#### ARCI-011-020 Medications and Prohibited Substances

Upon a finding of a violation of these medication and prohibited substances rules, the stewards shall consider the classification level of the violation as listed in at the time of the violation in the Uniform Classification Guidelines of Foreign Substances as promulgated by the Association of Racing Commissioners International and impose penalties and disciplinary measures consistent with the recommendations contained therein. The stewards shall also consult with the official veterinarian to determine if the violation was a result of the administration of a therapeutic medication as documented in a veterinarian's Medication Report Form received per ARCI-011-010 (C). The stewards may also consult with the laboratory director or other individuals to determine the seriousness or the laboratory finding or the medication violation Penalties for all medication and drug violations shall be investigated and reviewed on a case by case basis. Extenuating factors include, but are not limited to:

- (1) The past record of the trainer, veterinarian and owner in drug cases;
- (2) The potential of the drug(s) to influence a horse's racing performance;
- (3) The legal availability of the drug;
- (4) Whether there is reason to believe the responsible party knew of the administration of the drug or intentionally administered the drug;
- (5) The steps taken by the trainer to safeguard the horse;
- (6) The probability of environmental contamination or inadvertent exposure due to human drug use;
- (7) The purse of the race;
- (8) Whether the drug found was one for which the horse was receiving a treatment as determined by the Medication Report Form;
- (9) Whether there was any suspicious betting pattern in the race, and;
- (10) Whether the licensed trainer was acting under the advice of a licensed veterinarian. As a result of the investigation, there may be mitigating circumstances for which a lesser or no penalty is appropriate for the licensee and aggravating factors, which may increase

#### A. Uniform Classification Guidelines

the penalty beyond the minimum.

The following outline describes the types of substances placed in each category. This list shall be publicly posted in the offices of the official veterinarian and the racing secretary.

# (1) Class 1

Opiates, opium derivatives, synthetic opioids, psychoactive drugs, amphetamines and U.S. Drug Enforcement Agency (DEA) scheduled I and II drugs. Also found in this class are drugs which are potent stimulants of the nervous system. Drugs in this class have no generally accepted medical use in the racehorse and their pharmacological potential for altering the performance of a race is very high.

#### (2) Class 2

Drugs in this category have a high potential for affecting the outcome of a race. Most are not generally accepted as therapeutic agents in the racehorse. Many are products

intended to alter consciousness or the psychic state of humans, and have no approved or indicated use in the horse. Some, such as injectable local anesthetics, have legitimate use in equine medicine, but should not be found in a racehorse. The following groups of drugs are in this class:

- (a) Opiate partial agonists, or agonist-antagonists;
- (b) Non-opiate psychotropic drugs, which may have stimulant, depressant, analgesic or neuroleptic effects;
- (c) Miscellaneous drugs which might have a stimulant effect on the central nervous system (CNS);
- (d) Drugs with prominent CNS depressant action;
- (e) Antidepressant and antipsychotic drugs, with or without prominent CNS stimulatory or depressant effects;
- (f) Muscle blocking drugs which have a direct neuromuscular blocking action;
- (g) Local anesthetics which have a reasonable potential for use as nerve blocking agents (except procaine); and
- (h) Snake venoms and other biologic substances, which may be used as nerve blocking agents.

### (3) Class 3

Drugs in this class may or may not have an accepted therapeutic use in the horse. Many are drugs that affect the cardiovascular, pulmonary and autonomic nervous systems. They all have the potential of affecting the performance of a racehorse. The following groups of drugs are in this class:

- (a) Drugs affecting the autonomic nervous system which do not have prominent CNS effects, but which do have prominent cardiovascular or respiratory system effects (bronchodilators are included in this class);
- (b) A local anesthetic which has nerve blocking potential but also has a high potential for producing urine residue levels from a method of use not related to the anesthetic effect of the drug (procaine);
- (c) Miscellaneous drugs with mild sedative action, such as the sleep inducing antihistamines;
- (d) Primary vasodilating/hypotensive agents; and
- (e) Potent diuretics affecting renal function and body fluid composition.

# (4) Class 4

This category is comprised primarily of therapeutic medications routinely used in racehorses. These may influence performance, but generally have a more limited ability to do so. Groups of drugs assigned to this category include the following:

- (a) Non-opiate drugs which have a mild central analgesic effect;
- (b) Drugs affecting the autonomic nervous system which do not have prominent CNS, cardiovascular or respiratory effects
  - (A) Drugs used solely as topical vasoconstrictors or decongestants
  - (B) Drugs used as gastrointestinal antispasmodics
  - (C) Drugs used to void the urinary bladder

- (D) Drugs with a major effect on CNS vasculature or smooth muscle of visceral organs.
- (E) Antihistamines which do not have a significant CNS depressant effect (This does not include H1 blocking agents, which are listed in Class 5);
- (c) Mineralocorticoid drugs;
- (d) Skeletal muscle relaxants:
- (e) Anti-inflammatory drugs--those that may reduce pain as a consequence of their anti-inflammatory actions, which include:
  - (A) Non-Steroidal Anti-Inflammatory Drugs (NSAIDs;
  - (B) Corticosteroids (glucocorticoids); and
  - (C) Miscellaneous anti-inflammatory agents.
- (f) Anabolic and/or androgenic steroids and other drugs;
- (g) Less potent diuretics;
- (h) Cardiac glycosides and antiarrhythmics including:
  - (A) Cardiac glycosides;
  - (B) Antirryhthmic agents (exclusive of lidocaine, bretylium and propanolol); and
  - (C) Miscellaneous cardiotonic drugs.
- (i) Topical Anesthetics--agents not available in injectable formulations;
- (j) Antidiarrheal agents; and
- (k) Miscellaneous drugs including:
  - (A) Expectorants with little or no other pharmacologic action;
  - (B) Stomachies; and
  - (C) Mucolytic agents.

# (5) Class 5

Drugs in this category are therapeutic medications for which concentration limits have been established as well as certain miscellaneous agents. Included specifically are agents, which have very localized action only, such as anti-ulcer drugs and certain anti-allergenic drugs. The anticoagulant drugs are also included.

#### B. Penalties

- (1) In issuing penalties against individuals found guilty of medication and drug violations a regulatory distinction shall be made between the detection of therapeutic medications used routinely to treat racehorses and those drugs that have no reason to be found at any concentration in the test sample on race day.
- (2) The stewards or the commission will use the Racing Medication and Testing Consortium's penalty category and schedule as a starting place in the penalty stage of the deliberations for a rule violation for any drug listed in the Association of Racing Commissioners International Uniform Classification Guidelines for Foreign Substances.
- (3) If a licensed veterinarian is administering or prescribing a drug not listed in the RCI *Uniform Classification Guide lines for Foreign Substances* or shown in the RMTC

- *Penalty Guideline Listing*, the identity of the drug shall be forwarded to the official veterinarian to be forwarded to the Racing Medication and Testing Consortium for classification.
- (4) Any drug or metabolite thereof found to be presenting a pre- or post-race sample which is not classified in the most current RCI *Uniform Classification Guidelines* for Foreign Substances shall be assumed to be a RCI Class 1 Drug and the trainer and owner shall be subject to those penalties as set forth in schedule "A" unless satisfactorily demonstrated otherwise by the Racing Medication and Testing Consortium, with a penalty category assigned.
- (5) The penalty categories and their related schedules, if applicable, shall be on the following criteria:
  - (a) Whether the drug is approved by the U.S. Food and Drug Administration for use in the horse:
  - (b) Whether the drug is approved by the U.S. Food and Drug Administration for use in any species;
  - (c) Whether the drug has any legitimate therapeutic application in the equine athlete:
  - (d) Whether the drug was identified as "necessary" by the RMTC Veterinary Advisory Committee;
  - (e) Whether legitimate, recognized therapeutic alternatives exist, and;
  - (f) The current RCI Classification of the drug.
- (6) The penalty categories "A", "B" and "C" and their related schedules for Trainers and Owners are shown in the following tables.

The following are recommended penalties for violations due to the presence of a drug carrying a Category "A" penalty and for violations of ARCI-011-015: Prohibited Practices:

LICENSED TRAINER:				
1 <sup>st</sup> offense	2 <sup>nd</sup> LIFETIME offense in any jurisdiction	3 <sup>rd</sup> LIFETIME offense in any jurisdiction		
Minimum one-year suspension absent	Minimum three-year suspension absent	Minimum five-year suspension absent mitigating		
mitigating circumstances. The presence of	mitigating circumstances. The presence of	circumstances. The presence of aggravating factors		
aggravating factors could be used to impose a	aggravating factors could be used to impose a	could be used to impose a maximum of license		
maximum of a three-year suspension.	maximum of license revocation with no	revocation with no reapplication for a five-year		
	reapplication for a three-year period.	period.		
AND	AND	AND		
<ul> <li>Minimum fine of \$10,000 or 10% of total</li> </ul>	<ul> <li>Minimum fine of \$25,000 or 25% of total</li> </ul>	Minimum fine of \$50,000 or 50% of total purse		
purse (greater of the two) absent mitigating	purse (greater of the two) absent mitigating	(greater of the two) absent mitigating circumstances.		
circumstances. The presence of aggravating	circumstances. The presence of aggravating	The presence of aggravating factors could be used to		
factors could be used to impose a maximum of	factors could be used to impose a maximum of	impose a maximum of \$100,000 or 100% of purse		
\$25,000 or 25% of purse (greater of the two).	\$50,000 or 50% of purse (greater of the two).	(greater of the two).		
AND	AND	AND		
<ul> <li>May be referred to the Commission for any</li> </ul>	<ul> <li>May be referred to the Commission for any</li> </ul>	May be referred to the Commission for any further		
further action deemed necessary by the	further action deemed necessary by the	action deemed necessary by the Commission.		
Commission.	Commission.			
LICENSED OWNER:				
1 <sup>st</sup> offense	2 <sup>nd</sup> LIFETIME offense in owner's stable	3 <sup>rd</sup> LIFETIME offense in owner's stable		
	in any jurisdiction	in any jurisdiction		
<ul> <li>Disqualification and loss of purse.</li> </ul>	<ul> <li>Disqualification and loss of purse.</li> </ul>	Disqualification, loss of purse and \$50,000 fine.		
		AND		
AND	AND			
<ul> <li>Horse shall be placed on the veterinarian's</li> </ul>	<ul> <li>Horse shall be placed on the veterinarian's list</li> </ul>	Horse shall be placed on the veterinarian's list for		
list for 90 days and must pass a commission-	for 120 days and must pass a commission-	180 days and must pass a commission-approved		
approved examination before becoming	approved examination before becoming eligible	examination before becoming eligible to be entered.		
eligible to be entered.	to be entered.	AND		
		Referral to the Commission with a recommendation		
TDI 0.11 : 1.1	as for violations due to the presence of a drug of	of a suspension for a minimum of 90 days.		

The following are recommended penalties for violations due to the presence of a drug carrying Category "B" penalty, for the presence of more than one NSAID in a plasma/serum sample, subject to the provisions set forth in ARCI-011-020 E.(1)(c) and for violations of the established levels for total carbon dioxide:

# LICENSED TRAINER:

1 <sup>st</sup> offense	2 <sup>nd</sup> offense (365-day period) in any jurisdiction	3 <sup>rd</sup> offense (365-day period) in any jurisdiction			
Minimum 15-day suspension absent	Minimum 30-day suspension absent mitigating	Minimum 60-day suspension absent mitigating			
mitigating circumstances. The presence of	circumstances. The presence of aggravating factors	circumstances. The presence of aggravating factors			
aggravating factors could be used to impose	could be used to impose a maximum of a 180-day	could be used to impose a maximum of a one-year			
a maximum of a 60-day suspension.	suspension.	suspension.			
AND	AND	AND			
<ul> <li>Minimum fine of \$500 absent mitigating</li> </ul>	Minimum fine of \$1,000 absent mitigating	Minimum fine of \$2,500 absent mitigating			
circumstances. The presence of aggravating	circumstances. The presence of aggravating factors	circumstances. The presence of aggravating factors			
factors could be used to impose a maximum	could be used to impose a maximum of \$2,500.	could be used to impose a maximum of \$5,000 or 5%			
of \$1,000.		of purse (greater of the two).			
		AND			
		May be referred to the Commission for any further     the Commission for any further			
		action deemed necessary by the Commission.			
LICENSED OWNER:	LICENSED OWNER:				
1 <sup>st</sup> offense	2 <sup>nd</sup> offense in stable (365-day period) in any	3 <sup>rd</sup> offense in stable (365-day period) in any			
	jurisdiction	jurisdiction			
<ul> <li>Disqualification and loss of purse [in the</li> </ul>	<ul> <li>Disqualification and loss of purse [in the</li> </ul>	Disqualification and loss of purse, and in the			
absence of mitigating circumstances]*.	absence of mitigating circumstances]*.	absence of mitigating circumstances a \$5,000 fine.*			
AND					
		AND			
	AND				
<ul> <li>Horse must pass a commission-approved</li> </ul>	Horse must pass a commission-approved	Horse shall be placed on the veterinarian's list for			
examination before becoming eligible to be	examination before becoming eligible to be	45 days and must pass a commission-approved			
entered.	entered.	examination before becoming eligible to be entered.			

<sup>\* (</sup>The RMTC recommendation called for loss of purse to happen in absence of mitigating circumstances the Joint Model Rules Committee has made loss of purse mandatory in their proposal)

The following are recommended penalties for violations due to the presence of a drug carrying a Category "C" penalty and overages for permitted NSAIDs and furosemide: (*All concentrations are for measurements in serum or plasma*.)

LICENSED TRAINER	Phenylbutazone (>2.0 - 5.0 2.1 4.9 mcg/ml)* Flunixin (>20 - 100 21 99 ng/ml) Ketoprofen (>10 - 50 11 49 ng/ml) Furosemide (>100 ng/ml) and no furosemide when identified as administered**	Phenylbutazone (≥≥5.0 mcg/ml) Flunxin (≥≥100 ng/ml) Ketoprofen (≥≥50 ng/ml) and CLASS C Violations
1 <sup>st</sup> Offense (365-day period) in any jurisdiction	Minimum fine of a written warning to a maximum fine of \$500\\$250 absent mitigating circumstances	Minimum fine of \$1,000500 absent mitigating circumstances
2 <sup>nd</sup> Offense (365-day period) in any jurisdiction	Minimum fine of a written warning to a maximum fine of \$750\$500 absent mitigating circumstances	Minimum fine of \$1,5001,000 and 15-day suspension absent mitigating circumstances
3 <sup>rd</sup> Offense (365-day period) in any jurisdiction	Minimum fine of \$500 to a maximum fine of \$1,000 and 15 day suspension absent mitigating circumstances	Minimum fine of \$2,500 and 30-day suspension absent mitigating circumstances
LICENSED OWNER	Phenylbutazone ( <u>&gt;2.0 - 5.02.1-4.9</u> mcg/ml) Flunixin ( <u>&gt;20 - 5021-99</u> ng/ml) Ketoprofen ( <u>&gt;10 - 5011-49</u> ng/ml)	Phenylbutazone (≥≤5.0 mcg/ml) Flunixin (≥≤100 ng/ml) Ketoprofen (≥≤50 ng/ml) AND CLASS C VIOLATIONS
	Furosemide (>100 ng/ml) and no furosemide when identified as administered**	
1 <sup>st</sup> Offense (365-day period) in any jurisdiction	no furosemide when identified as	Loss of purse. Horse must pass commission-approved examination before being eligible to run
1 <sup>st</sup> Offense (365-day period) in any jurisdiction  2 <sup>nd</sup> Offense (365-day period) in any jurisdiction	no furosemide when identified as administered**  Horse may be required to pass commission-approved examination before being	

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eligible to run

\*If the trainer has not had more than one violation within the previous two years, the Stewards/Judges are encouraged to issue a warning in lieu of a fine provided the reported level is below 3.0 mcg/ml, absent of aggravating factors.

After a two-year period, if the licensee has had no further violations, any penalty due to an overage in the 2.0-5.0 category will be expunged from the licensee's record for penalty purposes.

- (7) The recommended penalty for a violation involving a drug that carries a Category "D" penalty is a written warning to the trainer and owner. Multiple violations may result in fines and/or suspensions
- (8) Any licensee of the commission, including veterinarians, found to be responsible for the improper or intentional administration of any drug resulting in a positive test may, after proper notice and hearing, be subject to the same penalties set forth for the licensed trainer.
- (9) The licensed owner, veterinarian or any other licensed party involved in a positive laboratory finding shall be notified in writing of the hearing and any resulting action. In addition their presence may be required at any and all hearings relative to the case.
- (10) Any veterinarian found to be involved in the administration of any drug carrying the penalty category of "A" shall be referred to the State Licensing Board of Veterinary Medicine for consideration of further disciplinary action and/or license revocation. This is in addition to any penalties issued by the stewards or the commission.
- (11) Any person who the stewards or the commission believe may have committed acts in violation of criminal statutes may be referred to the appropriate law enforcement agency. Administrative action taken by the stewards or the commission in no way prohibits a prosecution for criminal acts committed, nor does a potential criminal prosecution stall administrative action by the stewards or the commission.

Procedures shall be established to ensure that a licensed trainer is not able to benefit financially during the period for which the individual has been suspended. This includes, but is not limited to, ensuring that horses are not transferred to licensed family members.

#### C. Medication Restrictions

- (1) A finding by the commission approved laboratory of a prohibited drug, chemical or other substance in a test specimen of a horse is prima facie evidence that the prohibited drug, chemical or other substance was administered to the horse and, in the case of a post-race test, was present in the horse's body while it was participating in a race. Prohibited substances include:
  - (a) Drugs or medications for which no acceptable threshold concentration has been established;
  - (b) Therapeutic medications in excess of established threshold concentrations;
  - (c) Substances present in the horse in excess of concentrations at which such substances could occur naturally; and
  - (d) Substances foreign to a horse at concentrations that cause interference with testing procedures.
- (2) Except as otherwise provided by this chapter, a person may not administer or cause to be administered by any means to a horse a prohibited drug, medication, chemical or other substance, including any restricted medication pursuant to this chapter during the 24-hour period before post time for the race in which the horse is entered.

## D. Medical Labeling

(1) No person on association grounds where horses are lodged or kept, excluding licensed veterinarians, shall have in or upon association grounds which that person occupies or has the right to occupy, or in that person's personal property or effects or vehicle in that person's care, custody or control, a drug, medication, chemical, foreign substance or other substance

- that is prohibited in a horse on a race day unless the product is labeled in accordance with this subsection.
- (2) Any drug or medication which is used or kept on association grounds and which, by federal or state law, requires a prescription must have been validly prescribed by a duly licensed veterinarian, and in compliance with the applicable state statutes. All such allowable medications must have a prescription label which is securely attached and clearly ascribed to show the following:
  - (a) The name of the product;
  - (b) The name, address and telephone number of the veterinarian prescribing or dispensing the product;
  - (c) The name of each patient (horse) for whom the product is intended/prescribed;
  - (d) The dose, dosage, duration of treatment and expiration date of the prescribed/dispensed product; and
  - (e) The name of the person (trainer) to whom the product was dispensed.

### E. Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)

- (1) The use of one of three approved NSAIDs shall be permitted under the following conditions:
  - (a) Not to exceed the following permitted serum or plasma threshold concentrations which are consistent with administration by a single intravenous injection at least 24 hours before the post time for the race in which the horse is entered:
    - (i) Phenylbutazone 2 micrograms per milliliter;
    - (ii) Flunixin 20 nanograms per milliliter;
    - (iii) Ketoprofen 10 nanograms per milliliter.
  - (b) These or any other NSAID are prohibited to be administered within the 24 hours before post time for the race in which the horse is entered.
  - (c) The presence of more than one of the three approved NSAIDs, with the exception of Phenylbutazone in a concentration below 0.5 microgram per milliliter of serum or plasma or any unapproved NSAID in the post-race serum or plasma sample is not permitted. The use of all but one of the approved NSAIDs shall be discontinued at least 48 hours before the post time for the race in which the horse is entered.
- (2) Any horse to which a NSAID has been administered shall be subject to having a blood and/or urine sample(s) taken at the direction of the official veterinarian to determine the quantitative NSAID level(s) and/or the presence of other drugs which may be present in the blood or urine sample(s).

### F. Furosemide

(1) Furosemide may be administered intravenously to a horse, which is entered to compete in a race. Except under the instructions of the official veterinarian or the racing veterinarian for the purpose of removing a horse from the Veterinarian's List or to facilitate the collection of a post-race urine sample, furosemide shall be permitted only after the official veterinarian has placed the horse on the Furosemide List. In order for a horse to be placed on the Furosemide List the following process must be followed.

- (a) After the horse's licensed trainer and licensed veterinarian determine that it would be in the horse's best interests to race with furosemide they shall notify the official veterinarian or his/her designee, using the prescribed form, that they wish the horse to be put on the Furosemide List.
- (b) The form must be received by the official veterinarian or his/her designee by the proper time deadlines so as to ensure public notification.
- (c) A horse placed on the official Furosemide List must remain on that list unless the licensed trainer and licensed veterinarian submit a written request to remove the horse from the list. The request must be made to the official veterinarian or his/her designee, on the proper form, no later than the time of entry.
- (d) After a horse has been removed from the Furosemide List, the horse may not be placed back on the list for a period of 60 calendar days unless it is determined to be detrimental to the welfare of the horse, in consultation with the official veterinarian. If a horse is removed from the official Furosemide List a second time in a 365-day period, the horse may not be placed back on the list for a period of 90 calendar days.
- (e) Furosemide shall only be administered on association grounds.
- (f) Upon the request of the regulatory agency designee, the veterinarian administering the authorized bleeder medication shall surrender the syringe used to administer such medication which may then be submitted for testing
- (2) The use of furosemide shall be permitted under the following circumstances on association grounds where a detention barn is utilized:
  - (a) Furosemide shall be administered at the direction of the official veterinarian no less than four hours prior to post time for the race for which the horse is entered.
  - (b) A horse qualified for furosemide administration must be brought to the detention barn within time to comply with the four-hour administration requirement specified above.
  - (c) The dose administered shall not exceed 500 mg. nor be less than 150 mg.
  - (d) Furosemide shall be administered by a single, intravenous injection.
  - (e) After treatment, the horse shall be required by the Commission to remain in the detention barn in the care, custody and control of its trainer or the trainer's designated representative under association and/or Commission security supervision until called to the saddling paddock.
- (3) The use of furosemide shall be permitted under the following circumstances on association grounds where a detention barn is not utilized:
  - (a) Furosemide shall be administered no less than four hours prior to post time for the race for which the horse is entered.
  - (b) The furosemide dosage administered shall not exceed 500 mg. nor be less than 150 mg.
  - (c) Furosemide shall be administered by a single, intravenous injection.
  - (d) The trainer of the treated horse shall cause to be delivered to the official veterinarian no later than one hour prior to post time for the race for which the horse is entered the following information under oath on a form provided by the Commission:
    - (A) The name of the horse, racetrack name, the date and time the furosemide was administered to the entered horse;

- (B) The dosage amount of furosemide administered to the entered horse; and
- (C) The printed name and signature of the attending licensed veterinarian who administered the furosemide.
- (4) Test results must show a detectable concentration of the drug in the post-race serum, plasma or urine sample.
  - (a) The specific gravity of post-race urine samples may be measured to ensure that samples are sufficiently concentrated for proper chemical analysis. The specific gravity shall not be below 1.010. If the specific gravity of the urine is found to be below 1.010 or if a urine sample is unavailable for testing, quantitation of furosemide in serum or plasma shall be performed;
  - (b) Quantitation of furosemide in serum or plasma shall be performed when the specific gravity of the corresponding urine sample is not measured or if measured below 1.010. Concentrations may not exceed 100 nanograms of furosemide per milliliter of serum or plasma

#### G. Bleeder List

- (1) The official veterinarian shall maintain a Bleeder List of all horses, which have demonstrated external evidence of exercise induced pulmonary hemorrhage from one or both nostrils during or after a race or workout as observed by the official veterinarian.
- (2) Every confirmed bleeder, regardless of age, shall be placed on the Bleeder List and be ineligible to race for the following time periods:
  - (a) First incident 14 days;
  - (b) Second incident within 365 day period 30 days;
  - (c) Third incident within 365 day period –180 days;
  - (d) Fourth incident within 365-day period barred for racing lifetime.
- (3) For the purposes of counting the number of days a horse is ineligible to run, the day the horse bled externally is the first day of the recovery period.
- (4) The voluntary administration of furosemide without an external bleeding incident shall not subject the horse to the initial period of ineligibility as defined by this policy.
- (5) A horse may be removed from the Bleeder List only upon the direction of the official veterinarian, who shall certify in writing to the stewards the recommendation for removal.
- (6) A horse which has been placed on a Bleeder List in another jurisdiction pursuant to these rules shall be placed on a Bleeder List in this jurisdiction.

### H. Anti-Ulcer Medications

The following anti-ulcer medications are permitted to be administered, at the stated dosage, up to 24 hours prior to the race in which the horse is entered.

- (1) Cimetidine (Tagamet®) 8-20 mg/kg PO BID-TID
- (2) Omeprazole (Gastrogard®) 2.2 grams PO SID
- (3) Ranitidine (Zantac®) 8 mg/kg PO BID

COMMITTEE NOTE: Consortium is currently discussing administration dead-line for Ranitidine.

### I. Environmental Contaminants and Substances of Human Use

COMMITTEE NOTE: Consortium says that potential substances identified in this section will be put through the same scientific review process in order to determine whether a threshold concentration can be established.

- (1) The following substances can be environmental contaminants in that they are endogenous to the horse or that they can arise from plants traditionally grazed or harvested as equine feed or are present in equine feed because of contamination during the cultivation, processing, treatment, storage or transportation phases:
- (2) The following drugs are recognized as substances of human use and addiction and which could be found in the horse due to its close association with humans:
- (3) Regulatory thresholds have been set for the following substances.
  - (a) Caffeine 100 nanograms of caffeine per milliliter of serum or plasma
- (4) If the preponderance of evidence presented in the hearing shows that a positive test is the result of environmental contamination or inadvertent exposure due to human drug use it should be considered as a mitigating factor in any disciplinary action taken against the affected trainer.

### J. Androgenic-Anabolic Steroids

- (1) No AAS shall be permitted in test sample collected from racing horses except for residues of the major metabolite of **stanozolol, nandrolone,** and the naturally occurring substances **boldenone** and testosterone at concentrations less that the indicated thresholds.
- (2) Concentrations of these AAS shall not exceed the following urine threshold concentrations for total (*i.e.*, free drug or metabolite and drug or metabolite liberated from its conjugates):
  - (a) 16β-hydroxystanozolol (metabolite of stanozolol (Winstrol)) 1 ng/ml in urine for all horses regardless of sex;
  - (b) Boldenone (Equipoise® is the undecylenate ester of boldenone) in male horses other than geldings – 15 ng/ml in urine. No boldenone shall be permitted in geldings or female horses.
  - (c) Nandrolone (Durabolin® is the phenylpropionate ester and Deca-Durabolin® is the decanoate ester)
    - (A) In geldings 1 ng/ml in urine
    - (B) In fillies and mares -1 ng/ml in urine
  - (d) Testosterone
    - (A) In geldings 20 ng/ml in urine
    - (B) In fillies and mares 55 ng/ml in urine
- (3) Any other anabolic steroids are prohibited in racing horses.
- (4) Post-race urine samples must have the sex of the horse identified to the laboratory.
- (5) Any horse to which an anabolic steroid has been administered in order to assist in the recovery from illness or injury may be placed on the veterinarian's list in order to monitor the concentration of the drug or metabolite in urine. After the concentration has fallen below the designated threshold for the administrated AAS, the horse is eligible to be removed from the list.

# K. Alkalinizing Substances

The use of agents that elevate the horse's TCO2 or Base excess level above those existing naturally in the untreated horse at normal physiological concentrations is prohibited. The following levels also apply to blood gas analysis:

- (1) The regulatory threshold for TCO2 is 37.0 millimoles per liter of plasma/serum or a base excess level of 10.0 millimoles, and;
- (2) The decision level to be used for the regulation of TCO2 is 37.0 millimoles per liter of plasma/serum plus the measurement uncertainty of the laboratory analyzing the sample, or a base excess level of 10.4 millimoles per liter of plasma/serum.

Adopted in Version 1.4 ARCI 8/27/02 NAPRA 10/2/02

Version 1.4 to 2.0 ARCI 4/26/03 NAPRA 4/14/03: Rule topic was renumbered to ARCI-011-023

Version 2.1 to 3.0 ARCI 4/3/04 NAPRA 4/3/04: Amended and modified new rule language

Version 3.2 to 3.3 ARCI 12/7/05: Added and modified rule language

Version 4.0 to 4.1 ARCI 4/26/07: Added new rule language

Version 4.1 to 4.15 ARCI Board of Directors meeting 12/5/2007: Amended rule language

Version 4.3 to 4.4 ARCI Board 12/10/08: Amended language

Version 4.4 to 4.5 ARCI 4/23.09: Amended language added Alkalinizing Substances

Version 4.7 to 4.8 ARCI Board 10/22/10 Amended language regarding Phenylbutazone level 5.0 to 2.0