



Curt E. Harper, Ph.D., F-ABFT

Chief Toxicologist

2019 California Impaired Driving Task Force



## Alabama Dept. Forensic Sciences

- 15 Toxicologists
- 1 Laboratory (Birmingham)
- 500 cases/mo, 6,000/yr
- DUI & DUI/D (40%)
- Postmortem (60%)
  - 66 County Coroners
  - 3 Medical Examiner Offices



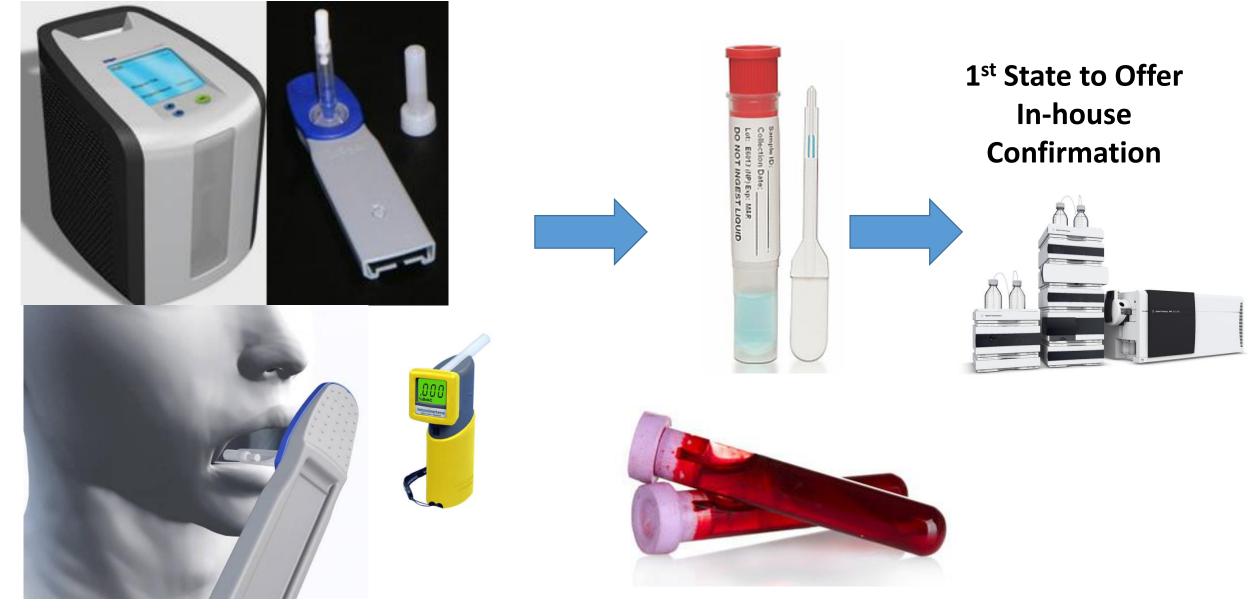
<0.08% (14%) [77% drug prev.] >0.08% (86%) [50% drug prev.]

5 years (2013-2017) **DUI – Traffic Stops** 6,355 blood tests >0.08% 81,039 breath tests 10% of <0.08% 1,135/11,345 <0.08%, **DUI – Traffic Crashes** Suspected DUI/D Involving Serious Injury/Deaths <0.08% (75%) [77% drug prev.] >0.08% (25%) [50% drug prev.]

## Oral Fluid Drug Testing

### **Roadside Screen (Probable Cause)**

**Confirmation (Evidentiary)** 



# SOFT/AAFS Oral Fluid Ad Hoc Committee

### Members

- Christine Moore (Chair)
- Curt Harper (Vice Chair)
- Marilyn Huestis
- Timothy Rohrig
- Jarrad Wagner
- Madeleine Swortwood
- Luke Rodda
- Chares LoDico
- Mandi Mohr
- Kristen Burke
- Nathalie Desrosiers

### Documents

- OF FAQ Document 2.0 (2017)
  - Advantages
  - Limitations
  - Specimen comparison

http://soft-tox.org/files/2017\_OF\_FAQ.pdf

- OF Pilot Project Guidelines
  - Key Stakeholders
  - Program Management
  - Program Protocol
  - Consent Form (example)

http://soft-tox.org/files/2014\_OF\_Pilot.pdf

# Step#1: Stakeholder's Meetings & Outreach

- Traffic Safety Resource Prosecutor (TSRP)
- DRE Coordinator & Program
- Prosecutor's Training (2016/2017/2018)
- Judge's Symposium (2016/2017)
- Collaboration with Vendors
- Alabama Impaired Driving Prevention Council (AIDPC)
- ADECA
- Step #2: Develop Study Design
  - Roadside screening, laboratory confirmation
  - No legislative change needed. Must modify ADFS Rules.

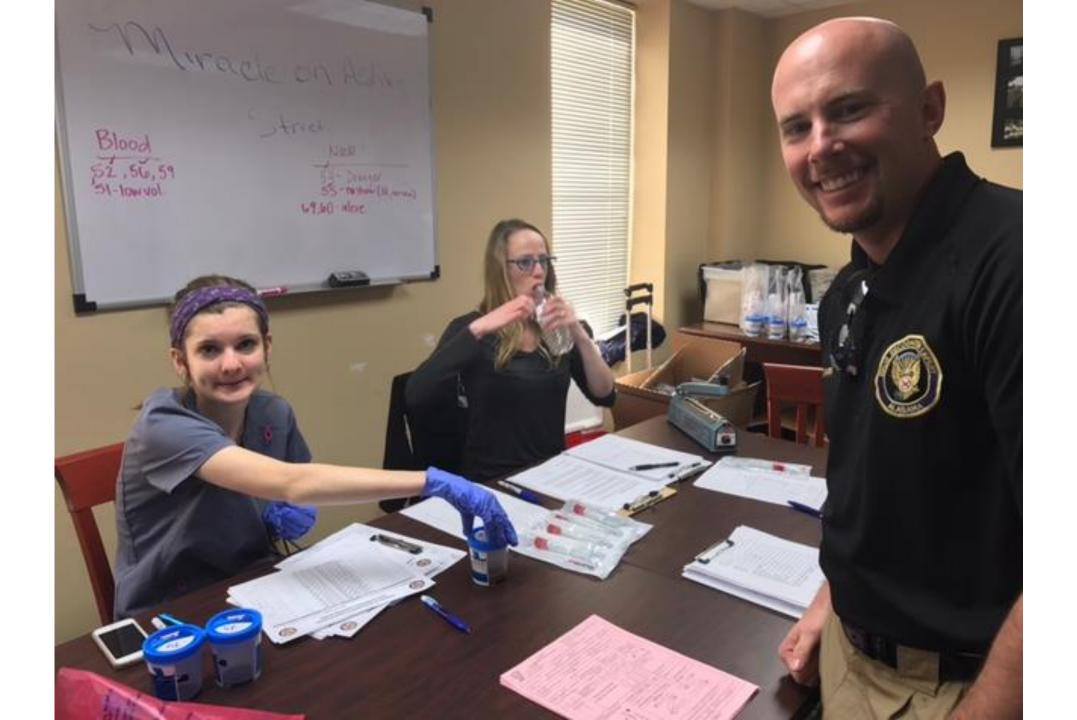
# ADFS OF Study Summary

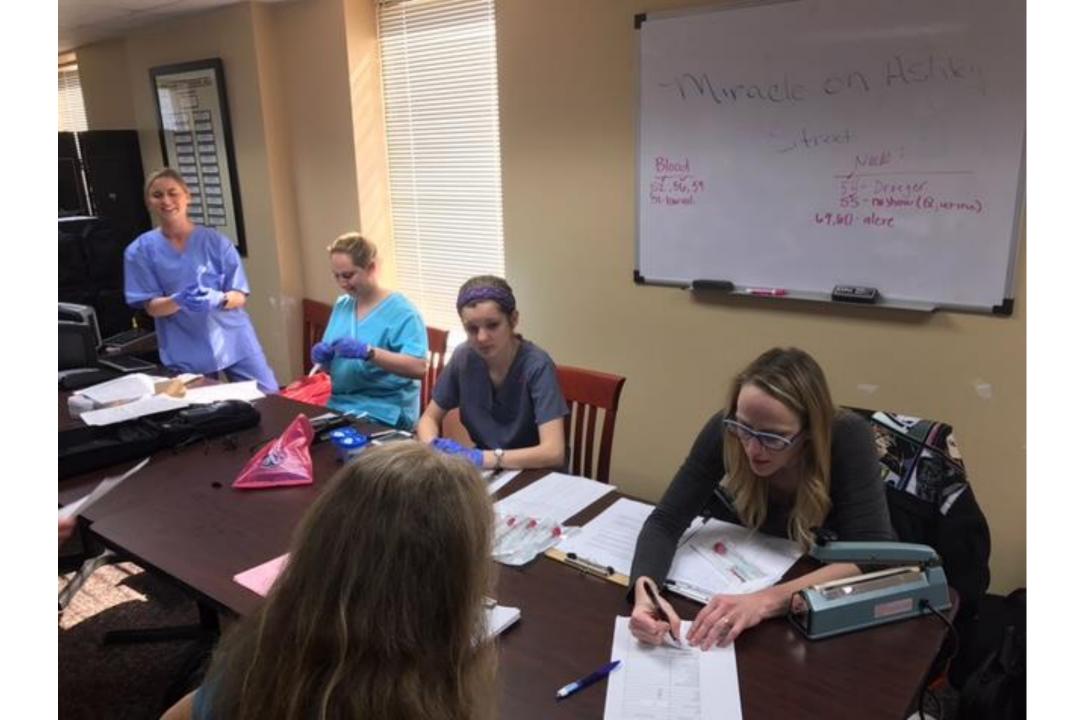
- Alabama Drug Recognition Expert (DRE), TSRP B. Lindsey
- Clara White Mission Jacksonville, FL
- Proof of principle study set forth to validate the use of :
  - Aim #1: OF screening in the field by officers
  - Aim #2: OF confirmation testing at ADFS
- 3 Oral Fluid Screening Devices
  - 1. SoToxa (fka Alere DDS2)
  - 2. Draeger DT5000
  - 3. Randox Evidence MultiSTAT
- Developed OF confirm methods at ADFS
- AL Code: Blood, Urine, or other bodily substance
  - There was no need to change state statue (huge advantage)
- Modified ADFS Rules and Regulations for Testing
  - In effect June 24, 2018
  - First oral fluid case: Summer 2018



### DUID Applications for Oral Fluid Drug Screening Devices











### Specification Comparison

	Alere DDS2	Draeger DT5000	Randox MultiSTAT
Time to complete (min)	5	10	17
Size	Small	Medium	Large
Number of targets	6	7	21

## Cutoffs (ng/mL)

Target	Alere SoToxa	Draeger DT5000	Randox MultiSTAT
Cocaine	30	20	20
ТНС	25	5	10
Opiates	40	20	10
Benzodiazepine	20	15	20
Methamphetamine	50	35	50
Amphetamine	50	50	50
Methadone	NA	20	4

### Parameters

- Sensitivity = TP/(TP + FN)
  - Ability to identify positive cases
- Specificity = TN/(TN+FP)
  - Ability to avoid false positives, identify negative cases
- Positive Predictive Value = TP/(TP+FP)
  - Ability to correctly label as positive
- Negative Predictive Value = TN/(TN+FN)
  - Ability to correctly label as negative
- Accuracy = (TP+TN)/(TP+FP+TN+FN)
  - Overall correctness

## Summary of Roadside Device Performance

	Alere SoToxa*	Draeger DT5000	Randox Multi-STAT
Specificity	>80	>80	>80
PPV	>80	>80	>80
NPV	>80	>80	>80
Sensitivity	>80	>80	>80
Accuracy	>80	>80	>80

Cocaine, THC, Meth, Opiates, Methadone\*

Specificity	98	100	100
PPV	75	75 100	
NPV	98	97	100
Sensitivity	75	57	100
Accuracy	96	97	100

Benzodiazepines

## SOFT/AAFS Oral Fluid FAQ 2.0: Should labs develop Qual or Quant OF confirmation

### The 3 No-no's

- Most drugs (e.g. smoked, vaped or snorted):
   <u>OF</u> drug concs <u>do not predict</u> concurrent <u>blood</u> drug concs
- 2. Not recommended to estimate drug concs in whole blood from OF drug concs or vice versa
- 3. <u>Not possible to correlate a quantitative drug conc.</u> in OF, blood or urine <u>directly to degree of impairment</u>

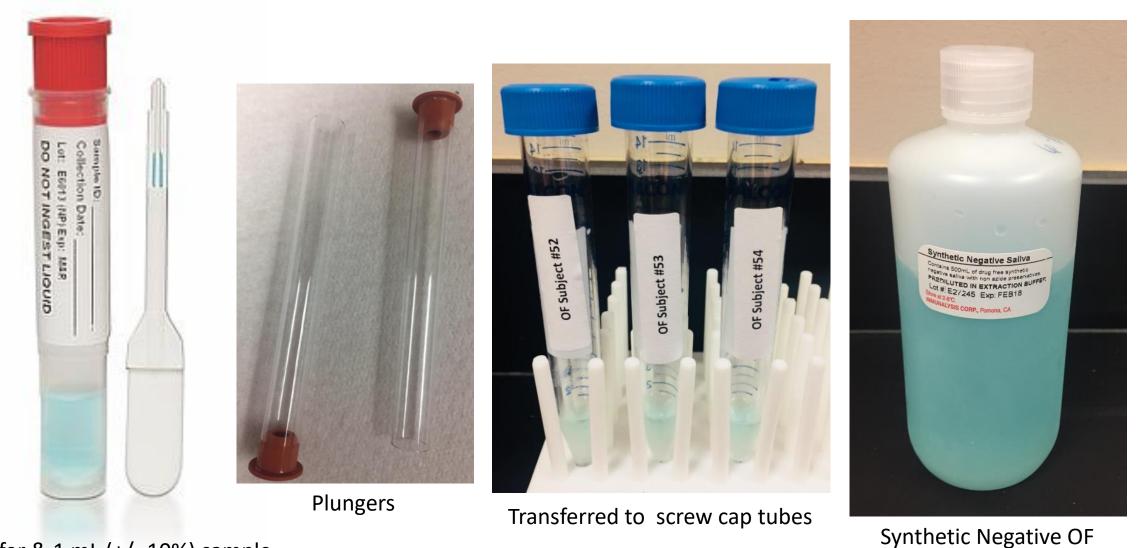
For these reasons....

## SOFT/AAFS Oral Fluid FAQ 2.0: Should labs develop Qual or Quant OF confirmation

### Recommended practice:

- Qualitative (present or ND) methods recommended since there is not a direct correlation b/t concs in OF and blood in most cases due to a variety of factors
  - Oral cavity contamination from recent use, unknown exact volume of confirmation fluid specimen, individual variability in PK and PD
- Quantitative measurement of drug concs for research purpose
  - Developing a better understanding of typical OF drug concs in various populations
  - Helps with the development of screening devices with the appropriate sensitivity

## Quantisal<sup>TM</sup> Oral Fluid Confirmation



3 mL buffer & 1 mL (+/- 10%) sample

## Quantisal 2.0<sup>TM</sup> Oral Fluid Confirmation

Sample ID: A Collection Date: Lot: E##### Exp: YYYY-MM-DD DO NOT INGEST LIQUID	
Sample ID:B Collection Date: Lot: E##### Exp: YYYY-MM-DD	

### Agilent 6460/6430 QQQ LC/MS/MS: Confirmation

2 Confirmation Methods at ADFS:

- 1. (19) Drugs of Abuse
- 2. (6) Cannabinoids

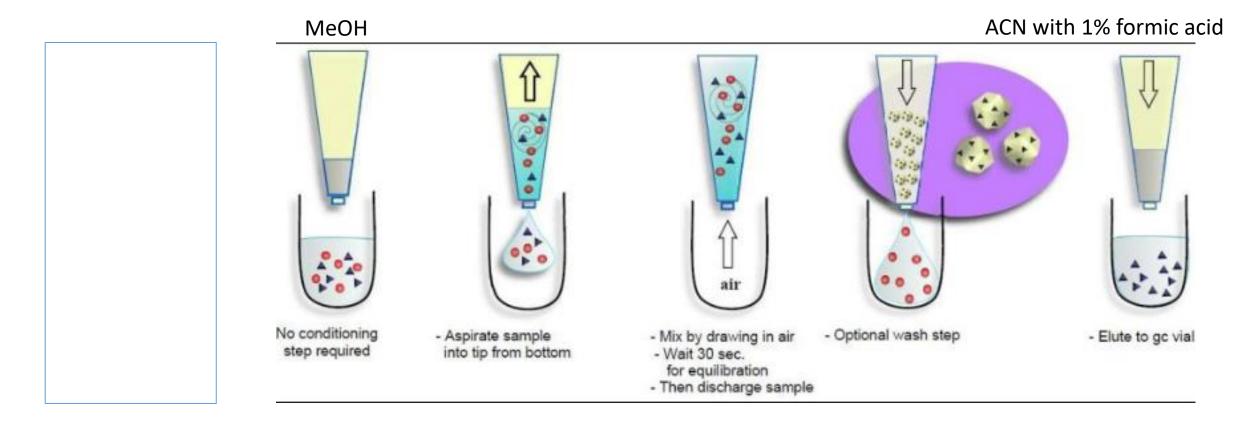
(25 Total Targets)

95% of DUID Targets



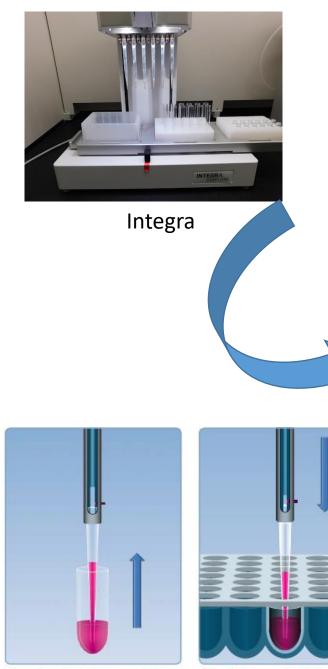
\*Recommendations for Toxicological Investigation of Drug-Impaired Driving and Motor Vehicle Fatalities. JAT 2017.

# ADFS 19 Target Method: DPX<sup>®</sup> Technology



### **Mobile Phases**

MPA- 0.1% formic acid in water MPB- 0.1% formic acid in ACN Gradient – 98:2 (0-8 min, 50:50 (8-9 min), 2:98 (9-12 min) Agilent Poroshell 120 EC-C18 (2.8 um, 2.1 x`100 mm) 5  $\mu$ L injection, Wash = IPA/MeOH/Water (33:33:33)

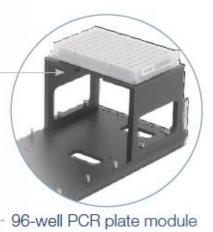


### Full Automation: The Wave of the <u>Future</u>





Hamilton Starlet



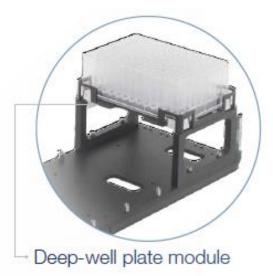


Figure 1: Aspiration from a sample tube

Figure 2: Dispense into a 96-well plate

	19 Targets: DPX Extraction OF	*NSC LOD (ng/m	L) Integra LOD (ng/mL)	Integra (Admin) LOD (ng/mL)	Hamilton Starlet LOD (ng/mL)
1	Amphetamine	20	1.0	20	20
2	Methamphetamine	20	1.0	20	20
3	Carisoprodol	100	1.0	100	100
4	Meprobamate	100	1.0	100	100
5	Diazepam	1.0	1.0	1.0	1.0
6	Nordiazepam	1.0	4.0*	4.0	4.0
7	Alprazolam	1.0	4.0*	4.0	4.0
8	Clonazepam	1.0	4.0*	4.0	4.0
9	Lorazepam	1.0	4.0*	4.0	10^
10	Cocaine	8	1.0	10	10
11	BE	8	1.0	10	10
12	Morphine	5	1.0	5	5
13	Hydrocodone	5	1.0	5	5
14	Oxycodone	5	1.0	5	5
15	Methadone	10	1.0	10	10
16	Zolpidem	10	1.0	10	10
17	6-MAM	2	0.5	0.5	0.5
18	Fentanyl	0.5	1.0	1.0	1.0
19	Buprenorphine	0.5	2.0	2.0	4.0^

## ADFS Cannabinoids LLE Method

### <u>Procedure</u>

- 1. Spike controls in 1 mL of oral fluid/buffer solution.
- 2. Add 50  $\mu$ L of IS Mix (0.1  $\mu$ g/mL) to 500  $\mu$ L of prepared sample.
- 3. Add 200  $\mu L$  of 5% formic acid in water.
- 4. Vortex.
- 5. Add 3 mL of 80:10:10 n-hexane/diethyl ether/ ethyl acetate
- 6. Rotate for 5 min.
- 7. Centrifuge for 10 min.
- 8. Transfer upper layer to a conical tube.
- 9. Dry down @ 45 C°
- 10. Reconstitute with 100  $\mu$ L of 50: 50 MPA/MPB

### Mobile Phases

- MPA- 5 mM Ammonium formate with 0.1% formic acid in water
- MPB- 0.1% formic acid in methanol
- Gradient 50:50 (0-5 min), 35:65 (5-8 min), 5-95 (8-10 min)
- Agilent Poroshell 120 EC-C18 (2.8 um, 2.1 x`100 mm)



## ADFS Cannabinoids LLE Method

Target	*NSC Recommended LOD (ng/mL)	LOD (ng/mL)	LOQs (as of March 2019)
ТНС	2	0.5	1.0
THC-OH	N/A	4.0	4.0
THC-COOH	N/A	1.0	4.0
Cannabinol	N/A	0.2	4.0
Cannabidiol	N/A	0.5	0.5
Cannabigerol	N/A	0.2	4.0

\*Recommendations for Toxicological Investigation of Drug-Impaired Driving and Motor Vehicle Fatalities. JAT 2017.

#### CHEMICAL TEST FOR INTOXICATION

CHAPTER 370-1-1

### The list of field sobriety alcohol screening devices approved for use in Alabama can be found by following the link on the Department's web page at <u>www.adfs.alabama.gov</u>.

Author: Dale A. Carpenter, Mark A. Pevey, Curt E. Harper, and Gregory L. Turner Statutory Authority: §32-6-49.13 <u>Code of Alabama</u>, 1975 as amended. History: Amended: Filed May 19, 2003; adopted July 7, 2003; effective August 11, 2003. Amended: Filed May 10<sup>th</sup>, 2018; effective June 24<sup>th</sup>, 2018.

#### 370-1-1-.06 Field Sobriety Drug Screening Devices

#### (1) Purpose.

This rule list the approved "FIELD BREATHALYZER or OTHER APPROVED DEVICE" as referred to by §32-6-49.13 <u>Code of Alabama</u>, 1975 as amended. Additionally, this rule describes training requirements and minimal operational criteria necessary for accurate and reliable results from oral fluid drug screening devices.

#### (2) Definitions and Abbreviations:

- (a) Drug. Any substance, when taken into the human body, which can impair the ability of a person to operate a vehicle safely.
- (b) Negative Result. A negative result indicate the samples is drug-free for the tested targets or below the cutoff level of the test.
- (c) Observation. For the purpose of this rule, use of the term observation shall mean to watch. Prior to the administration of a field sobriety screening test employing an approved field sobriety drug screening device a subject must be under the observation of the operator of the device or other law enforcement officer for a period of not less than ten minutes.
- (d) Oral Fluid. A clear, tasteless fluid comprised of saliva produced by multiple salivary glands, and other constituents inside the mouth.
- (e) Positive Result. A positive result indicates presence of the drug, its metabolites, or cross-reacting substance but does not indicate level of intoxication, administration route or concentration in the oral fluid. A positive test result should be confirmed by a second test method such as GC/MS (gas chromatography-mass spectrometry) or LC/MS (liquid chromatography-mass spectrometry).

#### CHEMICAL TEST FOR INTOXICATION

#### CHAPTER 370-1-1

(3) Approved Training. Training afforded by the manufacturer of an approved device and/or training received from the Alabama Department of Forensic Sciences shall constitute approved training.

(4) Operation of Devices.

- (a) Officers shall use the device according to the manufacturer's operational procedure.
- (b) Every subject must be under observation by an officer for a period of ten minutes before the screening device test is administered.
- (c) The subject should not eat, drink, or smoke ten minutes prior to giving a sample.
- (5) Quality Control Tests and Maintenance.
- (a) Quality control (QC) tests and annual maintenance shall be conducted per manufacturer's operational procedure.
- (b) The device is working properly if the QC test(s) pass.
- (c) QC test(s) should be conducted at the time of testing or within 24 hours of the subject test.

(6) Training and Maintenance Records. It shall be the responsibility of each law enforcement agency to maintain permanent records documenting the training of each officer in the use of approved field sobriety screening devices and the annual maintenance results on each device in use by the law enforcement agency.

(7) Approved Field Sobriety Drug Screening Device List.

**NOTE:** For the purpose of this rule, variations or enhancements that do not have any bearing on the drug detecting capability of the instrument, such as the addition of a modem, external printer or passive sampling systems are approved.

The list of field sobriety drug screening devices approved for use in Alabama can be found by following the link on the Department's web page at www.adfs.alabama.gov.

Author: Curt E. Harper Statutory Authority: §32-6-49.13 Code of Alabama, 1975 as amended. History: Filed May 10<sup>th</sup>, 2018; effective June 24<sup>th</sup>, 2018.



#### ALABAMA DEPARTMENT OF FORENSIC SCIENCES TOXICOLOGY SPECIMEN COLLECTION - ANTEMORTEM (Living Subjects)

Hoover/Birmingham Regional Laboratory 2026 Valleydale Road, Hoover, AL 35244 Tel (205)-982-9292 Fax (205) 403-2025 https://adfs.alabama.gov/

#### NOTE

The following are recommendations for collecting and submitting specimens for Toxicological Analyses. This kit is intended for the collection of specimens from LIVING SUBJECTS. It is considered best practice to collect both blood and oral fluid. They complement each other and will provide a more complete picture of recent drug use. Blood collection should be witnessed by the investigating officer or by his/her representative who can authenticate the specimens. Oral fluid should be collected by the investigating officer or by his/her representative as close to the arrest or crash as possible (e.g. at roadside).

#### ORAL FLUID SPECIMEN COLLECTION

- 1. Collect the Quantisal oral fluid sample in this order of preference:
  - a. At Roadside (after 10 minute observation period)
  - b. Prior to DRE evaluation (if applicable)
  - c. After DRE evaluation (if applicable)
  - d. At the same time as the blood draw
- 2. Check expiration date on Quantisal packaging and ensure subject has refrained from smoking and consumption of food or beverage for 10 minutes prior to specimen collection.
- Fill out Specimen Security Seal or label with subject's name, date & time of collection, and collector's initials. з.
- Peel back and open package to remove collector. Have subject move tongue side to side to accumulate oral fluid in his/her 4. mouth before starting to speed up the collection. Keep the tip of the device pointed down.
- Instruct subject to position collector (oral absorbent swab) under tongue and close mouth. Keep head down to allow gravity 5. to help with oral fluid collection. Wait until indicator turns BLUE or 10 minutes has elapsed. Note on submission form if indicator turned blue. Collection time may take from 2-10 minutes to collect approximately 1 mL of oral fluid.
- Hold red-capped tube with blue liquid in an upright position and uncap by pushing up with thumb(s). Instruct subject to 6. insert collector (oral absorbent swab) into the uncapped transport tube and replace the cap.
- Snap cap firmly into tube for transport. 7.
- 8. Mix saturated collector (oral absorbent pad) with the blue liquid by gently shaking tube.
- 9. Seal top of collector with evidence tape or specimen security seal. Initial and date seal.

#### Do's:

- Ensure deprivation period of 10 minutes before oral fluid collection. a)
- b) It is recommended to use gloves during sample collection for hygienic purposes.
- C) Collect both oral fluid and blood (seek warrant if necessary).
- d) Ship samples to laboratory as soon as possible.
- Store unused collectors at room temperature. Avoid prolonged exposure to heat/sunlight. e)

#### Don'ts:

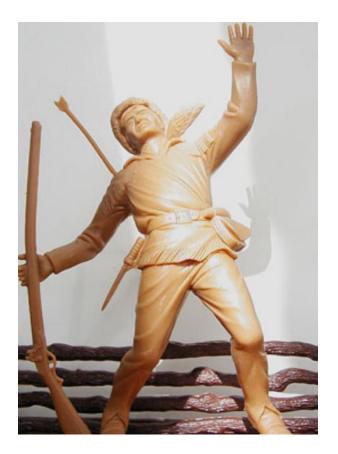
- Ensure subject does not chew or suck on pad, talk, or remove collector from mouth.
- b) Do not stand tube on table. Do not spill or empty liquid from tube.
- Do not consume buffer in the tube or place collector into mouth after it has been in buffer. C)
- d) Collector and subject should not touch the absorbent pad with fingers or environment.

Video Training: https://immunalysis.com/products/oral-fluid/quantisal/collector-training/

KIT SHIPPING SEAL CAT NO BEKIN send sample to laboratory for testing Quantisal® **Gral Fluid Collection Device** with Volume Adequacy Indicator NI LON SECURITY SEAL 

## 1 Year Status Update

- OF/Blood cases received = 63
- # of OF only = (~10%)
- # of Roadside Devices in the field = ~6-8 devices
- # of Roadside Tests = 33%
- # Kits distributed to LEAs = ~300
- 4 Subpoenas, No Testimony
- Officer Feedback
  - Dry mouth ... Doesn't Turn Blue!
  - Consent
  - Cost of Roadside Cartridges, devices



### An Increase in THC Concentrations Over the Years in Alabama

Year	Number of Cases (DUIs)	Average THC Concentration (ng/mL)	Number of Cases (Traffic Fatalities)	Average THC Concentration (ng/mL)
2010	98	4.1	53	8.2
2011	77	3.9	32	8.3
2012	53	3.8	26	7.1
2013	40	3.8	37	6.5
2014	88	4.0	43	7.0
2015	166	4.20	39	13
2016	246	4.8	67	10
2017	233	4.6	88	9.3
2018	208	6.8	66	7.8
2019*	275	5.2	57	7.1

## **OF THC Concentrations**

- Lee 2012
  - THC after use = 68 10,284 ng/mL
  - THC at 6 hrs = 1.3-11.9 ng/mL
  - Carboxy-THC = 0.02 0.76 ng/mL !!!
- Hartman 2015
  - After 15 minutes: Blood (33-68 ng/mL), OF (764-952 ng/mL)
  - After 2 hours: Blood (2-3 ng/mL), OF (33-91 ng/mL)
  - OF THC > 1600 ng/mL indicated within last 1.4 hours
  - OF THC > 600 ng/mL indicated within last 2.3 hours
- Lee 2011
  - Detection Window = 48 hours for 2 ng/mL
  - CBD, CBN, (CBG) useful markers for recent use
- Oral Cavity Contribution, XXX Contamination (Marker of recent use)

### Cannabinoid Concentrations in Blood vs Oral Fluid

	Target: TH	С	Target: Carboxy THC			Target: Hydroxy THC		
	Blood Concentration	OF Concentration		Blood Concentration	OF Concentration		Blood Concentration	OF Concentration
Mean	7.0	113	Mean	80	ND	Mean	4.4	ND
Median	5.6	37	Median	37	ND	Median	3.9	ND
Max	21.0	943	Max	400	ND	Max	11	ND
Min	1.0	0.7	Min	6.2	ND	Min	1.0	ND
(+) Rate	14/20 = 70%	22/24 = 92%	(+) Rate	16/20 = 80%	0/24 = 0%	(+) Rate	14/20 = 70%	0/24 = 0%
Т	arget: Cannak	pinol	Target: Cannabigerol		-	Farget: Cannak	oidiol	
	Blood Concentration	OF Concentration		Blood Concentration	OF Concentration		Blood Concentration	OF Concentration
Mean	1.0	7.1	Mean	5.5	5.5	Mean	1.8	3.7
Median	1.0	2.6	Median	1.6	1.6	Median	1.8	3.7
Max	1.1	25	Max	2.1	24	Max	2.3	3.7
Min	0.9	1.0	Min	0.6	0.8	Min	1.3	3.7
(+) Rate	2/20 = 10%	14/24 = 58%	(+) Rate	6/20 = 30%	13/24 = 54%	(+) Rate	2/20 = 10%	1/24 = 4%

### Methamphetamine & Amphetamine Concentrations in Blood vs Oral Fluid

Та	arget: Methamph	etamine			Target: Ampheta	mine
	Blood Concentration	OF Concentration			Blood Concentration	OF Concentration
Mean	555	1397		Mean	60	336
Median	385	650		Median	51	130
Max	2,200	4,800		Max	210	920
Min	34	12		Min	13	12
(+) Rate	13/16 = 81%	14/15 = 93%	]	(+) Rate	13/16 = 81%	14/15 = 93%

### Alprazolam Concentrations in Blood vs Oral Fluid

Target: Alprazolam					
	Blood Concentration OF Concentration				
Mean	96	20			
Median	79	13			
Max	190	44			
Min	<10	1.2			
(+) Rate	5/6 = 83%	8/8 =100%			

## Anticipated Challenges to OF Drug Testing

- OF conc. correlated to blood conc. and/or impairment?
- Appropriate window of detection?
- Environmental contamination?
- Passive exposure to THC = +OF?
- CBD ingestion = +OF THC?
- Roadside OF screen creates bias to DRE Evaluation?

### Result

But I'm a DRE...and don't' want my Opiates opinion to appear biased. Screened Cannabis Screened Amphetamine Screened Methamphetamine Screened Methadone

## Drugs Detected



06/14/19

005

### **Case Studies**





### Curt E. Harper, Ph.D., F-ABFT

Toxicology Discipline Chief



## Case Study #1 Crystal Meth Wreck

- White, Male, 38 year old
- Fiancé died 4 mo. ago, no one with Narcan to "watch after him"
- Admitted drug use:
  - 10 AM ("half a bag" of meth mixed with "little bit" of heroin)
- Time of Crash: 2:10 PM
- DRE Evaluation/OF Sample: 4:10 PM (~2 hours post-crash)
- Time of Blood Draw: 5:04 PM (~3 hours post-crash)
- DRE Evaluation: Narcotic Analgesic, CNS Stimulant

fiance died Ymonths ago

#### ALABAMA DRUG INFLUENCE EVALUATION

King         29747         18-17-017         Heat           Recorder/Witness         Crash:         None         Arresting	ting Officer's Agency
Recorder/Witness	ting Officer's Agency
Crash, Indie	
	over PO
Arrestee's Name (Last First Middle) Pate of Birth Sex Race Arrestin 19/80 M W	ting Officer (Name_ID#)
Date Examined / Time /Location Date Examined / Time /Location Date Examined / Time /Location Date Examined / Time /Location Miranda Warping Given Miranda Miranda Warping Given Miranda Miranda Miranda	-0506 Chemical Test: Urine Blood Coral Fluid Test or tests refused
Given by: Ning UNO Jurkey Senduich (150	What have you been drinking? How much? Time of last drink? Dr Pegger/Water 1602, 1030
Y Time now/ Actual When did you last sleep? How long? Are you sick or injured?	d? Are you diabetic or epileptic?
1600-1800 1608 Last night 3/13 DYes 10 No	TYes XNo
Y Ves ZNo Ves ZNo DYes ZNo DYes	under the care of a doctor or dentist? $S \square No f / 5 u b \theta \times e^{n}$
Image: Speech:       Are you taking any medication or drugs?       Attitude:         Image: Speech:       Speech:       Cooperation or drugs?	Coordination: Norme
Speech: Cheo-C Breath odor: Bad	Face:
	Blindness: Tracking:
Glasses Contacts, if so Hard Soft Normal Bloodshot Watery	None Left Right Equal Unequal
Pupil Size:     Pupil Contacts, if so     Inalu Soft       Pupil Size:     Pupil Contacts, if so     Inalu Contacts, if so       Pupil Size:     Pupil Contacts, if so     Inalu Contacts, if so       Pupil Size:     Pupil Contacts, if so     Inalu Contacts, if so       Pupil Size:     Pupil Contacts, if so     Inalu Contacts, if so       Pupil Size:     Pupil Contacts, if so     Inalu Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Size:     Pupil Contacts, if so     Pupil Contacts, if so       Pupil Contacts, if so     Pupil Contacts, if so     Pupil Contacts, if soft       Pupil Contacts, if soft     Pupil Contacts, if soft     Pupil Contacts, if soft	Able to follow stimulus Eyelids Normal

			140				- /
	Finger to Nose	PUPIL	Room light	Darkness	Direct	Nasal area:	2
	(Draw lines to spots touched)	SIZE	(2.5 - 5.0)	(5.0 - 8.5)	(2.0 - 4.5)		
S.		Left Eye	2,0	2.0	2.0	Oral cavity:	JeAt
du Bass		Right Eye	2.0	2.0	2.0	cleo r	dry north
Jan 1	2			nd Dilation: es DNo		Nonee	action to Light:
	(A) A SIN		RIGHT	r ARM bilawh		LEFT AF	Meelle track
	5 A	Ę	edle f	11	- ,		- Fa
	head coning down slow Risid	5wolter orr	6140	the puncture	•	With	Swollen
	Blood Pressure Temperature 140/87 98.3 °F Muscle Tone:	ficm needle	- Auto			$\geq$	Covared by bandaid
	Normal Flaccid Rigid						
	What drugs or medications have you been using? Heroin / Meth	How much?	ixed joed	heroin J	Time of use? 1000 om	100	the drugs used? (Location)
	Date / Time of arrest: Time DRE was notified: 7/10/18/15/1 15/5	Evaluation sta	rt time: Éval	luation completion 1704		Subject refused entire Subject stopped parti	e evaluation cipating during evaluation
	Officer's Signature:		roved by / date				DRE #
l	Opinion of Evaluator: Not Impaired Alcoho	ol Depressant	CNS S		Dissociat	ive Anesthetic Analgesic	<ul> <li>Inhalant</li> <li>Cannabis</li> </ul>

UPDATED 3/2018 JP

# Case Study #1: Toxicology Results

Target	Blood (ng/mL)	Oral Fluid (ng/mL)
Methamphetamine	780	>1000 (4,700)
Amphetamine	51	900
Fentanyl	12	52
Morphine	ND*	37
6-MAM	ND	P

\* Qualifier out of tolerance. ~ 5 ng/mL



DRE Evaluation: Narcotic Analgesic, CNS Stimulant

## Case Study #2 Watch the Fence

- White, Male, 24 year old
- Subject drove his vehicle off the road and into a chain link fence.
- Time of Crash: 10:00 PM
- DRE Evaluation/OF Sample: 11:30 PM (1.5 hours post-crash)
- Time of Blood Draw: 12:43 AM (2.75 hours post-crash)
- Alere DDS2 THC, Stimulant, Benzo, Opiates
- Draeger DT5000 THC, Stimulant, \*Invalid
- DRE Evaluation: Cannabis + CNS Stimulant

Comon l	1		ALA'	BAMA				EVA	ALUATOR:		
	D	DRUG INFLUENCE EVALUATION					IACP# 025222 ROLLING LOG#: 16-073				
	CASE NUN	MBERS:						SCR	LIBE: Non	<u>.</u>	
	TYPE EVA	ALUATION:	EN	FORCEM	ENT			WIT	WITNESS:		
ARRESTEE'S NA	ME (Last, Firs	t, Middle)		Date of Bir	th Age	Sex	Race	Arres	ting Officer (Na	me, ID#)	
				2/19/9	2 24	m	W				
Date Examined / T	ime /Location			Breath Resu	ults:	Test F	Refused 🗆			Chemical Test	
12/21/16	12330/1	1c Jail		Results: , (	200	Instru	.menit #: /	099	328 .		Test or tests refused
Miranda Warning		S Yes	What hav	ve you eaten to	oday? Wł				een drinking?	How much?	Time of last drink?
Given By: M. N.	lelson		Mexi	ican	Binner	-	water	<u> </u>	8-10 6.	Hler	notsure
Time now/ Actual		hen did you last			Are you si	ck or inju				tic or epileptic?	
After 2 / 233		6300-04			Yes [		A .		🔲 Yes 🖼 1		
Do you take insuli		<u> </u>	Do y	ou have any p	hysical de	fects?			Are you under	the care of a doo	ctor or dentist?
🔲 Yes 🏽 No				Yes No					🛐 Yes 🗆 N	o normal j	
Are you taking any				Attitud		r				Coordination	
🗃 Yes 🗆 No	Xamx, Traz	Adames Advai	1, Maria	man Coe	Nentive	C	onfor	<u>ed</u>	۹. 	Poor, St	asseria, Stunplinz
Speech:			Breat	ath Odor:				F	ace:		1
Low, this	ck, and sh	vired		A or ma				:		'n./	
Corrective Lenses:	: 🗆 None	Prescription I	think	Eyes: 🗆 Re					Blindness: 1+		Tracking:
	Contacts, if so			□ Normal	Bloo	dshot 2	Watery	8	None 🗖 Left	Right	Equal 🖸 Unequal
Pupil Size: 🖉	Equal					cal Nysta		A	Able to follow st		Eyelids 🖸 Normal
-	Unequal (expla					Yes 🔛	No	1	🖀 Yes 🗖	l No	Droopy

Pulse and time HG	N	Right Eye	Left Eye	Conve	rgence	ONE LE	EG STAND no
1. 132 / 2.344 Lac	k of Smooth Pursuit	Normal	Accoul	Collive	angenee	· @	
	ximum Deviation	normal	normal	(3)		ົ	UR 1
3. 134 / 0014 Ang	gle of Onset	None	none	Right eve	Left.eve		
Romberg Balance Wa	alk and turn test		Cannot keen	balance <u>5</u> ×		LR	
	Aid not	AHempt	Califor Keep			🇊 🖀 Sways wh	nile balancing
	CORECTED C		Starts too soo	n <u>no</u>		🖬 🖪 Uses arms	s to balance
$\downarrow^{1}$ $\downarrow$ $\downarrow$ $\downarrow^{2}$	ا	in the second		I# Nine	2 <sup>nd</sup> Nine	😫 🍘 Hopping	
	TER BER	Torrate	D Stops walkin	<sup>B</sup> IX		🗖 🖀 Puts foot	
			Misses heel-t		1 7	Tremors	Tremons
Ro	led Pante Les UN	° ~ ~	cl Steps off line			<6"	Down IX
2x - symmistic Den	constructe two 2x	Ticmors	Raises arms	<u> </u>		SWAY Arms	
naved formal JEFA /V. 1 "10	nsway hice turn	neine			1/	Hoping	
	AL Feet Algionint		Actual steps	<sup>taken</sup> <b>9</b> /1:	z /		
LLC Internal clock De	scribe Turn		Cannot d	o test (explair	n)	Type of footwea	r:
46 estimated as 30 seconds	NONE			ver, caild		Sock F	cet
Draw lines to spots to	ouched	PUPIL SIZE	Room light	Darkness 5.0 – 8.5	Direct 2.0 - 4.5	Nasal area:	
		Left Eye	2.5 5.0	5.0-0.5	4.9-9.5	Clear	
		Etti Eye	6.0	7.5	4.5	Oral cavity:	1.1
	)) 🕰 🛛 🗆	Right Eye			1	Raixed tost	
	(/ L		6.0	1.5	4,5	Green Catin	
1,0000	1a. I.	and the second	1	REBOUND D		REACTION	N TO LIGHT:
	KIV6 D	pupillary U	arest		Yes 📱 No	2104	

Draw lines to spots touched	PUPIL SIZE Room light 2.5 5.0		rect Nasal are	ea: Elcer
	Left Eye Right Eye 6.0 pupillary Unrest	7,5 4 REBOUND DILATIO	9 Oral cavi Raixe	
	RIGHT	ARM		ARM
Blood pressure 130 / 96 Temperature 180 / 96				
130         96         97.6         0           Muscle tone:         □         Normal         □         Flaccid         Image: Rigid           Comments:         □         Signal         □         Flaccid         Image: Rigid			-	
What drugs or medications have you been using? How $\mathcal{O}_{\mathcal{O}}_{\mathcal{O}_{\mathcal{O}_{\mathcal{O}_{\mathcalO}}}}}}}}}}$	much? c Line, Xunapometer		Chome	gs used? (Location)
Date / Time of arrest: Time DRE was notified:		Evaluation completion tin		
12/22/16 2317 2310	2330	0030	500	
Opinion of Evaluator: Depressant Hallucin Stimulant Dissocia	ogen INarcotic Ana tive Anesthetic Inhalant	algesic 🖉 Cannabis	Alcohol Rule Out	Unable to Determine Category No Impairment
Officer's Signature: Felony O	ffense: NONE	Misdemeanor Offense: $D \circ J$		Reviewed/approved by / date:
				Revised. 06/10

# Case Study #2: Toxicology Results

Target	Blood (ng/mL)	Oral Fluid		
THC	ND	315		
THC-OH	ND	ND		
THC-COOH	18	ND		
Cannabigerol (CBG)	ND	14		
Cannabinol (CBN)	ND	19		
Cannabidiol (CBD)	ND	ND		
Methamphetamine	560	>1000		
Amphetamine	55	>1000		
Cocaine	ND	66		
BE	75	456		
Alprazolam	150	>1400		
Morphine	ND*	1470		

### Specimen taken proximate to time of driving: How close?

- Roadside immediately after arrest
- Prior to DRE evaluation
- After DRE evaluation



• After search warrant, simultaneously with blood at hospital

- Considerations
  - Consent
  - Implied Consent Law
  - Search Warrant
    - Birchfield v. North Dakota
  - Case Law
  - Discuss with your TSRP or Local DA



### Case Study #3 Know your Limitations

- White, Male, 28 year old
- Struck a drive-thru while operating vehicle
- Time of Crash: 11:15 PM
- DRE Evaluation/OF Sample: 12:25 AM (~1 hours post-crash)
- Time of Blood Draw: 11:53 PM (0.75 hours post-crash)
- Alere DDS2 Negative
- Draeger DT5000 Negative
- DRE Evaluation: Depressant

# Case Study #3 - Toxicology Results

Target	Blood (ng/mL)	Oral Fluid (ng/mL)
Diazepam	180	ND*
Nordiazepam	97	4*
Zolpidem	390	130

\* Alere DDS2 & Draeger DT5000 were negative.

\* Highlights limitations of OF roadside screening and confirmation testing.



#### **Our Future Aims**

- Hamilton Starlet Extraction Automation = OF, Blood methods
- In the process of evaluating DrugWipe
- Newly designed BSKits = 2,500 per year.
- Quantitative results for cannabinoids
- Postmortem:
  - Traffic Fatalities, Suspected ODs

Journal of Analytical Toxicology, 2019;1–8 doi: 10.1093/jat/bkz032 Article

#### Article

Oral Cavity Fluid as an Investigative Approach for Qualitative and Quantitative Evaluations of Drugs in Postmortem Subjects

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#### Abstract

A relatively overlooked aspect of forensic science is the potential of oral cavity fluid for contributing to a forensic diagnosis. Although traditional specimens, like blood and urine, are routinely evaluated for forensic toxicology testing, fluid from the oral cavity has not been investigated as a matrix in postmortem cases. Our laboratory developed and validated qualitative and quantitative analytical methods for determining 47 medicinal and illicit drugs from oral cavity fluid. These developed methods aimed to compare results from liquid chromatography coupled with tandem mass spectrometry (LC-MS/MS) analyses of oral cavity fluid to those of traditional matrices collected from the same postmortem subjects. Of 34 cadavers studied, 32 (including two decomposed and two drowned subjects) had detectable and quantifiable drugs in the oral cavity fluid and/or blood, urine, bile, vitreous fluid and/or liver tissue. The most significant finding was that 6-acetylmorphine (6-AM) was detected more frequently in oral cavity fluid (11 cases) than in blood and urine combined (6 cases). Compounds with a short window of detection, like the heroin metabolite, 6-AM and even heroin, could be detected more readily in oral cavity fluid than in urine. In 2017, the incidence of heroin-related overdose deaths increased to 15,958. Those data have shed light on the practicality of testing oral cavity fluid postmortem and its significance in forensic toxicology. In conclusion, this study showed that oral cavity fluid could be useful for detecting and quantifying drugs in postmortem subjects; moreover, oral cavity fluid may be particularly suitable when other matrices are limited or difficult to collect, due to body condition or putrefaction