Differential Diagnosis in Small Animal Medicine

By

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To Naomi and Abigail, for their patience and support, and Mac, for a lifetime of companionship.

Introduction

This book was written to fill what I felt was a gap in the market. While working up difficult medical cases, I have often wanted a single ready reference to help me formulate a differential list from the clinical information I have available. Unfortunately, I found myself frequently having to consult multiple textbooks to bring all the information I needed together. I decided therefore to write a book that would serve as a ready reference for differential diagnoses of the majority of presentations that are encountered in practice, including both common and uncommon conditions. This text should be of use to veterinary students, general practitioners, university interns, residents and anyone who, like me, cannot fully carry these lists around in their heads. I hope other clinicians find it as useful as I do.

The differential diagnosis list is one of the most important aspects of the problemoriented approach to clinical diagnosis. For those who are not familiar with the problemoriented approach, a brief outline follows.

As the name implies, problem-oriented medical management (POMM), concentrates on the individual problems of a patient. A differential diagnosis list should be made for each and every problem that is found in a patient, whether in the history, the physical examination, imaging or clinicopathological tests. Although superficially this may not sound very 'holistic', in fact, if all the patient's problems are considered individually, the whole patient will have been evaluated, without falling into the trap of presuming that all of the findings are caused by a single condition.

The problem-oriented approach starts with a detailed history, and it is important to discover what the owners perceive to be the main problems – after all, they usually know their animal better than the clinician does. However, there may be relevant historical signs that the owners had not considered significant, so failing to systematically ask all the questions which could be of importance in a case can lead to overlooking important information. Using a checklist or form, such as the one in Appendix A can be useful as an aide-memoire.

In every case, a detailed physical examination should be carried out, including body systems that are not apparently of immediate concern. Again, a checklist or form, such as the one in Appendix B, can help ensure a systematic approach.

Once the history has been taken and the physical examination has been completed, the clinician should list every problem that has been discovered. Problems may include such findings as exercise intolerance, pruritus, pyrexia or a heart murmur. A differential diagnosis list should then be created for every problem. The list should be appropriate to that animal. There is no point listing feline leukaemia virus as a likely diagnosis in a dog!

An attempt should also be made to categorise the conditions in order of likelihood, or at least into common and uncommon. Although the more common conditions have been indicated in this book with an asterisk (*), there are few objective data regarding the true incidence of conditions, and the estimate of incidence is largely subjective and influenced by the author's geographical location and caseload. Familiarity with how

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common conditions are and their local incidence will help prioritise differential lists. The clinician can then select diagnostic tests in a rough order of probability, although rarer but life-threatening conditions, such as hypoadrenocorticism, should also be ruled out early in the course of investigations. Some authorities rightly point out that emphasis should be placed on historical and physical signs, and that 'over-investigating' can be expensive and potentially detrimental to the patient (Chesney, 2003).

It is this author's opinion, however, that it is possible to place too much importance on probabilities and how commonly a condition occurs. The newly-qualified veterinary surgeon will often look for the rare but exciting and memorable condition they learned about at college, while the experienced practitioner will often remind them that 'common things are common', and suggest they restrict their investigations only to commonly-encountered conditions. The ideal approach is probably somewhere in between.

Although it is self-evidently true that common things are common, it is also true that uncommon things are encountered relatively often. To take a hypothetical example: if a common problem is caused by common conditions A and B with a frequency of 80%, and by rare conditions C to Z the rest of the time, with conditions C to Z occurring with equal frequency, then each individual condition C to Z will be responsible for the problem approximately 0.9% of the time, making each individual condition quite uncommon. However, 1 in 5 presentations of this problem will be caused by an uncommon condition, and so uncommon conditions will be diagnosed commonly, provided they are looked for. The problem-oriented approach ensures that these uncommon conditions are not overlooked.

Some authorities prefer to categorise the initial approach to a case differently, and describe the subjective and objective assessment of a patient as part of the SOAP approach (Subjective, Objective, Assessment, Plan). The principle is the same however, in that a detailed history or physical examination is the basis of the initial differential list.

Once the differential diagnosis list has been formulated, the clinician is in a position to select appropriate tests to aid in making a definitive diagnosis. Prioritising the selection of diagnostic tests helps avoid placing undue financial strain on the client and inappropriate or unnecessary testing on the patient. Tests may be prioritised on such factors as: the number of conditions which will be ruled in and out; the sensitivity and specificity of the tests; the risk/benefit to the patient ratio; the financial cost/benefit to the client ratio; the incidence or prevalence of the condition being tested for; the importance of the condition being tested for (e.g. hypoadrenocorticism is uncommon, but the consequences of failing to diagnose it may be serious).

After the results of initial testing have been obtained the clinician may be in a position to make a definitive diagnosis. Often, however, it is necessary to refine the differential list and select further appropriate testing. The differential list may be reformulated as often as is necessary until a single diagnosis for that problem is made. Often, a single diagnosis will tie in all the problems satisfactorily. However, in many cases, particularly in geriatric patients, concurrent disorders will require multiple diagnoses.

For problem cases in which a clear diagnosis is not made, or the patient fails to respond to treatment as expected, returning to the beginning with the history and physical examination, with the condition often having progressed, can be helpful. However, very few tests are 100% sensitive and specific, and many 'definitive' diagnoses in fact leave room for some doubt. The clinician should never be afraid to revise the initial diagnosis if further evidence comes to light. Those who are concerned that failing to make the correct diagnosis in every case is somehow a sign of inferior clinical abilities

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should take heart from a recent study from the School of Veterinary Medicine at the University of California (Kent et al., 2004). In this paper, clinical and post mortem diagnoses of 623 dogs treated between 1989 and 1999 at the Veterinary Teaching Hospital were compared. It was found that the post mortem diagnosis, presumed to be the correct diagnosis, differed from the clinical diagnosis in approximately ¹/₃ of cases.

This book is organised into seven parts. Part 1 deals with signs likely to be uncovered during history taking. Part 2 deals with signs encountered at the physical examination. Part 3 deals with imaging findings, Part 4 with clinicopathological findings, and Part 5 electrophysiological findings. Part 6 outlines the techniques involved in some common diagnostic procedures and Part 7 contains some algorithms to aid in the diagnosis of common clinical presentations. Four appendices, containing checklists for diagnostic investigations, and a bibliography follow.

The individual lists are categorised as I felt was logical, for example by the DAMNIT-V organisation. DAMNIT-V is a mnemonic for remembering the various pathological processes that may cause a disease:

D - degenerative

A – anomalous (usually listed as congenital in this book)

M - metabolic

N – nutritional, neoplastic

I – inflammatory, infectious, immune-mediated, iatrogenic, idiopathic

T – traumatic, toxic

V - vascular

This categorisation is not appropriate in all cases, however. The individual lists are largely organised alphabetically. The more common conditions are labelled with an asterisk, but, as stated above, whether or not a condition is considered to be common is largely a matter of subjective opinion. Those conditions that are predominantly or exclusively found only in dogs are marked with a (D) and those in cats are marked with a (C).

Sources for the information in this book are wide ranging. A large number of text-books, listed in the bibliography, were consulted, but in most cases it was necessary to expand the lists found in these sources, using information from veterinary journals and conference proceedings.

Although there are undoubtedly omissions from some of the lists, encompassing as this book does virtually the whole of small animal veterinary medicine, I have tried to make it as comprehensive as possible. I would be happy to hear of any omissions, corrections or comments on the text, which can be e-mailed with any supporting references to alex.gough@btconnect.com.

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xvi Introduction

Key

- * = more common condition
- (D) = condition seen exclusively or predominantly in dogs
- (C) = condition seen exclusively or predominantly in cats
- q.v. = more information can be found on this condition elsewhere in this book see Index

References

Chesney, C. (2003) Overdiagnosis in the veterinary field? JSAP, 44:421.

Kent, M. S., et al. (2004) Concurrence between clinical and pathologic diagnoses in a veterinary medical teaching hospital: 623 cases (1989 and 1999). *JAVMA*, 224:403–406.

PART 1 HISTORICAL SIGNS

1.1 General, systemic and metabolic historical signs

1.1.1 Polyuria/polydipsia

Physiological

Exercise

High environmental temperature

Diet

Increased salt intake Very low protein diet

Electrolyte disorders

Hypercalcaemia *q.v.* Hypokalaemia *q.v.* Hypernatraemia *q.v.*

Endocrine disease

Acromegaly Diabetes mellitus* Diabetes insipidus

- Central
- Nephrogenic

Hyperadrenocorticism*

Hyperthyroidism* (C)

Hypoadrenocorticism (D)

Insulinoma

Phaeochromocytoma

Primary hyperaldosteronism

Primary hyperparathyroidism

Hepatobiliary disease, e.g.

Hepatic neoplasia* q.v. Hepatitis/cholangiohepatitis* q.v.

Infectious disease, e.g.

Toxaemia, e.g.

• Pyometra

Miscellaneous

Congenital lack of ADH receptors Hypothalamic disease Pericardial effusion Polycythaemia Psychogenic

Neoplasia*

Renal disorders

Acute renal failure* q.v. Chronic renal failure* q.v. Glomerulonephritis After urethral obstruction Primary renal glycosuria Pyelonephritis Renal medullary washout

Drugs/toxins

Aminophylline Corticosteroids Delmadinone acetate Diuretics Ethylene glycol Indomethacin



Fig. 1.1 Dorsal T1 weighted MR scan of the adrenal glands of a dog with pituitary-dependent hyperadrenocorticism, showing mild bilateral enlargement. Reproduced with permission of Downs Referrals, Bristol.

Lithium

NPK fertilisers

Paraquat

Phenobarbitone

Potassium bromide

Primidone

Proligestone

Terfenadine

Theophylline

Vitamin D rodenticides

Note: Polyuria and polydipsia are considered together here, since one will lead to the other, with only a few exceptions. These include polydipsia in the face of obstructive lower urinary tract disease or oliguric renal failure, and polyuria which is not matched by fluid intake, in which case dehydration will rapidly follow. None of these scenarios are encountered commonly in practice.

References

Garrett, L. D. (2003) Insulinomas: A review and what's new. *Proceedings*, ACVIM, 2003.

Lunn, K. F. (2005) Avoiding the water deprivation test. *Proceedings*, ACVIM, 2005.

Tobias, et al. (2002) Pericardial disorders: 87 cases of pericardial effusion in dogs (January 1, 1999 to December 31, 2001). *Proceedings, ACVIM*, 2002.

1.1.2 Weight loss

Decreased nutrient intake

Anorexia q.v.

Diet

- Poor-quality diet
- Underfeeding

Dysphagia q.v.

Increased nutrient loss

Burns

Chronic blood loss

- Epistaxis q.v.
- Haematemesis q.v.
- Haematuria q.v.
- Melaena q.v.

Diabetes mellitus*

Effusions q.v.

Fanconi syndrome (D)

Intestinal parasites*

Neoplasia

Protein-losing enteropathy*

Protein-losing nephropathy

Increased nutrient use

Endocrine, e.g.

Hyperthyroidism* (C)

Neoplasia*

Physiological

Cold environment

Exercise

Fever q.v.

Lactation

Pregnancy*

Malassimilation

Cardiac failure*

Exocrine pancreatic insufficiency

Hepatic failure/bile salt deficiency* q.v.

Hypoadrenocorticism (D)

Neoplasia*

Renal failure* q.v.

Small intestinal disease* q.v.

Regurgitation and vomiting q.v.

Reference

Rutz, G. M., et al. (2001) Pancreatic acinar atrophy in German Shepherds. Compend Contin Educ Pract Vet, 23:347–56.

1.1.3 Weight gain

Fluid accumulation

Ascites* q.v.

Peripheral oedema q.v.

Pleural effusion

Increased body fat

Overeating

Boredom

Excessive appetite (normal in some breeds)*

High-calorie diets

Overfeeding*

Endocrinopathies

Acromegaly

Hyperadrenocorticism*

Hypogonadism

Hypothyroidism* (D)

Insulinoma

Increased organ size

Hepatomegaly* *q.v.* Renomegaly *q.v.*

Splenomegaly* q.v.

Uterine enlargement q.v.

- Pregnancy*
- Pyometra*

Neoplasia

Large abdominal mass (often associated with poor body condition)* Drugs, e.g.

Corticosteroids

References

Garrett, L. D. (2003) Insulinomas: A review and what's new. *Proceedings*, ACVIM, 2003.

Peterson, M. E., et al. (1990) Acromegaly in 14 cats. JVIM, 4:192-201.

1.1.4 Polyphagia

Behavioural/psychological

Normal in some breeds* Boredom

Physiological

Cold environment Increased exercise Lactation* Pregnancy*

Malassimilation*

Increased nutrient loss

Increased nutrient use

Diet

Highly-palatable food* Poor-quality food

Endocrine

Diabetes mellitus*
Hyperadrenocorticism*
Hyperthyroidism* (C)
Insulinoma

Miscellaneous

Peritoneopericardial diaphragmatic hernia

Drugs/toxins

Aminophylline

Benzodiazepines

Cannabis

Cyproheptadine

Delmadinone acetate

Glucocorticoids

Phenobarbitone

Potassium bromide

Primidone

Proligestone

References

Garrett, L. D. (2003) Insulinomas: A review and what's new. Proceedings, ACVIM, 2003.

Rexing, J. F. & Coolman, B. R. (2004) A peritoneopericardial diaphragmatic hernia in a cat. *Vet Med*, 99:314–18.

1.1.5 Anorexia/inappetence

Difficulty with prehension

Blindness q.v.

Myopathy, e.g.

Masticatory myositis

Tetanus

Pain on opening jaw, e.g.

Mandibular or maxillary fracture

Retrobulbar abscess

Skull fractures

Soft tissue trauma

Temporo-mandibular joint disease

Trigeminal nerve disease, e.g.

Neoplasia

Trigeminal neuritis

Difficulty with mastication

Dental disease*

Lingual disease

Oral neoplasia*

Oral ulceration, e.g.

Ingestion of caustic or acidic substances*

· Renal disease

Difficulty with swallowing

Pharyngeal disease

Foreign body*

Neoplasia

Neurological disease

Ulceration

Oesophageal disease, e.g.

Foreign body*

Neoplasia

Ulceration

Megaoesophagus

Stricture

Vascular ring anomaly

Primary anorexia

Intracranial disease, e.g.

• Hypothalamic neoplasia

Secondary anorexia

Anosmia

- Chronic rhinitis q.v.
- Nasal neoplasia
- Other nasal disease
- Neurological disease

Endocrine disease, e.g.

- Diabetic ketoacidosis
- Hypoadrenocorticism (D)

Fever* q.v.

Gastrointestinal disease q.v., e.g.

- Gastritis
- Inflammatory bowel disease*

Heart disease, e.g.

· Cardiac failure*

Hepatic disease* q.v.

Infection*

Metabolic abnormalities, e.g.

- Hypercalcaemia q.v.
- Hypokalaemia q.v.

Pain*

Pancreatic disease*, e.g.

• Pancreatitis

Respiratory disease, e.g.

- Airway disease* q.v.
- Diaphragmatic hernia
- Pleural effusion* q.v.
- Pneumonia q.v.

Renal disease* q.v.

Drugs

- Acetazolamide
- Amiodarone
- Amphotericin B
- Bethanechol
- Bromocriptine
- Butorphanol
- Cardiac glycosides
- Chlorambucil
- Diazoxide
- Doxorubicin
- Fentanyl

- Hydralazine
- Itraconazole
- Ketoconazole
- Melphalan
- Methimazole
- Mitotane
- Nicotinamide
- Oxytetracycline (C)
- Penicillamine
- Theophylline
- Trimethoprim/sulphonamide (C)

Diet

Recent dietary changes* Unpalatable diet*

Psychological/behavioural* factors

Altered schedule New family members New house New pets

Reference

Forman, M. A., et al. (2004) Evaluation of serum feline pancreatic lipase immunoreactivity and helical computed tomography versus conventional testing for the diagnosis of feline pancreatitis. *JVIM*, 18:807–15.

1.1.6 Failure to grow

With good body condition

Chondrodystrophy (normal in many breeds)* (D)

Endocrine disorders

- Congenital hyposomatotropism (pituitary dwarfism)
- Congenital hypothyroidism
- Hyperadrenocorticism

With poor body condition

Dietary intolerance Exocrine pancreatic insufficiency*

Inadequate nutrient intake

Anorexia *q.v.*Poor-quality diet
Underfeeding

Cardiac disorders, e.g.

Congenital Endocarditis

Hepatic disorders, e.g.

Hepatitis q.v.

Portosystemic shunt

Oesophageal disorders, e.g.

Megaoesophagus q.v.

Vascular ring anomaly (e.g. persistent right aortic arch)

Gastrointestinal disease, e.g.

Histoplasmosis

Obstruction, e.g.

- Foreign body*
- Intussusception*

Parasites*

Renal disease

Congenital kidney disease Glomerulonephritis Pyelonephritis

Inflammatory disease

Endocrine disease

Diabetes insipidus

Diabetes mellitus*

Hypoadrenocorticism (D)

Reference

Chastain, C. B., et al. (2001) Combined pituitary hormone deficiency in German shepherd dogs with dwarfism. *Sm Anim Clin Endocrinol*, 11:1–4.

1.1.7 Syncope/collapse (see Table 1.1)

Cardiovascular dysfunction

Myocardial failure Myocardial infarction Shock *q.v.*

Bradyarrhythmias q.v., e.g.

High grade second degree heart block Sick sinus syndrome (D) Third degree heart block

Tachyarrhythmias q.v.

Supraventricular tachycardia* Ventricular tachycardia*

Table 1.1 Differentiating seizures from syncope. This table is a guide to the differentiation of generalised seizures from syncopal episodes. However, there is a lot of overlap between the two: syncopal episodes may involve convulsions; seizures may occur on exercise; tonic-clonic motions may not always be observed with seizures.

	Syncope	Seizure (generalised)
Precipitating event/ timing	Exercise, excitement, stress, cough, urination, defecation	Often at rest or on waking
Pre-event	Acute weakness, staggering, vocalisation	Anxiety, attention-seeking
Event	Usually flaccid limbs but may be rigid	Jaw motions, hypersalivation, tonic-clonic limb motion or limb rigidity
	Duration less than 1 minute	Duration often greater than 1 minute
	Rarely urination/defecation Usually retain consciousness, but may lose consciousness	Urination and/or defecation Loss of consciousness
	Abnormal heart rhythm or rate may or may not be palpatated/auscultated	Often sinus tachycardia
Post-event	Rapid recovery	Slow recovery Prolonged post-event disorientation

Obstruction to flow

Congenital, e.g.

- Aortic stenosis (D)
- Pulmonic stenosis (D)

Hypertrophic obstructive cardiomyopathy

Pericardial effusion* (D)

Pulmonary hypertension

Arterial obstruction, e.g.

- Neoplasia
- Thrombosis

Hypoxaemic disease

Carboxyhaemoglobinaemia Methaemoglobinaemia

Respiratory disease

Upper airway, e.g.

- Brachycephalic obstructive airway syndrome
- Laryngeal paralysis
- Tracheal collapse
- Tracheal obstruction

Lower airway, e.g.

- Pneumonia
- Small airway disease

Ventilation-perfusion mismatch, e.g.

• Lung collapse

Pleural/thoracic disorders, e.g.

- Pleural effusion
- Pneumothorax
- Rib fractures

Right-to-left cardiac shunt, e.g.

Reverse-shunting patent ductus arteriosus Severe anaemia

Neurological dysfunction

Brainstem disease Glossopharyngeal neuralgia Micturition-related collapse Narcolepsy/cataplexy Seizures *q.v.* Swallowing-related collapse

Diffuse cerebral dysfunction, e.g.

Encephalopathy
Haemorrhage
Hydrocephalus
Inflammation
Oedema
Space occupying lesion
Trauma

Lower motor neurone disorders

Endocrine neuropathies, e.g.

- Diabetes mellitus*
- Hyperadrenocorticism
- Hypothyroidism* (D)

Lumbosacral disease

Paraneoplastic neuropathies, e.g.

• Insulinoma

Peripheral nerve neoplasia

Polyneuropathy

Polyradiculoneuropathy

Neuromuscular junction disorders

Botulism Myasthenia gravis

Upper motor neurone disorders

Central vestibular disease Cerebellar disease Cerebral disease Peripheral vestibular disease Spinal disease

Miscellaneous

Carotid sinus stimulation, e.g.

- Neoplasia
- Tight collar

Hyperventilation

Postural hypotension

Tussive syncope

Metabolic disorders

Diabetic ketoacidosis

Hypercalcaemia/hypocalcaemia q.v.

Hypernatraemia/hyponatraemia q.v.

Hyperthermia/hypothermia q.v.

Hypoglycaemia q.v.

Hypokalaemia q.v.

Severe acidosis q.v.

Severe alkalosis q.v.

Myopathies

Corticosteroid myopathy

Exertional myopathy

Hypocalcaemic myopathy

Hypokalaemic myopathy

Malignant hyperthermia

Mitochondrial myopathy

Muscular dystrophy

Polymyopathy

Polymyositis

Protozoal myopathy

Skeletal/joint disorders

Bilateral cranial cruciate disease

Bilateral hip disease

Discospondylitis

Intervertebral disc disease

Multiple myeloma

Osteoarthritis

Panosteitis

Patellar luxation

Polyarthritis

Drugs

Anti-arrhythmics, e.g.

- Atenolol
- Digoxin
- Propranolol
- Quinidine

Sedatives, e.g.

• Phenothiazines

Vasodilators, e.g.

- ACE inhibitors
- Hydralazine
- Nitroglycerine

References

Berendt, M. (2001) The diagnosis of epilepsy: seizure phenomenology and classification. *Proceedings of the World Small Animal Veterinary Association World Congress*, 2001.

Shelton, G. D. (1998) Myasthenia gravis: lessons from the past 10 years. *JSAP*, 39:368–72.

Ware, W. A. (2002) Syncope. Proceedings, Waltham/OSU Symposium, Small Animal Cardiology, 2002.

Wray, J. (2005) Differential diagnosis of collapse in the dog. 1. Aetiology and investigation. *In Practice* 27:16–28.

1.1.8 Weakness

Metabolic disease

Renal failure* q.v.

Hepatic failure* q.v.

Hypoglycaemia q.v.

Electrolyte disorders*

- Hypercalcaemia*/hypocalcaemia q.v.
- Hyperkalaemia/hypokalaemia* q.v.
- Hypernatraemia/hyponatraemia q.v.

Acid-base disorders

- Acidosis *q.v.*
- Alkalosis *q.v.*

Infectious diseases*

Bacterial

Viral

Fungal

Rickettsial

Protozoal

Other parasitic diseases

Immune-mediated/inflammatory diseases

Chronic inflammatory conditions*

Immune-mediated haemolytic anaemia* q.v.

Immune-mediated polyarthritis

Haematological diseases

Anaemia* q.v.

Hyperviscosity syndrome

Endocrine diseases

Diabetes mellitus*

Hyperadrenocorticism

Hyperparathyroidism

Hypoadrenocorticism (D)

Hypoparathyroidism

Hypothyroidism* (D)

Insulinoma

Cardiovascular diseases

Bradyarrhythmias q.v., e.g.

- High grade second degree heart block
- Sick sinus syndrome (D)
- Third degree heart block

Congestive heart failure*

Pericardial effusion* q.v.

Hypertension* q.v.

Hypotension* q.v.

Tachyarrhythmias q.v., e.g.

• Ventricular tachycardia*

Respiratory diseases

Airway obstruction, e.g.

- Feline asthma* (C)
- Foreign body*
- Neoplasia *

Intrathoracic neoplasia*

- Pleural effusion*
- Pulmonary hypertension
- Pulmonary oedema* q.v.
- Pulmonary thromboembolism

Severe pulmonary parenchymal disease

Neuromuscular diseases

Epilepsy* q.v.

Myasthenia gravis

Myopathies

Vestibular disease* q.v.

Intracranial disease, e.g.

Cerebrovascular accident

Infection

Inflammation

Space-occupying lesions

Spinal cord disease q.v., e.g.

Infection

Inflammation

Intervertebral disc disease* (D)

Neoplasia

Trauma*

Peripheral polyneuropathies

Endocrine disorders, e.g.

- Diabetes mellitus*
- Hyperadrenocorticism
- Hypothyroidism* (D)

Polyradiculoneuritis

Paraneoplastic disorders

Drugs/toxins, e.g.

- Cisplatin
- Lead
- Vincristine

Infections

Botulism

Tick paralysis

Systemic disorders

Dehydration*

Fever* q.v.

Neoplasia*

Nutritional disorders

Cachexia, e.g.

Heart failure*

Neoplasia*

Inadequate calorie intake, e.g.

Anorexia* q.v.

Poor-quality diet

Specific nutrient deficiencies, e.g.

Minerals

Vitamins

Physiological factors

Over-exercise

Pain*

Stress/anxiety*

Drugs/toxins

Alphachloralose

Anticoagulant rodenticides

Anticonvulsants

Antihistamines

Blue-green algae

Cannabis

Diclofenac sodium

Glucocorticoids

Hypotensive agents, e.g.

- Beta-blockers
- Vasodilators

Ibuprofen

Insulin overdosage

Iron salts

Mistletoe

Opioids

Organophosphates
Petroleum distillates
Phenoxy acid herbicides
Pyrethrin/pyrethroids
Rhododendron
Salbutamol
Sedatives

References

Sadek, D. & Schaer, M. (1996) Atypical Addison's disease in the dog: a retrospective survey of 14 cases. *JAAHA*, 32:159–63.

Shelton, G. D. (1998) Myasthenia gravis: lessons from the past 10 years. *JSAP*, 39:368–72.

1.2 Gastrointestinal/abdominal historical signs

1.2.1 Ptyalism/salivation/hypersalivation

Physiological factors

Appetite stimulation* Fear* Stress*

Oral cavity disease

Dental disease* Foreign body* Neoplasia*

Inability to close mouth, e.g.

Mandibular trauma*

Trigeminal nerve disease, e.g.

- Idiopathic trigeminal neuritis
- Infiltrating neoplasia, e.g.
 - Lymphoma
 - Nerve sheath tumours

Ulceration*, e.g.

Immune-mediated disease Ingestion of irritant substance Renal failure*

Inflammation*

Faucitis*
Gingivitis*
Glossitis*
Oesophagitis*
Stomatitis*

Neurological disease

Cataplexy/narcolepsy
Hepatic encephalopathy
Intracranial neoplasia
Partial seizures

Nausea/regurgitation/vomiting q.v.

Salivary gland disease q.v.

Salivary gland necrosis/sialadenitis Salivary mucocoele Sialadenosis

Normal breed variation, e.g.

St Bernards

Drugs/toxins

Adder bites

Alphachloralose

Baclofen

Batteries

Benzodiazepines

Bethanechol

Blue-green algae

Cannabis

Carbamate

Chocolate/theobromine

Cotoneaster

Cyanoacrylate adhesives

Daffodil

Dieffenbachia

Dinoprost tromethamine

Glyphosphate

Horse chestnut

Ivermectin

Ketamine

Laburnum

Levamisole (C)

Loperamide

Metronidazole

Mistletoe

NPK fertilisers

Organophosphates

Paracetamol

Paraquat

Phenoxy acid herbicides

Plastic explosives

Pyrethrin/pyrethroids

Pyridostigmine

Rhododendron

Rowan

Terfenadine

Toads

Trimethoprim/sulphonamide (C)

Xylazine

References

Patterson, E. E., et al. (2003) Clinical characteristics and inheritance of idiopathic epilepsy in Vizslas. *JVIM*, 17:319–25.

Schroeder, H. & Berry, W. L. (1998) Salivary gland necrosis in dogs: a retrospective study of 19 cases. *JSAP*, 39:121–25.

Sozmen, M., et al. (2000) Idiopathic salivary gland enlargement (sialadenosis) in dogs: a microscopic study. *JSAP*, 41:243–47.

1.2.2 Gagging/retching

Congenital disease

Achalasia, e.g.

• Cricopharyngeal achalasia (D)

Cleft palate

Hydrocephalus

Neuromuscular disease

Brainstem disease

Cranial nerve defects (V, VII, IX, XII)

Encephalitis

Laryngeal paralysis*

Muscular dystrophy

Myasthenia gravis

Immune-mediated and infectious disease

Asthma* (C)

Bacterial encephalitis

Fungal disease

• Granuloma complex

Idiopathic glossopharyngitis

Laryngitis*

Pharyngitis*

Rabies

Rhinitis*

Sialadenitis

Viral encephalitis

Systemic disorders

Hypocalcaemia

Renal failure*

Trauma

Foreign body*

Pharyngeal haematoma

Styloid apparatus trauma Tracheal rupture

Neoplasia

Central nervous system

Epiglottis

Inner ear

Nasal

Pharyngeal

Tonsillar

Nutrition

Food texture and size

Respiratory disease (expectoration), e.g.

Bronchitis*

Haemorrhage

Pulmonary oedema*

Toxic

Botulism

Ingestion of irritant chemical

Smoke

Reference

Schroeder, H. & Berry, W. L. (1998) Salivary gland necrosis in dogs: a retrospective study of 19 cases. *JSAP*, 39:121–25.

1.2.3 Dysphagia

Infectious/inflammatory disease

Oral disease

Dental disease*

Osteomyelitis of jaw

Periodontitis*

Pharyngitis*

Rabies

Retrobulbar abscess

Severe gingivitis*

Tooth root abscess*

Ulceration, e.g.

- Ingestion of irritant substance
- · Renal disease*

Obstruction

Foreign body*

Granuloma

Neoplasia Sialocoele

Trauma

Fracture*
Haematoma
Laceration*

Temporomandibular joint disease

Neuromuscular disease

Cricopharyngeal achalasia Myasthenia gravis

Myopathy, e.g.

Masticatory myopathy

Trigeminal nerve disease, e.g.

- Intracranial disease
- Trigeminal neuritis

References

Meomartino, L., et al. (1999) Temporomandibular ankylosis in the cat: a review of seven cases. *JSAP*, 40:7–10.

Preifer, R. M. (2003) Cricopharyngeal achalasia in a dog. Can Vet J, 44:993-5.

1.2.4 Regurgitation

Salivary gland disease

Sialadenitis

Oesophageal disease

Foreign body*

Megaoesophagus

- Idiopathic
- Acquired

Neoplasia

Oesophageal diverticulum

Oesophageal fistula

Oesophageal inclusion cysts

Oesophagitis*

Stricture

Vascular ring anomaly, e.g.

• Persistent right aortic arch

Gastric disease

Gastric dilatation-volvulus* (D)

Hiatal hernia

Pyloric outflow obstruction, e.g.

- Foreign body*
- Neoplasia
- Pyloric stenosis

Neuromuscular disease

Peripheral neuropathies, e.g.

Giant cell axonal neuropathy (D)

Lead poisoning

Polyneuritis

Polyradiculoneuritis

Central nervous system disease, e.g.

Brainstem disease

Infection

Inflammation

Intracranial space occupying lesion

Trauma

Neuromuscular junctionopathies, e.g.

Acetylcholinesterase toxicity

Botulism

Myasthenia gravis

Tetanus

Immune-mediated disease

Dermatomyositis (D)

Polymyositis

Systemic lupus erythematosus

Endocrine disease

Hypoadrenocorticism (D)

Hypothyroidism* (D)

References

Han, E., et al. (2003) Feline esophagitis secondary to gastroesophageal reflux disease: clinical: signs and radiographic, endoscopic and histopathological findings. *JAAHA*, 39:161–7.

Hodges, J., et al. (2004) Recurrent regurgitation in a young cat with an unknown history. *Vet Med*, 99:244–51.

Schroeder, H. & Berry, W. L. (1998) Salivary gland necrosis in dogs: a retrospective study of 19 cases. *ISAP*, 39:121–5.

White, R. N., et al. (2003) Vascular ring anomaly with coarctation of the aorta in a cat. *JSAP*, 44:330–34.

1.2.5 Vomiting

ACUTE VOMITING

Dietary

Dietary indiscretion*

Dietary intolerance*

Sudden change in diet*

Gastrointestinal disease

Colitis*

Constipation/obstipation* q.v.

Foreign body*

Gastric dilatation/volvulus*

Gastric or duodenal ulceration*

Gastritis/enteritis*

Haemorrhagic gastroenteritis*

Infection, e.g.

- Bacterial*
- Parasites*
- Viral*

Inflammatory bowel disease*

Intestinal volvulus

Intussusception

Neoplasia*

Endocrine disease, e.g.

Diabetic ketoacidosis*

Hypoadrenocorticism (D)

Metabolic/systemic disease

Hypercalcaemia/hypocalcaemia q.v.

Hyperkalaemia/hypokalaemia* q.v.

Hyperthermia* q.v.

Liver disease* q.v.

Pancreatitis*

Peritonitis*

Prostatitis*

Pyometra* (D)

Renal disease* q.v.

Septicaemia*

Urinary obstruction*

Vestibular disease*

Miscellaneous conditions

Central nervous system disease

Diaphragmatic hernia

Motion sickness

Psychogenic

Drugs/toxins

Acetazolamide

Adder bite

Allopurinol

Alpha-2 agonists

Aminophylline

Amphotericin B

Apomorphine

Aspirin

Atipamezole

Atropine