

Acute Complications of Hemodialysis

新光急診 張志華醫師

Intradialytic hypotension

- **Definition:** A decrease in systolic BP ≥ 20 mm Hg or a decrease in MAP ≥ 10 mm Hg associated with symptoms
- **Complication:** cardiac arrhythmias, coronary and/or cerebral ischemic events
- **Long-term side effects:** volume overload due to suboptimal ultrafiltration, LVH, and interdialytic hypertension
- **A third of dialysis patients**

Risk Factors of Dialysis Hypotension

- Low body mass
- Poor nutritional status and hypoalbuminemia
- Severe anemia
- Advanced age (Age > 65 years old)
- Cardiovascular disease
- Large interdialysis weight gain
- Low blood pressure (predialysis systolic BP <100 mm Hg)

Etiology of Dialysis Hypotension (I)

- **Excessive rate and degree of ultrafiltration**
- **Inappropriate peripheral venodilation**
- **Autonomic dysfunction**
- **Inadequate vasoconstrictor secretion**

Etiology of Dialysis Hypotension (II)

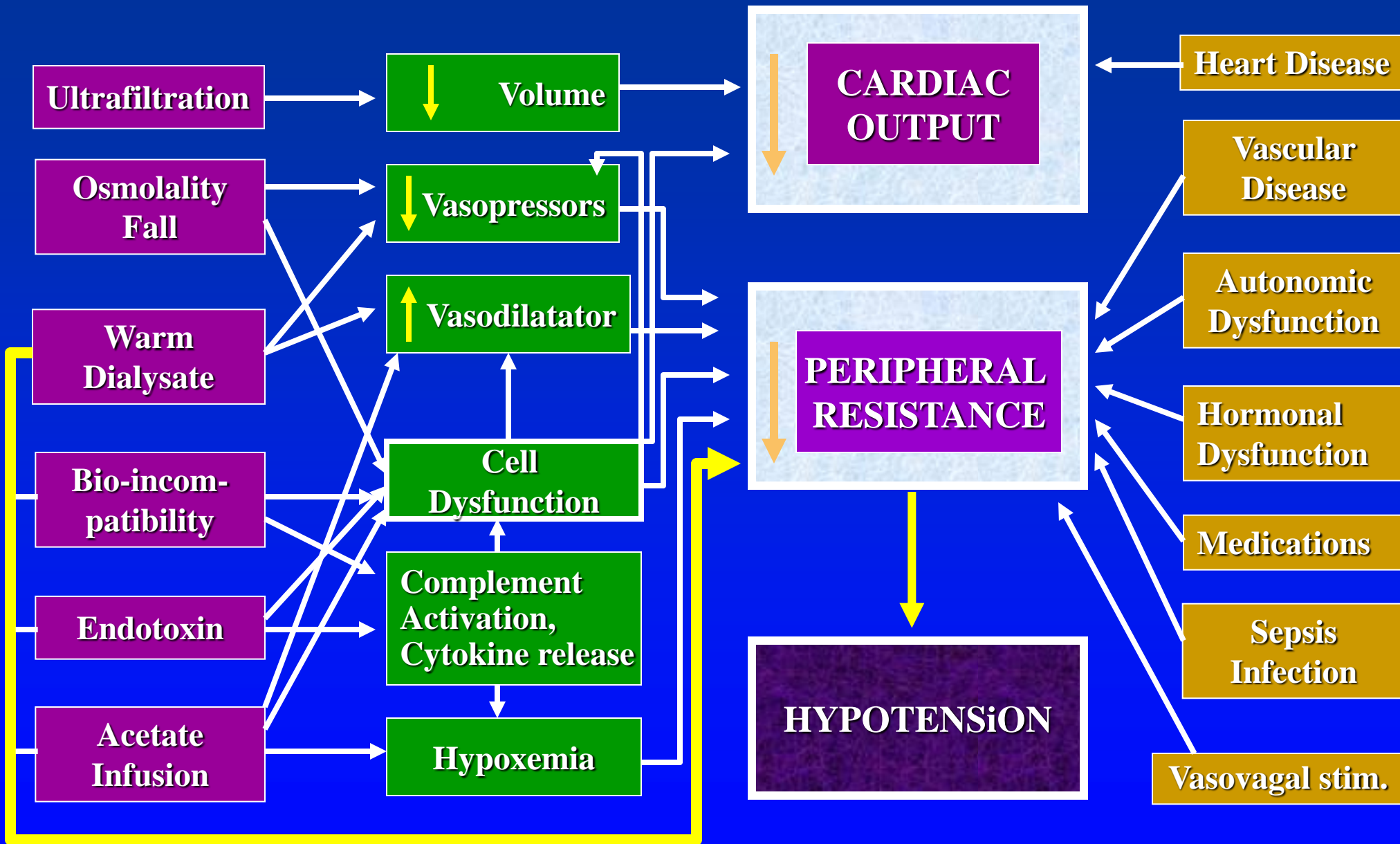
- **Acetate dialysate**
- **Low calcium dialysate**
- **Eat shortly before dialysis**
- **Antihypertensive medications**
- **LV dysfunction**

PATHOGENESIS

MEDIATORS

PATHOPHYSIOLOGY

PATIENT



Prevention and Management of Dialysis Hypotension (I)

- Limiting sodium intake
- Minimize **interdialytic weight gain** by education
- Blood sugar control
- Slow ultrafiltration
- Sodium modeling
- Raise dialysate calcium
- Lower dialysate temperature

Prevention and Management of Dialysis Hypotension (II)

- Switch to CAPD
- Hyperoncotic albumin
- Nasal oxygen
- Mannitol infusion

Prevention and Management of Dialysis Hypotension (III)

- L-Carnitine therapy
- Sertraline
- Midodrine (midorine 2.5 mg)
1# Bid ~ 2# Tid, max: 40 mg/d
- Blood transfusion or erythropoietin therapy
- Volume expansion
- Vasoconstrictor

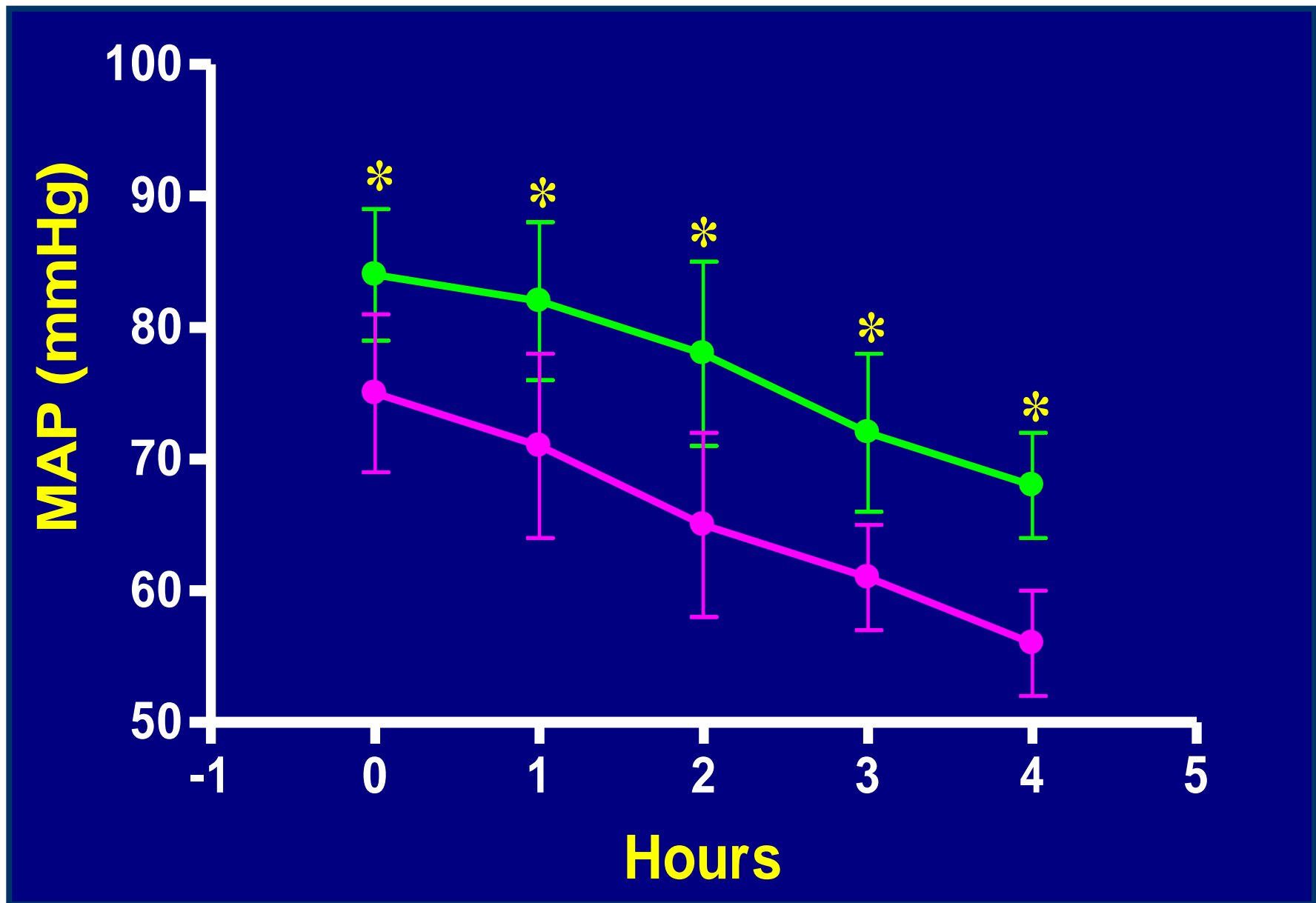


Figure. Serial changes in MAP HD before (●) and after (●)midodrine therapy.

Muscle Cramps

- **35-86%** of hemodialysis patients
- **Lower extremities**
- **Mechanisms: Rapid ultrafiltration, **Intradialytic hypotension**, tissue hypoxia**
- **Treatment: Quinine, Vit E, L-carnitine, Creatine monohydrate, sodium modeling, hypertonic solution**

Acute Allergic Reaction

- First use syndrome
- Burning retrosternal pain
- Diffuse heat, cold perspiration, **urticaria**, **pruritus**, laryngeal stridor, bronchospasm,
loss of consciousness
- Polyurethane function as a reservoir for ethylene oxide

Uremic Pruritus (I)

- 50-90% of dialysis patients
- Risk: male, high serum BUN, Ca, P, β 2-microglobulin, duration of dialysis
- Diagnositc criteria

 - 1 Pruritus appears shortly before the onset of dialysis, or at any time, without evidence of any other active disease that could explain the pruritus.
 - 2 more than or equal to three episodes of itch during a period of <2 weeks, with the symptom appearing a few times a day, lasting at least few minutes, and troubling the patient.
 - 3 Appearance of an itch in a regular pattern during a period of 6 months, but less frequently than listed above.

Causes of itching in ESRD

(1) *Uremia related*

- (a) Uremic itching
- (b) Xerosis
- (c) Anemia of chronic kidney disease
- (d) Secondary hyperparathyroidism

(2) *Uremia unrelated*

- (a) Drug-induced hypersensitivity
 - (b) Senility
 - (c) Hepatitis
 - (d) Diabetes mellitus
 - (e) Hypothyroidism
 - (f) Iron-deficiency anemia
 - (g) Lymphoproliferative/solid tumors
 - (h) Hypercalcemic states
-

Uremic Pruritus (II)

- Optimize the dialysis dose
- Treat anemia
- Treat 2nd hyperparathyroidism
- Ultraviolet B phototherapy
- **Topical emollients**
- Capsaicin
- **Antihistamine**
- **Anti-serotonin agents**

Arrhythmia (I)

- 30-48% of dialysis patients
- Risk factor:
 - ▲ Compromised myocardium: CAD, Intermycardiocytic fibrosis, Pericarditis
 - ▲ Increased **QT interval** or dispersion

Arrhythmia (II)

- ▲ Electrolyte imbalance: **hypokalemia**, **hyperkalemia**, hypercalcemia, hypermagnesemia
- ▲ Anemia
- ▲ Increased LV mass
- ▲ Advanced age
- ▲ Acetate dialysate

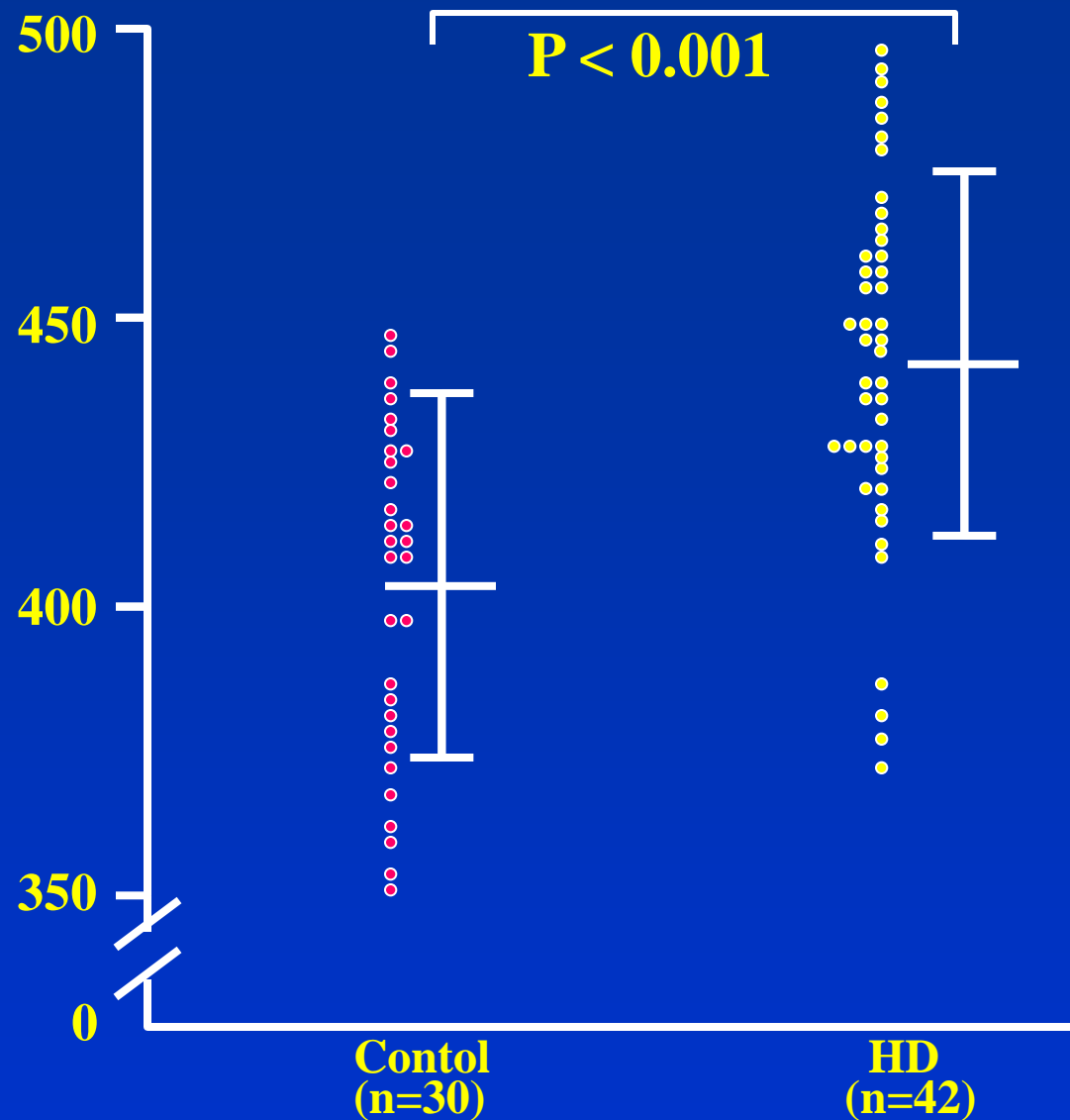


Fig. Distribution of QTc values among hemodialysis patients and controls. The mean value of QTc was significantly increased in hemodialysis patients (432.6 ± 24.9 ms) compared controls (402.0 ± 21.0 ms) ($p < 0.01$)

Results of 24-Hour Holter ECG Monitoring

Arrhythmias Seen	No. of Tapes (%)
Ventricular ectopic beats (> 20/hr)	15 (24)
Ventricular ectopic beats (> 100/hr)	2 (3)
Episodes of ventricular tachycardia	5 (8)
Episodes of supraventricular tachycardia	2 (3)
Episodic atrial fibrillation	7 (11)
Heart block (intermittent)	1 (1.6)

Bleeding During Dialysis (I)

- Platelet dysfunction
- Impaired dense granule release of ATP and serotonin
- Reduced synthesis of thromboxane A2
- Elevated platelet cytosolic cAMP and calcium
- Impaired aggregation response

Bleeding During Dialysis (II)

- Altered adhesive fibrinogen and vWf
- Impaired fibrinogen receptor (GPIIbIIIa)
function
- Uremic toxin or inhibitors
- Erythropoietin augments GPIIbIIIa

Bleeding During Dialysis (III)

- Pack RBC
- Cryoprecipitate, FFP(VIII/vWF)
- Desmopressin (DDAVP) 4mcg/ml/Amp
0.3mcg/kg in N/S 100ml over 15~30 min
- Estrogen

Air Embolism

- 1 ml/kg air may be fatal
- Occlude RV outflow tract and pulmonary vascular bed
- Thromboxane B₂, endothelin
- Trendelenburg position with left side down
- Withdrawal of air from RA
- **Hyperbaric oxygen**

Dialysis Pericarditis I

- **Uremic pericarditis:**
pericarditis before RRT or within 8 wks of its initiation
- **Dialysis pericarditis:**
 ≥ 8 wks after initiation of RRT
- **Incidence of dialysis pericarditis: 2-12%**
- **Etiology: inadequate dialysis, volume overload, infection, autoimmune, drugs**

Dialysis Pericarditis II

- Precordial pain, hypotension, dyspnea, fever, weight gain
- Heparin free dialysis
- Intensive dialysis
- NSAID
- Subxiphoid pericardiostomy

Dialysis Disequilibrium (I)

- **Headache, vomiting, seizure, delirium**
- Rapid correction of marked azotemia
- Cerebral swelling
- Reverse urea effect
- Acidosis of the CSF

Dialysis Disequilibrium (II)

- Shorten the duration
- Lower dialyzer blood flow
- Less efficient dialyzer
- Osmotic agents, high sodium
- IV diazepam

Hypokalemia

- Loss into dialysate, **alkali therapy**
- Renal or extrarenal losses
- **Arrhythmia**, hypotension, fatigue, weakness, **paralysis**
- **CAD**, **digitalis**, **hypercalcemia**, **hypomagnesemia**,
meta alkalosis
- Adjust **dialysate potassium** and buffer

Hyperkalemia

- Dietary intake
- GI bleeding
- Overheated or hypotonic dialysate
- Medications
- Metabolic acidosis

Hypophosphatemia

- Intensive dialysis
- Phosphorus binders
- Reduced intake
- Dysfunction of erythrocytes, CNS, skeletal and cardiac muscle
- Phosphorus rich food

Hypercalcemia (I)

- Liberation of calcium from bone
- Intradialytic gain
- Phosphorus binders
- Widespread use of calcitriol
- Aluminum poisoning

Hypercalcemia (II)

- Low dialysate calcium
- Phosphorus binders during meals
- Discontinue vitamin D Therapy
- Treat aluminum toxicity
- Pamidronate

Endotoxin

- **Bacterial infections**
- **Header sepsis syndrome:**
waterborne *Xanthomonas*–induced fevers
- **Pyrogens**

Hypertensive Emergencies

- Paradoxical, hypertensive response
- Rise in plasma catecholamine
- Activation of renin-angiotensin system
- Antihypertensive withdrawal
- Tx: Sublingual captopril and nifedipine

Bowel Ischemia

- Abdominal pain, acute diarrhea
- Dialysis **hypotension**
- Digitalis, β blockers
- Occlusive and **non-occlusive** infarction (25~60%)
- Heart: Congestive heart failure, arrhythmia (**Af**)
- Hyperkalemia, acidemia, leukocytosis
- Dx: **Inappropriate pain**, elevated LDH and CPK

THANK YOU