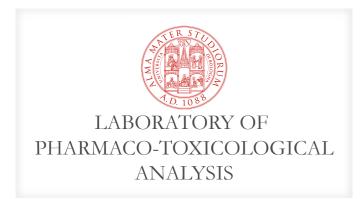
THERAPEUTIC DRUG MONITORING (TDM) BY MEANS OF NOVEL SAMPLING AND EXTRACTION PROCEDURES: A COMPARATIVE STUDY

LAURA MERCOLINI¹, MICHELE PROTTI¹, LAWRENCE J. ALBERS² AND CHRISTOPHER REIST²

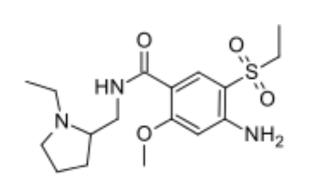


¹ Laboratory of Pharmaco-Toxicological Analysis, Department of Pharmacy and Biotechnology, Alma Mater Studiorum - University of Bologna, Via Belmeloro 6, 40126 Bologna, (Italy) ² Mental Health, Veterans Affairs Medical Center, 5901 East Seventh Street, 90822 Long Beach, CA (USA) email: laura.mercolini@unibo.it



THERAPEUTIC DRUG MONITORING (TDM)

The therapeutic drug monitoring (TDM) is designed to study important correlations between the administered drug dosage and the related levels found in biological matrices, to better understand therapeutic effects, side effects and toxicity. In this work TDM is applied to psychiatric patients undergoing therapy with central nervous system drugs, particularly with atypical antipsychotics. The biological matrices most commonly used for TDM purpose are blood, plasma and serum but some crucial issues concerning their sampling and pretreatment are still present

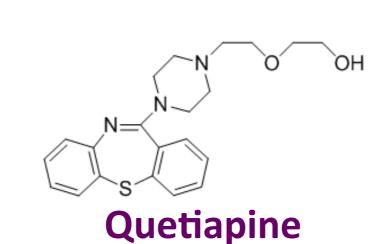


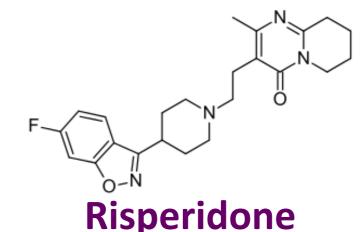
Amisulpride





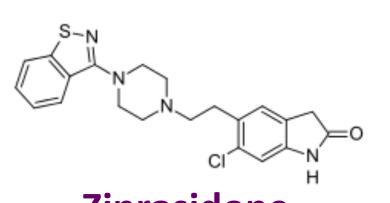






PLASMA

SERUM



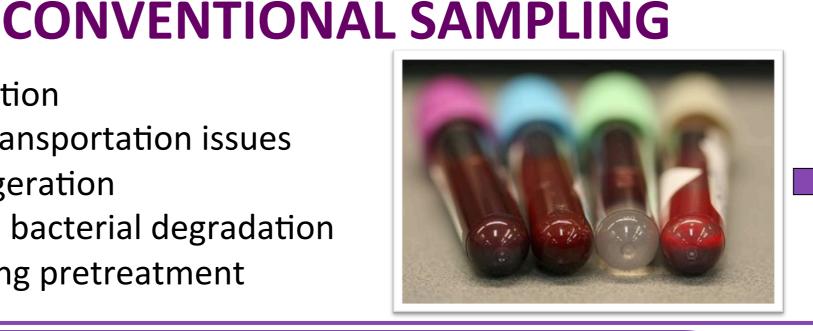
Ziprasidone



BLOOD

Invasive collection

- Storage and transportation issues
- Need for refrigeration
- Enzymatic and bacterial degradation
- Time consuming pretreatment



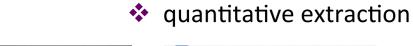
- After-sampling processes
- Careful handling
- Biological hazard
- Risk for sample contamination
- High cost process



DRIED MATRIX SPOTS

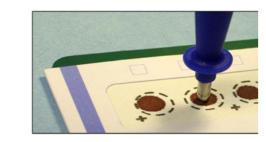
Paper-like support issuse related to:

matrix effect









VOLUMETRIC

BLOOD: Capillary blood from a finger prick blotted onto a card PLASMA/SERUM: Collection of a sample fixed volume of $(10 \mu L)$

AIM OF THE RESEARCH

This study proposed an innovative and alternative approach for the TDM of psychiatric patients treated with some atypical antipsychotic drugs



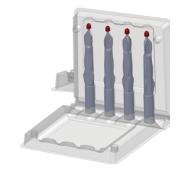
10 μL) of matrix by means of

VOLUMETRIC ABSORPTIVE MICROSAMPLING and **DRIED MATRIX SPOTS**

VOLUMETRIC ABSORPTIVE MICROSAMPLING







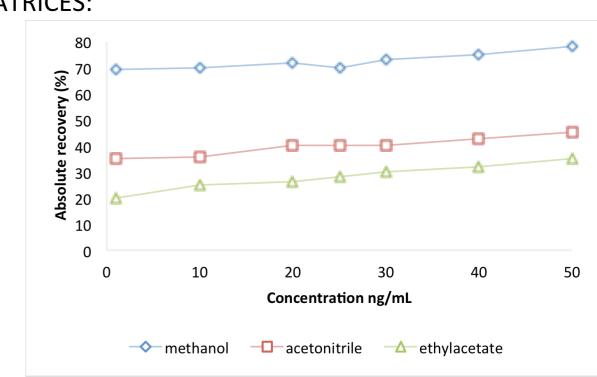
Polymeric tip which absorbs an accurate sample volume of (10 μL) by wicking for 2 seconds and drying at RT

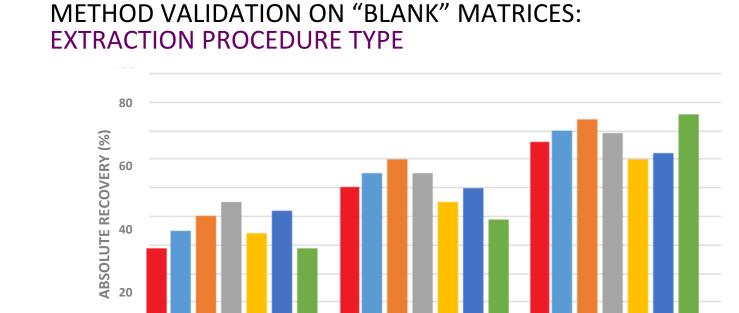
METHOD VALIDATION ON "BLANK" MATRICES: **ABSOLUTE RECOVERY (%)**

DRIED MATRIX SPOTS	BLOOD (DBS)	PLASMA (DPS)	SERUM (DSS)
AMISULPRIDE	65	65	64
ARIPIPRAZOLE	69	67	66
CLOZAPINE	64	65	64
OLANZAPINE	65	62	63
QUETIAPINE	60	62	61
RISPERIDONE	61	60	62
7IPRASIDONE	62	63	60

ABSORPTIVE MICROSAMPLING	FROM BLOOD	FROM PLASMA	FROM SERUM
AMISULPRIDE	67	70	70
ARIPIPRAZOLE	70	71	70
CLOZAPINE	73	72	72
OLANZAPINE	69	70	69
QUETIAPINE	60	60	61
RISPERIDONE	63	64	55
ZIPRASIDONE	77	76	74

METHOD VALIDATION ON "BLANK" MATRICES: **EXTRACTION SOLVENT TESTED EXTRACTION SOLVENTS:** methanol acetonitrile





USA = ultrasound agitation (5 min) MA = microwaves (15 sec) COMBO = USA (5 min) + MA (15 sec)

- AMISULPRIDE
- **ARIPIPRAZOLE CLOZAPINE**
- **OLANZAPINE**
- QUETIAPINE RISPERIDONE
- ZIPRASIDONE

COMPARATIVE STUDY ON "BLANK" MATRICES:

CONVENTIONAL SAMPLING = liquid-liquid extraction **DRIED MATRIX SPOTS** = DBS, DPS, DSS **VOLUMETRIC ABSORPTIVE**

ABSOLUTE RECOVERY (%)

ethylacetate

MICROSAMPLING = from blood, plasma, serum Conventional sampling **Dried Matrix Spots**

COMPARATIVE STUDY ON "REAL" SAMPLES: TDM OF PSYCHIATRIC PATIENTS

CONVENTIONAL SAMPLING	BLOOD	PLASMA	SERUM
PATIENT 1 amisulpride	320 ng/mL	318 ng/mL	320 ng/mL
PATIENT 2 aripiprazole	120 ng/mL	124 ng/mL	119 ng/mL
PATIENT 3 clozapine	400 ng/mL	410 ng/mL	397 ng/mL
PATIENT 4 olanzapine	22 ng/mL	23 ng/mL	21 ng/mL
PATIENT 5 quetiapine	110 ng/mL	112 ng/mL	111 ng/mL
PATIENT 6 risperidone	23 ng/mL	22 ng/mL	22 ng/mL
PATIENT 7 ziprasidone	140 ng/mL	138 ng/mL	139 ng/mL

DMS	BLOOD (DBS)	PLASMA (DPS)	SERUM (DSS)
PATIENT 1 amisulpride	321 ng/mL	320 ng/mL	319 ng/mL
PATIENT 2 aripiprazole	123 ng/mL	124 ng/mL	120 ng/mL
PATIENT 3 clozapine	410 ng/mL	415 ng/mL	403 ng/mL
PATIENT 4 olanzapine	22 ng/mL	24 ng/mL	23 ng/mL
PATIENT 5 quetiapine	115 ng/mL	112 ng/mL	113 ng/mL
PATIENT 6 risperidone	24 ng/mL	22 ng/mL	24 ng/mL
PATIENT 7 ziprasidone	148 ng/mL	141 ng/mL	141 ng/mL

VAMS	FROM BLOOD	FROM PLASMA	FROM SERUM
PATIENT 1 amisulpride	323 ng/mL	321 ng/mL	320 ng/mL
PATIENT 2 aripiprazole	125 ng/mL	124 ng/mL	119 ng/mL
PATIENT 3 clozapine	408 ng/mL	412 ng/mL	400 ng/mL
PATIENT 4 olanzapine	21 ng/mL	21 ng/mL	20 ng/mL
PATIENT 5 quetiapine	116 ng/mL	112 ng/mL	114 ng/mL
PATIENT 6 risperidone	22 ng/mL	22 ng/mL	24 ng/mL
PATIENT 7 ziprasidone	144 ng/mL	139 ng/mL	143 ng/mL

CONCLUSION

The present method was validated on "blank" matrices and then applied to "real" samples from psychiatric patients treated with atypical antipsychotics. The TDM has been performed on dried fluids obtained as dried matrix samples and volumetric absorptive microsamples (blood, plasma and serum).

The use of dried microvolumes has proved to be able to grant the golden standard features for drug determination, therapeutic and side effect correlations, but allowing to overcome issues and weaknesses due to traditional analyses.

