ID (name/tag#)	Species	Breed	Age
see multiple animal form	domestic goats	-	adult
Sex	Animal Weight		
-	-		
Location of Lesion	No. in group	No. Dead	No. Sick
N/A	N/A	N/A	N/A
No. on Premises	Duration of Problem		
N/A	N/A		

* Was animal euthanized? If so, what method?

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

M. ovipneumoniae qPCR

Please save remaining DNA isolations and call Maggie for pick up.

Bill to ADRU-ARS-USDA acct #RSA 2540-1080

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's
 or Owner's Signature:

Condition(s)
 Suspected:

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: P.O. Box 647034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 E-Mail: waddl@vetmed.wsu.edu
 Shipping address: Bustad Hall, Rm. 155-N Pullman, WA. 99164-7034
 FAX: (509) 335-7424
 Web Site: <http://waddl.vetmed.wsu.edu>

2016-7117
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routing: .md

06/02/16

Owner: ADRU-ARS-USDA

Veterinarian: Highland

TEST(S) REQUESTED: M. ovipneumoniae qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	e
1	13_A	26		51		76	
2	13_B	27		52		77	
3	13_C	28		53		78	
4	13_D	29		54		79	
5	15_A	30		55		80	
6	15_B	31		56		81	
7	15_C	32		57		82	
8	15_D	33		58		83	
9		34		59		84	
10		35		60		85	
11		36		61		86	
12		37		62		87	
13		38		63		88	
14		39		64		89	
15		40		65		90	
16		41		66		91	
17		42		67		92	
18		43		68		93	
19		44		69		94	
20		45		70		95	
21		46		71		96	
22		47		72		97	
23		48		73		98	
24		49		74		99	
25		50		75		100 *	

* For over 100 samples, please copy this form and continue numbering.

Washington Animal Disease Diagnostic Lab

**P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424**

**Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF**

**Case#: 2016-7117
Report Date: 06/07/16**

Pullman, WA 99164-6630

Submittal Date: 06/02/16
Owner: USDA-ARS-ADRU

Species: Domestic Goat

Age: Adult
Sex:

Final Report:

Molecular Diagnostics- Reported on 06/07/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
13_A	Nasal swab	Not detected
13_B	Nasal swab	Not detected
13_C	Nasal swab	Not detected
13_D	Nasal swab	Not detected
15_A	Nasal swab	Not detected
15_B	Nasal swab	Not detected
15_C	Nasal swab	Not detected
15_D	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

8 nasal swabs
- dropped off
by MAH

2016-7117
Ref Vet: Highland, Margaret
Owner: USDA-ARS-ADRU
Breed: Domestic Goat
Routed: .md

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms:	Opened by:
Samples Received Via:	<input type="checkbox"/> US Mail	<input type="checkbox"/> FedEx	<input checked="" type="checkbox"/> Drop off	<input type="checkbox"/> Other:	<input checked="" type="checkbox"/> Yes	<input type="checkbox"/> No
	<input type="checkbox"/> UPS	<input type="checkbox"/> FedEx-R			Explain below: <i>not</i>	

Comments for Case Tracking:



06/02/16
notes: 1 page

Sample Label ✓

MAH

ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University
 Web Site: <http://waddl.vetmed.wsu.edu>

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 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
 Bustad Hall, Rm.155-N
 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

Ref Vet: Highland, Margaret
 Owner: Highland, Margaret
 Breed: Domestic Goat
 Routed: md

2016-7913

06/20/16
 form: 3 pages

Please type or use black ink and print clearly.

Veterinarian or Last Case Coordinator Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF 3033		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99164	
Phone: 509-335-6327	Fax: 509-335-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☐ Owner ☒ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.
Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted:		Date Collected: June 2016	
nasal swabs		Date Shipped:	
(Please use WADDL Animal ID Sheet for multiple animals.)			
Tests Requested:	<input type="checkbox"/> Necropsy <input type="checkbox"/> Histopathology <input type="checkbox"/> Toxicology	<input type="checkbox"/> Virology <input type="checkbox"/> Serology <input type="checkbox"/> Fungal Culture	<input type="checkbox"/> Bacteriology <input type="checkbox"/> Mycoplasma culture <input type="checkbox"/> Parasitology
		<input type="checkbox"/> IHC <input checked="" type="checkbox"/> PCR <input type="checkbox"/> Other:	
Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.			
Animal ID (name/tag#)	Species	Breed	Age
see multiple animal form	domestic goats	multiple	multiple
Sex	Animal Weight	Location of Lesion	No. in group
		N/A	N/A
No. Dead	No. Sick	No. on Premises	Duration of Problem
N/A	N/A		N/A

* Was animal euthanized? If so, what method?

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

M.ovipneumoniae qPCR on each sample.

Please save remaining DNA isolation and call Maggie for pick up or may request further testing (sequencing) be performed by WADDL, depending on the results of qPCR analyses.

Bill to ADRU-ARS-USDA acct #RSA 2540-1080

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's or Owner's Signature: <i>Maggie Highland</i>	Condition(s) Suspected: <i>N/A (surveillance)</i>
--	---

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University

Mailing address:

P.O. Box 647034

Pullman, WA. 99164-7034

Phone: (509) 335-9696

E-Mail: waddl@vetmed.wsu.edu

Web Site: http://waddl.vetmed.wsu.edu

Shipping address:

Bustad Hall, Rm.155-N

Pullman, WA. 99164-7034

FAX: (509) 335-7424

2016-7913

Ref Vet: Highland, Margaret

Owner:

Breed: Domestic Goat

Routing: md

06/20/16

Owner: ADRU-ARS-USDA

Veterinarian: Highland

TEST(S) REQUESTED: Mycoplasma ovipneumoniae qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
1	24_A	26	1_S	51		76	5_B
2	24_B	27	1_T	52		77	5_C
3	24_C	28	1_U	53		78	5_D
4	3_A	29	1_V	54		79	5_E
5	3_B	30	1_W	55		80	5_F
6	3_C	31	1_X	56		81	5_G
7	3_D	32	1_Y	57		82	5_H
8	1_A	33	1_Aa	58		83	5_I
9	1_B	34	1_Bb	59		84	5_J
10	1_C	35	1_Cc	60		85	5_K
11	1_D	36	1_Dd	61		86	5_L
12	1_E	37	1_Ee	62		87	5_M
13	1_F	38	1_Ff	63		88	5_N
14	1_G	39	1_Gg	64		89	5_O
15	1_H	40	1_Hh	65		90	5_P
16	1_I	41	1_Ii	66		91	4_A
17	1_J	42	1_Jj	67		92	4_B
18	1_K	43	1_Kk	68		93	2_A
19	1_L	44	1_Ll	69		94	2_B
20	1_M	45	1_Mm	70		95	2_C
21	1_N	46	1_Nn	71		96	2_D
22	1_O	47	1_Oo	72	4_A	97	2_E
23	1_P	48		73	4_B	98	2_F
24	1_Q	49		74	4_C	99	2_G
25	1_R	50		75	5_A	100*	2_H

* For over 100 samples, please copy this form and continue numbering.

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: Shipping address:
 P.O. Box 647034 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 FAX: (509) 335-7424
 E-Mail: waddl@vetmed.wsu.edu
 Web Site: http://waddl.vetmed.wsu.edu

2016-7913

06/20/16

Ref Vet: Highland, Margaret
 Owner:
 Breed: Domestic Goat
 Routing: md

Owner: ADRU-ARS-USDA

Veterinarian: HIGHLAND

TEST(S) REQUESTED: M. ovipneumoniae qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
1	5_A	26	2_A	51		76	
2	5_B	27	2_B	52	6_G	77	
3	5_C	28	2_C	53	13_A	78	
4	5_D	29	2_D	54	13_B	79	
5	5_E	30	3_A	55	13_C	80	
6	5_F	31	3_B	56	13_D	81	
7	5_G	32	3_C	57	13_E	82	
8	5_H	33	3_D	58	13_F	83	
9	5_I	34	4_A	59	2-b-H	84	
10	5_J	35	4_B	60	2-A	85	
11	5_K	36	4_C	61	2-B	86	
12	5_L	37	1_A	62	2-C	87	
13	5_M	38	1_B	63	2-D	88	
14	5_N	39	21_A	64	2-E	89	
15	5_O	40	5_A	65	2-F	90	
16	5_P	41	5_B	66	2-G	91	
17	4_A	42	5_C	67	2-H	92	
18	4_B	43	5_D	68	2-I	93	
19	1_A	44	5_E	69	2-A	94	
20	1_B	45	6_A	70	2B	95	
21	1_C	46	6_B	71	2-C	96	
22	1_D	47	6_C	72	2-D	97	
23	1_E	48	6_D	73	7-A	98	
24	7_A	49	6_E	74	7-B	99	
25	7_B	50	6_F	75		100 *	

* For over 100 samples, please copy this form and continue numbering.

Washington Animal Disease Diagnostic Lab

**P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424**

**Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF**

**Case#: 2016-7913
Report Date: 07/01/16**

Pullman, WA 99164-6630

Submittal Date: 06/20/16
Owner:

Species: Domestic Goat

Age:
Sex:

Final Report:

Molecular Diagnostics- Reported on 07/01/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
24_A	Nasal swab	Not detected
24_B	Nasal swab	Not detected
24_C	Nasal swab	Not detected
03_A	Nasal swab	Not detected
03_B	Nasal swab	Not detected
03_C	Nasal swab	Not detected
03_D	Nasal swab	Not detected
11_A	Nasal swab	Not detected
11_B	Nasal swab	Not detected
11_C	Nasal swab	Indeterminate
11_D	Nasal swab	Not detected
11_E	Nasal swab	Not detected
11_F	Nasal swab	Not detected
11_G	Nasal swab	Not detected
11_H	Nasal swab	Detected
11_I	Nasal swab	Not detected
11_J	Nasal swab	Not detected
11_K	Nasal swab	Not detected
11_L	Nasal swab	Not detected
11_M	Nasal swab	Not detected
11_N	Nasal swab	Not detected
11_O	Nasal swab	Not detected
11_P	Nasal swab	Not detected
11_Q	Nasal swab	Not detected
11_R	Nasal swab	Not detected
11_S	Nasal swab	Not detected
11_T	Nasal swab	Not detected
11_U	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
1.V	Nasal swab	Not detected
1.W	Nasal swab	Not detected
1.X	Nasal swab	Not detected
1.Y	Nasal swab	Not detected
1.Aa	Nasal swab	Not detected
1.Bb	Nasal swab	Detected
1.Cc	Nasal swab	Not detected
1.Dd	Nasal swab	Not detected
1.Ee	Nasal swab	Detected
1.Ff	Nasal swab	Detected
1.Gg	Nasal swab	Indeterminate
1.Hh	Nasal swab	Not detected
1.Ii	Nasal swab	Indeterminate
1.Jj	Nasal swab	Detected
1.Kk	Nasal swab	Not detected
1.Ll	Nasal swab	Indeterminate
1.Mm	Nasal swab	Not detected
1.Nn	Nasal swab	Indeterminate
1.Oo	Nasal swab	Detected
4.A	Nasal swab	Not detected
4.B	Nasal swab	Not detected
4.C	Nasal swab	Not detected
5.A	Nasal swab	Not detected
5.B	Nasal swab	Not detected
5.C	Nasal swab	Not detected
5.D	Nasal swab	Not detected
5.E	Nasal swab	Not detected
5.F	Nasal swab	Not detected
5.G	Nasal swab	Not detected
5.H	Nasal swab	Not detected
5.I	Nasal swab	Not detected
5.J	Nasal swab	Not detected
5.K	Nasal swab	Not detected
5.L	Nasal swab	Not detected
5.M	Nasal swab	Not detected
5.N	Nasal swab	Not detected
5.O	Nasal swab	Not detected
5.P	Nasal swab	Indeterminate
4.A	Nasal swab	Not detected
4.B	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
2.E	Nasal swab	Not detected
2.F	Nasal swab	Not detected
2.G	Nasal swab	Not detected
2.H	Nasal swab	Not detected
5.A	Nasal swab	Not detected
5.B	Nasal swab	Not detected
5.C	Nasal swab	Not detected
5.D	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

Animal	Specimen	Result
T.5.E	Nasal swab	Indeterminate
T.5.F	Nasal swab	Not detected
T.5.G	Nasal swab	Not detected
T.5.H	Nasal swab	Not detected
T.5.I	Nasal swab	Not detected
T.5.J	Nasal swab	Not detected
T.5.K	Nasal swab	Not detected
T.5.L	Nasal swab	Not detected
T.5.M	Nasal swab	Not detected
T.5.N	Nasal swab	Not detected
T.5.O	Nasal swab	Detected
T.5.P	Nasal swab	Indeterminate
Y.4.A	Nasal swab	Not detected
Y.4.B	Nasal swab	Not detected
A.1.A	Nasal swab	Not detected
A.1.B	Nasal swab	Not detected
A.1.C	Nasal swab	Not detected
A.1.D	Nasal swab	Not detected
A.1.E	Nasal swab	Not detected
L.7.A	Nasal swab	Not detected
L.7.B	Nasal swab	Indeterminate
L.2.A	Nasal swab	Not detected
L.2.B	Nasal swab	Not detected
L.2.C	Nasal swab	Indeterminate
L.2.D	Nasal swab	Not detected
L.3.A	Nasal swab	Not detected
L.3.B	Nasal swab	Not detected
L.3.C	Nasal swab	Not detected
L.3.D	Nasal swab	Not detected
L.4.A	Nasal swab	Not detected
L.4.B	Nasal swab	Not detected
L.4.C	Nasal swab	Not detected
L.1.A	Nasal swab	Not detected
L.1.B	Nasal swab	Not detected
P.21.A	Nasal swab	Not detected
R.5.A	Nasal swab	Not detected
R.5.B	Nasal swab	Not detected
R.5.C	Nasal swab	Indeterminate
R.5.D	Nasal swab	Indeterminate
R.5.E	Nasal swab	Not detected
L.6.A	Nasal swab	Not detected
L.6.B	Nasal swab	Not detected
L.6.C	Nasal swab	Not detected
L.6.D	Nasal swab	Not detected
L.6.E	Nasal swab	Not detected
L.6.F	Nasal swab	Not detected
L.6.G	Nasal swab	Not detected
L.6.H	Nasal swab	Not detected
L.13.A	Nasal swab	Not detected
L.13.B	Nasal swab	Not detected
L.13.C	Nasal swab	Not detected
L.13.D	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.03.17

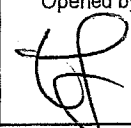
Animal	Specimen	Result
[REDACTED] 13_E	Nasal swab	Not detected
[REDACTED] 13_F	Nasal swab	Not detected
[REDACTED] 2_A	Nasal swab	Not detected
[REDACTED] 2_B	Nasal swab	Not detected
[REDACTED] 2_C	Nasal swab	Not detected
[REDACTED] 2_D	Nasal swab	Not detected
[REDACTED] 2_E	Nasal swab	Not detected
[REDACTED] 2_F	Nasal swab	Not detected
[REDACTED] 2_G	Nasal swab	Not detected
[REDACTED] 2_H	Nasal swab	Detected
[REDACTED] 2_I	Nasal swab	Not detected
[REDACTED] 2_A	Nasal swab	Not detected
[REDACTED] 2_B	Nasal swab	Not detected
[REDACTED] 2_C	Nasal swab	Not detected
[REDACTED] 2_D	Nasal swab	Not detected
[REDACTED] 7_A	Nasal swab	Not detected
[REDACTED] 7_B	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

149 dry swabs

2016-7913
Ref Vet: Highland, Margaret
Owner:
Breed: Domestic Goat
Routed: md

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No Explain below:	Opened by: 
	Samples Received Via: <input type="checkbox"/> US Mail <input type="checkbox"/> FedEx <input checked="" type="checkbox"/> Drop off <input type="checkbox"/> UPS <input type="checkbox"/> FedEx-R <input type="checkbox"/> Other:					


Comments for Case Tracking:



pages: 1 page

06/20/16

Sample Label <input checked="" type="checkbox"/>
--



ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University

Web Site: <http://waddl.vetmed.wsu.edu>

US Postal Service mailing address:
 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
 Bustad Hall, Rm.155-N
 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

Ret Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routed: md

2016 - 10050

08/04/16
 form: 3 pages

Please type or use black ink and print clearly.

Veterinarian or Case Coordinator: Last Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF 3033		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99164	
Phone: 509-335-6327	Fax: 509-335-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☐ Owner ☒ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.

Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted:		Date Collected: July 2016	
(Please use WADDL Animal ID Sheet for multiple animals.)		Date Shipped: n/a	
<div style="text-align: center; font-size: 1.5em; font-weight: bold;">nasal swabs</div>			
Tests Requested: <input type="checkbox"/> Necropsy <input type="checkbox"/> Histopathology <input type="checkbox"/> Toxicology	<input type="checkbox"/> Virology <input type="checkbox"/> Serology <input type="checkbox"/> Fungal Culture	<input type="checkbox"/> Bacteriology <input type="checkbox"/> Mycoplasma culture <input type="checkbox"/> Parasitology	<input type="checkbox"/> IHC <input checked="" type="checkbox"/> PCR <input type="checkbox"/> Other:
Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.			
Animal ID (name/tag#)	Species	Breed	Age
see multiple animal form	domestic goats	multiple	multiple
Location of Lesion	No. in group	No. Dead	No. Sick
N/A		N/A	N/A
Sex			
Animal Weight			
No. on Premises			
Duration of Problem			
N/A			

* Was animal euthanized? If so, what method? N/A

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

M. ovipneumoniae qPCR on each sample

Please save remaining DNA isolations and call Maggie for pick up or may request further testing (sequencing) be performed by WADDL, depending on the results of qPCR analysis.

Please bill to ADRU-ARS-USDA account #RSA 2540-1080

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's or Owner's Signature: *Maggie Highland*

Condition(s) Suspected: *N/A screening study*

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 College of Veterinary Medicine, Washington State University
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2016 - 10050
 Ret Vet: Highland, Margaret
 Owner: USDA - ARS - ADRI
 Breed: Domestic Goat
 Routing: md

08/04/16

Owner: Highland, Maggie

Veterinarian: Highland, Maggie

TEST(S) REQUESTED: Movi qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
1	8_A	26	5_O	51	4_B	76	1_E
2	8_B	27	5_P	52	4_C	77	1_F
3	4_A	28	2_A	53	4_D	78	1_G
4	4_B	29	2_B	54	1_A	79	1_A
5	4_C	30	2_C	55	1_B	80	1_B
6	4_D	31	2_D	56	1_A	81	1_C
7	4_E	32	6_A	57	1_B	82	1_D
8	4_F	33	1_A	58	1_C	83	1_E
9	4_G	34	2_B	59	1_D	84	1_F
10	1_A	35	2_C	60	6_A	85	1_G
11	1_B	36	2_A	61	6_B	86	5_A
12	5_A	37	2_B	62	6_C	87	5_B
13	5_B	38	2_C	63	2_A	88	5_C
14	5_C	39	2_D	64	2_B	89	5_D
15	5_D	40	25_A	65	2_C	90	5_E
16	5_E	41	25_B	66	2_D	91	5_F
17	5_F	42	25_C	67	2_E	92	5_G
18	5_G	43	25_D	68	2_F	93	5_H
19	5_H	44	25_E	69	2_G	94	5_I
20	5_I	45	1_A	70	2_H	95	5_J
21	5_J	46	1_B	71	2_I	96	5_K
22	5_K	47	1_C	72	1_A	97	5_L
23	5_L	48	1_D	73	1_B	98	5_M
24	5_M	49	1_E	74	1_C	99	5_N
25	5_N	50	4_A	75	1_D	100 *	5_O

* For over 100 samples, please copy this form and continue numbering.

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

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College of Veterinary Medicine, Washington State University

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Shipping address:
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Pullman, WA. 99164-7034
FAX: (509) 335-7424

2016-10050
Ref Vet: Highland, Margaret
Owner: USDA - ARS - ADRI
Breed: Domestic Goat
Routing: md

08/04/16

Owner: Highland, Maggie

Veterinarian: Highland, Maggie

TEST(S) REQUESTED: Movi qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
101	5_P	126	1_E	51		76	
102	5_Q	127	7_A	52		77	
103	5_R	128	7_B	53		78	
104	5_S	129	7_C	54		79	
105	5_T	130	7_D	55		80	
106	5_U	131	2_A	56		81	
107	5_V	132	2_B	57		82	
108	5_W	133	2_C	58		83	
109	5_X	34		59		84	
110	5_Y	35		60		85	
111	3_A	36		61		86	
112	3_B	37		62		87	
113	1_A	38		63		88	
114	1_B	39		64		89	
115	1_C	40		65		90	
116	1_D	41		66		91	
117	1_E	42		67		92	
118	1_F	43		68		93	
119	1_G	44		69		94	
120	5_A	45		70		95	
121	5_B	46		71		96	
122	1_A	47		72		97	
123	1_B	48		73		98	
124	1_C	49		74		99	
125	1_D	50		75		100 *	

* For over 100 samples, please copy this form and continue numbering.

Washington Animal Disease Diagnostic Lab

P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424

Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF

Case#: 2016-10050
Report Date: 08/19/16

Pullman, WA 99164-6630

Submittal Date: 08/04/16
Owner: USDA-ARS-ADRU

Species: Domestic Goat

Age:
Sex:

Final Report:

Molecular Diagnostics- Reported on 08/19/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
18.A	Nasal swab	Not detected
18.B	Nasal swab	Not detected
14.A	Nasal swab	Indeterminate
14.B	Nasal swab	Not detected
14.C	Nasal swab	Indeterminate
14.D	Nasal swab	Indeterminate
14.E	Nasal swab	Not detected
14.F	Nasal swab	Indeterminate
14.G	Nasal swab	Not detected
11.A	Nasal swab	Not detected
11.B	Nasal swab	Not detected
15.A	Nasal swab	Not detected
15.B	Nasal swab	Not detected
15.C	Nasal swab	Not detected
15.D	Nasal swab	Not detected
15.E	Nasal swab	Not detected
15.F	Nasal swab	Not detected
15.G	Nasal swab	Not detected
15.H	Nasal swab	Not detected
15.I	Nasal swab	Indeterminate
15.J	Nasal swab	Not detected
15.K	Nasal swab	Not detected
15.L	Nasal swab	Not detected
15.M	Nasal swab	Not detected
15.N	Nasal swab	Indeterminate
15.O	Nasal swab	Not detected
15.P	Nasal swab	Not detected
12.A	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
6.A	Nasal swab	Not detected
1.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
25.A	Nasal swab	Not detected
25.B	Nasal swab	Not detected
25.C	Nasal swab	Not detected
25.D	Nasal swab	Not detected
25.E	Nasal swab	Not detected
1.A	Nasal swab	Not detected
1.B	Nasal swab	Not detected
1.C	Nasal swab	Not detected
1.D	Nasal swab	Not detected
1.E	Nasal swab	Not detected
4.A	Nasal swab	Not detected
4.B	Nasal swab	Not detected
4.C	Nasal swab	Not detected
4.D	Nasal swab	Not detected
11.A	Nasal swab	Not detected
11.B	Nasal swab	Not detected
11.A	Nasal swab	Not detected
11.B	Nasal swab	Not detected
11.C	Nasal swab	Not detected
11.D	Nasal swab	Not detected
6.A	Nasal swab	Not detected
6.B	Nasal swab	Not detected
6.C	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
2.E	Nasal swab	Not detected
2.F	Nasal swab	Not detected
2.G	Nasal swab	Not detected
2.H	Nasal swab	Indeterminate
2.I	Nasal swab	Not detected
Y.1.A	Nasal swab	Not detected
Y.1.B	Nasal swab	Not detected
Y.1.C	Nasal swab	Not detected
Y.1.D	Nasal swab	Not detected
Y.1.E	Nasal swab	Not detected
Y.1.F	Nasal swab	Not detected
Y.1.G	Nasal swab	Not detected
Y.1.A	Nasal swab	Not detected
Y.1.B	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
1.C	Nasal swab	Not detected
1.D	Nasal swab	Not detected
1.E	Nasal swab	Not detected
1.F	Nasal swab	Not detected
1.G	Nasal swab	Not detected
5.A	Nasal swab	Not detected
5.B	Nasal swab	Not detected
5.C	Nasal swab	Not detected
5.D	Nasal swab	Not detected
5.E	Nasal swab	Not detected
5.F	Nasal swab	Not detected
5.G	Nasal swab	Not detected
5.H	Nasal swab	Not detected
5.I	Nasal swab	Not detected
5.J	Nasal swab	Not detected
5.K	Nasal swab	Not detected
5.L	Nasal swab	Not detected
5.M	Nasal swab	Not detected
5.N	Nasal swab	Not detected
5.O	Nasal swab	Not detected
5.P	Nasal swab	Not detected
5.Q	Nasal swab	Not detected
5.R	Nasal swab	Not detected
5.S	Nasal swab	Not detected
5.T	Nasal swab	Not detected
5.U	Nasal swab	Not detected
5.V	Nasal swab	Not detected
5.W	Nasal swab	Not detected
5.X	Nasal swab	Not detected
5.Y	Nasal swab	Not detected
3.A	Nasal swab	Not detected
3.B	Nasal swab	Not detected
1.A	Nasal swab	Not detected
1.B	Nasal swab	Not detected
1.C	Nasal swab	Not detected
1.D	Nasal swab	Not detected
1.E	Nasal swab	Not detected
1.F	Nasal swab	Not detected
1.G	Nasal swab	Not detected
5.A	Nasal swab	Not detected
5.B	Nasal swab	Not detected
1.A	Nasal swab	Not detected
1.B	Nasal swab	Not detected
1.C	Nasal swab	Not detected
1.D	Nasal swab	Not detected
1.E	Nasal swab	Not detected
7.A	Nasal swab	Not detected
7.B	Nasal swab	Not detected
7.C	Nasal swab	Not detected
7.D	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
2-C	Nasal swab	Not detected
1.H	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

133 nasal swabs
- dropped off by
Maggie Highland

2016 - 10050
Ref Vet: Highland, Margaret
Owner: USDA - ARS - ADPU
Breed: Domestic Goat
Routed: md

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms: <input type="checkbox"/> Yes <input type="checkbox"/> No Explain below:	Opened by: MA
	Samples Received Via:					
	<input type="checkbox"/> US Mail	<input type="checkbox"/> FedEx	<input checked="" type="checkbox"/> Drop off			
	<input type="checkbox"/> UPS	<input type="checkbox"/> FedEx-R	<input type="checkbox"/> Other:			

Comments for Case Tracking:



08/04/16
Page 1 of 1

Sample Label ✓

ACCESSION FORM FOR GENERAL DIAGNOSTICS
Washington Animal Disease Diagnostic Laboratory

College of Veterinary Medicine, Washington State University

Web Site: <http://waddl.vetmed.wsu.edu>

US Postal Service mailing address:
 PO Box 647034
 Pullman, WA. 99164-7034

UPS, FedEx or Courier shipping address:
 Bustad Hall, Rm.155-N
 Pullman, WA. 99164-7034

Phone: (509) 335-9696
 FAX: (509) 335 7424
 E-Mail: waddl@vetmed.wsu.edu

2016-12311
 Ref Vet: Highland, Margaret
 Owner: USDA - ARS - ADRU
 Breed: Domestic Goat
 Routed: md

09/21/16
 form 2 pages

Please type or use black ink and print clearly.

Veterinarian or Last Case Coordinator Name: Highland		First Name: Maggie	
Clinic: ADRU-ARS-USDA			
Street address: ADBF 3033		Mailing Address or PO Box:	
City: Pullman	State: WA	Zip: 99164	
Phone: 509-335-6327	Fax: 509-335-8328	E-mail: mah@vetmed.wsu.edu	
Owner: Last Name first: same as above		Guardian Name: (if owner is under 18)	
Farm Name:		First Time Submitter? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Street address:		Mailing Address or PO Box:	
City:	State:	Zip:	
Phone:	Fax:	E-mail:	

Billing: ☐ Owner ☒ Clinic ☐ 3rd Party (preapproval required) Please note: WADDL policy is to bill the clinic if provided, unless prepaid.
Reporting Preference: ☐ Mail ☐ Fax ☒ Web access - register on web site at <http://waddl.vetmed.wsu.edu>

Please fill out completely as possible:

Specimen(s) Submitted: (Please use WADDL Animal ID Sheet for multiple animals.)		nasal swabs-frozen (-20C)		Date Collected: Aug-Sept 2016	
				Date Shipped: n/a	
Tests Requested: <input type="checkbox"/> Necropsy <input type="checkbox"/> Virology <input type="checkbox"/> Bacteriology <input type="checkbox"/> IHC <input type="checkbox"/> Histopathology <input type="checkbox"/> Serology <input type="checkbox"/> Mycoplasma culture <input checked="" type="checkbox"/> PCR <input type="checkbox"/> Toxicology <input type="checkbox"/> Fungal Culture <input type="checkbox"/> Parasitology <input type="checkbox"/> Other:		Note: WADDL reserves the right to modify the tests requested for more efficient case work-up and / or to send specimens to outside laboratories to perform testing not done at WADDL.			
Animal ID (name/tag#)	Species	Breed	Age	Sex	Animal Weight
see multiple animal form	domestic goats	multiple	multiple		
Location of Lesion	No. in group	No. Dead	No. Sick	No. on Premises	Duration of Problem
N/A		N/A	N/A		N/A

* Was animal euthanized? If so, what method? N/A

Additional History: Vaccinations, signs, stress factors, treatments, post mortem findings, pertinent feed or feed additives, clinical lab results, previous WADDL Case Numbers. (Attach additional sheets as necessary.)

M. ovipneumoniae qPCR on each sample

Please save remaining DNA isolations and call Maggie for pick up or may request further testing (sequencing) be performed by WADDL, depending on the results of qPCR analysis.

Please bill to ADRU-ARS-USDA account #RSA 2540-1094

WADDL is an official brucellosis testing laboratory. All serology for brucellosis, including abortion screens, requires identification of animals, date of sample collection, and signature of an accredited veterinarian attesting to the following statement:

"I certify that the specimens submitted with this form were collected by me from the animal(s) described on the date indicated."

Veterinarian's, Clinician's or Owner's Signature:

Condition(s) Suspected:

IDENTIFICATION SHEET FOR MULTIPLE ANIMALS

(To accompany WADDL Accession form, if needed)

Washington Animal Disease Diagnostic Laboratory
 College of Veterinary Medicine, Washington State University
 Mailing address: Shipping address:
 P.O. Box 647034 Bustad Hall, Rm. 155-N
 Pullman, WA. 99164-7034 Pullman, WA. 99164-7034
 Phone: (509) 335-9696 FAX: (509) 335-7424
 E-Mail: waddl@vetmed.wsu.edu
 Web Site: http://waddl.vetmed.wsu.edu

2016-12311

09/21/16

Ref Vet: Highland, Margaret
 Owner: USDA-ARS-ADRU
 Breed: Domestic Goat
 Routing: md

Owner: Highland, Maggie

Veterinarian: Highland, Maggie

TEST(S) REQUESTED: Movi qPCR

Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name	Tube	Animal # or Name
1	2_A	26	2_F	51	4_B (3)	76	1_J (2)
2	2_B	27	2_G	52	4_C (3)	77	1_L (2)
3	2_C	28	2_H	53	4_D (3)	78	1_N (2)
4	2_D	29	2_I	54	4_E (3)	79	1_O (2)
5	9_A	30	2_J	55	4_F (3)	80	1_P (2)
6	9_B	31	3_A	56	4_G (3)	81	1_Q (2)
7	9_C	32	3_B	57	4_H (3)	82	1_R (2)
8	9_D	33	3_C	58	4_I (3)	83	1_S (2)
9	9_E	34	4_A	59	4_J (3)	84	1_T (2)
10	5_A	35	4_B	60	4_K (3)	85	1_U (2)
11	5_B	36	4_C	61	4_L (3)	86	1_V (2)
12	5_C	37	4_D	62	4_M (3)	87	1_W (2)
13	5_D	38	4_E	63	4_N (3)	88	1_X (2)
14	5_E	39	3_A	64	4_O (3)	89	1_Y (2)
15	5_F	40	3_B	65	4_S (3)	90	1_HH (2)
16	5_G	41	3_C	66	9_F (2)	91	1_II (2)
17	5_H	42	3_D	67	9_G (2)	92	1_KK (2)
18	5_I	43	3_E	68	17_J (2)	93	1_LL (2)
19	5_J	44	3_F	69	17_K (2)	94	1_MM (2)
20	5_K	45	3_G	70	1_A (2)	95	1_NN (2)
21	2_A	46	3_H	71	1_B (2)	96	1_SS (2)
22	2_B	47	A_3_I	72	1_D (2)	97	1_ZZ (2)
23	2_C	48	26_A	73	1_E (2)	98	1_BC (2)
24	2_D	49	26_B	74	1_F (2)	99	2_H (4)
25	2_E	50	4_A (3)	75	1_G (2)	100 *	

* For over 100 samples, please copy this form and continue numbering.

Washington Animal Disease Diagnostic Lab

P.O. Box 647034
Pullman, WA 99164-7034
Telephone : (509) 335-9696
Fax : (509) 335-7424

Dr. Margaret Highland
USDA-ARS-ADRU
WSU - 3003 ADBF

Case#: 2016-12311
Report Date: 10/05/16

Pullman, WA 99164-6630

Submittal Date: 09/21/16
Owner: USDA-ARS-ADRU

Species: Domestic Goat

Age:
Sex:

Final Report:

Molecular Diagnostics- Reported on 10/05/16 Authorized by Daniel Bradway, Lab Manager

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
9.A	Nasal swab	Not detected
9.B	Nasal swab	Not detected
9.C	Nasal swab	Not detected
9.D	Nasal swab	Not detected
9.E	Nasal swab	Not detected
5.A	Nasal swab	Not detected
5.B	Nasal swab	Not detected
5.C	Nasal swab	Not detected
5.D	Nasal swab	Not detected
5.E	Nasal swab	Not detected
5.F	Nasal swab	Not detected
5.G	Nasal swab	Not detected
5.H	Nasal swab	Not detected
5.I	Nasal swab	Not detected
5.J	Nasal swab	Not detected
5.K	Nasal swab	Not detected
2.A	Nasal swab	Not detected
2.B	Nasal swab	Not detected
2.C	Nasal swab	Not detected
2.D	Nasal swab	Not detected
2.E	Nasal swab	Not detected
2.F	Nasal swab	Not detected
2.G	Nasal swab	Not detected
2.H	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
2.I	Nasal swab	Not detected
2.J	Nasal swab	Not detected
3.A	Nasal swab	Not detected
3.B	Nasal swab	Not detected
3.C	Nasal swab	Not detected
4.A	Nasal swab	Not detected
4.B	Nasal swab	Not detected
4.C	Nasal swab	Not detected
4.D	Nasal swab	Not detected
4.E	Nasal swab	Not detected
3.A	Nasal swab	Not detected
3.B	Nasal swab	Not detected
3.C	Nasal swab	Indeterminate
3.D	Nasal swab	Indeterminate
3.E	Nasal swab	Not detected
3.F	Nasal swab	Indeterminate
3.G	Nasal swab	Not detected
3.H	Nasal swab	Not detected
3.I	Nasal swab	Not detected
26.A	Nasal swab	Not detected
26.B	Nasal swab	Not detected
4.A (3)	Nasal swab	Not detected
4.B (3)	Nasal swab	Not detected
4.C (3)	Nasal swab	Indeterminate
4.D (3)	Nasal swab	Indeterminate
4.E (3)	Nasal swab	Not detected
4.F (3)	Nasal swab	Not detected
4.G (3)	Nasal swab	Not detected
4.H (3)	Nasal swab	Not detected
4.I (3)	Nasal swab	Not detected
4.J (3)	Nasal swab	Indeterminate
4.K (3)	Nasal swab	Not detected
4.L (3)	Nasal swab	Indeterminate
4.M (3)	Nasal swab	Indeterminate
4.N (3)	Nasal swab	Not detected
4.O (3)	Nasal swab	Not detected
4.S (3)	Nasal swab	Indeterminate
9.F (2)	Nasal swab	Not detected
9.G (2)	Nasal swab	Not detected
17.J (2)	Nasal swab	Indeterminate
17.K (2)	Nasal swab	Not detected
1.A (2)	Nasal swab	Not detected
1.B (2)	Nasal swab	Not detected
1.D (2)	Nasal swab	Not detected
1.E (2)	Nasal swab	Not detected
1.F (2)	Nasal swab	Not detected
1.G (2)	Nasal swab	Not detected
1.J (2)	Nasal swab	Not detected
1.L (2)	Nasal swab	Not detected
1.N (2)	Nasal swab	Not detected
1.O (2)	Nasal swab	Not detected
1.P (2)	Nasal swab	Not detected

Washington Animal Disease Diagnostic Lab

PCR-Mycoplasma ovipneumoniae SOP: 501.40RT.2016.07.18

Animal	Specimen	Result
1-Q (2)	Nasal swab	Not detected
1-R (2)	Nasal swab	Not detected
1-S (2)	Nasal swab	Not detected
1-T (2)	Nasal swab	Not detected
1-U (2)	Nasal swab	Not detected
1-V (2)	Nasal swab	Indeterminate
1-W (2)	Nasal swab	Not detected
1-X (2)	Nasal swab	Not detected
1-Y (2)	Nasal swab	Not detected
1-HH (2)	Nasal swab	Indeterminate
1-II (2)	Nasal swab	Not detected
1-KK (2)	Nasal swab	Not detected
1-LL (2)	Nasal swab	Not detected
1-MM (2)	Nasal swab	Indeterminate
1-NN (2)	Nasal swab	Not detected
1-SS (2)	Nasal swab	Not detected
1-ZZ (2)	Nasal swab	Not detected
1-BC (2)	Nasal swab	Not detected
2-H (4)	Nasal swab	Not detected

PCR-Mycoplasma ovipneumoniae test comment: This assay detects only Mycoplasma ovipneumoniae. Culture is available at WADDL to detect other species of Mycoplasma if desired. Fees for culture are available on our website. Please contact the lab if Mycoplasma culture or other testing is desired.

Quantity/Description/Routing of Samples

99 nasal swabs → MD
per MAH

by 2 M. Highland
→ 250B

Sample Condition:	<input type="checkbox"/> Room Temp.	<input type="checkbox"/> On ice	<input checked="" type="checkbox"/> Frozen	<input type="checkbox"/> Fixed	Contents match forms: <input type="checkbox"/> Yes <input type="checkbox"/> No Explain below:	Opened by: <i>not</i>
	Samples Received Via:					
	<input type="checkbox"/> US Mail	<input type="checkbox"/> FedEx	<input checked="" type="checkbox"/> Drop off			
	<input type="checkbox"/> UPS	<input type="checkbox"/> FedEx-R	<input type="checkbox"/> Other:			

Comments for Case Tracking:

unk
↓
MD to verify

2016-12311
Ref Vet: Highland, Margaret
Owner: USDA - ARS - ADRI
Breed: Domestic Goat
Routed: md



09/21/16
notes: 1 page

Sample Label