

GUIDE TO CONGENITAL AND HERITABLE DISORDERS IN DOGS*

Before you decide whether to buy a dog from a breeder or a pet shop, please assess the impact this will have on the current serious problem of dog overpopulation. In our country alone, millions of dogs are killed each year simply because there are too many for the number of homes available. Purchasing dogs from commercial sources ensures the continued death of other dogs waiting for homes in our animal control facilities or shelters. *Adopting a dog from an animal control facility or shelter will save a life.* Dogs from these sources will not cost any more to feed and are unlikely to require any greater care than purebred dogs. More importantly, they will be equally as loving, lovable and compatible with your family.

If you are set on having a particular breed of dog, there are many options available. There are many active breed rescue groups which concentrate on finding homes for existing dogs of virtually any breed. Also, at many of the animal control facilities or shelters, a substantial percentage of the dogs available are purebreds.

There is another consideration with respect to purebred dogs, particularly those from irresponsible breeders or puppy mills and that is the issue of congenital or genetically transmitted disorders. Purebred dogs of many breeds and even mixed breed dogs are prone to specific abnormalities which may be familial or genetic in nature. Often, these health problems are unapparent to the average person, but could result in misery for both the dog and her or his guardians.

This booklet is intended to make you aware of the potential problems associated with various purebred dogs. If you do decide to purchase a purebred dog, it would be prudent to ask the seller if any of the relatives have been affected by the conditions listed for that breed. Furthermore, you should ask the seller who will be responsible for the veterinary costs if a puppy is afflicted with a heritable disease which may not manifest itself until later in life.

*Includes Genetic Predisposition to Diseases

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Directory

Section I

Provides a list of 166 more commonly known purebred dog breeds, each of which is accompanied by a number or series of numbers that correspond to the congenital and heritable diseases identified and described in Section II.

Section II

This section provides an alphabetical listing of congenital and genetically transmitted diseases that may occur in purebred dogs. Each disease is assigned an identification number, and some diseases are followed by the names of the breeds known to be subject to those diseases.

How to use this book

If you are thinking about acquiring a purebred dog, refer to Section I to find out the number of congenital and genetically transmitted diseases associated with the breed or breeds in which you are interested. Refer to Section II to find the names and definitions of those diseases.

Disclaimer:

This report was designed to catalogue the identified congenital and hereditary disorders found in purebred dogs and to describe other common conditions that are thought to be genetic in origin because they appear with higher-than-expected frequency in certain breeds.

The knowledge of and experience with animal diseases is always changing, and new diseases are discovered each year. As a result, this catalogue cannot be complete. Every attempt has been made, nevertheless, to provide the latest information from published articles in scientific and popular journals, from review chapters written by veterinary and other professionals, and from breed-club literature provided by designated representatives of individual breed clubs.

The conditions listed in the Guide to Congenital and Heritable Disorders in Dogs occur with different degrees of frequency and severity from one breed to the next. Hip dysplasia, umbilical hernias, and allergies, for example, are common conditions found in many breeds, whereas serious malformations and life-threatening diseases are typically found with less frequency. The impact of a particular disorder on the health and longevity of the breed or breeds that it affects will depend on the nature and severity of that disorder. Therefore, no attempt has been made to assign priority among the diseases identified in this catalogue. This sort of information should be sought from veterinarians and other medical professionals with relevant experience and from national breed-club representatives, whose names can be obtained from the American Kennel Club, 51 Madison Avenue, New York, NY 10010.

Section I:

Aberdeen Terrier: 324

Affenpinscher: 12, 55, 98, 218, 235, 236, 330

Afghan Hound: 14, 42, 65, 96, 114, 121, 135, 145, 147, 166, 192, 206a, 211, 221, 221a, 239, 245, 256, 269, 270, 312, 330

Airedale Terrier: 7, 9, 48, 65, 88, 103, 122, 140, 146, 148, 165, 166, 168, 192, 206a, 230, 256, 269, 270, 273a, 312, 314, 318, 330

Akbash: 166, 192, 312, 318

Akita: 9, 10, 11, 27, 31, 43, 65, 71, 103, 114, 115, 131, 135, 137, 140, 146, 152, 156, 166, 172, 178a, 192, 193a, 199, 206, 221, 221a, 239, 256, 258a, 270, 273a, 291, 311a, 312, 318, 329, 330

Alaskan Malamute: 13, 42, 59, 65, 67, 80, 85, 89, 120, 121, 122, 128a, 135, 144, 147, 148, 150, 152, 166, 206a, 213, 221, 221a, 250b, 256, 266, 299b, 330, 334

American Bulldog: 42, 80, 95, 103, 152, 166, 214, 330

American Cocker Spaniel: 1, 2, 10, 12, 18, 26, 27, 38, 38a, 42, 43, 54, 55, 64, 65, 69, 72, 73, 88, 94, 94a, 95, 103, 107, 109, 121, 123, 135, 146, 147, 148, 149, 150, 152, 154, 166, 171, 173, 179, 181, 186, 188, 192, 193a, 197, 220, 221, 221a, 226, 228, 235, 236, 242, 245, 250, 254, 256, 266, 270, 275, 276, 286, 307, 311a, 312, 318, 319, 320, 330

American Eskimo: 10, 21, 22, 81, 156, 166, 239, 263, 303a

American Foxhound: 78, 199, 290, 311

American Staffordshire Terrier: 42, 54, 55, 88, 103, 143a, 152, 166, 193a, 204a, 214, 221, 221a, 256

American Water Spaniel: 42, 150, 270

Antarctic Husky: 103, 147

Australian Cattle Dog (Australian Blue Heeler): 42, 78, 109, 121, 124b, 147, 149, 152, 166, 171, 177, 186, 193a, 200, 214, 221, 228, 245, 256, 270, 318, 319

Australian Kelpie: 58, 199, 203, 256

Australian Shepherd: 42, 52, 55, 58, 78, 86, 89, 152, 166, 176a, 193a, 199, 203, 214, 221, 221a, 245, 256, 269, 270, 287, 318, 328, 329, 330

Australian Terrier: 85, 185, 256, 270

Basenji: 27, 56, 59, 66, 124c, 146, 166, 171, 172, 192, 245, 256, 263, 268, 270, 318

Basset Hound: 5, 9, 9a, 15, 24a, 27, 31, 61a, 70, 94, 103, 105, 109, 114, 120, 121, 131, 135, 136, 140, 146, 147, 157, 159a, 166, 168, 169, 170, 171, 173, 174, 186, 190, 193, 196, 221, 221a, 222, 226, 231, 235, 245, 249, 250, 256, 273a, 274, 291, 299, 311, 318, 330, 332

Beagle: 10, 11, 21, 34, 37, 42, 43, 54, 55, 65, 72, 80, 88, 94a, 109, 114, 120, 121, 135, 136, 146, 147, 150, 157, 166, 168, 173, 182, 188a, 192, 193a, 202, 204, 212, 220, 227, 242, 245, 256, 260, 267, 270, 275, 280, 310, 312, 327, 330

Bearded Collie: 9a, 27, 42, 65, 146, 152, 159a, 166, 192, 239, 245, 256, 269, 270, 286, 303, 311a, 312

Bedlington Terrier: 2, 23, 42, 52a, 64, 88, 94, 184, 199, 210, 223, 256, 265, 266, 269, 270

Belgian Malinois: 109, 152, 166, 256

Belgian Sheepdog: 42, 109, 152, 166, 204a, 230, 256, 269, 270

Belgian Tervuren: 42, 109, 166, 221, 221a, 230, 245b, 256

Bernese Mountain Dog: 20, 42, 46, 47, 54, 55, 95, 103, 149, 149d, 152, 153a, 166, 194b, 221, 221a, 255, 256, 258a, 269, 318

Bichon Frise: 42, 53, 65, 103, 109, 110, 122, 148, 235, 250, 269, 270, 322a, 331a

Black and Tan Coonhound: 94, 103, 122, 148, 152, 221, 221a, 252

Bloodhound: 31, 94, 103, 114, 131, 152, 166, 179, 181, 195, 221, 221a, 245, 324

Blue Tick Hound: 136, 193a, 221, 221a

Border Collie: 2, 52, 58, 64, 65, 86, 109, 149, 152, 186, 214, 221, 221a, 256

Border Terrier: 17, 39, 42, 58, 68, 70, 126, 145, 186, 196, 217, 235, 248, 256, 270, 324, 327

Borzoi (Russian Wolfhound): 31, 36, 42, 118, 131, 152, 155, 166, 192, 199, 200, 230, 245, 256, 270, 312, 330

Boston Terrier: 10, 12, 17, 22, 39, 42, 54, 55, 65, 67, 68, 71, 78, 80, 88, 90, 103, 112, 114, 135, 145, 151, 154, 159, 166, 171, 174, 179, 181, 196, 197, 235, 236, 237, 248, 256, 262, 275, 295, 304, 308

Bouvier des Flandres: 27, 42, 55, 74, 90, 94, 100, 103, 131, 135, 152, 166, 184a, 221, 221a, 300, 318, 330

Boxer: 3, 6, 8, 10, 22, 24, 38, 38a, 42, 67, 72, 75, 80, 83, 88, 94a, 99, 103, 113, 114, 119, 121, 131, 134, 139, 149, 153, 156, 166, 192, 193, 196, 214a, 221, 221a, 242a, 250, 256, 277, 293, 294, 297, 300, 304a, 312, 317, 325, 330

Briard: 42, 61d, 105, 166, 231, 256, 330

Brittany: 42, 55, 61b, 61d, 88, 109, 121, 147, 149c, 166, 186, 221, 221a, 256, 270

Brussels Griffon: 1, 42, 54, 55, 88, 152, 154, 185, 235, 256, 278, 281

Bullmastiff: 3, 27, 31, 51, 55, 88, 103, 114, 131, 135, 152, 166, 192, 193, 221, 221a, 245, 256, 270, 273, 280, 312, 325

Bull Terrier: 7, 12, 21, 27, 37b, 70, 78, 80, 94, 103, 130, 143a, 149b, 171, 186, 192, 196, 201, 221, 221a, 235, 250b, 261, 264a, 266a, 294, 312, 318, 333

Cairn Terrier: 1, 42, 48, 68, 75, 85, 121, 122, 135, 136, 147, 148, 149, 149c, 166, 171, 186, 193a, 199a, 252a, 256, 263, 270, 330

Cardigan Welsh Corgi: 61a, 75, 90, 103, 135, 169, 173, 186, 245, 256, 270

Cavalier King Charles Spaniel: 37b, 42, 65, 85, 88, 103, 114a, 124d, 166, 179, 199, 201, 235, 249, 256, 270, 305, 311a

Chesapeake Bay Retriever: 42, 88, 103, 114, 152, 166, 192, 221, 221a, 256, 270, 312, 330

Chihuahua: 55, 57, 65, 70, 87, 103, 121, 135, 147, 149, 154, 160, 162, 166, 175, 179, 181, 186, 201, 206, 214, 221, 221a, 235, 256, 260, 313

Chinese Crested: 21, 85, 143a, 166, 271

Chinese Shar-Pei: 10, 11, 22, 29, 72a, 80, 94, 95, 103, 128, 129, 130, 135, 152, 166, 168, 172, 186, 187, 202a, 221, 221a, 226, 235, 250, 256, 270, 276, 277a, 295, 319, 326

Chinook: 22, 98a, 166.

Chow Chow: 9b, 27, 31, 42, 48, 55, 61, 80, 88, 94, 95, 98, 103, 130, 135, 140, 152, 160a, 165, 166, 172, 192, 193, 197, 204, 221, 221a, 230, 239, 245, 250, 256, 273a, 280, 312

Clumber Spaniel: 94, 103, 152, 309, 319, 324

Collie: 33, 37, 42, 45, 52, 58, 59, 60, 65, 73, 78, 80, 82, 86, 88, 89, 103, 109, 121, 126, 131, 140, 147, 152, 166, 171, 176, 176a, 192, 199, 203, 208, 209, 220, 221, 221a, 236, 238, 239, 245, 250, 256, 269, 270, 273a, 312, 318, 330

Coton de Tulear: 21, 78, 115, 152, 166, 192, 200, 235, 312

Curly-Coated Retriever: 42, 88, 94, 103, 137, 193a, 256, 299b

Dalmatian: 2, 10, 22, 32, 38, 64, 78, 80, 81, 83, 88, 103, 129, 130, 135, 136, 140, 143a, 152, 166, 184a, 192, 193a, 199, 214, 221, 221a, 230, 250b, 256, 258a, 268, 273a, 275, 294, 312, 321, 322, 322a

Dandie Dinmont Terrier: 3, 42, 67, 87, 97, 103, 135, 149, 152, 166, 173, 235, 245, 281

Doberman Pinscher: 2, 3, 4, 6, 7, 9a, 27, 35, 38, 42, 51, 52a, 53, 59, 60a, 64, 68, 80, 103, 105, 121, 127, 129, 138a, 140, 143a, 146, 147, 152, 161, 166, 170, 173, 182, 192, 199, 206a, 221, 221a, 225, 231, 239, 243, 245, 250, 251, 256, 266, 266a, 267, 270, 273a, 292, 303a, 304a, 312, 328, 330

English Bulldog: 1, 3, 6, 19, 42, 54, 55, 57, 80, 88, 90, 94, 98, 103, 116, 128, 129, 130, 145, 152, 154, 164, 166, 179, 181, 193, 196, 201, 205, 217, 242, 245, 250, 260, 261, 278, 280, 287, 308, 313, 325, 330

English Cocker Spaniel: 42, 70, 88, 94, 103, 119, 135, 146, 147, 149b, 150, 166, 177, 186, 214, 221, 221a, 226, 236, 245, 256, 259, 270, 304, 330

English Foxhound: 78, 290

English Setter: 27, 31, 42, 61, 68, 78, 91, 94, 103, 121, 124b, 147, 152, 160, 166, 177, 181, 188a, 192, 193a, 214, 221, 221a, 256, 258a, 312, 323, 330

English Springer Spaniel: 9a, 10, 12, 18, 26, 27, 42, 43, 54, 55, 59, 65, 69, 72, 88, 94, 94a, 95, 103, 107, 109, 121, 123, 124, 129b, 135, 140, 146, 147, 148, 149, 150, 152, 154, 159a, 160a, 166, 171, 173, 188, 193a, 197, 206, 206a, 213, 220, 221, 221a, 226, 228, 235, 236, 242, 245, 245a, 254, 256, 264a, 266, 270, 273a, 275, 276, 286, 307, 318, 319, 320, 330

English Toy Spaniel (King Charles and Ruby Blenheim Spaniels): 42, 55, 65, 85, 103, 143, 235, 249, 270, 311a, 318

Field Spaniel: 14, 42, 166, 256, 270

Finnish Spitz: 85, 166

Flat-Coated Retriever: 42, 88, 94, 103, 152, 153a, 166, 256, 286a

French Bulldog: 42, 54, 55, 88, 90, 98, 103, 119, 121, 122, 145, 147, 148, 172, 173, 261, 330

German Pinscher: 27, 166

German Shepherd: 10, 21, 27, 36, 38, 42, 43a, 44, 54, 55, 59, 65, 72, 72b, 75, 79a, 81, 83, 86, 89, 94a, 95, 102, 103, 105, 109, 112, 114, 121, 122, 129a, 130, 221a, 131, 137, 140, 143a, 147, 148, 149c, 152, 166, 168, 171a, 180, 186, 190a, 191, 192, 193a, 194, 202b, 208, 214a, 218a, 220, 221, 221a, 225, 226, 229, 230, 231, 236, 238, 241, 243, 247, 250, 250b, 256, 258a, 266, 270, 273a, 276, 283, 299b, 300, 306, 312, 316, 320, 327a, 330

German Shorthaired Pointer: 8, 9a, 42, 65, 103, 114, 124b, 125, 131, 150, 152, 159a, 166, 177, 188a, 191, 193a, 196, 197, 206, 214, 221, 221a, 230, 256, 300, 311, 330

German Wirehaired Pointer: 42, 103, 148, 152, 166, 221, 221a, 270, 302, 330

Giant Schnauzer: 42, 89, 105, 135, 146, 152, 155, 158, 166, 190a, 192, 218a, 221, 221a, 231, 256, 269, 270, 276, 311a, 312, 327b

Golden Retriever: 7, 9, 10, 21, 22, 27, 38, 42, 49a, 52a, 59, 65, 81, 85, 88, 94, 95, 103, 121, 129, 129a, 130, 140, 143a, 146, 147, 149, 152, 166, 178, 192, 193, 193a, 204a, 206, 220, 221, 221a, 245, 250, 250b, 256, 258a, 262, 266a, 273a, 300, 308a, 312, 328, 329

Gordon Setter: 31, 42, 45, 103, 128a, 131, 152, 166, 221, 221a, 256, 179, 181, 270, 307

Great Dane: 6, 7, 9a, 31, 36, 38, 42, 48, 50, 51, 61, 75, 78, 80, 83, 88, 94, 103, 114, 131, 135, 144, 152, 153, 155, 158, 159a, 166, 176, 192, 198, 199, 201, 204a, 211, 221, 221a, 225, 243, 250, 255, 256, 270, 292, 298, 312, 330

Great Pyrenees: 5, 9, 16, 31, 42, 78, 80, 94, 103, 122, 124, 131, 148, 152, 166, 195, 221, 221a, 244, 245, 256, 303a, 304, 311a, 325

Greyhound: 14, 42, 65, 72, 88, 90, 94a, 109, 112, 121, 143a, 147, 155, 166, 186, 190a, 218a, 220, 221, 221a, 225, 230, 237, 245, 256, 279, 326, 330

Harrier: none reported

Havanese: 10, 42, 140, 149, 166, 245, 256, 269, 273a

Ibizan Hound: 10, 14, 42, 70, 166, 270, 311a

Irish Red and White Setter: 37a, 166, 192, 312

Irish Setter: 7, 10, 22, 27, 31, 37a, 40, 42, 61, 65, 79, 81, 88, 103, 109, 121, 129, 130, 131, 132, 136a, 138a, 140, 146, 147, 152, 166, 168, 172, 186, 191, 192, 198, 206a, 220, 221, 221a, 225, 243, 245, 250, 256, 258a, 262, 264, 273a, 275, 276, 311a, 312, 320, 324, 328, 329

Irish Terrier: 75, 85a, 204a, 256

Irish Water Spaniel: 42, 152, 166, 167, 195, 245, 256, 330

Irish Wolfhound: 10, 31, 38, 42, 103, 131, 149, 152, 155, 158, 166, 221, 221a, 225, 250, 330

Italian Greyhound: 14, 42, 61, 65, 70, 109, 135, 143a, 166, 220, 243, 256, 311a

Japanese Spaniel (Japanese Chin): 42, 57, 70, 88, 103, 110, 137, 235, 256, 313

Keeshond: 1, 9b, 42, 63, 71, 85, 94, 109, 135, 149, 156, 165, 166, 180, 197, 201, 220, 253a, 260, 266, 274, 310, 327, 330

Kerry Blue Terrier: 42, 47, 88, 103, 124, 141, 146, 150, 166, 179, 181, 207, 256, 311a, 320, 330

Komondor: 42, 103, 152, 166, 285

Kuvasz: 42, 103, 152, 166, 221, 221a, 303a, 330

Labrador Retriever: 3, 7, 9, 9a, 10, 21, 22, 40, 42, 52a, 59, 68, 75, 77, 78, 85, 88, 89, 94, 95, 103, 109, 114a, 121, 122, 129a, 143a, 146, 147, 148, 149, 152, 158, 159a, 160, 166, 192, 193, 194b, 197, 204a, 206a, 221, 221a, 244, 245, 256, 257, 258, 269, 270, 276, 282, 304a, 312, 315, 330

Lakeland Terrier: 42, 70, 88, 166, 186, 245, 319, 320, 330

Lhasa Apso: 1, 10, 22, 42, 65, 81, 88, 94, 103, 140, 166, 171, 173, 179, 181, 189, 235, 256, 266, 266a, 273a, 275, 330

Leonberger: 9, 9a, 10, 21, 27, 152, 159a, 166, 192, 221, 221a, 312, 330

Lowchen: 42, 235, 256

Maltese: 1, 30, 57, 70, 78, 88, 110, 135, 140, 146, 149, 152, 160, 166, 192, 235, 256, 270, 273a, 311a, 312, 313, 330, 331a

Mastiff: 27, 31, 65, 80, 94, 103, 131, 166, 193, 221, 221a, 245, 256, 270, 325

Miniature Bull Terrier: 103, 166, 186

Miniature Dachshund: 4, 5, 42, 43a, 54, 55, 61, 65, 70, 72, 75, 78, 80, 85, 94a, 129, 146, 156, 161, 166, 173, 176, 178, 187, 196, 199, 206a, 214, 216, 224, 228, 230, 237, 239, 250, 256, 263, 267, 275, 296, 297, 311a, 326, 328, 330

Miniature Pinscher: 42, 65, 85, 87, 103, 128a, 140, 171, 179, 181, 185, 202b, 230, 235, 256, 273a, 299b

Miniature Poodle: 5, 10, 22, 26, 27, 42, 49, 57, 70, 78, 81, 85, 88, 92, 93, 103, 109, 110, 111, 121, 135, 136, 140, 144, 146, 147, 156, 165, 166, 173, 175, 184, 186, 192, 193a, 199, 206, 206a, 220, 221, 221a, 223, 226, 230, 235, 236, 250, 256, 269, 273a, 275, 294, 311a, 312, 313, 322a, 327, 330

Miniature Schnauzer: 10, 22, 24a, 42, 70, 71, 76, 85, 88, 103, 112, 121, 144a, 146, 147, 149, 149a, 156, 157, 157c, 166, 179, 181, 185, 192, 206a, 214, 221, 221a, 256, 259, 260, 266a, 270, 271, 284, 301, 311a, 312, 322a, 330

Neapolitan Mastiff: 42, 83, 94, 103, 152, 158, 166, 221, 245, 256

Newfoundland: 9a, 21, 25, 31, 38, 42, 75, 81, 83, 94, 95, 103, 114, 129a, 131, 146, 152, 166, 183, 192, 204a, 206, 221, 221a, 236, 239, 300, 308a, 311a, 312, 320, 327, 330

Norfolk Terrier: 109, 149, 166, 201, 304a

Norwegian Dunkerhound: 78, 199

Norwegian Elkhound: 42, 88, 103, 135, 152, 166, 180, 186, 256, 266, 275, 276, 302

Norwich Terrier: 65, 166, 186, 330

Nova Scotia Duck Tolling Retriever: 9a, 42, 159a, 166, 256

Old English Sheepdog: 9a, 27, 38, 42, 44a, 45, 47, 80, 88, 103, 122, 129, 140, 146, 148, 149, 152, 159a, 161, 166, 172, 176a, 192, 199, 221, 221a, 226, 250, 256, 269, 270, 273a, 283, 292, 311a, 312, 328, 330

Otter Hound: 119, 152, 166, 221, 221a, 249, 274, 311, 330

Papillon: 12, 42, 65, 78, 103, 128a, 166, 235, 256, 330

Parson Russell Terrier: 20, 78, 123, 186, 206, 235, 330

Pekingese: 42, 57, 88, 94, 103, 116, 128, 146, 162, 166, 171, 173, 179, 181, 184, 186, 199, 230, 246, 256, 277, 278, 304, 311a, 313, 317, 318

Pembroke Welsh Corgi: 27, 42, 50, 61a, 65, 72, 75, 82, 90, 109, 166, 173, 186, 193a, 206a, 214, 236, 245, 256, 270, 330

Petit Basset Griffon Vendeen: 21, 42, 65, 109, 166, 192, 245, 270, 312

Pharaoh Hound: 10, 166, 220, 311a

Pointer: 8, 10, 36, 42, 44a, 65, 80, 89, 103, 109, 149c, 152, 166, 178, 213, 214a, 215, 230, 231, 239, 242a, 256, 258a, 318, 330

Polish Lowland Sheepdog (PONS): 166, 192, 193a, 214, 312, 318

Pomeranian: 9b, 42, 70, 73, 87, 88, 103, 110, 136, 137, 140, 149, 162, 165, 166, 184, 186, 192, 210, 219, 235, 236, 256, 273a, 312, 313

Portuguese Water Dog: 9a, 38, 42, 88, 128a, 147, 159a, 166, 188a, 193a, 199, 245, 256, 273a, 299b

Pug: 9a, 10, 22, 50, 54, 57, 65, 67, 76, 80, 81, 85, 88, 90, 98, 98a, 103, 109, 116, 128, 143, 145, 149, 150, 152, 166, 173, 179, 181, 185, 195, 196, 230, 235, 246, 256, 259, 293, 295, 304a, 308, 309, 317

Puli: 27, 42, 152, 256, 270

Rhodesian Ridgeback: 42, 45, 51, 84, 103, 143a, 146, 152, 166, 190a, 192, 218a, 221, 221a, 245, 256, 312, 330

Rottweiler: 9a, 27, 42, 85, 88, 94, 95, 103, 105, 129, 129a, 146, 149c, 152, 159a, 161, 166, 172, 190a, 192, 193, 218a, 221, 221a, 225, 231, 245, 250b, 256, 258a, 269, 270, 300, 311a, 312, 326, 328, 330

Saint Bernard: 27, 31, 38, 42, 60, 72, 83, 88, 94, 94a, 103, 109, 114, 118, 121, 122, 128, 131, 133, 140, 147, 148, 149, 152, 155, 166, 188, 198, 221, 221a, 225, 262, 273a, 298, 325, 328, 329, 330

Saluki: 14, 27, 42, 65, 103, 146, 166, 214, 245, 256, 269, 270, 311a, 330

Samoyed: 9b, 24, 42, 65, 85, 88, 89, 103, 121, 135, 140, 146, 147, 149, 149b, 152, 166, 192, 204a, 221, 221a, 240, 242, 245, 256, 260, 269, 270, 273a, 274, 303a, 311a, 312, 328, 330

Schipperke: 42, 85, 88, 103, 166, 185, 202b, 207, 239, 245, 256

Scottish Deerhound: 31, 42, 105, 131, 158, 166, 221, 221a

Scottish Terrier: 5, 10, 22, 37, 42, 68, 75, 78, 80, 81, 122, 129, 140, 148, 149a, 166, 186, 193, 197, 206, 245, 256, 272, 273a, 294, 313a, 324, 330

Sealyham Terrier: 22, 42, 81, 135, 166, 186, 245, 256, 269, 270

Shetland Sheepdog: 5, 37, 42, 52, 52a, 58, 59, 65, 82, 86, 88, 108, 121, 122, 129, 147, 148, 149a, 151, 152, 157, 166, 176a, 192, 220, 236, 245, 256, 270, 306, 312, 313a, 328, 329, 330

Shiba Inu: 9, 10, 21, 22, 27, 42, 43, 115, 152, 157a, 166, 235, 324a, 329

Shih Tzu: 1, 42, 54, 55, 83, 88, 89, 94, 103, 110, 146, 149, 166, 173, 179, 181, 182, 187, 256, 266, 266a, 269, 311a, 317, 322a, 330

Shiloh Shepherd: 61c, 166, 229, 231a, 266, 290a

Siberian Husky: 41, 42, 65, 86, 103, 104, 121, 128a, 135, 147, 149, 152, 157c, 166, 184a, 186, 192, 221, 221a, 230, 245, 256, 270, 312, 324a, 328, 329, 330, 334

Silky Terrier: 42, 57, 70, 85, 154, 185, 188a, 193a, 235, 245, 256, 311a, 313

Skye Terrier: 27, 88, 101, 111, 163, 166, 183, 186, 192, 206, 312, 316, 330

Sloughi: 256

Smooth Fox Terrier: 3, 10, 22, 42, 78, 81, 87, 88, 112, 135, 138, 166, 185, 186, 206, 221, 221a, 243, 260, 288, 330

Soft Coated Wheaten Terrier: 9a, 10, 22, 42, 81, 159a, 166, 172, 220, 245, 253, 256, 258a, 258b, 266a, 270, 330

Spinone Italiano: 91, 103

Staffordshire Bull Terrier: 42, 109, 143a, 166

Standard Dachshund: 1, 4, 5, 8, 42, 43a, 54, 55, 61, 65, 70, 72, 75, 78, 80, 83, 85, 94a, 103, 115, 129, 135, 140, 146, 156, 161, 166, 173, 176, 179, 181, 187, 192, 196, 199, 214, 214a, 216, 220, 224, 228, 230, 237, 239, 242a, 245, 250, 256, 263, 267, 273a, 275, 296, 297, 311a, 312, 326, 328, 330

Standard Manchester Terrier: 42, 72, 109, 135, 166, 185, 186, 237, 256, 330

Standard Poodle: 9a, 10, 21, 22, 27, 31, 42, 61, 81, 88, 103, 109, 110, 121, 124a, 135, 140, 144, 146, 147, 152, 159a, 166, 175, 184, 186, 192, 199, 220, 221, 221a, 223, 230, 245, 256, 269, 273a, 311a, 312, 330

Standard Schnauzer: 9a, 23, 42, 62, 121, 147, 166, 192, 193, 221, 221a, 240, 260, 270, 286, 299a, 312, 330

Sussex Spaniel: 38, 42, 88, 103, 270

Swiss Mountain Dog: 221, 221a, 249, 311, 311a

Tibetan Mastiff: 27, 95, 152, 158, 166, 192, 221, 245, 312, 330

Tibetan Terrier: 14, 42, 78, 103, 157d, 186, 193a, 199, 214, 245, 250b, 256, 270

Tosu Inu: 10, 27, 157a, 166

Toy Fox Terrier: 80, 235, 185, 330

Toy Manchester Terrier: 42, 166, 186, 237, 256, 330

Toy Poodle: 5, 10, 22, 26, 27, 42, 49, 57, 70, 78, 81, 85, 88, 92, 93, 103, 109, 110, 111, 121, 124a, 135, 136, 140, 144, 146, 147, 149, 156, 165, 166, 173, 175, 184, 186, 193a, 199, 220, 221, 221a, 223, 226, 230, 235, 236, 256, 269, 273a, 275, 294, 311a, 313, 322a, 327, 330

Vizsla: 27, 42, 68, 80, 103, 109, 117, 121, 140, 143a, 147, 152, 161, 166, 193, 221, 221a, 256, 273a, 289, 296, 305, 318, 319, 326

Weimaraner: 27, 31, 43a, 61, 65, 80, 88, 103, 105, 114, 121, 131, 140, 142, 147, 150, 152, 158, 160a, 165, 166, 168, 169, 170, 178a, 196, 206, 250, 256, 273a, 289, 296, 305, 311b, 318, 319, 326

Welsh Springer Spaniel: 42, 135, 152, 245, 256

Welsh Terrier: 42, 135, 166, 186, 190a, 218a, 330

West Highland White Terrier: 2, 4, 9a, 10, 21, 22, 37, 42, 52a, 64, 68, 71, 80, 81, 106, 130, 136, 156, 159a, 171, 181, 185, 186, 199, 245, 250, 263, 270, 276, 313a, 331, 331a

Whippet: 42, 61, 70, 80, 103, 143a, 166, 186, 221, 221a, 234, 237, 256, 330

Wire Fox Terrier: 3, 22, 37, 42, 78, 87, 88, 103, 112, 135, 138, 166, 185, 186, 206, 243, 256, 260, 286, 288, 310, 313a, 330

Wirehaired Pointing Griffon: 152, 206a, 226

Yorkshire Terrier: 42, 57, 70, 71, 80, 88, 103, 149, 154, 160, 162, 166, 179, 181, 185, 199b, 235, 236, 245, 252a, 256, 269, 270, 276, 313, 330

Section II:

- 1. Aberrant cilia:** eyelashes growing abnormally, such as rubbing against the eyeball. (also see #88.)
- 2. Abnormal copper metabolism:** an inability to utilize and store copper properly, resulting in liver disease and other problems. * Bedlington Terrier or Doberman Pinscher
- 3. Abnormal dentition:** abnormal placement, number and development of teeth.
- 4. Acanthosis nigricans:** a skin disease where the skin becomes thickened and dark, primarily affecting the axillae (armpits). *Dachshund
- 5. Achondroplasia:** abnormal development of cartilage leading to dwarfism (seen aberrantly in most breeds, but what makes a Basset Hound and other achondroplastic breeds long and low).
- 6. Acne:** same thing as in humans, affects the muzzle and lip areas.
- 7. Acral lick dermatitis:** a skin disease caused by an animal's licking a localized area excessively, especially on the legs and paws.
- 8. Acral mutilation:** a progressive self-mutilation of the feet and legs. Also known as peripheral sensory neuropathy in pointing breeds, because they are born without pain sensation.
- 9. Acute moist dermatitis:** known as "hot spots," a localized area of a severely itchy, inflamed and oozing dermatitis exacerbated by the animal's intense licking and chewing at the spot.
- 9a. Addison's disease (hypoadrenocorticism):** a disease characterized by inadequate secretion of cortisone from the adrenal glands. *Old English Sheepdog, Standard Poodle and Bearded Collie (also see #159a.)
- 9b. Adrenal sex hormone dermatosis:** also called "alopecia-x," this disorder is caused by overproduction of adrenal sex hormones and results in patchy or generalized hair loss over the trunk. *Young male Pomeranian, Keeshond, and Samoyed, as well as other breeds
- 10. Allergies:** same as in humans. Dogs can be allergic to things they come in contact with, eat or inhale.
- 11. Amyloidosis:** a condition where abnormal deposits of proteinaceous material called amyloid are laid down in tissues and impair their function. * Akita and Chinese Shar-Pei.
- 12. Anasarca:** a condition where neonatal puppies have an abnormal accumulation of fluids in their tissues. *English Bulldog
- 13. Anemia with chondrodysplasia:** a condition of Alaskan Malamutes where there is malformation and maldevelopment of cartilage and red blood cells. This condition is also called stomatocytosis because of the mouth-like shape of red blood cells.
- 14. Anesthetic idiosyncrasy:** a condition where an individual has an abnormal response to commonly used anesthetics sometimes leading to death. Idiosyncratic means there is no good explanation or way to predict this.
- 15. Anomaly of third cervical vertebra:** a malformation of one of the neck bones in the spinal column.
- 16. Anophthalmia:** a condition where an animal is born without any eyes.

- 17. Aortic body tumors:** a cancer that arises from a small organ at the base of the aorta, near the heart.
- 18. Apocrine gland tumor:** a cancer arising from glands which secrete fluids (such as mammary glands).
- 19. Arteriovenous fistula:** an abnormal connection that forms between arteries and veins.
- 20. Ataxia:** incoordination associated with a variety of central and peripheral nervous system abnormalities, spinal cord instability or inner ear disorders. This condition is seen in many breeds and with episodes of spasmodic muscle activity in the Parson Russell Terrier.
- 21. Atopic dermatitis:** a skin disease caused by a dog's reaction to an inhalant allergy. (also see #81.)
- 22. Atopy:** an allergy caused from things dogs inhale.
- 23. Atresia of nasolacrimal puncta:** a condition where the openings on the inside of the lower eyelids (puncta) are too small or closed so tears spill over the lid instead of draining to the nose.
- 24. Atrial septal defects:** a malformation of the dividing wall between two chambers of the heart, usually resulting in a hole which then causes blood flow abnormalities.
- 24a. Avian tuberculosis:** an often fatal infection in immune deficient animals due to avian tuberculosis. *Basset Hound and Miniature Schnauzer
- 25. Avulsion fractures:** fractures of the bones caused by a piece being ripped away, usually because of its strong attachment to a ligament or tendon.
- 26. Basal cell tumor:** a cancer arising from a type of skin cell.
- 27. Behavioral abnormalities:** a whole range of abnormal behavior patterns, such as aggression, panic disorders, etc., obsessive compulsive disorders like "spinners" or tail chasing, and pica seen in the Bull Terrier.
- 28. Bithoracic ectromelia:** a condition where the front leg bones are very small or absent.
- 29. Blepharospasm:** an abnormal tightening of the muscles around the eyes, which causes repetitive blinking.
- 30. Blindness:** an inability to see due to a large variety of causes.
- 31. Bloat:** a condition where a dog's stomach produces excessive gas and enlarges severely enough to cause death without immediate treatment. Bloat is usually associated with gastric torsion. (also see #131.)
- 32. Blue eyes:** an adverse reaction to certain vaccines containing canine hepatitis virus (adenovirus 1) which produces a bluish discoloration to the cornea.
- 33. Bullous pemphigoid:** an autoimmune (i.e., the body attacking itself) disease associated with the formation of painful blisters.
- 34. Bundle branch block:** an abnormality of the electrical conduction mechanism of the heart.
- 35. Bundle of His degeneration:** a condition where a part of the electrical system of the heart deteriorates.
- 36. Calcinosis circumscripta:** the development of lumps of hard calcium deposits in the skin.

37. Cancer, bladder: transitional cell carcinoma (TCC) of the bladder is prevalent in the Scottish Terrier and West Highland White Terrier. Herbicide exposure increases the risk of TCC, whereas increased consumption of green leafy and yellow orange vegetables decreases the disease risk.

37a. Canine leukocyte adhesion deficiency (CLAD): a condition where the leukocytes fail to adhere normally, leading to recurrent infections. *Irish setter and Irish Red and White Setter

37b. Cardiac valvular disease: weakness of heart valves producing heart murmurs and leading to cardiac failure. *Cavalier King Charles Spaniel

38. Cardiomyopathy: a disease of weakened heart muscles, associated with taurine deficiency in the Golden Retriever. *giant breeds, Boxer, American Cocker Spaniel and Doberman Pinscher

38a. Carnitine deficiency: low concentrations of this quaternary amine are associated with reduced contraction and relaxation of cells, especially heart muscle, because it is involved in long-chain fatty acid transport within cells to generate metabolic energy. Deficiency of carnitine is thereby linked to dilated cardiomyopathy. *Boxer and American Cocker Spaniel (also see #38.)

39. Carotid body tumors: cancer arising from a small organ located on the carotid in the neck.

40. Carpal subluxation: a condition where the "wrist" bones are loose and out of alignment.

41. Castration responsive dermatosis: a skin condition characterized by loss of hair, thickened skin and inflammation which responds to castration (i.e., hormonally dependent).

42. Cataract: as in humans, a change in structure of the lens of the eye leading to cloudiness and usually to blindness.

43. Cataract with microphthalmia: a condition where a dog has cataracts along with abnormally small eyes.

43a. Cell-mediated immunodeficiency: a deficiency of T-lymphocyte function causing impaired immunity, chronic recurrent infections and stunted growth. * Weimaraner and Dachshund (also see #311b.)

44. Cellulitis (folliculitis and furunculosis): inflammation and infection of the cells of the skin including the hair follicles and deeper structures.

44a. Cerebellar ataxia: an x-chromosome-linked disorder of pointers. Affected puppies have episodes of dysmetria, nystagmus and disorientation, which progresses to marked ataxia by 16 months of age.

45. Cerebellar cortical abiotrophy: malformation of the neurons in the cerebellum, a part of the brain.

46. Cerebellar degeneration: a condition where a part of the brain deteriorates.

47. Cerebellar and extrapyramidal abiotrophy: a condition where the neurons in the cerebellum part of the brain and parts of the spinal cord are malformed and eventually malfunction.

48. Cerebellar hypoplasia: a condition where the cerebellum, a part of the brain, is poorly formed (too small or absent) and doesn't function properly or at all.

- 49. Cerebrospinal demyelination:** a condition where the neurons of the brain and spinal cord are malformed, lacking a specialized sheath, which causes malfunction.
- 50. Cervical disc disease:** a degeneration or malformation of the cushioning discs between the spinal column bones (vertebrae) in the neck.
- 51. Cervical vertebral malformation or instability:** a malformation of the vertebrae in the neck usually leading to nerve damage. This is commonly seen in the Doberman Pinscher and causes a hackneyed gait (high stepping).
- 52. Choroidal hypoplasia:** the technical name for collie eye anomaly (see #58), this autosomal recessive disorder causes improper development of the choroid vascular layer under the retina of the eye in young puppies. There is no treatment or cure.
*Collie, Border Collie, Shetland Sheepdog, Australian Shepherd and Lancashire Heeler.
- 52a. Chronic progressive hepatitis:** an uncommon disease sometimes called chronic active hepatitis. *Doberman Pinscher, Dalmatian, Labrador Retriever, Golden Retriever, and Shetland Sheepdog (also see #2, 62.)
- 53. Ciliary dyskinesia:** congenital trait in several breeds where all ciliated cells (those with hairs or moving appendages) are deformed and rigid. This is also called immotile cilia syndrome and Kartagener's syndrome, and causes chronic pneumonia and sterility.
*Bichon Frise and Doberman Pinscher
- 54. Cleft lip:** a condition where the two halves of the upper lip do not join together. Cleft palate and cleft lip are often seen together.
- 55. Cleft palate:** a condition where the roof of the mouth is not closed and the inside of the nose opens into the mouth.
- 56. Coliform enteritis:** an inflammation of the bowel tract caused by a certain group of bacteria.
- 57. Collapsed trachea:** a condition where the cartilage rings that make up the trachea are malformed and tend to collapse easily.
- 58. Collie eye anomaly:** detected by ophthalmologic examination between 5-8 weeks of age. This is seen worldwide in several breeds, as well as Rough and Smooth Collies. In mild disease, vision may not be impaired, but mildly affected dogs can produce severely affected offspring. In severe form, colobomas and retinal detachment can occur by about 2 years of age, although total blindness is rarely seen. Genetic testing from a blood sample can distinguish normal, carrier, and affected dogs.
- 59. Coloboma:** an abnormal development of the eye, usually seen in the Collie, which can lead to blindness. (See #58, 60, 203.)
- 60. Colobomas with aphakia:** same as above but with congenital absence of the lens.
- 60a. Color dilution alopecia:** a form of follicular dysplasia causing various degrees of hair loss in middle-aged blue and fawn Doberman Pinschers.
- 61. Color mutant alopecia:** a condition where certain pigmented areas of a dog's skin grow less or no fur. *Yorkshire Terrier and fawn Irish Setter
- 61a. Combined immunodeficiency:** a severe combined deficiency of cell-mediated immunity (T-cell function) and low concentrations of serum immunoglobulins (IgA, IgG, and variably IgM). Affected puppies usually die from viral infections by 12-16 weeks of age. *Basset Hound. (See #43a, 168, 169, 170.)

61b. Complement deficiency: a deficiency in serum concentrations of the third component of complement which impairs neutrophil function and causes recurrent infections. *Brittany Spaniel

61c. Compressive myelopathy: a condition of Shiloh Shepherds with malformed spinal vertebral processes which compress the spinal cord and cause spastic paraparesis, acute progressive pelvic limb weakness and incoordination (ataxia). (also see #290a.)

61d. Congenital stationary night blindness: a condition at birth where affected pups cannot see at night. This is seen in the Briard

62. Conjunctivitis: an inflammation of the conjunctival membrane of the eye.

63. Conus septal defect: a developmental abnormality of the right ventricle of the heart.

64. Copper storage abnormality in liver: (also see #2.)

65. Corneal dystrophy: an abnormality of the cornea usually characterized by shallow pits in the surface.

66. Corneal leukomas: an abnormal accumulation of a white material in or on the cornea.

67. Corneal ulcer, superficial: an erosion of the outer membrane and outer surface of the cornea.

68. Craniomandibular osteopathy: an abnormal development of the bones of the face and the jaw. *West Highland White and Cairn Terrier, among other breeds.

69. Cranioschisis: abnormal development of the skull characterized by openings between or in the bones.

69a. Cricopharyngeal dysfunction: delayed swallowing times causing dysphagia (also see #89a) observed on videofluoscopy. *Golden Retriever

70. Cryptorchidism: a condition where one testicle does not descend into the scrotal sac.

71. Cushing's disease (hyperadrenocorticism): a disease characterized by an excess secretion of corticosteroids from the adrenal glands. (also see #156.)

72. Cutaneous asthenia: a condition where the skin lacks its normal strength, elasticity and sensation. Also called Ehlers-Danlos syndrome. *English Springer Spaniel and Boxer (also see #94a.)

72a. Cutaneous mucinosis: (See #202a.)

72b. Cutaneous vasculopathy: a disease of autosomal recessive inheritance in which young puppies exhibit footpad swelling and depigmentation, with crusting and ulceration of the tips of ears and tail. *German Shepherd

73. Cyclic neutropenia: a condition characterized by periodic lowering of neutrophils, a type of white blood cell. *Grey Collie

74. Cystic ovaries: a condition where the ovarian follicles become cystic (fluid filled) leading to hormonal imbalances and other problems.

75. Cystinuria: an abnormal excretion of a substance (cystine) in the urine.

76. Cystitis and cystic calculi: infection of the bladder which often leads to formation of abnormal mineral deposits (bladder stones).

77. Dacryocystitis: inflammation of a tear sac.

78. Deafness: an inability to hear, due to many different causes.

79. Deformed tail: a congenital condition where the tail is malformed.

- 79a. Degenerative myelopathy:** progressive disorder primarily in German Shepherd dogs where the spinal cord degenerates and causes rear limb weakness and incoordination. Can end up with cauda equine syndrome, where the nerve roots at the end of the spinal column become atrophied.
- 80. Demodicosis:** a kind of skin disease (mange) caused by microscopic *Demodex canis* mites living within the skin layers and producing an immunodeficiency syndrome.
- 81. Dermatitis, atopic:** inflammation and subsequent infection of the skin due to atopy. (also see #21, 22.)
- 82. Dermatomyositis:** a disease affecting the skin and muscles, usually in the Collie or Shetland Sheepdog
- 83. Dermoid cyst:** a small growth composed of skin-like structures.
- 84. Dermoid sinus:** similar to a dermoid cyst but usually larger. *Rhodesian Ridgeback
- 85. Diabetes mellitus:** a metabolic disease caused by insulin deficiency and characterized by the inability to utilize sugars normally.
- 85a. Digital hyperkeratosis:** a condition of Irish Terrier puppies that causes marked thickening of the foot pads. Affected feet crack, become infected and painful.
- 86. Discoid lupus erythematosus:** a form of autoimmune disease affecting the skin.
- 87. Dislocation of shoulder:** a condition where the bones of the shoulder joint are out of proper position.
- 88. Distichiasis:** abnormally growing eyelashes.
- 89. Dwarfism:** an abnormality of the normal growth pattern resulting in an undersized individual.
- 89a. Dysphagia:** inability to swallow properly. (also see #69a.)
- 90. Dystocia:** complications of the birth process (difficult birth).
- 91. Eclampsia:** convulsions usually seen around the time of parturition (whelping).
- 92. Ectodermal defects:** any of a multitude of abnormalities arising from maldevelopment of the fetal ectoderm (e.g., skin, nervous system, eyes).
- 93. Ectopic ureters:** the ureters (tubes leading from the kidneys to the bladder) do not empty into the bladder in the normal location.
- 94. Ectropion:** an abnormal rolling out of the eyelids.
- 94a. Ehlers-Danlos syndrome:** a connective tissue disease characterized by loose, hyperextensible and very fragile skin that tears easily. (also see #72.)
- 95. Elbow dysplasia:** an abnormal development of the elbow joint.
- 96. Elbow joint malformation:** (See #95.)
- 97. Elbow subluxation:** a condition where the elbow joint is loose and out of alignment.
- 98. Elongated soft palate:** the soft palate is abnormally long and causes breathing disorders.
- 98a. Encephalitis:** an inflammatory condition of the brain causing signs of central nervous system dysfunction and epilepsy (seizures). A unique form of encephalitis is prevalent in the Pug and is called "Pug Dog Encephalitis." (also see #109.)
- 99. Endocardial fibroelastosis:** an abnormal condition of scarring of the muscles of the heart.
- 100. Endometritis:** inflammation of the internal layer of the uterus.
- 101. Enlarged foramen magnum:** a condition in which the opening in the skull where the vertebral column begins is too large.
- 102. Enostosis:** a bony growth within the hollow part of a bone.

- 103. Entropion:** an abnormal rolling in of the eyelid.
- 104. Eosinophilic granuloma:** an allergic reactive syndrome characterized by the plaque-like accumulation of eosinophils, a type of white blood cell.
- 105. Eosinophilic panosteitis:** a painful inflammatory bone disease of young, rapidly growing dogs, often characterized by increased eosinophils in the blood. (also see #231.)
- 106. Epidermal dysplasia:** abnormal development of the outer layer of the skin. Common in the West Highland White Terrier and begins in as a puppy. (also see #331.)
- 107. Epidermoid cyst:** a small growth consisting of tissues of the outer layer of the skin. (also see #274.)
- 108. Epidermolysis bullosa:** an abnormal looseness to the skin characterized by large, deep, blister-like lesions.
- 109. Epilepsy:** a disease characterized by convulsions (seizures) and/or disturbances of consciousness.
- 110. Epiphora:** abnormal draining of tears often due to overproduction.
- 111. Epiphyseal dysplasia:** abnormal development of the epiphysis, a part of the long bones.
- 112. Esophageal achalasia:** a functional stricture or spasm of the muscles of the esophagus where it joins the stomach.
- 113. Esophageal dilatation:** an abnormally large and usually flaccid esophagus.
- 114. Eversion of nictitating membrane:** a condition where the third eyelid is protruding.
- 114a. Exercise-induced collapse:** seen in Cavalier King Charles Spaniel puppies associated with exercise- or excitement-induced muscle hypertonicity (similar to startle disease in people). *Also seen in young adult field trial Labrador Retrievers.
- 115. Eye abnormality:** any of a number of problems with the eye.
- 116. Facial fold dermatitis:** an infection of the facial skin caused by unusual or excessive skin folds (seen in dogs such as the Pekingese or Chinese Shar-Pei).
- 117. Facial nerve paralysis:** a decrease or cessation of function of the facial nerve leading to a drooping of the affected side of the face.
- 118. Factor I deficiency or hypofibrinogenemia:** a rare deficiency of a clotting factor (fibrinogen), which causes excessive bleeding.
- 119. Factor II deficiency or hypoprothrombinemia:** a rare deficiency of prothrombin, a clotting factor needed to control bleeding.
- 120. Factor VII deficiency:** a mild bleeding disease primarily of Beagle dogs.
- 121. Factor VIII deficiency or hemophilia A:** the most common severe inherited clotting disorder of humans and animals. The disorder is inherited as a sex-linked recessive trait (carried by females and manifested in males). Affects most dog breeds.
- 122. Factor IX deficiency or hemophilia B:** same as hemophilia A, but rarer and involves a different clotting factor.
- 123. Factor X deficiency:** a rare clotting disorder primarily of the American Cocker Spaniel and an autosomal trait (affects both sexes).
- 124. Factor XI deficiency:** a rare clotting disorder; protracted bleeding from surgical procedures is a feature. Both sexes are affected.

124a. Factor XII deficiency: a clotting factor deficiency that rarely produces clinical signs, which is usually diagnosed incidentally during blood testing for potential bleeding disorder. *Toy and Standard poodle, and occasionally in other breeds

124b. Familial amaurotic idiocy: deposits of fatty pigments in the brain produce loss of vision, stupor, and seizures. *English Setter, German Shorthaired Pointer and Australian Cattle Dog
(also see #177, 193a, 214.)

124c. Fanconi syndrome: a kidney tubular dysfunction of the Basenji which leads to glycosuria. (also see #268.)

124d. Femoral artery occlusion: a rather common disorder of genetic predisposition and probable weakness in the femoral artery wall of the Cavalier King Charles Spaniel

125. Fibrosarcoma: a cancer arising from certain types of fibrous cells.

126. Fibrous histiocytoma: a type of fibrous tumor arising from cells called histiocytes.

127. Flank sucking: a behavioral problem common in the Doberman Pinscher and exhibited as a continually wet patch on the flank (from sucking the skin).

128. Fold dermatitis: an inflammation of skin folds especially in dogs with loose skin (e.g., Chinese Shar-Pei).

128a. Follicular dysplasia: malformation of skin follicles.

129. Folliculitis: an infection of the hair follicles.

129a. Fragmented coronoid process: osteochondrosis of the elbow joint. (also see #221a.)

129b. Fucosidosis: an autosomal recessive, fatal disease caused by deficiency of alpha-fucosidase enzyme resulting in accumulation of fucose-containing metabolites in cells throughout the body. Neurologic signs predominate. *English Springer Spaniel (also see #193a.)

130. Furunculosis: an infection of the deeper structures of the skin.

131. Gastric torsion: a condition where the stomach twists, thereby impeding input and output. (also see #31.)

132. Generalized myopathy: a condition affecting all the muscles of the body which produces weakness.

133. Genu valgum: malformation of the knee joint ("knock-kneed").

134. Gingival hyperplasia: overgrowth of the gum tissues.

135. Glaucoma: abnormally high pressure in the eye.

136. Globoid cell leukodystrophy: abnormal development and/or function of certain types of white globoid cells in the brain. (also see #193a.)

136a. Gluten-sensitive enteropathy: also called wheat-sensitive enteropathy of the Irish Setter. Intolerance to foods containing glutes, affected dogs primarily have chronic diarrhea and weight loss. (also see #258a.)

137. Glycogen storage disease: characterized by an inability to store and utilize carbohydrates. (also see #193a.)

138. Goiter: a swelling of the thyroid gland.

138a. Granulocyte dysfunction or adhesion defect: an impairment of neutrophil function or adhesion which causes chronic recurring infections, stunted growth and secondary increase in immune globulins (hypergammaglobulinemia). *Irish Setter and Doberman Pinscher

- 139. Granulomatous colitis:** a type of chronic inflammation of the colon characterized by reactive tissue growths.
- 140. Granulomatous sebaceous adenitis:** a disease of sebaceous (sweat) skin glands characterized by reactive tissue growth and autoimmune destruction of the sebaceous glands. Hair loss occurs and is poorly responsive to treatment. *Standard Poodle, Akita, Samoyed and Vizsla
- 141. Hair follicle tumors:** abnormal growths of the hair follicles.
- 142. Hairlessness:** also called alopecia or loss of hair.
- 143. Hanging tongue:** a syndrome where the tongue does not retract into the mouth properly, due to neurologic or anatomic defects. *Cavalier King Charles Spaniel
- 143a. Hemangiosarcoma:** a cancer of blood vessels involving liver, spleen or skin.
- 144. Hemeralopia:** inability to see in daylight.
- 144a. Hemorrhagic gastroenteritis:** an acute disorder characterized by bloody diarrhea, elevated hematocrit and shock. *Miniature Schnauzer
- 145. Hemivertebra:** a particular kind of malformation of the vertebra where only half of the structure is formed.
- 146. Hemolytic anemia:** anemia caused by the destruction of the red blood cells by an autoimmune process. *Cocker Spaniel and Old English Sheepdogs, as well as several other breeds.
- 147. Hemophilia A:** a blood clotting disorder due to deficiency of coagulation factor VIII (this is the most common type of hemophilia in dogs). (also see #121.)
- 148. Hemophilia B:** a blood clotting disorder due to lack of coagulation factor IX. (also see #122.)
- 149. Hepatic portosystemic shunt or arteriovenous fistula:** a malformation of blood vessels in the liver or an abnormal communication between the arteries and veins in the liver.
- 149a. Hepatic lipidosis:** an abnormal accumulation of lipids in the liver which leads to liver failure. *Miniature Schnauzer and Shetland Sheepdog
- 149b. Hereditary nephritis:** also called "Samoyed hereditary glomerulopathy," a sex-linked disease of young males. Affected dogs have renal glomerular disease which rapidly progresses to kidney failure and death. Female carriers have abnormal glomerular basement membrane, as well, but usually remain healthy until later in life when renal failure may occur.
- 149c. Hereditary spinal muscular atrophy:** an autosomal dominant degenerative disease of motor neurons characterized by weakness and muscle atrophy with a typical gait, and progressing to dangling of the head and a drooping, paralyzed tail. Severely affected dogs become paralyzed and die by 3-4 months of age. *Brittany Spaniel
- 149d. Hepatocerebellar degeneration:** a syndrome of progressive cerebellar and hepatic disease of 6-8-week-old Bernese Mountain Dogs with lesions of cerebellar abiotrophy and coexistent hepatic lesions. Autosomal recessive inheritance.
- 150. Hermaphroditism:** a syndrome where the individual has anatomical features of both sexes.
- 151. Heterochromia, iris:** the presence of different colors in the same or both irises.
- 152. Hip dysplasia:** a developmental malformation or subluxation of the hip joints.
- 153. Histiocytoma:** a common benign tumor of certain skin tissue cells (i.e., histiocytes).

- 153a. Histiocytosis:** the most prevalent cancer of Bernese Mountain Dogs, usually leading to early death.
- 154. Hydrocephalus:** a condition where there is an abnormal accumulation of fluid in the ventricles of the brain.
- 155. Hygroma:** a fluid-filled sac usually occurring on the elbows of large breed dogs such as the Great Dane or Irish Wolfhound.
- 156. Hyperadrenocorticism or Cushing's disease:** a disease where the adrenal glands are overactive. (also see #71.)
- 157. Hypercholesterolemia:** a disease where the animal has too much cholesterol in the blood system. This is commonly associated with hypothyroidism.
- 157a. Hyperkalemia:** a benign condition of large Japanese dog breeds (Akita, Shiba Inu, Tosu Inu) in which the red blood cell membrane has an altered metabolism and leaks potassium into the serum making concentrations very high. The condition is aggravated by ingesting onions.
- 157b. Hyperlipidemia:** an idiopathic disorder of the Miniature Schnauzer in which blood lipid levels become very high and predispose to pancreatitis. (also see #149a.)
- 157c. Hyperphosphatasemia:** a benign familial condition in humans and the Siberian Husky in which serum alkaline phosphatase concentrations are very high.
- 157d. Hypertrophic neuropathy:** a form of polyneuropathy seen in the Tibetan Terrier (also see #250a.)
- 158. Hypertrophic osteodystrophy:** a condition of rapidly growing giant breeds where there is an abnormal inflammation of bones with pain and development of excessive bony growths.
- 159. Hypertrophy of membrana nictitans gland:** a condition where the gland of the third eyelid is abnormally large.
- 159a. Hypoadrenocorticism:** a disease where autoimmune or other causes of destruction of the adrenal glands produces a deficiency of corticosteroids. (also see #9a.)
- 160. Hypoglycemia:** a syndrome where the animal has abnormally low blood glucose levels.
- 160a. Hypomyelinogenesis:** failure of the nervous system to form myelin, seen at birth.
- 161. Hypopigmentation, lips and nose:** a condition where an animal lacks pigment (color) in areas where it is usually present. (also see #328.)
- 162. Hypoplasia of dens:** a condition where part of the second vertebra fails to develop fully and leads to instability.
- 163. Hypoplasia of larynx:** a condition where the larynx (cartilage of the "voice box") fails to develop fully.
- 164. Hypoplasia of trachea:** a trachea that fails to develop fully.
- 165. Hyposomatotropism:** failure of the body growth hormones (somatomedins) to develop fully. This is also known as growth hormone responsive dermatosis. *Pomeranian
- 166. Hypothyroidism:** a common endocrine disease where the body produces an abnormally low amount of thyroid hormones. This is an autoimmune destruction of the thyroid gland which affects more than 50 dog breeds. (also see #192, 312.)

- 167. Hypotrichosis:** a condition where there is an abnormally small amount of hair growth.
- 168. Immunoglobulin A deficiency:** a condition where concentrations of secretory immune globulins are low. *Chinese Shar-Pei and Beagle (also see #187.)
- 169. Immunoglobulin G deficiency:** a condition where circulating antibody concentrations are low, which produces immune deficiency and susceptibility to infections.
- 170. Immunoglobulin M deficiency:** a condition where antibodies produced in early stages of an immune response are low, producing susceptibility to infection. *Doberman pinscher
- 171. Inguinal hernia:** a break in the muscular layer of the body wall occurring at the inguinal canal (where the back leg meets the body).
- 171a. Inherited ventricular tachycardia:** a condition of young German Shepherds with very rapid heart rates, ventricular arrhythmias, and sudden death.
- 172. Intestinal malabsorption:** a disease where the intestinal tract does not absorb nutrients properly. This is also known as protein-losing enteropathy as a consequence of inflammatory bowel disease. In Irish Setters, there is also a wheat-sensitive enteropathy.
- 173. Intervertebral disc disease:** a disease where the discs between the vertebrae are abnormal and prone to rupture and misplacement.
- 174. Intussusception:** a serious condition where the intestinal tract telescopes in on itself.
- 175. Iris atrophy:** a condition where the iris (the colored part of the eye) shrinks and becomes non-functional.
- 176. Iris heterochromia:** a condition where one iris is a different color from the other or has more than one color to it.
- 176a. Ivermectin sensitivity:** a prevalent condition of the Collie where a mutation of the MDRI gene produces susceptibility to ivermectin toxicity.
- 177. Juvenile amaurotic idiocy:** a syndrome characterized by early onset blindness and low mental capacity.
- 178. Juvenile cellulitis:** an inflammation of cells (usually skin cells) occurring in the young animal.
- 178a. Juvenile polyarthritis:** a form of arthritis affecting multiple joints of young Akitas, and usually occurring within 1-4 weeks of vaccination. Cases typically occur at 3-4 months of age after the second or subsequent booster vaccination and respond poorly to therapy. Affected dogs may progress to develop amyloidosis (#11) and renal failure.
- 179. Keratitis sicca:** a condition where one or both eyes do not produce a normal amount or type of tears.
- 180. Keratoacanthoma:** a small growth, usually on the face, filled with keratin material.
- 181. Keratoconjunctivitis sicca:** (See #179.)
- 182. Kidney aplasia, unilateral:** a developmental abnormality where one kidney fails to develop, also called renal agenesis.
- 183. Kinked tail:** a developmental abnormality where the tail has a pronounced kink.
- 184. Lacrimal duct atresia:** a condition where the duct draining tears from the eye is too small or not formed.

184a. Laryngeal paralysis: a progressive paralysis of the larynx of young Bouvier des Flandres, Siberian Huskies, and Dalmatians. In the Dalmatian, the condition is usually linked to polyneuropathy (#250a). Affected dogs have an unusual bark and are prone to aspiration pneumonia.

185. Legg-Perthes disease: disease in which the blood vessels feeding the femoral head (top part of the thigh bone) shrink, leading to starvation and death of the femoral head (the ball of the ball-and-socket joint of the hip). Also called Legg-Calve'-Perthes disease.

186. Lens luxation: a condition where the lens in the eye is displaced into an abnormal position.

187. Linear IgA dermatosis: a type of skin disease resulting from an abnormality of the secretory immune system. *Chinese Shar-Pei

188. Lip fold dermatitis: a skin infection caused by redundant skin folds around the mouth.

188a. Lipidosis: a form of lysosomal 'storage' disease where lipids accumulate in nerves. Called GM-1 gangliosidosis in the Portuguese Water Dog (also see #193a.)

189. Lissencephaly: an abnormal brain development where the surface lacks gyri (the grooves).

190. Lung torsion: a condition where one or more lung lobes twist upon themselves.

190a. Lupoid onchodystrophy: (also see #218a.)

191. Lymphedema: a disorder where valvular blockage of lymph flow or twisted lymphatic ducts causes an accumulation of fluid to swell tissues with edema.

192. Lymphocytic thyroiditis: an autoimmune disease causing inflammation and destruction of the thyroid gland, which becomes infiltrated with lymphocytes (white blood cells) and leads to hypothyroidism. This is the most common endocrine disease of the dog and has an inherited predisposition. (also see #166, 312.)

193. Lymphosarcoma: a cancerous condition involving the lymphatic system. This is one of the more common canine cancers.

193a. Lysosomal 'storage' diseases: a group of progressive multifocal neurologic disorders caused by specific enzyme deficiencies leading to death of nerve cells and accumulation of their respective enzyme substrates in cells. (also see #299a.)

194. Malabsorption syndrome: (See #172.)

194a. Malignant histiocytosis: the more aggressive, rapidly fatal systemic form of histiocytosis. Heritable in the Bernese Mountain Dog, with no cure. (also see #153a.)

194b. Malignant hyperthermia: an autosomal dominant trait of the black Labrador Retriever. Very high body temperatures develop in response to gaseous anesthesia.

195. Malocclusion: a condition where the teeth do not meet properly.

196. Mastocytoma: a rare cancer developing from a type of tissue cell known as a mast cell.

197. Melanoma: a rare cancer developing from the type of skin cell which produces pigment (melanin).

197a. "Merle" eye anomaly: (also see #52, 58.) The breeding of two merle colored parents can produce some offspring with whiter coat color. These puppies typically inherit a variety of anomalies of the back (fundus) of the eye, which can be confused with choroidal hypoplasia.

- 198. Metabolic bone disease:** any of a number of diseases affecting the bones due to an abnormality of metabolism.
- 199. Microphthalmia:** a condition where one or both eyes are too small.
- 199a. Microvascular dysplasia:** (See #252a.)
- 200. Missing teeth:** a condition where there are too few teeth.
- 201. Mitral valve defects:** a group of abnormalities of the mitral valve of the heart.
- 202. Mononephrosis:** a condition where only one kidney is present.
- 202a. Mucinosis:** a common skin disorder of the Chinese Shar-Pei characterized by generalized pitting edematous folds, variable itching, and severe puffiness and wrinkling of the head and extremities. Vesicles may be present and rupture draining clear, stringy fluid. This is often associated with hypothyroidism and IgA deficiency. (also see #166, 168.)
- 202b. Mucopolysaccharidosis:** an inborn metabolic error of several types leading to storage disease and debilitation. *Schipperke (type IIIb or Sanfilippo syndrome) *Miniature Pinscher (type VI) *German Shepherd (type VII). (also see #299a.)
- 203. Multiple colobomas:** a developmental abnormality of the structures of the eye.
- 204. Multiple epiphyseal dysplasia:** a condition where many of the long bones develop abnormally due to changes in the growth plates.
- 204a. Muscular dystrophy:** a congenital and often inherited form of generalized muscle dysfunction which causes signs such as poor growth, weakness, abnormal gait, difficulty eating and swallowing, and muscle atrophy. Affected animals have serious health problems and may die or be euthanatized. Inheritance is sex-linked in the Golden Retriever, Irish Terrier, Samoyed and Belgian Shepherd.
- 205. Muzzle pyoderma:** an infectious skin disease on the muzzle of an animal.
- 206. Myasthenia gravis:** a syndrome characterized by muscle fatigue due to an autoimmune disease which produces chemical abnormalities of the muscles and nerves. An enlarged esophagus called megaesophagus can result and causes regurgitation of food.
- 206a. Myotonia congenita:** a condition present at birth characterized by tonic muscle contractions and twitching. *Miniature Schnauzer
- 206b. Narcolepsy:** a neurological disorder characterized by falling asleep suddenly (collapse) which can occur during periods of activity and last for various lengths of time. *Doberman Pinscher and Labrador Retriever
- 207. Narrow palpebral fissure:** an abnormally small opening between the upper and lower eyelids.
- 208. Nasal pyoderma:** a skin infection of the nose.
- 209. Nasal solar dermatitis:** a skin disease of the nose and muzzle which is greatly affected by exposure to sunlight. *Collie
- 210. Nasolacrimal puncta atresia:** (See #23.)
- 211. Necrotizing myelopathy:** a condition where the spinal cord gradually dies.
- 212. Necrotizing panotitis:** a severe infection of the ear and surrounding tissues.
- 213. Neuromuscular atrophy:** a condition where the muscles waste away due to lack of proper nerve supply.
- 214. Neuronal ceroid lipofuscinosis:** a congenital disease where fatty pigments are deposited in the brain and cause brain dysfunction. (also see #193a.)

- 215. Neurotropic osteopathy:** a disease of the bones due to abnormalities of the nerves.
- 216. Nodular panniculitis:** a skin disease characterized by nodules of inflammation under the skin.
- 217. Oligodendroglioma:** a cancer arising from a type of cell found in the brain and spinal cord.
- 218. Oligodontia:** an abnormally small number of teeth.
- 218a. Onchodystrophy:** painful symmetrical nail bed disorder causing the nails to fall off; cause unknown. *Greyhound, Rottweilers and several other breeds. (also see #190a.)
- 219. Open fontanel:** a condition where the suture lines between bones of the skull do not fuse together properly.
- 220. Optic nerve hypoplasia:** a condition where the optic nerve going from the eye to the brain is too small.
- 221. Osteochondritis dissecans:** a specific form of inflammation of the cartilage of certain joints which causes arthritis. (also see #221a.)
- 221a. Osteochondrosis:** a group of developmental diseases resulting in abnormal formation of joint cartilage. Commonly involves the shoulder, stifle, hock or elbow. (also see #221.)
- 222. Osteodystrophy:** any of a number of diseases involving the development of the bones.
- 223. Osteogenesis imperfecta:** imperfect development of the structure and/or mineralization of the bones.
- 224. Osteopetrosis:** a condition where the bones are abnormally dense and hard.
- 225. Osteosarcoma:** a cancer arising from the cells of the bones.
- 226. Otitis externa:** an infection of the external structures of the ear.
- 227. Otocephalic syndrome:** a developmental abnormality where the animal lacks a lower jaw and the ears meet below the face.
- 228. Overshot jaw:** a condition where the upper jaw is too long for the lower jaw.
- 229. Pancreatic insufficiency:** a condition where the pancreas does not produce the proper enzymes for digesting food.
- 230. Pannus:** an immunologic eye disease characterized by abnormal growth of tissue over the cornea.
- 231. Panosteitis:** (See #105.)
- 232. Parosteitis:** inflammation of tissue around a bone.
- 233. Parotitis:** inflammation of the parotid salivary gland, also called parotiditis.
- 234. Partial alopecia:** some loss of the normal hair coat.
- 235. Patella luxation:** a condition where the knee caps slide in and out of place.
- 236. Patent ductus arteriosus:** failure of the vessel remnant joining the aorta and pulmonary artery in fetal life to close properly at birth, thereby shunting blood away from the lungs.
- 237. Pattern alopecia or baldness:** hair loss occurring in certain patterns. *Dachshund
- 238. Pemphigus erythematous:** one of many skin diseases caused by an autoimmune mechanism.
- 239. Pemphigus foliaceus:** another skin disease caused by autoimmune destruction of tissues.

- 240. Perianal adenoma:** a cancer arising from a cell of a gland found near the anus.
- 241. Perianal fistulas:** a condition characterized by abnormal communications from deeper tissues to the skin surrounding the anus.
- 242. Perianal gland tumor:** (See #240.)
- 242a. Peripheral sensory neuropathy:** a disorder of young puppies born without pain sensation of peripheral tissues. (also see #8.)
- 243. Persistent right aortic arch:** a developmental abnormality where one of the fetal blood vessels near the heart does not atrophy as it should.
- 244. Persistent hyaloid artery:** as #243, however, involving a blood vessel inside the eye.
- 245. Persistent pupillary membrane:** a developmental abnormality where the membrane forming the iris does not form properly.
- 245a. Phosphofruktokinase deficiency:** a deficiency of a specific red blood cell enzyme in the English Springer Spaniel. Causes chronic anemia, exercise-induced acute hemolytic crises and enlarged spleen.
- 245b. Physiologic leukopenia:** a condition of most healthy adult Belgian Terverun whereby the total white blood cell count (neutrophils, lymphocytes, and monocytes) is below 6,000/ul. Because of their ancestral relationship to the other Belgian Shepherd dogs (Belgian Sheepdog and Belgian Malinois), these breed/varieties may also show the same phenomenon.
- 246. Pigmentary keratitis:** an inflammatory condition of the cornea characterized by abnormal pigmentation.
- 247. Pituitary dwarfism:** a developmental abnormality resulting in an undersized animal due to a defective pituitary gland.
- 248. Pituitary tumor:** a cancer arising from the pituitary gland.
- 249. Platelet disorder:** a group of abnormalities of small blood cells necessary to control bleeding. (also see #311, 311a.)
- 250. Pododermatitis:** a skin infection of the paws.
- 250a. Polycystic kidney disease:** malformation of kidneys where the renal pelvis is cystic. This disease may be associated with heart valvular disease in the Bull Terrier.
- 250ba. Polyneuropathy:** a progressive polyneuropathy of young dogs leading to neuromuscular atrophy, variable demyelination, paraparesis, exercise intolerance and hyperesthesia. *Alaskan Malamute, Dalmatian, Golden Retriever, Rottweiler and German Shepherd (also see #213.)
- 251. Polyostotic fibrous dysplasia:** a type of bone disease where the bones are composed of improper fibrous tissues.
- 252. Polyradiculoneuritis:** an acute inflammatory disease of several groups of nerves causing fever.
- 252a. Portosystemic shunt:** a congenital anomaly of blood vessels supplying the liver, causing varying degrees of liver dysfunction or failure. This condition can also be manifested as microvascular dysplasia. *Yorkshire and Cairn Terrier, but can occur in any breed. (also see #149, 199a.)
- 253. Posterior retinal atrophy:** a deterioration of the part of the eye which translates light to electric impulses (the retina) and produces night blindness. (also see #256.)

253a. Primary hyperparathyroidism: overactive production of parathyroid hormone, most often caused by parathyroid gland tumor, and leading to renal failure from secondary hypercalcemia. *Keeshond

254. Primary peripheral retinal dystrophy: a certain type of developmental disease affecting the retina.

255. Progressive ataxia: a condition where the animal's sense of coordination deteriorates.

256. Progressive retinal atrophy: a disease where the retina slowly deteriorates, producing night blindness.

257. Prolapsed rectum: a condition where the inside of the rectum protrudes outside the anus.

258. Prolapsed uterus: a condition where the uterus protrudes into the vaginal canal or through the vaginal opening.

258a. Protein-losing enteropathy: a relatively common genetically predisposed condition also called inflammatory bowel disease. Vomiting, diarrhea, and weight loss are the common signs. *Seen in many breeds, but mostly in Soft Coated Wheaten Terrier (in conjunction with #258b), Rottweiler, Bernese Mountain Dog, German Shepherd, Golden Retriever, Dalmatian, Akita, Irish Setter and English Setter

258b. Protein-losing nephropathy: a condition where protein is lost through the kidney. Affected dogs have excessive thirst and urination, which progresses to peripheral edema and renal failure. *Soft Coated Wheaten Terrier (in conjunction with #258a).

259. Pseudohermaphroditism (pseudohermaphroditism): a condition where the animal has the gonads of one sex, but the appearance is ambiguous or is of the opposite sex.

260. Pulmonic stenosis: a condition where one of the valves of the heart does not open properly.

261. Pyloric stenosis: a condition where the opening leading from the stomach does not function properly.

262. Pyometra: a bacterial infection of the uterus where it fills with pus.

263. Pyruvate kinase deficiency: a deficiency of a specific red blood cell enzyme. *Basenji, Beagle and Cairn Terrier

264. Quadriplegia with amblyopia: a syndrome characterized by weakness of all four limbs, as well as of vision.

264a. Rage syndrome: sudden unprovoked aggression of serious nature. *English Springer Spaniel (also see #27.)

265. Recessive retinal dysplasia: a developmental disorder resulting in an abnormal retina, carried by a recessive gene.

266. Renal cortical hypoplasia: a condition where the cortex of the kidney(s) develops incompletely.

266a. Renal dysplasia: a condition where the kidneys form abnormally. Renal failure develops with protein loss in urine.

267. Renal hypoplasia: a condition where the kidney(s) do not develop completely.

268. Renal tubular dysfunction: a condition where the tubules of the kidneys (the filtering structures) do not function properly. (also see #124c.) *In the Basenji, glycosuria develops and is called Fanconi syndrome.

- 269. Retinal detachment:** where the retina is unattached to the back of the eye.
- 270. Retinal dysplasia:** a condition where the retina is malformed.
- 271. Schnauzer comedo syndrome:** a skin disease of schnauzers where the skin forms comedones ("blackheads").
- 272. Scotty cramp:** a condition found in the Scottish Terrier where the animal has periodic, generalized cramping of the muscles.
- 273. Screw tail:** a birth defect where the tail is twisted tightly on itself.
- 273a. Sebaceous adenitis:** (See #140.)
- 274. Sebaceous cyst:** a small mass in the skin with a secretory lining and filled with a yellow waxy-like material. (also see #107.)
- 275. Sebaceous gland tumor:** a tumor arising from sebaceous glands of the skin.
- 276. Seborrhea:** a skin disease with excess scaling of the skin and often an excess of sebum (oil-like substance) and odor.
- 277. Sertoli cell tumor:** a tumor of the testicles which secretes estrogen and causes feminization.
- 277a. Shar-Pei fever syndrome:** similar to familial Mediterranean fever of humans, affected Shar-Pei have waxing and waning high fevers, and swelling of the tarsus joints (swollen hock syndrome), which can progress to renal or hepatic amyloidosis.
- 278. Short skull:** a skull that is abnormally short for the breed in question.
- 279. Short spine:** a spine that is abnormally short for the breed in question.
- 280. Short tail:** a tail that is abnormally short for the breed in question.
- 281. Shoulder abnormalities:** a group of disorders of the shoulder joint due to malformation or subluxation.
- 282. Shoulder dysplasia:** a looseness of the shoulder joint.
- 283. Silica uroliths:** stones which are composed primarily from silicone that forms in the bladder.
- 284. Sinoatrial syncope:** a condition where the electrical impulses of the heart are abnormal and the animal has episodes of syncope (fainting).
- 285. Skin disorders:** any of a number of abnormalities of the skin.
- 286. Skin neoplasms:** any number of tumors arising from cells of the skin.
- 286a. Soft tissue cancers:** prevalent and inherited in the Flat Coated Retriever.
- 287. Spina bifida:** a developmental abnormality where some vertebra are malformed thereby exposing the spinal cord.
- 288. Spinal cord demyelination (ataxia):** an abnormality of the nervous tissue of the spinal cord leading to incoordination.
- 289. Spinal dysraphism:** a developmental abnormality where the spinal cord does not form completely. (also see #305.)
- 290. Spinal osteochondrosis:** a specific type of developmental abnormality of the vertebrae.
- 290a. Spinal process (vertebral) malformation:** (See #61c.)
- 291. Splenic torsion:** a condition where the spleen twists upon itself.
- 292. Spondylolisthesis (Wobbler's syndrome):** a condition where the vertebrae of the neck slip out of joint and are malformed causing progressive incoordination of the rear legs. *Doberman Pinscher (also see #51, 332.)
- 293. Spondylosis:** a malformation of the vertebrae.
- 294. Squamous cell carcinoma:** a cancer arising from the squamous type of skin cell.

- 295. Stenotic nares:** a condition where the openings of the nose (nares) are too small.
- 296. Sterile pyogranuloma syndrome:** a disease of the deeper layers of the skin characterized by formation of abnormal tissues, with no infectious organisms involved.
- 297. Sternal callus:** a thickened, hairless area forming on the chest of an animal.
- 298. Stockard's paralysis:** a degeneration of parts of the spinal cord causing paralysis.
- 299. Stomach torsion:** (See #131.)
- 299a. Stomatocytosis:** disorder where red blood cells have a mouth-like shape leading to frequent bouts of hemolytic anemia, and increased red cell osmotic fragility. (also see #13.)
- 299b. 'Storage' disease:** (See #193a, 188a.)
- 300. Subaortic stenosis:** a tightening of the outflow opening for blood to go from the heart into the aorta. *Golden Retriever and Newfoundland
- 301. Subcorneal pustular dermatosis:** a skin inflammation occurring between certain layers of the skin.
- 302. Subcutaneous cysts:** small fluid-filled masses accumulating under the skin.
- 303. Subvalvular aortic stenosis:** as #300, but the tightening occurs below the aortic valve.
- 303a. Sulfonamide sensitivity:** a condition in genetically predisposed breeds where metabolism of potentiated sulfonamides is impaired and adverse side-effects are seen, including liver dysfunction, dry eye (#181), rheumatoid arthritis and bone marrow failure (red blood cell and/or platelet destruction). *Doberman Pinscher, Samoyed, American Eskimo, Kuvasz and Great Pyrenees
- 304. Swimmer puppies:** a developmental defect which causes a flattening of the body so that newborn pups are unable to place their feet under them for proper locomotion.
- 304a. Syncope:** a brief period of fainting or collapse. (also see #206a.)
- 305. Syringomyelia:** developmental abnormalities causing cavities within the spinal cord, probably just an effect of #289. *Rhodesian Ridgeback
- 306. Systemic lupus erythematosus:** an autoimmune disease where antibodies form against the nuclear protein of cells, characterized by skin lesions as well as other organ dysfunctions and blood abnormalities.
- 307. Tail abnormalities:** any number of problems associated with the tail.
- 308. Tail fold dermatitis:** a skin infection caused by abnormal tissue folds around the tail.
- 308a. Taurine-deficient cardiomyopathy:** a reversible dilated cardiomyopathy caused by taurine deficiency in the Golden Retriever and Newfoundland. (also see #38.)
- 309. Teeth abnormalities:** any number of problems of the teeth.
- 310. Tetralogy of Fallot:** a specific four-way developmental abnormality of the structures of the heart and associated great vessels.
- 311. Thrombocytopathy:** a functional abnormality of small blood cells (thrombocytes or platelets) which are needed to control bleeding. (also see #249.)
- 311a. Thrombocytopenia:** a reduced number of platelets in the blood which causes pinpoint hemorrhages in the skin and mucosa and often accompanies #146 as an autoimmune syndrome called Evans syndrome. (also see #249.) *Cavalier King Charles and English Toy Spaniel (mild to moderately severe familial thrombocytopenia can be an incidental finding in clinically normal animals).

311b. Thymic atrophy: a deficiency of cell-mediated immunity expressed by decreased T-cell function and low concentrations of growth hormone. *Weimaraner (also see #165.)

312. Thyroiditis: an autoimmune inflammatory disease of the thyroid gland. (also see #166, 192.)

313. Tracheal collapse: (See #57.)

313a. Transitional cell carcinoma: a form of bladder cancer especially common in the Scottish Terrier and West Highland White Terrier (also see #37.)

314. Trembling of the hindquarters: a condition where the rear legs tremble due to muscle weakness or other pathologies.

315. Type II muscle fiber deficiency: a deficiency in form and/or function of a specific type of muscle fiber.

316. Ulcerative colitis: an autoimmune inflammation of the lining of the colon characterized by formation of ulcers.

317. Ulcerative keratitis: an inflammation of the cornea characterized by the formation of ulcers.

318. Umbilical hernia: a break in the abdominal muscle wall at the point where the umbilical cord enters the body.

319. Undershot jaw: a condition where the lower jaw is too long for the upper jaw.

320. Ununited anconeal process: a developmental abnormality of one of the bones of the elbow joint causing pain. (also see #221a.)

321. Uric acid calculi: bladder stones which are formed primarily from urates.

*Dalmatian

322. Uric acid excretion abnormalities: an abnormality in the process of the excretion of the uric acid formed during metabolism. *Dalmatian

322a. Urolithiasis: stone formation in the urinary tract.

323. Uterine eclampsia: (See #91.)

324. Uterine inertia, primary: a condition where the uterus does not have the muscular strength to proceed with the birth process, and not due to any acquired problems (e.g., malnutrition).

324a. Uveodermatologic syndrome: (See #329.)

325. Vaginal hyperplasia: an overgrowth of tissues of the vagina.

326. Vasculitis: an inflammatory condition of the blood vessels.

327. Ventricular septal defect: an abnormality (usually a hole) in the wall between the two chambers of the heart.

327a. Ventricular tachycardia: a condition where a ventricle of the heart beats too rapidly, leading to varying degrees of cardiac irregularity or syncope.

327b. Vitamin B12-responsive malabsorption: a disease of young giant Schnauzer in which there is selective inability to absorb vitamin B12 from the bowel. Affected puppies have chronic non-regenerative anemia, low white blood cell counts, low serum vitamin B12, metabolites (methylmalonic acid) in the urine, and failure to thrive.

328. Vitiligo: a lack of pigment in the skin (called vitiligo in man and hypopigmentation in animals). *Rottweiler, Doberman Pinscher, Old English Sheepdog and Dachshund (also see #161.)

329. Vogt-Koyanagi-Harada-like syndrome: an autoimmune disease common in the Akita and the "sled" dog breeds where the eyes, blood and other tissues are progressively destroyed leading to blindness and death, also called uveodermatologic syndrome.

330. von Willebrand's disease: a type of bleeding disorder caused by defective blood platelet function (an autosomal trait affecting both sexes). *Occurs in 59 dog breeds, but most often in the Doberman Pinscher

331. Westie armadillo syndrome: a condition of the West Highland White Terrier where the skin becomes very thickened. This condition is related to atopic (inhalant) allergies. (also see #106.)

331a. White dog shaker syndrome: a disorder mainly of white dogs having muscular tremors over entire body, incoordination and rapid eye movements. Episodes occur with stress or excitement.

332. Wobbler's syndrome: (See #51, 292.)

333. Zinc deficiency: can be caused by dietary problems, but also from an inability to utilize and store zinc properly and is seen as a lethal problem called acrodermatitis in the Bull Terrier.

334. Zinc-responsive dermatosis: a condition where the skin is abnormal (scaly, hair loss, etc.) but which responds to the administration of zinc in the diet.