### PART 1: PK IN PATIENTS REQUIRING HEMODIALYSIS

Arthur J. Atkinson, Jr., M.D.

**Adjunct Professor** 

Department of Molecular Pharmacology and Biochemistry Feinberg School of Medicine Northwestern University

# FIRST DESCRIPTION OF HEMODIALYSIS IN ANIMALS\*

Copy of the face page of an article from Abel JJ, et al. J Pharmacol Exp Ther 1914:5:275-317.

Dr. John J. Abel did pioneering work in hemodialysis.

## WILLEM J. KOLFF, M.D. (1911 - )

Photograph of Dr. Willem J. Kolff, developer of the first functioning artificial kidney (1943).

### **ELIMINATION BY DIFFERENT ROUTES**

Chart showing elimination by renal, hepatic and dialysis routes using blood flow, afferent concentration, efferent concentration and eliminated drug as measurements.

## IMPACT OF CL<sub>D</sub>

Formula showing that CLR, CLNR and CLD are additive.

### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE
MECHANISTIC – RENKIN APPROACH
EMPIRICAL

EMPIRICAL
FICK EQUATION
RECOVERY CLEARANCE
CLINICAL STUDIES OF DIALYSIS PK
MODEL PROSPECTIVE STUDY
TREATMENT OF DRUG TOXICITY
PHYSIOLOGIC CHANGES DURING DIALYSIS
USE OF KINETIC METHODS FOR ANALYSIS
PATHOPHYSIOLOGIC CONSEQUENCES

### EUGENE RENKIN PROFESSOR EMERITUS AT UC DAVIS

Photograph of Eugene Renkin.

## **RENKIN DIALYSIS EQUATION\***

Equation showing dialyzer blood flow and permeability-surface area product of dialysis membrane.

Neglects: Boundary effects, ultrafiltration.

\* From Renkin EM. Tr Am Soc Artific Organs 1956;2:102-5

# DETERMINANTS OF PERMEABILITY TERM (P or P $\cdot$ S)

- DIALYZER MEMBRANE CHARACTERISTICS
  - MEMBRANE SURFACE AREA
  - MEMBRANE THICKNESS
  - MEMBRANE POROSITY
- DRUG BINDING TO PLASMA PROTEINS
- SOLUTE SIZE AND DIFFUSIVITY

## DIALYZER PERMEABILITY VS. FREE WATER DIFFUSION COEFFICIENTS

#### **PROCAINAMIDE/NAPA**:

RATIO OF DIALYZER PERMEABILITY COEFFICIENTS\*  $1.29 \pm 0.22$ 

RATIO OF FREE WATER
DIFFUSION COEFFICIENTS 1.23

<sup>\*</sup> From Gibson TP et al. Clin Pharmacol Ther 1976;20:720-6.

# DIALYSIS CLEARANCE VS. DIALYZER BLOOD FLOW\*

Chart showing dialysis clearance vs. dialyzer blood flow and the impact of P.S. values for urea (high), creatine, phosphate, and phenol red (low).

### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE
MECHANISTIC - RENKIN APPROACH
EMPIRICAL
FICK EQUATION
RECOVERY CLEARANCE
CLINICAL STUDIES OF DIALYSIS PK
MODEL PROSPECTIVE STUDY
TREATMENT OF DRUG TOXICITY
PHYSIOLOGIC CHANGES DURING DIALYSIS
USE OF KINETIC METHODS FOR ANALYSIS
PATHOPHYSIOLOGIC CONSEQUENCES

## DATA SOURCES FOR FICK EQUATION

Illustration of these sources in a dialysis machine.

## FICK EQUATION FOR CLEARANCE

Q = DIALYZER BLOOD FLOW

**A = CONCENTRATION IN BLOOD COMING TO DIALYZER** 

**V = CONCENTRATION IN BLOOD LEAVING DIALYZER** 

E = EXTRACTION RATIO

### **EXTRACTION RATIO**

The Renkin Equation and the Fick Equation terms for extraction ratio.

### **RECOVERY CLEARANCE**

The gold standard equation for clearance.

**U = DIALYSATE CONCENTRATION** 

V = DIALYSATE VOLUME

t = DIALYSIS TIME

**P = MEAN PLASMA CONCENTRATION** 

### TWO DIALYSIS MYTHS

- NEED TO USE BLOOD CONCENTRATIONS WHEN CALCULATING BLOOD CLEARANCE

BUT PLASMA CONCENTRATIONS PROPORTIONAL TO BLOOD CONCENTRATIONS, SO MAKES NO DIFFERENCE IN A/ [A + V] RATIO

- NEED TO USE PLASMA FLOW WHEN CALCULATING PLASMA CLEARANCE

### PLASMA VS. BLOOD CLEARANCE

Equation showing recovery and equation showing the Fick approach.

### NAPA IN RBC IS DIALYZED

Chart comparing flow parameters.

\* QEFF = [(1 - Hct) + (RBC/P) (HCT)] QMEAS

# DIALYSIS SATURATION VS. RECOVERY CLEARANCE

Formula for dialysis saturation and formula for recovery clearance.

### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE
MECHANISTIC - RENKIN APPROACH
EMPIRICAL
FICK EQUATION
RECOVERY CLEARANCE
CLINICAL STUDIES OF DIALYSIS PK
MODEL PROSPECTIVE STUDY
TREATMENT OF DRUG TOXICITY
PHYSIOLOGIC CHANGES DURING DIALYSIS
USE OF KINETIC METHODS FOR ANALYSIS
PATHOPHYSIOLOGIC CONSEQUENCES

# DATA SOURCES FOR RIGOROUS PK ANALYSIS

Chart illustrating this with a dialysis machine. Measurement of recovered drug is essential.

# KINETIC MODEL USED TO ANALYZE HEMODIALYSIS DATA\*

Chart of 3-compartment model.

\* From Stec GP, et al. Clin Pharmacol Ther 1979;26:618-28.

# KINETIC MODEL USED TO ANALYZE HEMODIALYSIS DATA\*

Chart showing 3-compartment model and dialysis machine.

\* From Stec GP, et al. Clin Pharmacol Ther 1979;26:618-28.

## FICK CLEARANCE EQUATION

## TWO PROBLEMS WITH FIXED-PARAMETER MODEL\*

Chart illustrating these two problems.

DURING DIALYSIS: [A] AND [V] DROP MORE
THAN EXPECTED FROM DRUG RECOVERY
AFTER DIALYSIS: CONCENTRATION
REBOUND IS LESS THAN EXPECTED

<sup>\*</sup> From Stec GP, et al. Clin Pharmacol Ther 1979;26:618-28.

# KINETIC MODEL USED TO ANALYZE HEMODIALYSIS DATA\*

Chart showing 3-compartment model and dialysis machine.

\* From Stec GP, et al. Clin Pharmacol Ther 1979;26:618-28.

# REDUCTION IN CL<sub>s</sub> DURING AND AFTER HEMODIALYSIS\*

Charts illustrating reduction in slow intercompartmental clearance.

\* From Stec GP, et al. Clin Pharmacol Ther 1979;26:618-28.

### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE
MECHANISTIC - RENKIN APPROACH
EMPIRICAL
FICK EQUATION
RECOVERY CLEARANCE
CLINICAL STUDIES OF DIALYSIS PK
MODEL PROSPECTIVE STUDY
TREATMENT OF DRUG TOXICITY
PHYSIOLOGIC CHANGES DURING DIALYSIS
USE OF KINETIC METHODS FOR ANALYSIS
PATHOPHYSIOLOGIC CONSEQUENCES

### **CASE HISTORY**

A 67 year-old woman became lethargic and confused and developed hypotension, renal insufficiency, junctional tachycardia and intraventricular conduction delay after ingesting an estimated 7gm of procainamide (PA). Plasma PA and NAPA concentrations were 57  $\mu$ g/mL and 55  $\mu$ g/mL, respectively.

## **CASE HISTORY (cont.)**

Hemodialysis was performed for 4 hr. By the end of the second hour BP was maintained in the range of 110/80 mm Hg without vasopressor therapy. At the end of dialysis, the patient was alert and oriented although only 340 mg of PA and 470 mg of NAPA had been removed by this procedure.

## **DIALYSIS CASE HISTORY (cont.)**

Fifteen hours after dialysis, PA and NAPA levels were 9.2  $\mu g/mL$  and 33  $\mu g/mL$ , respectively. The patient had returned to normal sinus rhythm with QRS = 0.12 sec.

## KINETIC ANALYSIS OF HEMODIALYSIS FOR PROCAINAMIDE TOXICITY\*

Chart illustrating this analysis and drug removal during dialysis.

\* From: Atkinson AJ Jr, et al. Clin Pharmacol Ther 1976;20:585-92.

### **CRITERION FOR DIALYSIS EFFICACY\***

Formula showing that dialysis clearance must be greater than 30% of total organ clearance to be effective.

\* Levy G. Am J Med 1977;62:461-5.

#### WAS DIALYSIS EFFICACIOUS?

- DIALYSIS INCREASED DRUG CLEARANCE

PA – TWO FOLD

NAPA – 3.8 FOLD

- BUT 4 hr OF DIALYSIS REMOVED < 1 gm of 7 gm DOSE

340 mg PA

470 mg NAPA

- HOWEVER, BLOOD LEVELS FELL SUBSTANTIALLY

PA: 25.7 μg/mL NAPA: 47.0 μg/mL

15.5 µg/mL

35.5 μg/mL

**AND PATIENT'S CONDITION STABILIZED** 

### PA & NAPA KINETICS IN TOXIC PATIENT

Chart showing a lower VD-beta in toxic patient.

### ESTIMATION OF $V_d$

Question: Why was distribution volume estimate so much lower in patient than in normal subjects?

Formulas comparing the usual with the dialysis estimates.

#### SEQUESTRATION OF DRUG IN SOMATIC TISSUES

Chart illustrating this effect with a 3-compartment model.

## EFFICACY OF EXTRACORPOREAL TREATMENT OF DRUG TOXICITY

- TOTAL EXTENT OF DRUG REMOVAL MAY BE COMPROMIZED BY  $\downarrow$  CLS.
- ↓ CLS FROM SOMATIC TISSUES CAN ACCELERATE ↓ IN DRUG CONCENTRATION TO WHICH VITAL ORGANS (CNS, HEART) ARE EXPOSED AND RESULT IN A BENEFICIAL CLINICAL RESPONSE > EXTENT OF DRUG REMOVAL.
- ↓ CLS FROM SOMATIC TISSUES ALSO ATTENUATES POST-DIALYSIS REBOUND.

#### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE MECHANISTIC - RENKIN APPROACH EMPIRICAL

EMPIRICAL

FICK EQUATION

RECOVERY CLEARANCE

CLINICAL STUDIES OF DIALYSIS PK

MODEL PROSPECTIVE STUDY

TREATMENT OF DRUG TOXICITY

PHYSIOLOGIC CHANGES DURING DIALYSIS

USE OF KINETIC METHODS FOR ANALYSIS

PATHOPHYSIOLOGIC CONSEQUENCES

#### WHY DOES $CL_S \downarrow$ DURING DIALYSIS?

**POSSIBILITIES:** 

CAPILLARY BLOOD FLOW DECREASES CAPILLARY P x S PRODUCT DECREASES BOTH DECREASE

### **RENKIN EQUATION\***

Q = capillary blood flow P = capillary permeability coefficient-surface area product (sometimes denoted P x S).

<sup>\*</sup> From Renkin EM. Am J Physiol 1953;183:125-36.

### MULTICOMPARTMENTAL MODEL OF INULIN AND UREA KINETICS\*

Illustration of this model.

<sup>\*</sup> From Atkinson AJ Jr, et al. Trends Pharmacol Sci 1991;12:96-101.

# BASIS FOR KINETIC HETEROGENETIY OF INTERSTITIAL FLUID SPACE

Chart comparing effective pore size with capillary structure and primary location in splanchnic and somatic tissues.

# ENDOTHELIAL FENESTRAE IN HEPATIC SINUSOIDS

Photomicrograph.

# INTERENDOTHELIAL CELL JUNCTION IN CONTINUOUS CAPILLARY

Photomicrograph.

## UREA (x) AND INULIN (▲) KINETICS DURING AND AFTER HEMODIALYSIS\*

Chart illustrating the kinetics during and after hemodialysis.

\* From Bowsher DJ, et al. J Lab Clin Med 1985;105:489-97.

#### EFFECT OF MOLECULAR WEIGHT (M) ON SOLUTE DIFFUSIVITY (D)\*

Chart illustrating this activity.

\* From Henderson LW: *In*: Brenner BM, Rector FC Jr. The Kidney. 1976, p. 1643-71.

#### RELATIONSHIP BETWEEN BLOOD FLOW (Q) AND $\operatorname{CL}_{\rm I}$

Chart illustrating this relationship.

\* From Bowsher DJ, et al. J Lab Clin Med 1985;105:489-97.

#### UREA AND INULIN KINETICS DURING AND AFTER HEMODIALYSIS

Chart showing the flow and permeability parameters before, during and after.

### RENIN-ANGIOTENSIN SYSTEM ACTIVATION DURING AND AFTER HEMODIALYSIS\*

Chart illustrating this system activation during and after hemodialysis.

\* From Bowsher DJ, et al. J Lab Clin Med 1985;105:489-97.

## DIFFERENT MICROCIRCULATORY ACTIONS OF ANGIOTENSIN II AND AVP\*

Chart illustrating actions of angiotensin II and AVP.

\* From Atkinson AJ Jr: The Pharmacologist 1989;31:229-34.

#### EFFECT OF ARGININE VASOPRESSIN (AVP) ON P• S\*

Chart illustrating this effect.

\* From Atkinson AJ Jr: The Pharmacologist 1989;31:229-34.

#### **GOALS OF DIALYSIS DISCUSSION**

DISCUSSION OF DIALYSIS CLEARANCE MECHANISTIC - RENKIN APPROACH EMPIRICAL

EMPIRICAL
FICK EQUATION
RECOVERY CLEARANCE
CLINICAL STUDIES OF DIALYSIS PK
MODEL PROSPECTIVE STUDY
TREATMENT OF DRUG TOXICITY
PHYSIOLOGIC CHANGES DURING DIALYSIS
USE OF KINETIC METHODS FOR ANALYSIS
PATHOPHYSIOLOGIC CONSEQUENCES

## HEMODIALYSIS-ASSOCATED SKELETAL MUSCLE CRAMPS

- COMPLICATE MORE THAN 20% OF HEMODIALYSIS SESSIONS
- OCCUR MORE FREQUENTLY IN SOME PATIENTS THAN OTHERS
- PATHOGENESIS UNCLEAR

- SYMPTOMATIC THERAPY: NaCI, MANNITOL

- PREVENTIVE THERAPY: NaCI INFUSION

## RESPONSE OF CRAMPING AND NONCRAMPING PATIENTS TO TILT\*

Chart illustrating this response.

\* Kaplan B et al.: Int J Clin Pharmacol Ther Toxicol 1992;30:173-80.

# ACTIONS OF ANGIOTENSIN II & SYMPATHETIC NERVOUS SYSTEM

Chart illustrating these actions.

### ONLY SOME PATIENTS HAVE DIALYSIS-ASSOCIATED SKELETAL MUSCLE CRAMPS\*

Chart comparing dialysis reaction with normal and cramper and noncramper.

\* Sidhom OA, et al. Clin Pharmacol Ther 1994;56:445-51

#### AUGUST KROGH 1920 NOBEL LAUREATE

Photo of August Krogh.

# CROSS SECTION OF MUSCLE SHOWING OPEN (O) & CLOSED (x) CAPILLARIES\*

Chart illustrating this cross section.

\*From Krogh A. Nobel Lecture, December 11, 1920.

# CAPILLARY DERECRUITMENT (OPEN (O) & CLOSED (x) CAPILLARIES)

8 OPEN CAPILLARIES IN MUSCLE CROSS SECTION.

### PATHOGENESIS OF DIALYSIS-ASSOCIATED SKELETAL MUSCLE CRAMPS

Chart illustrating the basis of dialysis-associated skeletal muscle cramps.

#### **CONCLUDING THOUGHT**

ALTHOUGH NON-COMPARTMENTAL ANALYSIS OF PK DATA IS CURRENTLY IN VOGUE, IT IS UNABLE TO PROVIDE INSIGHT INTO SOME IMPORTANT PHENOMENA:

- IMPACT OF DIALYSIS-ASSOCIATED HEMODYNAMIC CHANGES (↓ CLS)
- IMPACT OF ↓ SPLANCHNIC BLOOD FLOW (↓ CLF) ON BIOAVAILABILITY

# RELATIONSHIP BETWEEN $\operatorname{CL_F}$ AND EXTENT OF NAPA ABSORPTION\*

Chart sowing this directly proportional relationship.