Guidelines for Veterinary Thermography

General Statement

This guideline was prepared by members of the American Academy Of Thermology (AAT) as a guide to aid the performance of medical infrared imaging in evaluating veterinary patients. It implies a consensus of those substantially concerned with its scope and provisions. The AAT guideline may be revised or withdrawn at any time. The procedures of the AAT require that action be taken to reaffirm, revise or withdraw this guideline no later than three years from the date of publication. Suggestions for improvement of this guideline are welcome and should be sent to the executive director of the American Academy of Thermology. No part of this guideline may be reproduced in any form, in an electronic retrieval system or otherwise, without the prior written permission of the publisher.

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Statement of Need

Heat is a cardinal sign of inflammation. Vasomotor tone and vasomotor capacitance plays a significant role in thermoregulation, clinical symptomatology and manifestations of systemic illness. In the Animal Kingdom the clinical manifestation of pain can be detrimental to survival. As such, changes in the previously mentioned vascular parameters may be the only clinical signs shown.

Infrared Thermal Imaging is the only non-invasive technology available to image and map circulatory changes associated with these disorders. It can play an important role in clinical diagnosis and enhance the clinical examination. But infrared imaging may also be valuable to document musculoskeletal stress caused by training as well as circulatory effects of therapeutic modalities. In addition, infrared imaging can be used as an aid in the regulation of the animal industry.

Other technologies such as Radiography, Ultrasonography, Scintigraphy, and MRI do not provide the physiologic information offered by Medical Thermal imaging. The clinical application of Infrared Thermal may be instrumental in understanding the pathophysiology associated with veterinary disease and can lead to improved patient outcomes.

The mission and bylaws of the American Academy of Thermology support the incorporation of thermal imaging into veterinary clinical medicine. The AAT recognizes a current and ongoing need to promulgate continuing education in the science and methods of thermal imaging and the clinical application of heat asymmetry patterns obtained from thermal imaging among both veterinarians and thermal technologists.

Purpose

Veterinary medicine is a unique branch of the health sciences that involves multiple species many of which are not domesticated. Regardless, any can be dangerous to the examiner. Furthermore, veterinary patients do not seek medical attention, rather the owner, rider, trainer, or caretaker seeks medical advice based on their observations. Thus, veterinary examinations must be very thorough with attention to basic clinical signs that provide insight to potential inflammatory conditions. Unfortunately, many veterinary patients can only be examined using anesthesia or sedation. The ability of infrared imaging to detect changes in the heat patterns of skin makes it an invaluable tool in the clinical assessment of veterinary patients with certain types of problems. The thermal examination can be performed from the cranium to the base of the spine, from the torso to the extremities, including the digits and may include the oral and abdominal cavities.
Common Indications

There are 4 common uses for infrared imaging in veterinary medicine (1-35)

1. As a diagnostic aid where changes in the thermal patterns of scanned areas suggest a regional or global diagnosis whereby anatomic imaging modalities can then be used to characterize the nature of the problem.

2. As a method to enhance clinical assessment of the veterinary patient to include:
   - Evaluation or follow-up of patients with known or suspected vasomotor instability.
   - Pre-procedure assessment to aid in the planning of interventional therapeutics or diagnostics
   - Follow-up to detect improvement, progression or spread of disease, which may reflect change in condition.
   - Evaluation of muscle and peripheral circulation
   - Evaluation of post-surgical swelling and wound healing
   - Evaluation of unexpected post-operative or post fracture pain.
   - Qualitative and quantitative assessment of the vasculature and blood flow to tissues
   - Evaluation of peripheral neuropathies.
   - Evaluation of inflammation associated with dental or periodontal disease
   - To determine if areas of palpable soreness are associated with changes in temperature
   - Monitoring of testicular temperature
   - Monitoring of extremities post cast application
   - Monitoring progression and appropriateness of exercise protocols through the post-injury rehabilitation process.
3. As a method to assess thermal, functional, ergonomic, musculoskeletal stress in animals during and after training, competition, daily routines, and travel & performance, including but not limited to:
   - Identification of the presence of subclinical inflammation to allow for appropriate intervention.
   - Assessment of foot, hoof and shoe balance.
   - Assessment of foot and hoof prints to evaluate corresponding pressure changes that the animal is applying versus other limbs.
   - Assessment of temperature changes associated with wounds or surgery.
   - Assessment of saddle & harness fit in horses.
   - Assessment of the effects of the rider on the horse’s back.
   - Evaluation of the ergonomic fitness of harnesses, thermal control/equipment vests and foot protection booties in performance and working animals.
   - Identification of the onset of hyperthermia.
   - Structural evaluation of thermal and functional conditions of animal facilities.

4. For regulatory medicine to determine welfare compliance in jumping horses and Tennessee Walking Horses (TWH) including but not limited to:
   - Assessment of limb temperature changes for limb sensitivity in show jumpers
   - Identification of abnormal cold on the distal limb of TWHs suggestive of foreign substance application
   - Assessment of thermal patterns on the palmar pastern of TWHs suggestive of abnormal scar formation
   - Assessment of thermal patterns of TWHs distal limbs consistent with application of “soring” agents
Contraindications and Limitations

Infrared thermal imaging shows only skin temperature and therefore is a reflection of the deeper tissues.

Infrared imaging is highly sensitive to environmental factors and an understanding of these factors is imperative if infrared imaging is to be successful.

- Infrared imaging is contraindicated if the thermal evaluator does not understand infrared radiation physics
- Imaging is contraindicated if bilaterally symmetrical images cannot be evaluated.
- Imaging is contraindicated if environmental factors cannot be controlled, including sunshine, ambient temperature, drafts, haircoat, topical moisture, topical liniments, and the presence of bandages or blankets or articles of harness.
- Imaging is contraindicated if the patient is uncooperative.

Guideline 1: Owner (Person Responsible) Communication and Veterinary Patient Preparation:

Communication- The examiner:

1.1 Explains the medical necessity for performing infrared imaging to the Person Responsible (PR) for the Patient.

1.2 Responds to questions and concerns from the PR about any aspect of the examination.

1.3 Advises the PR about risk factors and the information hoped to be obtained by infrared imaging. Obtains informed consent either written or orally from the PR to proceed with infrared imaging.

1.4 Refers specific diagnostic, treatment or prognosis questions to the Veterinarian.

Preparation:

1.5 The Patient should be as clean as possible and groomed to remove any debris from the body surface. Patient should not have contact with any object if that body part is being imaged. All leg wraps, bandages, blankets, or any other object having contact with the area of anatomy to be examined should be removed for a minimum of 20 minutes prior to the commencement of the Thermal Exam. Animals should not be touched or allowed to sit or lay down.

1.6 Avoid placing any material of any kind on the patient’s skin, such as any skin lotions, liniments, fly spray, topical external parasite treatments, etc. The haircoat must be dry.
1.7 The PR should declare the presence of any vasoactive topical or systemic substances to the examiner. Whenever possible steroids, sympathetic agonists and antagonists, vasoactive medications, opiates and transdermal patches should be avoided for 24 hours prior to testing. Exceptions should always be recorded in the record.

1.8 Avoid any therapy or exposure that is applied to the skin, goes through the skin or may affect skin for 24 hours before imaging. The same is true for any electrodiagnostic technique that may affect skin. It should be noted within the record whenever exceptions are made and why.

1.9 In the absence of extenuating circumstances, diagnostic studies using neurolytic blocks should be avoided for 3 days prior to testing.

1.10 Controlled exercise such as riding, lunging or treadmill for horses, play, running or treadmill for dogs are useful dynamic examinations part of the thermographic evaluation but must be performed in a controlled manner. Intense exercise should be avoided in all animals for at least 2 hours prior to a resting examination.

**Guideline 2: Patient Assessment**

Patient assessment should be performed before infrared imaging.

2.1 Obtain a complete, pertinent history by interview of the PR and/or review of the patient’s medical record. A pertinent history includes:
   a. Current medical status, especially regarding pain, lameness, and possible vascular issues.
   b. Presence of any signs or symptoms of vasomotor or other autonomic dysfunction.
   c. Relevant risk factors for injury, age, use, breed, and weight
   d. Symptoms of odontalgia
   e. Recent surgery
   f. Pathology/Laboratory investigation values.
   g. Current medication or therapies
   h. Results of other diagnostic Imaging modalities, radiographic, sonographic, scintigraphic, thermographic, computed tomography or magnetic resonance studies
   i. Results of prior treatments

2.2 Veterinarians should complete a limited, focused, detailed or extensive physical examination, which includes assessment of all structures under study. The extent of the examination may be dependent on the tractability of the veterinary patient. Veterinary assistants should follow policy and procedure of the interpreting or attending veterinarian.

2.3 Compliance with Federal, State or other applicable regulatory rules and regulations should be maintained.
Guideline 3: Examination Guidelines

3.1 Infrared imaging measures and maps the degree and distribution of IR thermal emission. Skin temperature is largely under the control of the autonomic nervous system and inflammatory processes. Sagittal symmetry is expected throughout the body. Asymmetric IR emission of 1°C or greater can be indicative of sympathetic nervous system (SNS) dysfunction or other pathology.

Infrared evaluations do not test structure, but rather correlates to sympathetic nervous system physiology as well as local or systemic inflammation. Therefore, when structural injury is suspected additional radiographic imaging or diagnostic studies may still need to be performed.

Due to the complex nature and etiology of painful conditions associated with skin temperature asymmetry patterns, only those veterinarians trained in the proper techniques required to perform and interpret infrared imaging should do so. When present, the pattern of asymmetry discovered by infrared examination should guide the treating veterinarian in determining the source or generator of the abnormality. Both response to treatment and additional examination or testing may still be required to complete this task.

3.2 The following minimum specifications should guide the selection and use of infrared hardware and software systems. These specifications apply to modern infrared imagers. They are not intended to reflect on systems used in the past. While recognizing that individual circumstances will vary, for the purposes of this document, the typical imager Field of View (FOV) is 25 degrees, patient to imager distance 3-8 feet (as needed to allow the region of interest to fill approximately 75% of the image) and optical quality is satisfactory to the vast majority of observers.

Emissivity is a fractional representation of the amount of energy radiated from a material versus the energy that would come from a true black body at the same temperature. Passive IR imaging (thermology) measures and maps the pattern of skin thermal radiance (the degree and distribution of skin temperature changes). If needed for the examination being performed, medical grade imagers should be calibrated against two black bodies having emissivity of 0.98, and spanning the physiologic temperature range.

Imager emissivity set to 0.98 (human skin). The emissivity is a fractional representation of the amount of energy radiated from a material versus the energy that would come from a black body at the same temperature.

Image detector spectral bandwidth: typically, 8 to 14 microns (micrometers).

- Preferred absolute detector resolution of ≥ 640 X 480 coupled with a suitable microbolometer and lens. Today, most medical imaging systems utilize uncooled focal plane array detectors found in the 320 X 240 sensor range or higher. When systems with 320 X 240 sensor arrays are coupled with a high-quality microbolometer, lens, and compensatory software
or firmware innovations, they can approach the image quality, spatial resolution, and spot measurement requirements found in 640 X 480 systems.

- Min. measurable spot size is 2.1x2.0 mm (3x3 or 9 pixels) at 40 cm distance.
- Spatial resolution quality at 8 feet (2.4 meters) equivalent to ≤ 2.6 mRad IFOV (Instantaneous Field of View) at 40 cm minimum focus.
- Thermal sensitivity of < 50 mK NETD (Noise Equivalent Temperature Difference) @ 30 °C.
- Ability to perform accurate quantitative differential temperature analysis with a precision of ≤ ± 0.05 °C (50mK).
- Repeatability and precision of ≤ ±0.05 °C (50mK) detection of temperature difference.
- Thermal drift (caused by internal heating of equipment during normal operation or by changes in external ambient temperature) to be strictly controlled by calibration to a known temperature standard if necessary for the study under consideration.
- Maintenance of detector uniformity and correction via calibration to a known temperature standard.
- Ability to render images in hi-resolution color and grayscale.
- High-resolution image visual display for interpretation.
- If video mode is used, it may incorporate real-time image focus and capture capability. While 10Hz, 20Hz, and 30Hz frame rates are capable of real-time imaging, having faster capability is preferred (i.e., 50Hz). For temperature analysis, radiometric video files are preferred.
- Precision Autofocus is recommended.
- Imager temperature range set to cover temperatures within the range of emissions (20-45 °C).
- Ability to archive images for future reference and image comparison at same patient positioning and distance from the imager.
- Software manipulation of the images should be maintained within strict parameters to ensure that the original qualities of the images are not compromised.
- Imaging software capable of identifying areas of temperature calculations and locations for reporting
- Appropriate infrared instrumentation, including real-time display, electronic static image capture, storage, post-capture annotation, or hard copy documentation capabilities, should be utilized.

Contact thermology devices that utilize single or multiple probes or sheets of thermally-sensitive liquid crystals for breast thermographic analysis are considered obsolete considering the current advances in non-contact digital infrared imaging. Thermographic scanning systems that cannot
acquire and display thermal differences of 0.05°C are also to be considered obsolete for medical purposes.

3.3 All studies should be performed in an environment in which, to the extent possible, ambient temperature is controlled, free from drafts and where there is no exposure to infrared rays, such as airflow, incandescent lights or sunlight that may result in heating or reflective artifacts. The imaging room should be comfortably cool to allow for dissipation of superficial heat which may produce artifact from the skin. Ideally the ambient temperature range for IR imaging suite should be between 20° to 25°. However, as long as it is below 30º C and sweating is not induced, thermal imaging may be performed.

3.4 Ventilation systems should be designed to avoid direct airflow onto the patient. The patient should not be placed near or under any light fixture that itself emits heat. Standard fluorescent and/or LED lights are appropriate. Ideally the floor should be made of a non-reflective material.

3.5 When possible, an equilibration time of fifteen – sixty minutes is deemed appropriate prior to obtaining the images. The use of a blanket and humidity can affect equilibration time. One way of knowing when the "blanket effect" is no longer in play is to assess whether pressure point areas such as the top of the withers, top of the shoulders, over the gluteals, and tailhead have all adjusted toward normal.

While infrared studies can usually be accomplished with one set of images, post exercise examination or saddle test examinations require more than one set of images. Images taken in these serial studies should have accompanying time stamps for clarity.

3.6 A standard exam protocol for each segment evaluated should be used. Each point of focus should include dorsal, palmar/plantar, medial, and/or lateral views, as required. Contralateral and dorsal views should be equidistant and fill seventy-five (75) percent of the image screen when possible. When possible, it is recommended that the contralateral extremity images should be captured in the same image as the extremity under examination. Obtaining additional images may be required for patients with specific, unique circumstances.

3.7 Neuro-musculoskeletal (NMSK) studies that wish to highlight vasomotor mapping and that are prepared for interpretation without post-acquisition radiometric image manipulation typically employ palettes of no less than ten colors and are formatted at 1°C per color. Many have found it beneficial to use a temperature span of more than 10°C (the 1°C per color format, however, is retained). The intent of using a broader temperature span is to ensure that no relevant radiometric image information is lost at the extremes of temperature maxima or minima across a wide range of regions of interest.

While vasomotor mapping may be more readily visible with a 1°C per color palette, gradient palettes that span less than 10°C are also commonly employed in NMSK studies that have a limited focus or region of interest and where visualization of vasomotor maps may not be necessary for that study’s intended use. Nonetheless it is the interpreting thermologist’s
responsibility to make sure that the absence of a vasomotor map is not clinically relevant for each study performed where the same is omitted.

The interpreting thermologist must also ensure that the range of temperature maxima and minima utilized does not exclude pertinent radiometric data. Individual protocols for acquisition that do not exclude relevant findings should be established. If a small temperature span is employed the interpreting physician should explain why it was required.

3.8 The patient’s physical and demeanor is assessed and monitored during the examination, with modifications made to image acquisition as necessary. Also, findings are analyzed throughout the course of the examination so that the veterinarian is provided sufficient data to direct patient management and render a thermographic impression.

3.9 Appropriate infrared instrumentation, which includes real time display, electronic static image capture, storage, post capture annotation, or hard copy documentation should be utilized. It is further recommended that static image files be available for viewing and storage, either as radiometric or JPEG images stored in a unique folder for each animal patient, or in Digital Imaging and Communications in Medicine (DICOM) format for Picture Archiving and Communication System (PACS) distribution.

3.10 Evaluate the patient’s physical and mental status prior to discharge. Additional discharge instructions to the PR may include recommendation to schedule follow up appointment with the attending veterinarian, and to resume all medication medical treatment that may have been discontinued prior to the infrared study.

Guideline 4: Review of the Infrared Thermography Examination (example templates are available for Members in the Knowledge Center within the AAT Member Portal)

4.1 The data acquired during the extremity and spinal infrared examination should be reviewed to ensure that a complete and comprehensive evaluation has been performed and documented. Any exceptions to the routine examination protocol (i.e., study omissions or revisions) should be noted and reasons given.

4.2 The examiner should record all technical findings required to complete the final interpretation so that the measurements can be classified according to the standard laboratory diagnostic criteria.

4.3 The examiner should complete the required documentation of the study in a timely manner.

4.4 The examiner should alert the medical director or other responsible veterinarian when immediate medical attention is indicated, based on the infrared examination findings.
**Guideline 5: Presentation of Exam Findings**

5.1 The examiner must provide preliminary results as provided for by internal policy based on examination findings.

5.2 The examiner must present the record of diagnostic images and when applicable, explanations for sub-optimal examination findings to the interpreting veterinarian for use in diagnosis and archival purposes.

**Guideline 6: Preparation and Storage of Exam Findings**

6.1 Images should be presented to the interpreting physician for use in analysis and archival purposes. Radiometric images in either radiometric image format or non-radiometric formats such as lossless JPEG, PNG, or DICOM are acceptable. A color-to-temperature Thermal Scale must accompany each image.

6.2 The imaging clinic should adhere to all established federal and state regulations. Archiving of image data and the analysis/report are to be maintained for no less than seven years.

**Guideline 7: Exam Time Recommendations**

High quality and accurate results are fundamental elements of the infrared study. A combination of direct and indirect exam components is the foundation for maximizing exam quality and accuracy.

7.1 Indirect exam components include pre-exam procedures:
   a) obtaining previous exam data, completing pre-exam paperwork,
   b) exam room and equipment preparation and
   c) patient assessment, history, and positioning.

7.2 Post exam procedures include:
   a) clean up consisting of compiling, processing, and reviewing data for preliminary and/or formal interpretation,
   b) Communication with the Person Responsible (PR)
   c) examination charge and billing activities where appropriate.

Recommended time: 30-60 minutes.

7.3 Direct exam components include:
   a) equipment optimization,
   b) patient positioning throughout the exam,
   c) the actual hands-on examination process.
Recommended time: 20-60 minutes.

**Guideline 8: Reporting**

8.1 A medical record report should be prepared within 24 hours of the study. As part of the imaging protocol, the thermographer should consider sending each PR a summary report within 30 days of the thermographic examination.

8.2 Report layout: The body of the Infrared Veterinary Thermographic report should clearly state that procedures that follow a peer reviewed, internationally accepted guideline was utilized. The set of images obtained for study should be documented. If a standard protocol for reading images is used then this should be stated as well.

Thermographic Findings should be documented and any abnormalities noted. Findings include asymmetry of ≥ 1 degree Centigrade in ≥ 25% of the surface area of any individual region of interest and localized hot or cold spots.

Thermographic impressions include classification according to an accepted naming system or summarization of the thermographic findings. When recognized patterns (thermal signatures) are seen, the thermographic impressions may include the description of that pattern (for example: a back thermographic pattern consistent with overriding vertebral spinous processes) however care should be taken not to make any statements about clinical diagnosis in this section of the report. That is these patterns have been associated with particular disease but the pattern does not prove the disease.

Clinical Impressions are not to be included in the Thermographic Findings paragraph but rather in a separately identifiable paragraph that speaks to the generator or etiology of those findings. Any discussion that is clinically relevant should be reserved for this paragraph.

**Guideline 9: Continuing Professional Education**

Certification is considered the standard of practice for infrared technology. It indicates an individual’s competence to perform medical technology at the entry level. After achieving certification, all Registered Infrared SSR Technologists are expected to keep current with:

9.1 Advances in diagnosis and treatment of pain syndromes with and without sympathetic nervous system dysfunction (vasomotor instability).

9.2 Changes in infrared examination protocols or published laboratory diagnostic criteria.

9.3 Advances in infrared technology used for veterinary examinations.

9.4 Advances in other technology used for infrared examinations.
Guideline 10: Emerging Technologies

10.1 Technology is constantly being introduced that can challenge existing guidelines or that do not necessarily conform to currently accepted practices. These technologies can span the entire spectrum of sophistication and therefore require different adaptive responses. On one end of the spectrum there are innovations based upon generally accepted medical scientific methodology that have gained regulatory acceptance and on the other end there are technologies that are intended for personal use only or that have applications in non-medical fields but have not been accepted as suitable for medical practice.

10.2 General industrial or personal use thermal imaging cameras that do not meet the specification guidelines contained herein are not intended for use in Medical Thermology. Add-on” thermal imagers that plug into a cellular phone are, at present, not adequate for medical thermology imaging.

10.3 Technologies not otherwise covered in these Guidelines that employ methodologies, hardware, or protocols that have gained Federal Regulatory listing for Medical Thermology may become available over time. In cases where these technologies are employed the body of the report should document which device(s) were used and why. Other components of the Guideline should still be followed.

References

10. Turner TA: Thermography as an aid to the clinical lameness evaluation, Vet Clin of

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