

Problem-based Nephrology Orientation

腎臟科 王國彥



基本住院需知

- Check I/O , body weight QD
- 病況穩定的病人，減少IV的量，洗腎病患以40cc/hr為上限
- Regular F/U renal function and electrolyte
- 避免腎毒性的藥物，或依腎功能調節劑量
- 初次血液透析病患，需確認解釋及permit之完成
- 使用3%NaCl需精密計算劑量，check serum Na至少q4h
- 減少不必要的輸血
- 腎功能不全的病人應儘量避免使用含鎂antacid, NSAID, aminoglycoside, and Demerol, Fleet enema, 除非病人已長期無尿。即使已經洗腎的病人若餘尿 >100cc/day 也禁用aminoglycoside and NSAID 除非主治醫師同意。



Introduction Case

- 54y/o female , underlying DM and chronic renal insufficiency with OPD control for more than 8 years
- Baseline renal function 3 months ago : BUN/Cr 45/2.4
- Complaint progressive nausea , vomiting and general weakness in recent 2 weeks
- Besides , decreased urine output was also noted
- PE : bilateral lower leg edema , basal rales(+)
- Lab : Hb 6.4gm/dl MCV 85 BUN/Cr 108/7.5

What's your next step ?



Admission order for this patient

Exam :

- CBC/DC/Plt
- MAR : 加測 **iCa** , **P** , **iPTH** , **Iron profile**
- EKG , CXR
- ABG
- U/R , S/R
- **Check retinopathy**
- 24hr urine for total protein , Cr
- Record I/O , BW QD
- Specific survey cause of renal failure :
 - Renal echo
 - **FeNa**

Medication :

- Avoid nephrotoxic agent
- **Dose adjustment according to GFR**
- Hypertension control
- Sugar control
- Diuretics
- **EPO ?**
- **Sodium bicarbonate ?**
- **Dialysis ?**



Why should we check retinopathy ?

IDDM

- Onset usually well known and progress stage by stage.
- HTN always after stage 3.
- **>90%** with retinopathy
- Non-DM renal disease was low

NIDDM

- Onset usually not known and may present in ant stage.
- HTN may precede nephropathy.
- **60%** with retinopathy.
- Non-DM renal disease was high.



Differential Dx of ARF

- DDx between acute renal failure & acute on chronic renal failure :
 - Renal anemia
 - Renal echo
 - Size : CRF may have **smaller** long axis(<8cm , exception : *DM , multiple myeloma , polycystic kidney disease , collagen storage disease*)
 - Echogenicity : **increased** in CRF and acute inflammation
 - Cortex width : thin(<2mm) in CRF
 - Structure abnormality
- Classification : Pre-renal , intrinsic and post-renal



Fractional Excretion of Na

- The FeNa evaluates only the fraction of filtrated Na that is excreted and is not affected by changes in water reabsorption.

$$FENa = \frac{UNa \times Pcr}{PNa \times Ucr} \times 100 \%$$



DDx in ARF

Type	U/A	UNa	FENa	BUN/Cr
Pre-renal	↑ SG	<20	<1%	>20:1
ATN	↓ SG	>40	>1%	<20:1
Vascular	Hematuria	>20	>1%	<20:1
GN	Proteinuria, RBC cast	>20	> 1%	<20:1
Interstitial nephritis	Hematuria, WBC cast	>20	>1%	<20:1
Post-renal	Hematuria or normal	>20	>1%	< 20:1



Dose Adjustment According to GFR

Drugs not requiring adjustment :

- Antibiotic: clindamycin, doxycycline, nafcillin
- Anticoagulant: heparin, warfarin
- Anticonvulsants: tegretal, phenytoin, deparkin
- Anti-fungal agent: ketoconazole, miconazole
- Anti-TB: rifampin, INH



Drugs Use in Renal Failure

■ Oral hypoglycemic agent :

■ Sulfonylurea: glibenclamide (Daonil), gliclazide (Diamicon), glimepiride (Amaryl) → ↓ elimination in renal failure *May Induce hypoglycemia in renal failure*

■ Gliquidone (Glurenorm), NovoNorm (Repaglinide) → hepatic metabolism → no accumulation in renal failure

■ Metformin (Bentomin) → *metabolic acidosis in severe renal failure*

• Antibiotics :

■ β -lactam : potent convulsant → *reduce dose*

■ Aminoglycoside & Vancomycin : nephrotoxicity → *never use unless starting dialysis*

■ Imipenem: watch for neurological complication



Drugs Use in Renal Failure

- Analgesic agent :
 - NSAID : nephrotoxicity → never use unless starting dialysis
 - Demerol : metabolite accumulate in brain may induce conscious change → avoid use in renal failure

- Others :
 - Muscle relaxant: baclofen (baclon) → contraindicated in uremia patients
 - Antiviral agents: acyclovir (zovirax), gancyclovir, valaciclovir (valtrex), amantadine (PK-Merz) → ↓ to 1/3 to 1/7 dosage



Diuretics

- Thiazide & spironolactone have only little effect when $C_{cr} < 30$ cc/min (serum Cr about 2-2.5 mg/dl)
 - Spironolactone → no use in hyperkalemia
 - Thiazide → no use in hypercalcemia
- **Loop diuretic** is drug of choice



ACEI or ARB

- ACEI or ARB (angiotensin II receptor blocker) → drug of choice in hypertension with proteinuria
- Hyperkalemia: a common side effect, especially in CRI or CRF
 - Serum Cr > 3 → use with caution
 - Serum Cr > 6 → stop use , unless dialysis
 - Serum Cr 3-6 → F/U potassium, kalimate / diuretics use



When to use EPO & NaHCO₃ ?

- EPO :

- Indication : Cr > 6.0mg/dl with Hct < 31% or Hb < 11gm/dl
- Target : Hct 33~35%
- Dosage :
 - Darbepoetin alfa (Aranesp , 25mg) 1Amp QW → used in pre-dialysis stage & PD
 - Erythropoietin beta (Recommen) → used in HD room (EPO as HD room routine)

- NaHCO₃ :

- Indication : Chronic renal failure with metabolic acidosis
- Reserve for ABG : PH < 7.2 , HCO₃ < 16



When is the time of blood transfusion ?

- Hct < 20%
- Hct < 24% + Recognized symptoms or signs due to anemia (angina , dyspnea , weakness , hemodynamics change)
- The Epoetin-resistant patient who has chronic blood loss



值班時 ~

- This patient is dyspnea with chest tightness
- Vital sign : T/P/R 37.1 °C /105/25
- ABG : Respiratory alkalosis & metabolic acidosis
- EKG : sinus tachycardia
- PE : basal rales(+)
- CXR : bilateral pulmonary edema

What's your next step ?



Dyspnea

- Differential Dx :
 - Cardiogenic
 - Pulmonary
 - Metabolic
 - Anemia
 - Psychiatric
- Routine check : *checked before you call for help*
 - PE : crackles , rales , rhonchi , murmur
 - ABG
 - CXR
 - EKG
 - Cardiac enzyme



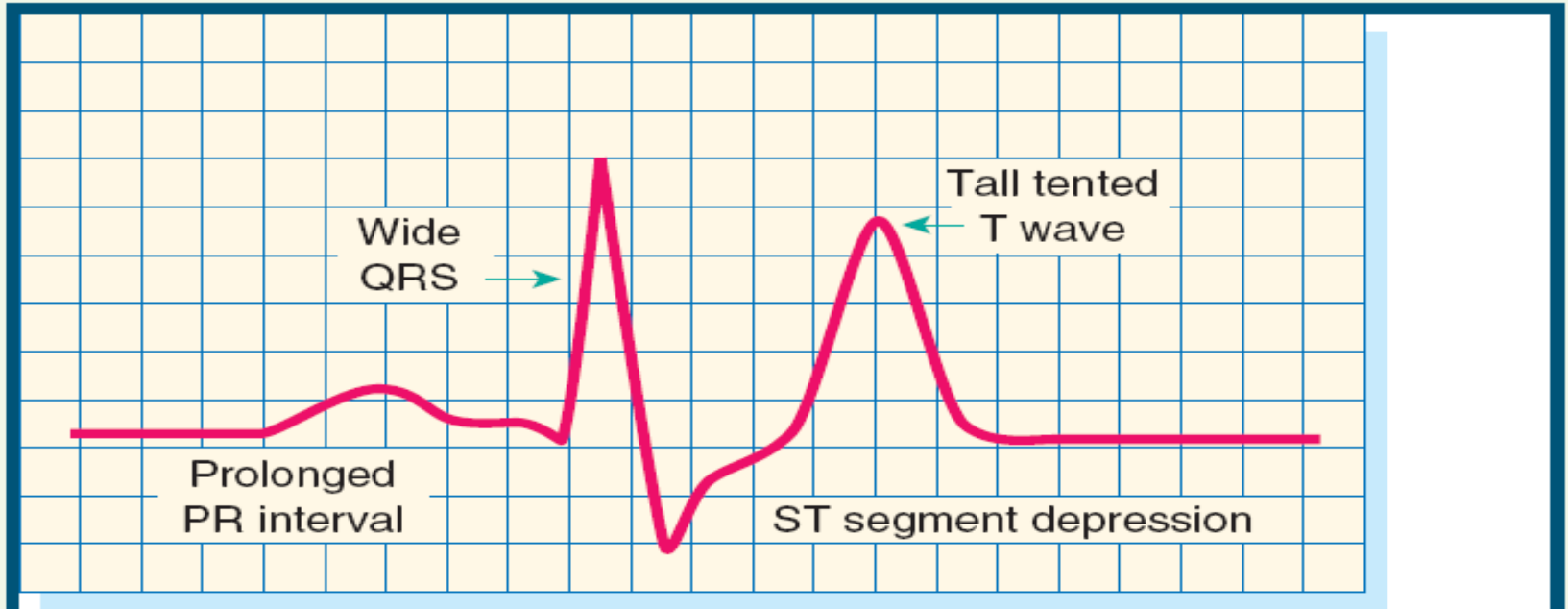
When to start dialysis ?

~ It's time to call CR

- Fluid overloading refractory to medication
- Hyperkalemia ($K > 6 - 6.5$ meq/L) : **medical treatment first**
- Severe metabolic acidosis
- Uremic symptoms (nausea, vomiting, conscious change, seizure)
- Uremic Pericarditis
- Uremic encephalopathy



ECG Change in Hyperkalemia



Electrocardiographic changes resulting from an elevated serum potassium level.

Medical Treatment of Hyperkalemia

- Ca gluconate
- NaHCO₃
- Insulin and 25-50% glucose
- β₂ agonist inhalation
- Diuretic
- Cation exchange resins (Kalimate)
- Dialysis



Medical Treatment

- Calcium gluconate
 - EKG有變化時
 - 10% 10cc (1支) 滴 2-3 分鐘
 - 5-10分鐘後如果沒效可再打一支
 - 藥效短 (30-60分鐘)
- Insulin + glucose
 - 10-20U RI + 25-50g glucose (50-100cc D50W)
 - 15-30分鐘開始作用, 可維持數小時
- Loop and thiazide diuretics

- NaHCO₃
 - 嚴重高血鉀合併酸血症
- β₂-agonist
 - 靜脈注射或吸入
 - 30分鐘作用, 持續2-4小時
- Kaysalate :
 - Remove K via GI tract
 - Kalimate 2pk po TID or 4pk in N/S 100cc enema

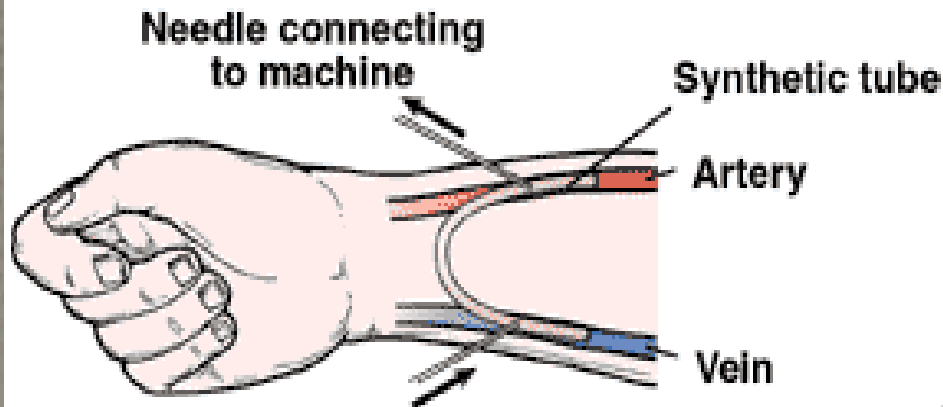


How to dialysis ?

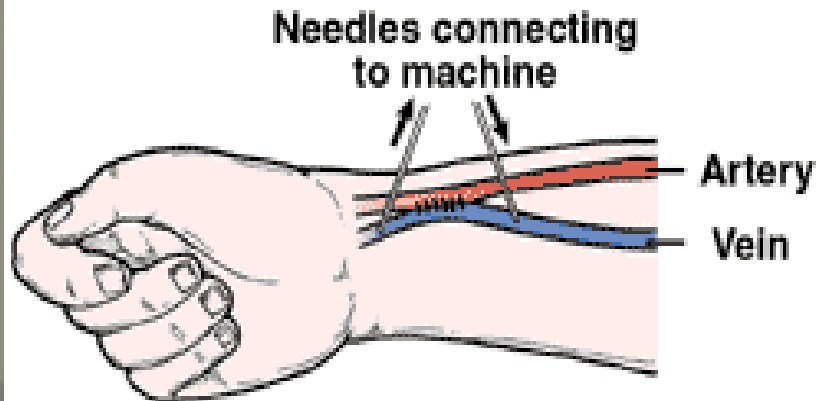
- Hemodialysis
 - AV graft : 2~3週可使用, 平均使用3~5年
 - AV fistula : 4~6週可使用, 平均使用5~7年
 - Perm-Cath : inserted by CVS , 馬上可用
 - Double lumen : inserted by CR , 馬上可用, 使用期為一個月
- Peritoneal dialysis : inserted by GS , 10天~2週可用
 - If PD started in < 10 days following catheter placement,
→do low-volume, supine dialysis.



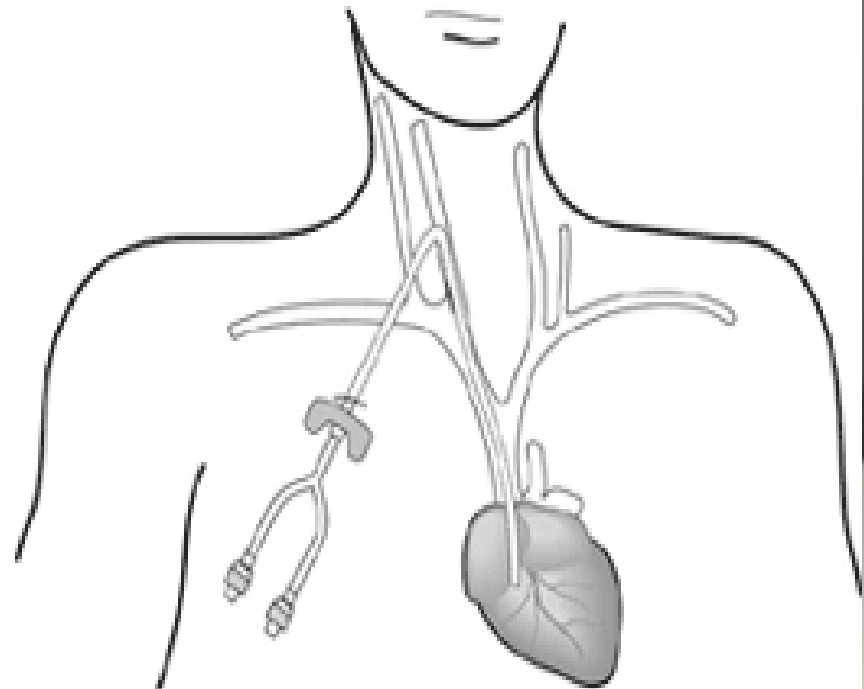
Types of Access for Dialysis



Graft

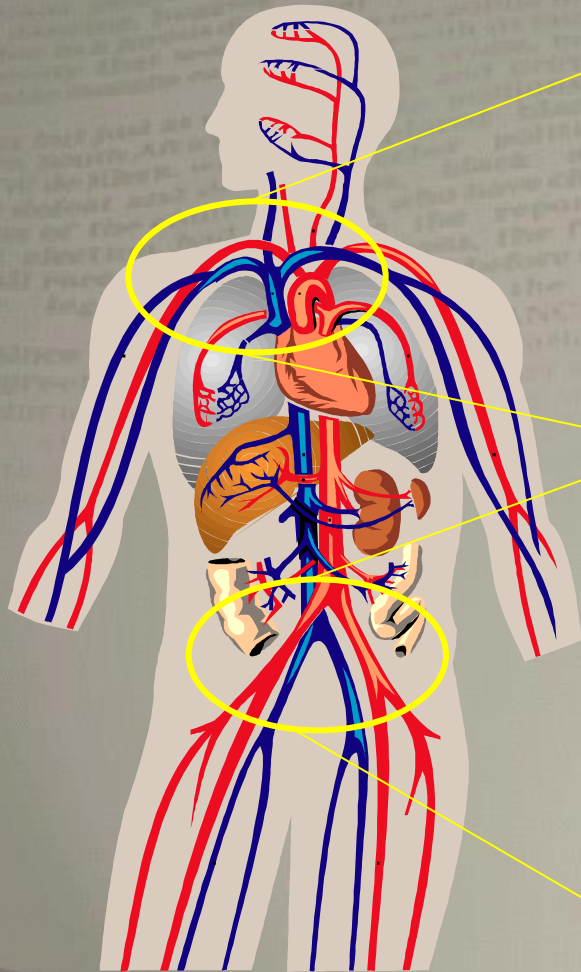


Fistula

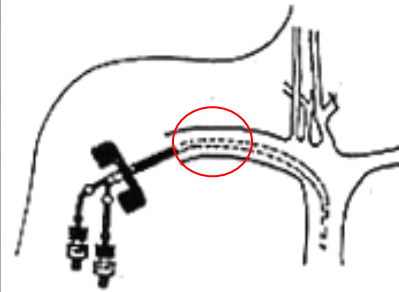


Catheter in neck

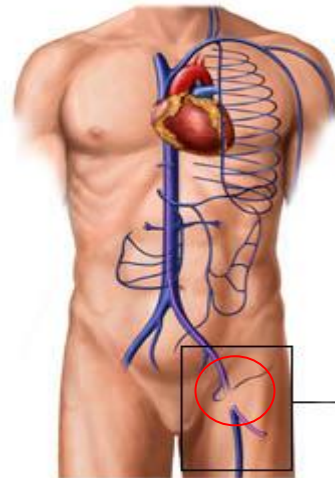
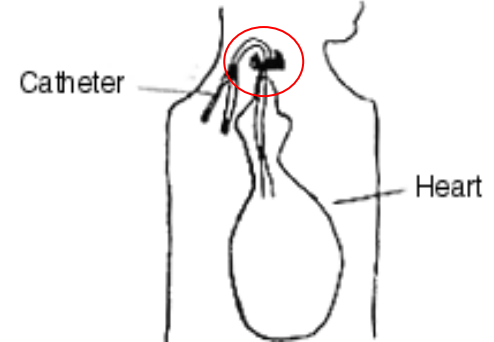
Temporary vascular access



Subclavian Catheter

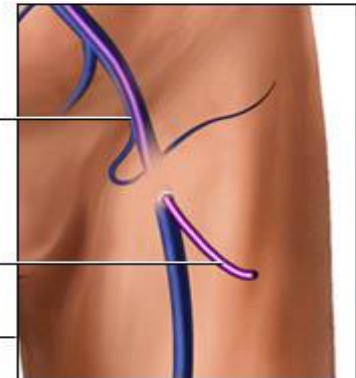


Internal Jugular Catheter

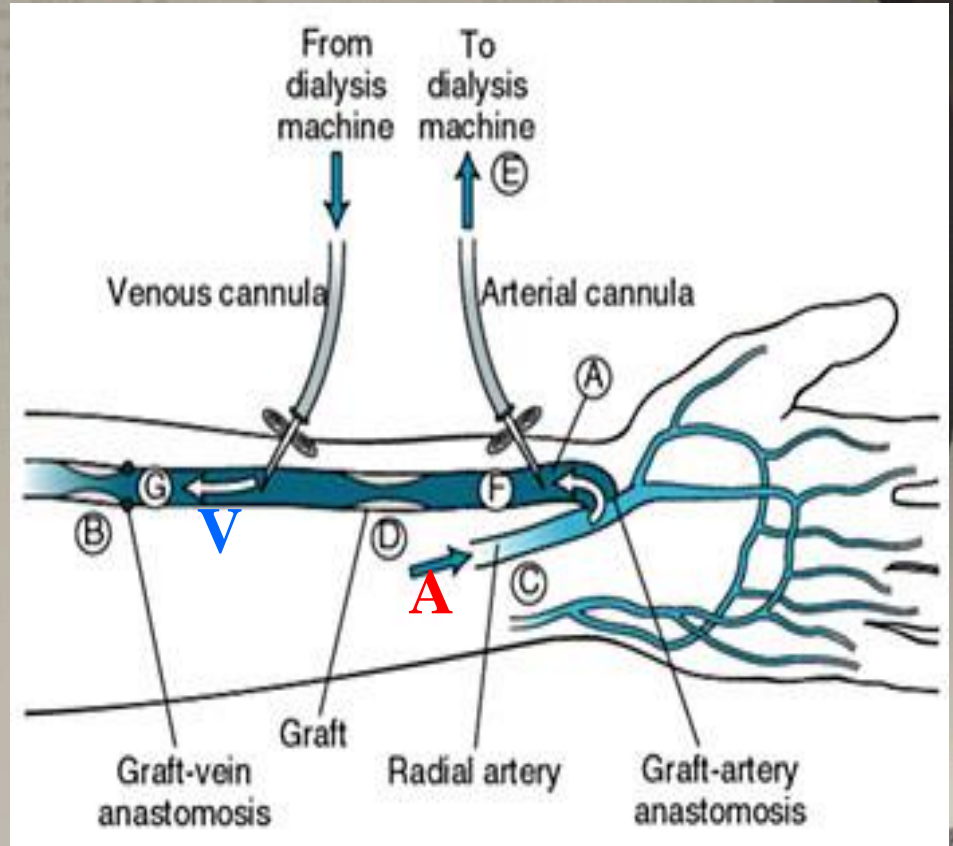
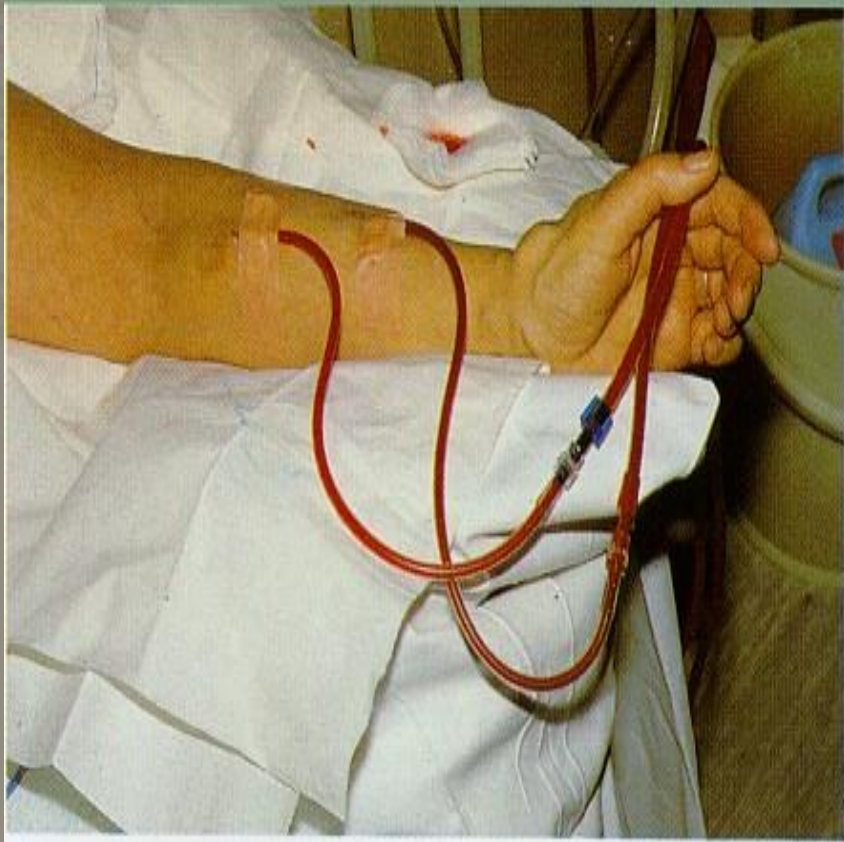


Femoral vein

Catheter



Standard of A-V Fistula



After HD ~

- Patient felt comfortable and no more dyspnea was complained
- Double lumen insertion site blood oozing even after compression for 30mins

What's your next step ?



Evaluation of vital sign

- Tachycardia +/- shock
 - Clinical evidence of volume depletion
 - Fluid challenge
 - Blood transfusion → use whole blood if shock is present
- No tachycardia , no shock



What is the cause of bleeding tendency ?

- Anti-coagulant use (local or systemic) during HD
 - Plt count normal , PT normal , aPTT prolong
 - Antagnoist use : Protamine , FFP transfusion
- Uremic bleeding
 - Plt count normal , PT normal , aPTT normal
 - Cause :
 - Platelet : abnormal Ca flux ↓ ADP and serotonin ,dense granula uremic toxin: iPTH
 - Platelet–vessel wall interaction : altered adhesion and vW Factor
 - Management



Therapeutic Strategies for Uremic Bleeding

TX	Dosage	Start	Peak	Recommend use
DDAVP	0.3 μ /kg	1 hour	2- 4 hours	4Amp in N/S 100cc drip >30min st
Cryoprecipitate	10 Units	1 hour	4 - 12 hours	2U TID
Platelet	12-24 Units			12-24 Units st
EPO	50-150 U/kg			As previous use
Estrogen	0.6/kg/d x 5 days	6 hour	5 - 7 days	Premarin 1# po TID



Sudden onset of conscious change ~

- After HD , she felt nausea and vomit some food substance
- Sudden onset of conscious change with coma status was noted by nurse

What's your next step ?



Step by step

- Always check vital sign at first
- DDX of conscious change un post-HD patient :
 - Shock → iv , inotropic agent use
 - Arrhythmia → ACLS
 - Stroke → hemorrhagic & non-hemorrhagic
 - Electrolyte unbalance → check Na , K , iCa , P
 - Dialysis Disequilibrium Syndrome(DDS)



Dialysis Disequilibrium Syndrome (DDS)

- Most in first dialysis with high BUN level
- Risk factors : old age or children , previous brain damage , severe metabolic syndrome
- Clinical features : neusea , headache , hypertension , conscious change , seizure , coma , death
- Pathophysiology : faster removal of BUN in blood & slower removal in CSF
 - Water shift to brain
 - Brain swelling or osmolytes accumulate
- Management : mannitol 100~250mg iv st



