

1837
2017
YEARS



Γ' ΠΑΘΟΛΟΓΙΚΗ ΚΛΙΝΙΚΗ ΙΑΤΡΙΚΗΣ ΣΧΟΛΗΣ
ΕΘΝΙΚΟΥ ΚΑΙ ΚΑΠΟΔΙΣΤΡΙΑΚΟΥ ΠΑΝΕΠΙΣΤΗΜΙΟΥ ΑΘΗΝΩΝ
ΔΙΕΥΘΥΝΤΗΣ: Καθηγητής Κων/νος Νικ. Συρίγος, MD, PhD, FCCP



ΟΓΚΟΛΟΓΙΚΑ ΕΠΕΙΓΟΝΤΑ

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Γ' Πανεπιστημιακή Παθολογική Κλινική,
Νοσοκομείο «Σωτηρία»

Ανδρας καπνιστής 62 ετών

**Αναφέρει οίδημα προσώπου, τραχήλου από ημερών,
άλγος στον θώρακα, ήπια δύσπνοια**

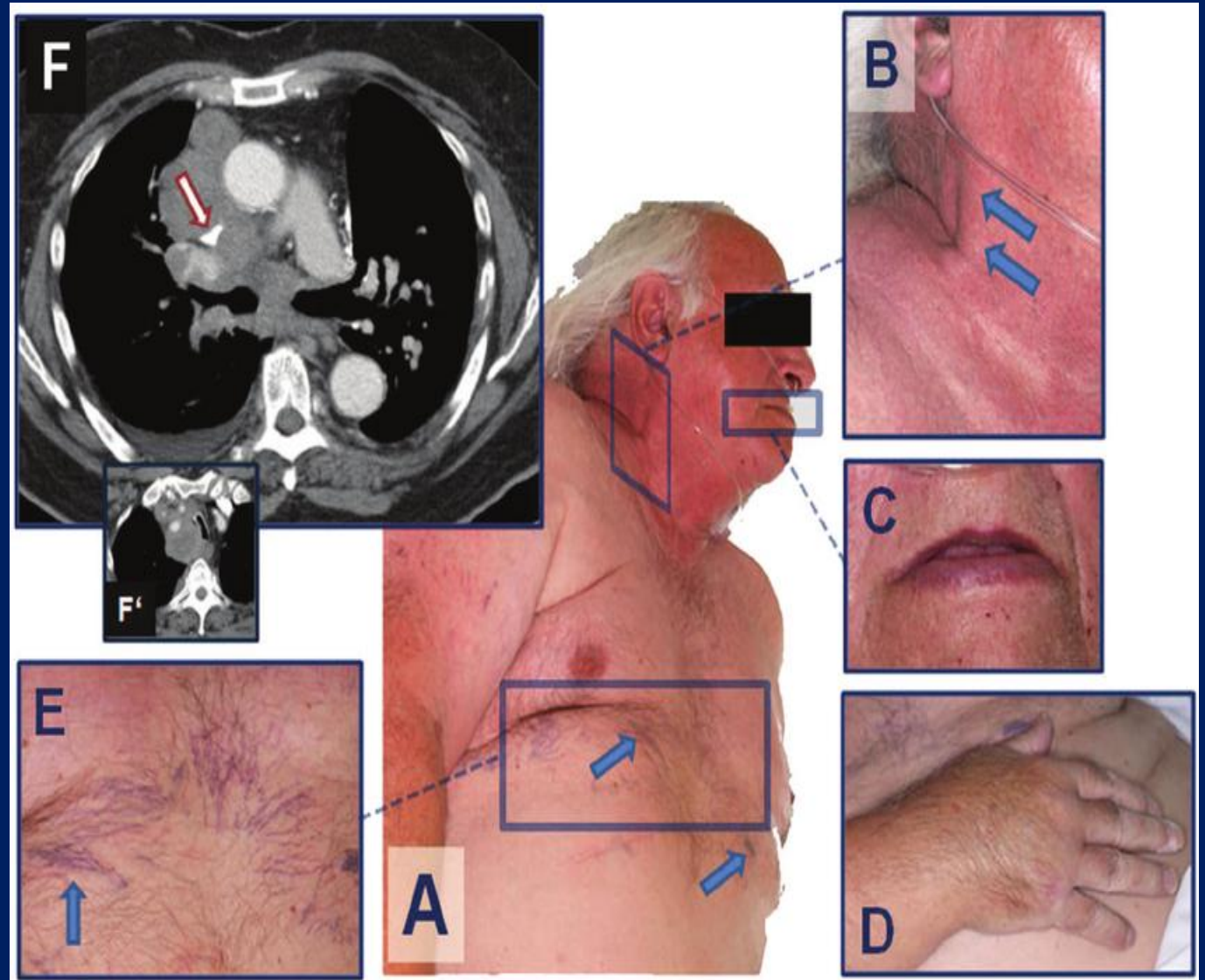
Διόγκωση στις σφαγίτιδες

Θωρακικό επίφλεβο

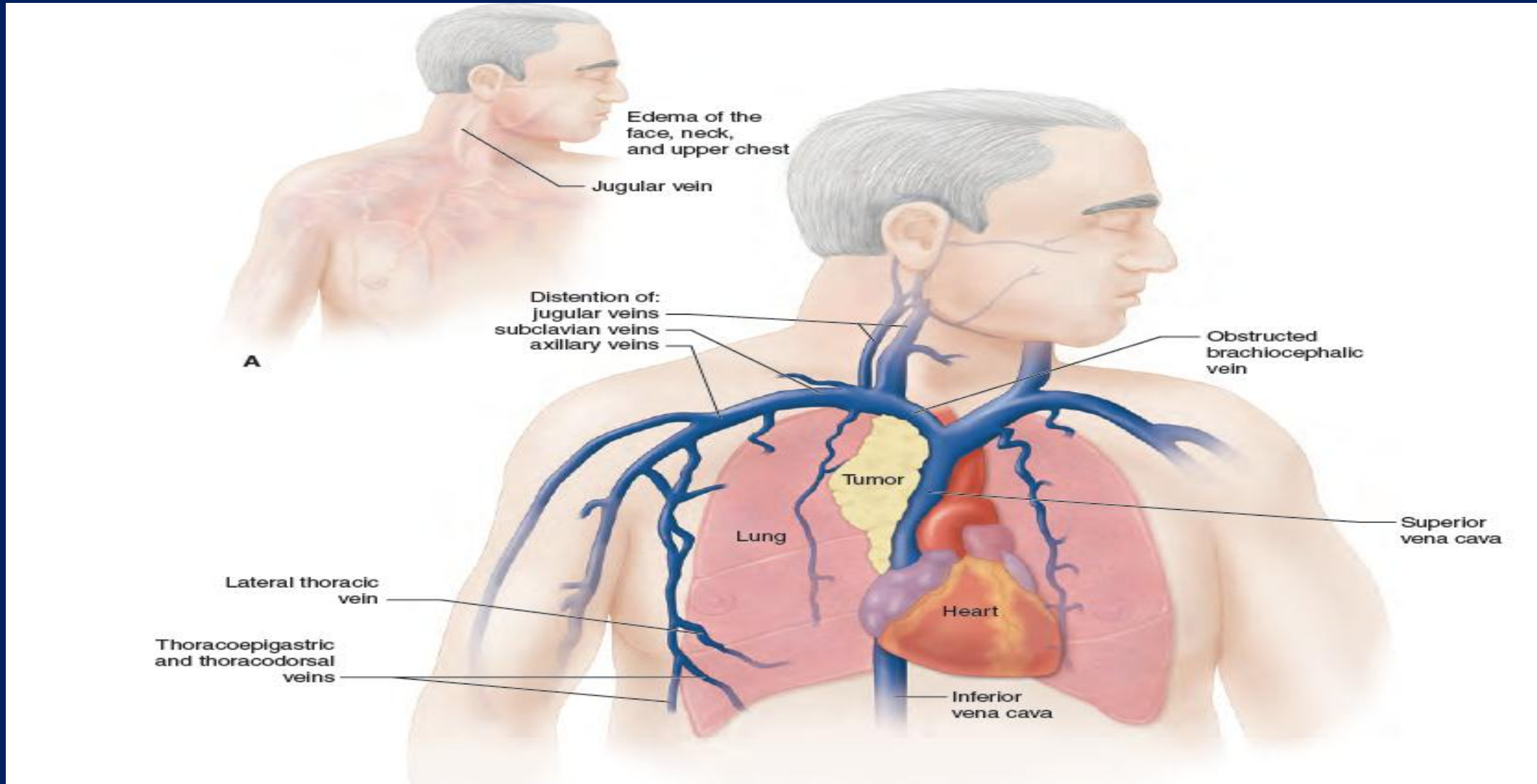
Συμπτώματα



Figure 3. Collateral circulation develops across the chest.



Σύνδρομο άνω κοίλης φλέβας



Βιοψία: μικροκυτταρικός καρκίνος πνεύμονα

Συμπτώματα

TABLE 119.1

Common Symptoms and Physical Findings of Superior Vena Cava Syndrome

Symptoms	Patients Affected ^a (%)	Physical Findings	Patients Affected ^a (%)
Dyspnea	63	Venous distention of neck	66
Facial swelling and head fullness	50	Venous distention of chest wall	54
Cough	24	Facial edema	46
Arm swelling	18	Cyanosis	20
Chest pain	15	Plethora of face	19
Dysphagia	9	Edema of arms	14

Αίτια

Primary Pathologic Diagnoses for Superior Vena Cava Syndrome

Histologic Diagnosis	Bell et al. ⁷ 159 Patients (%)	Schraufnagel et al. ¹⁰ 107 Patients (%)	Parish et al. ⁹ 86 Patients (%)	Yellin et al. ¹¹ 63 Patients (%)	Rice et al. ¹² 78 Patients (%)
Lung cancer	129 (81)	67 (63)	45 (52)	30 (48)	36 (46)
Lymphoma	3 (2)	10 (9)	8 (9)	13 (21)	6 (8)
Other malignancies (primary or metastatic)	4 (3)	14 (13)	14 (16)	8 (13)	5 (6)
Nonneoplastic	2 (1)	16 (15)	19 (22)	11 (18)	31 (40)
Undiagnosed	21 (13)	—	—	—	—

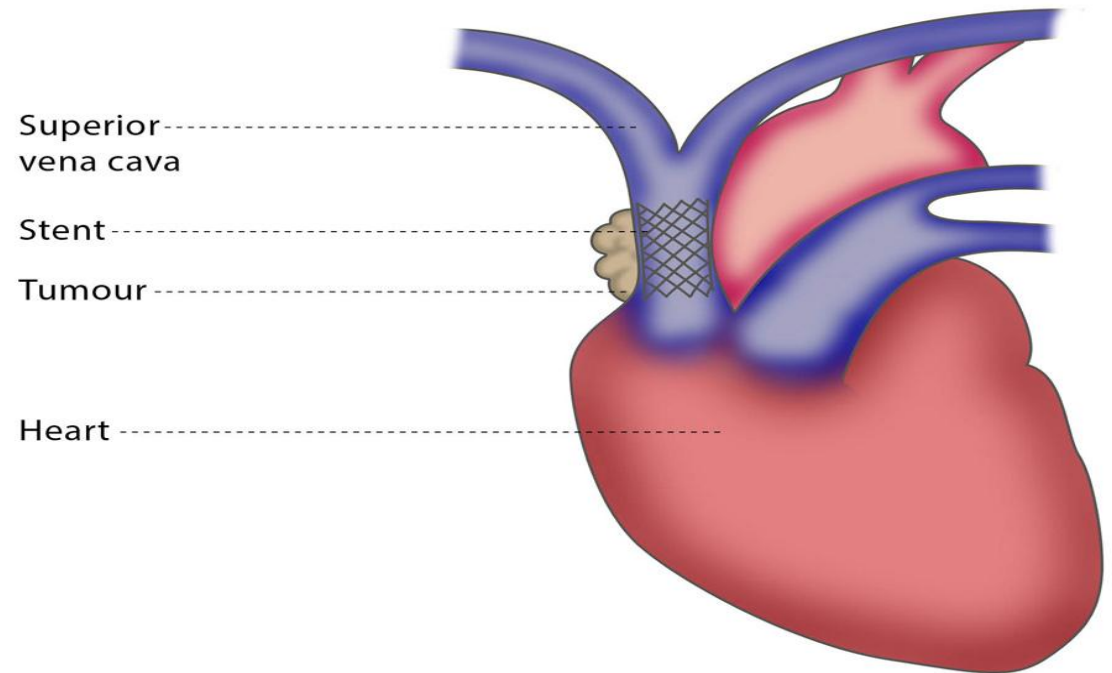
Αντιμετώπιση

Ακτινοθεραπεία

Χημειοθεραπεία

Stent

Superior vena cava (SVC) after stent placement

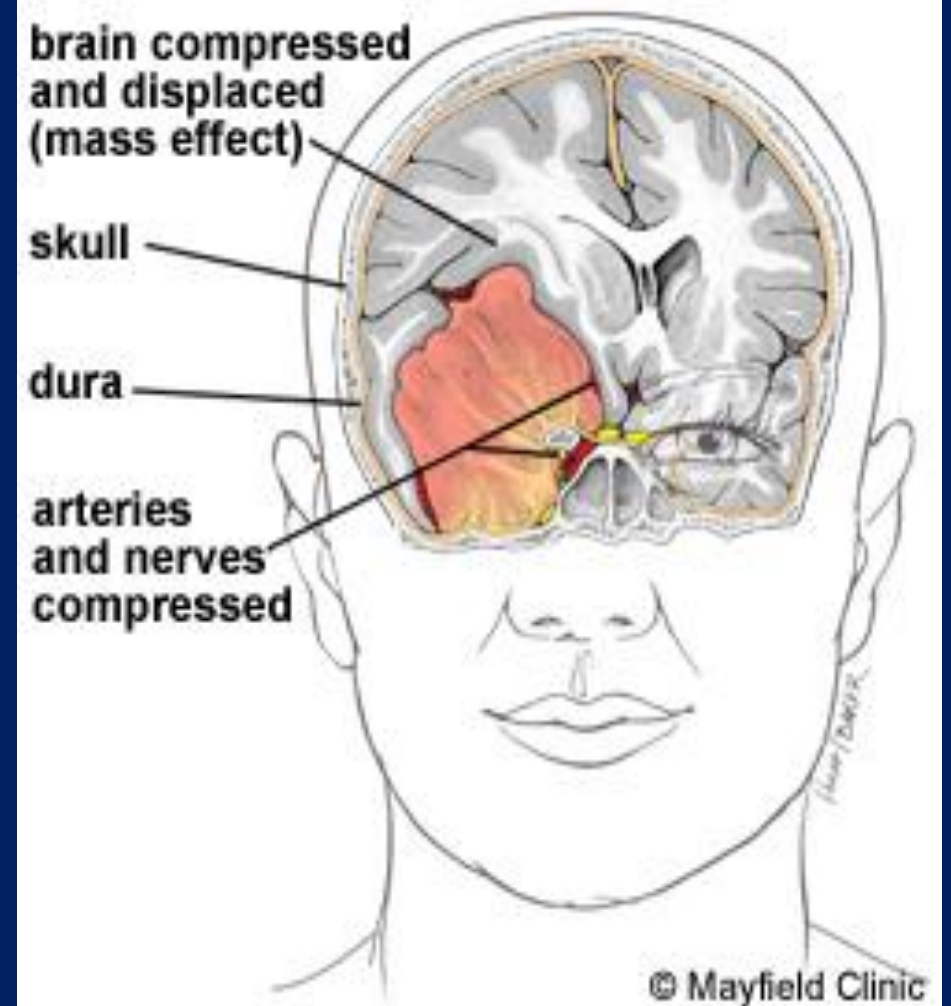


Γυναίκα 55 ετών

Διάγνωση μελανώματος ράχης πρό 4 ετίας

Χειρουργείο + ανοσοθεραπεία για 3 έτη

Αναφέρει ναυτία-εμέτους, κεφαλαλγία από ημερών



INCREASED INTRACRANIAL PRESSURE (ICP)

"CUSHING'S TRIAD"

(Symptoms of ICP are OPPOSITE of Shock)

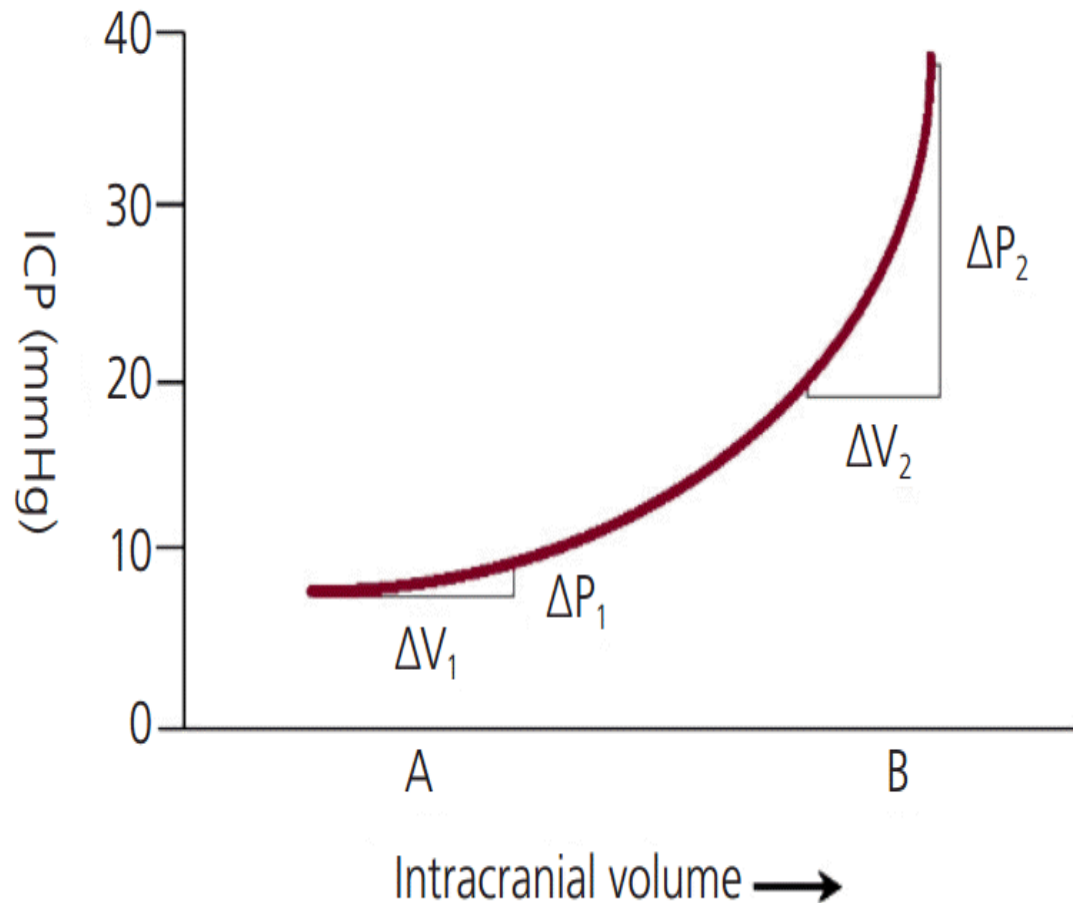
ICP

↑
↓
↓
Systolic B/P
Pulse
Respirations



Shock

↓
↑
↑
B/P
Pulse
Respirations



Intracranial Volume-Pressure Curve

- Fig. 54.3 p. 1612
- Stage 1: good compensation for volume change with no $\uparrow$ ICP
- Stage 2: good compensation, but $\uparrow$ risk of ICP
- Stage 3: steep rise in curve; loss of autoregulation; Cushing's Triad
- Stage 4: ICP at terminal levels; herniation

Complications

- ❑ **Increased intracranial pressure**
- ❑ **Increased intraocular pressure**
- ❑ **Increased intragastric pressure**
- ❑ **Aspiration due to decreased gag reflex**
- ❑ **Malignant hyperthermia**
- ❑ **Dysrhythmias**
- ❑ **Hypoxemia**
- ❑ **Airway trauma**
- ❑ **Failure to intubate / failure to ventilate**
- ❑ **DEATH**

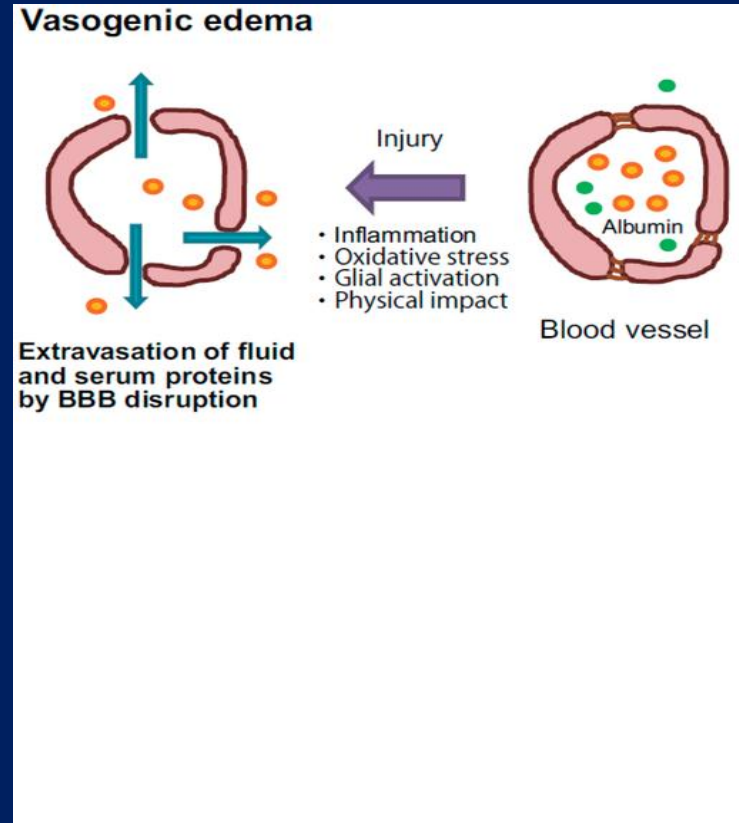


Αντιμετώπιση

Δεξαμεθαζόνη

Μαννιτόλη

Ακτινοθεραπεία



Γυναίκα 46 ετών

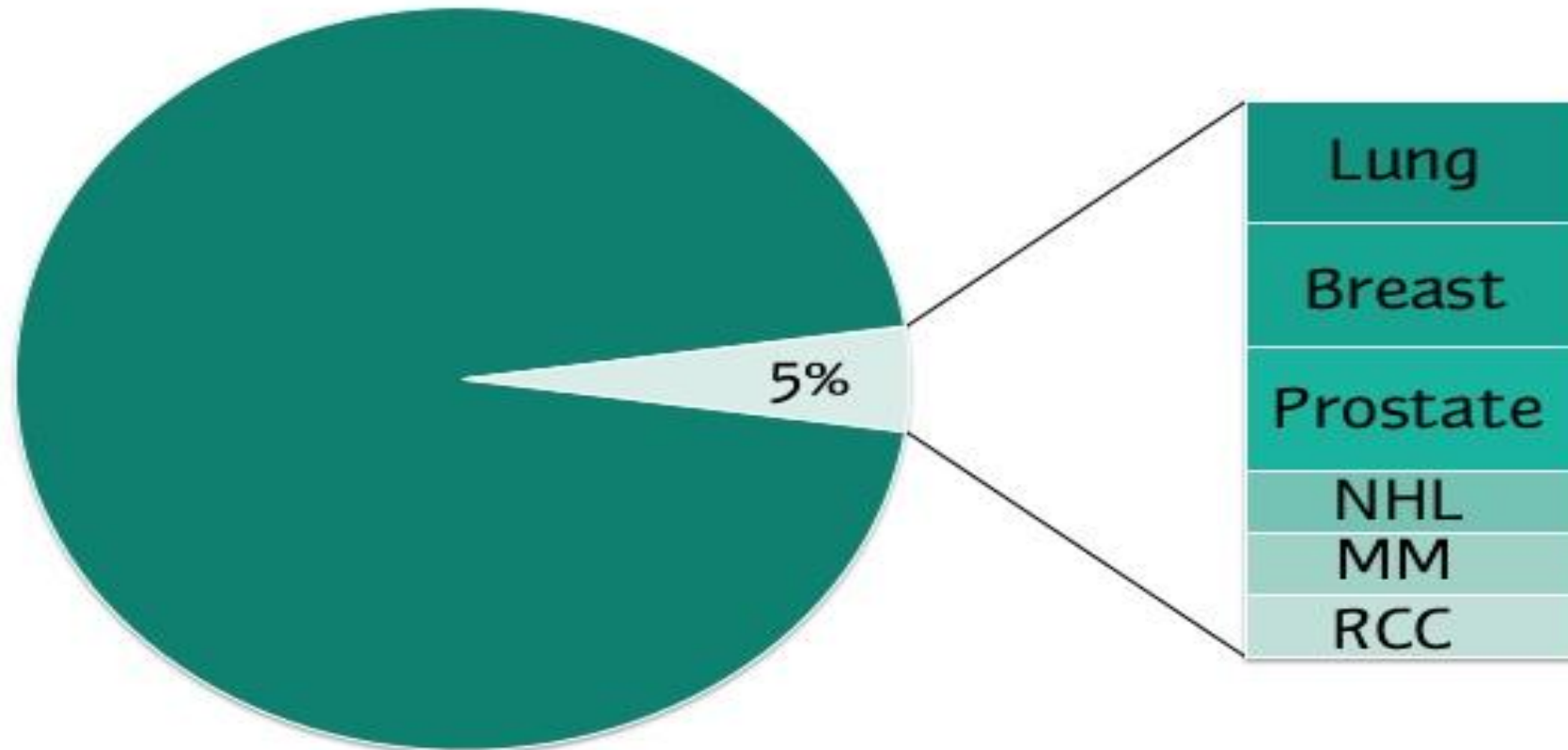
Ca μαστου σταδίου IV

Άλγος στην οσφύ

Άδυναμία-αιμωδίες κάτω άκρων

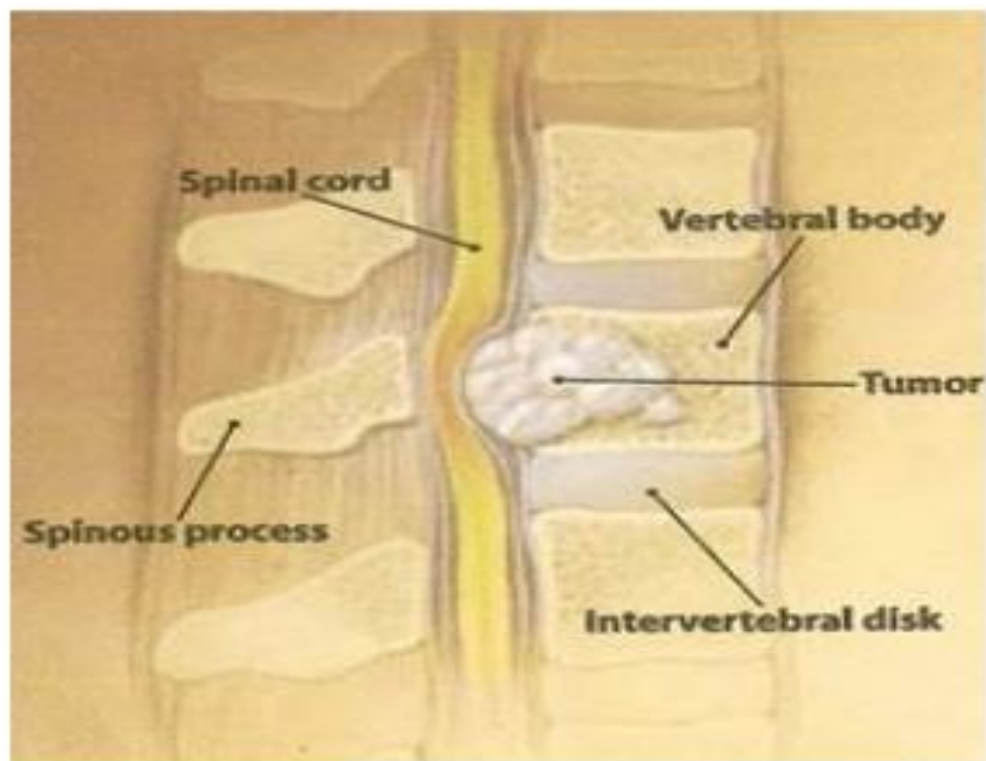
EPIDEMIOLOGY & PROGNOSIS

Incidence of MSCC in terminal CA patients

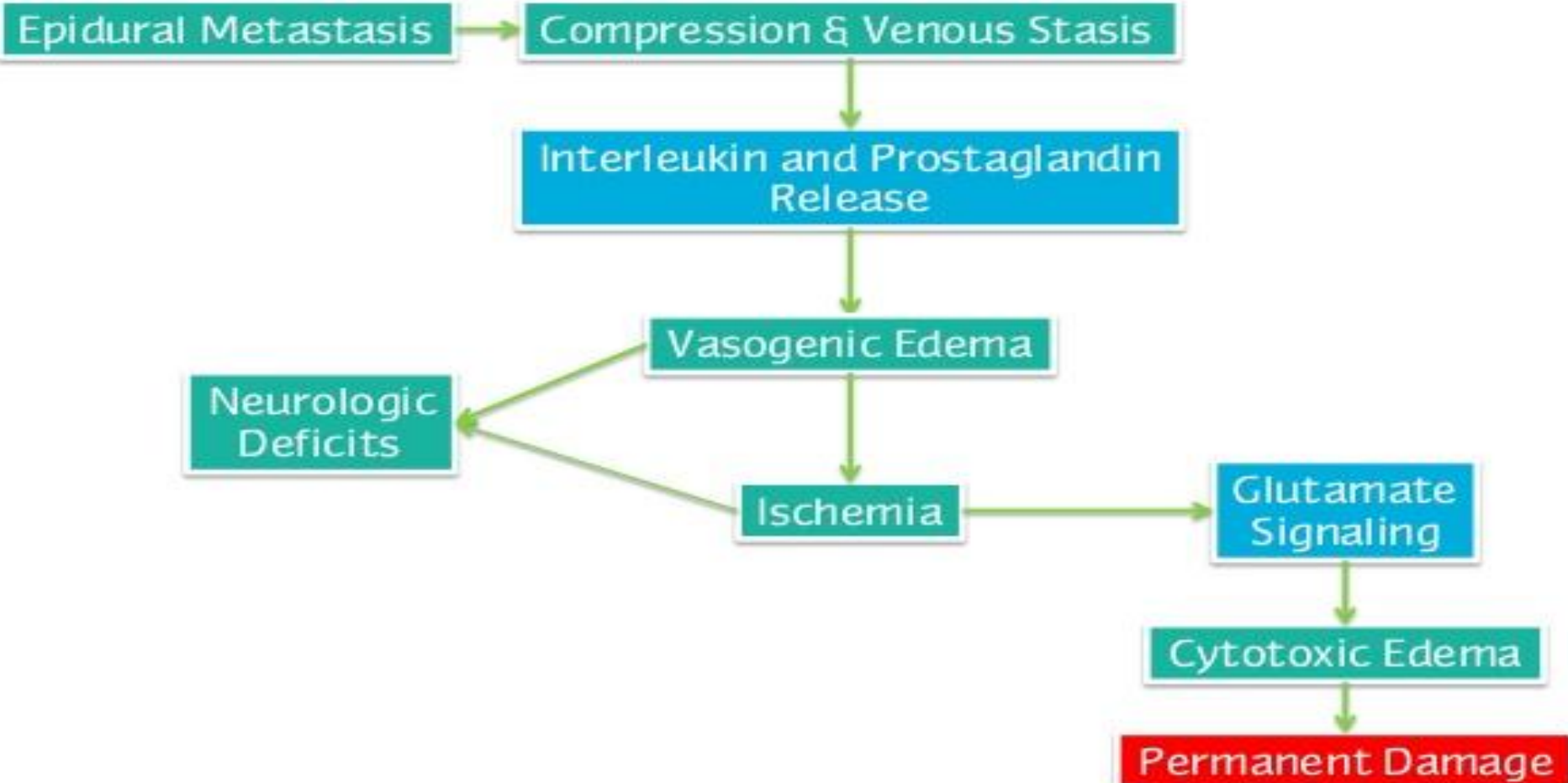


Median Survival after Diagnosis < 6 months

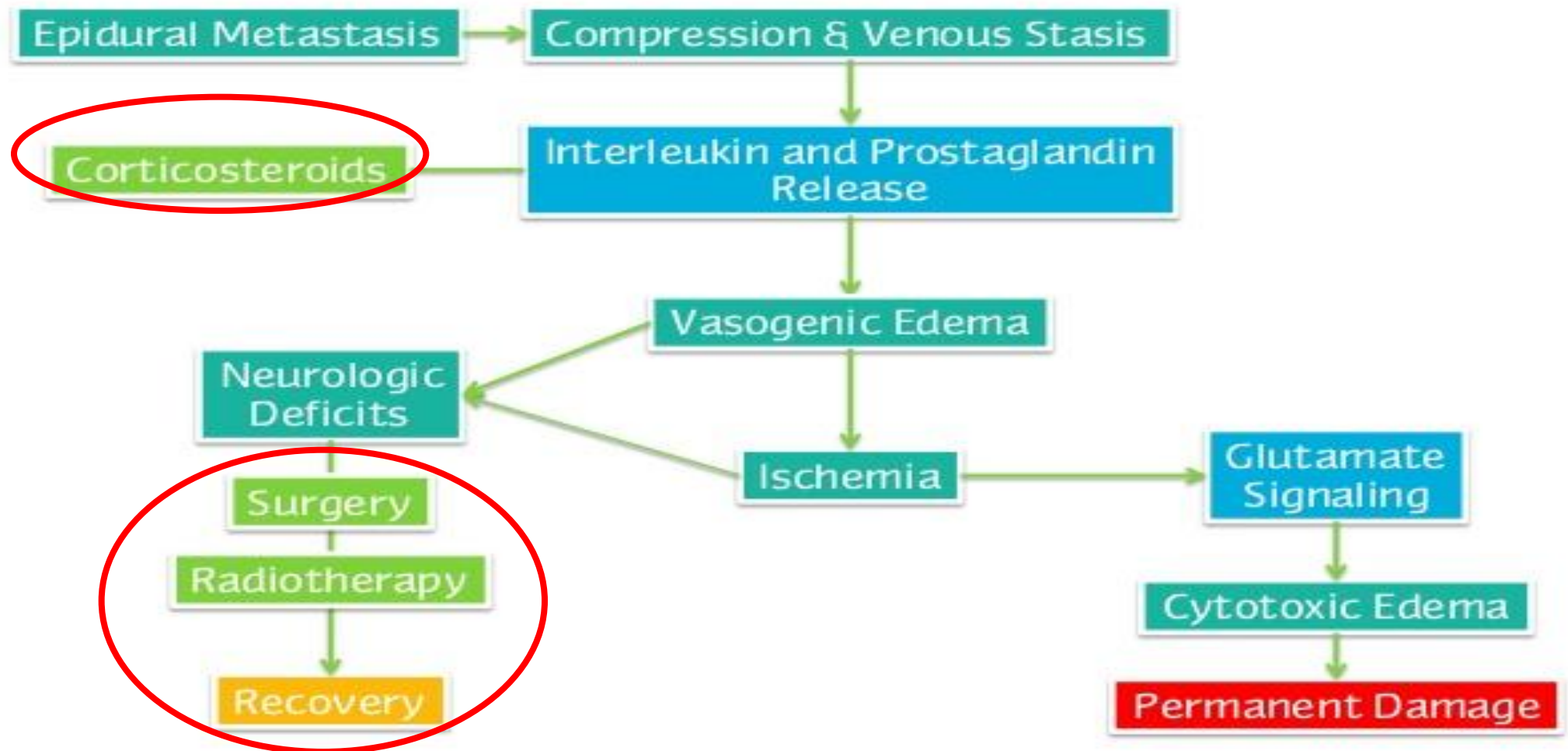
PATHOPHYSIOLOGY



PATHOPHYSIOLOGY



TREATMENT



CORTICOSTEROIDS

High Dose Dexamethasone	Standard Dose Dexamethasone
96 mg IV bolus, 24mg PO q6h x 3 days , then taper	10 mg IV bolus, then 4 mg IV q6h, then taper

Adverse
Effects

Benefits

GI perforation

Infection

Psychosis

Severe Neuro
deficits



RADIATION and SURGERY

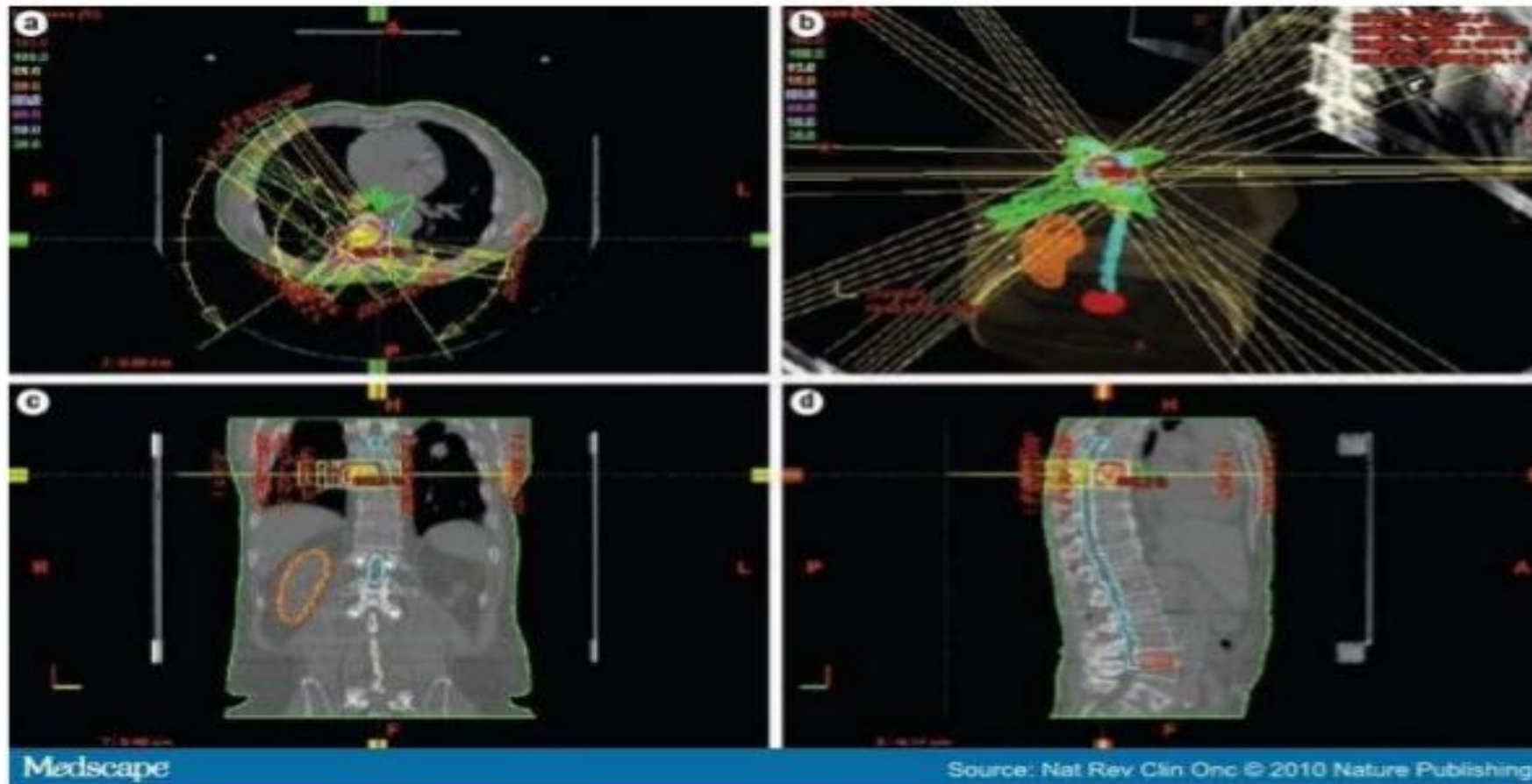
MSCC best treated with COMBINATION therapy

Radiation*
Shrink the tumor!

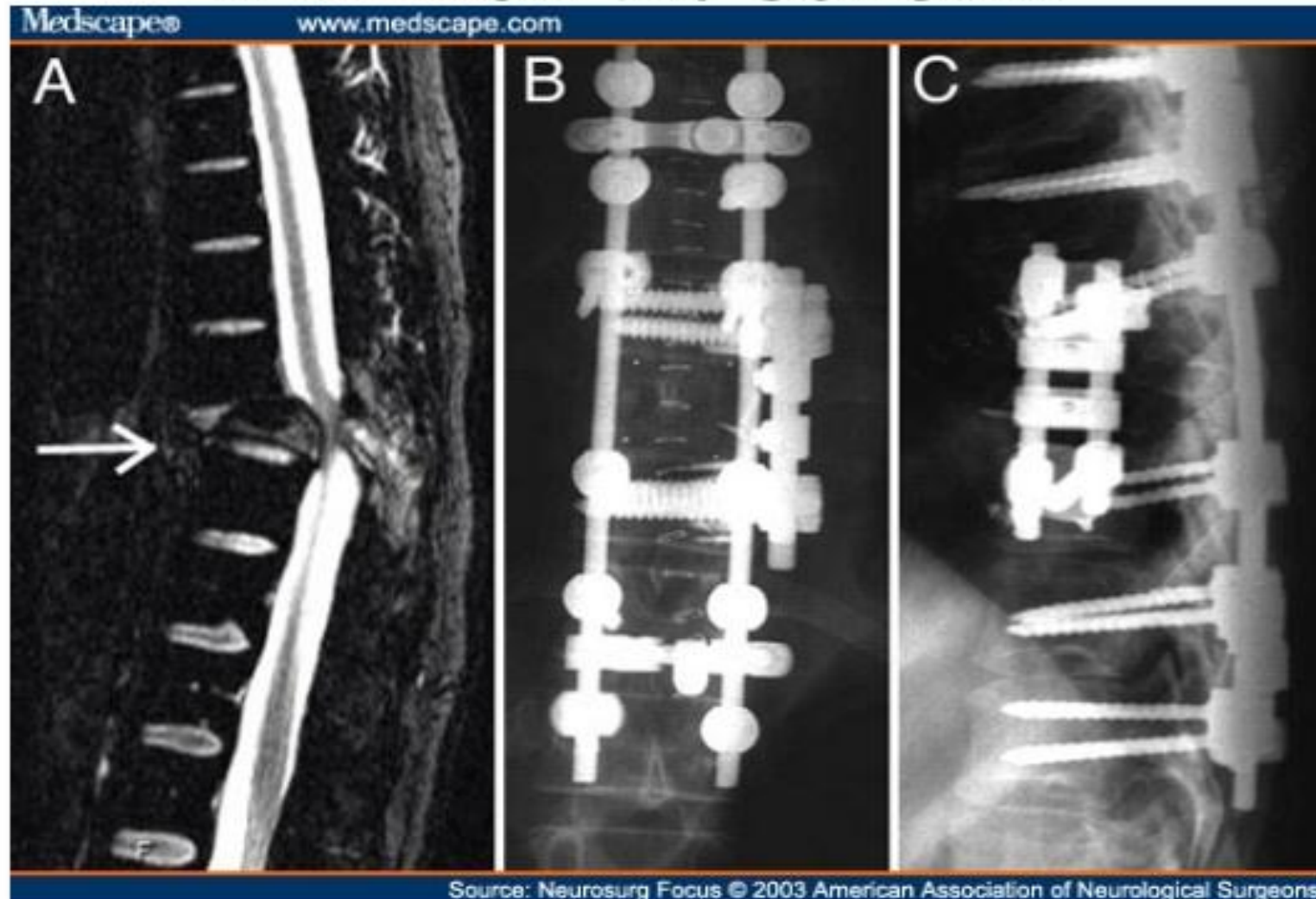
Surgery
Stabilize the spine
and/or resect tumor

*Chemotherapy may also be appropriate if the tumor is chemotherapy sensitive

Treatment



RADIATION and SURGERY



Άνδρας 55 ετών

Ca προστάτη

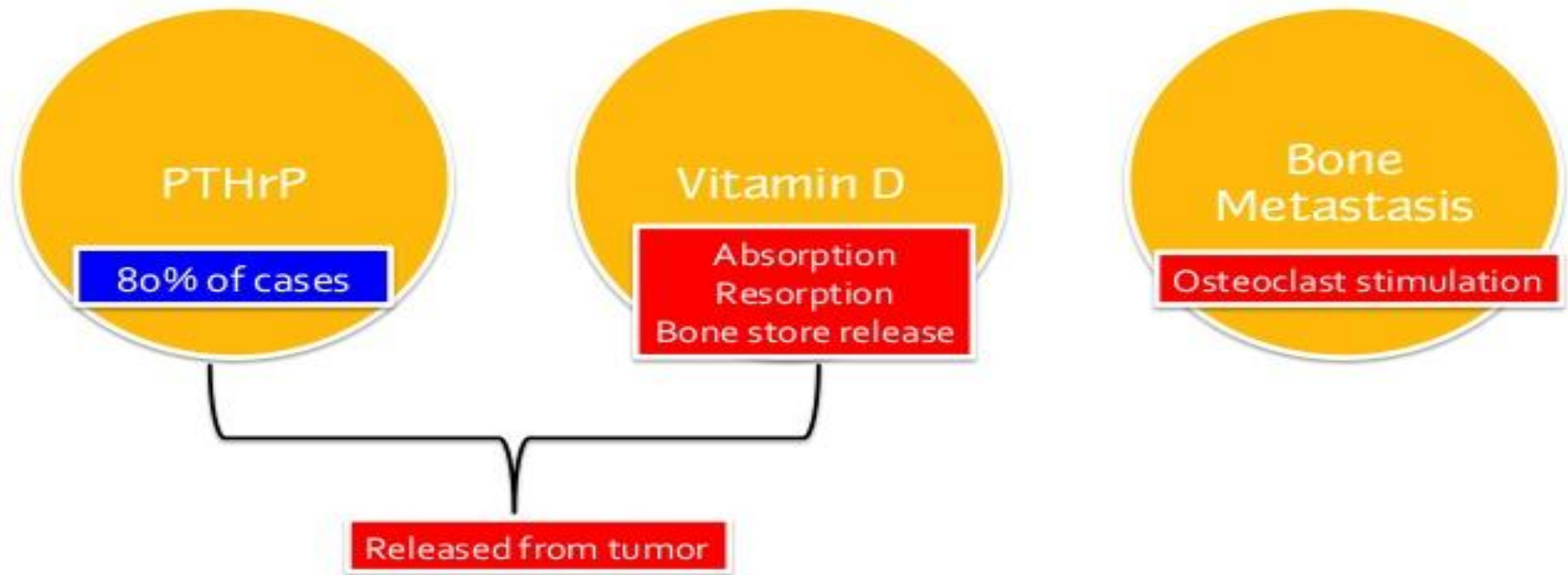
Κρεατινίνη: 1.67

Κοιλιακό άλγος

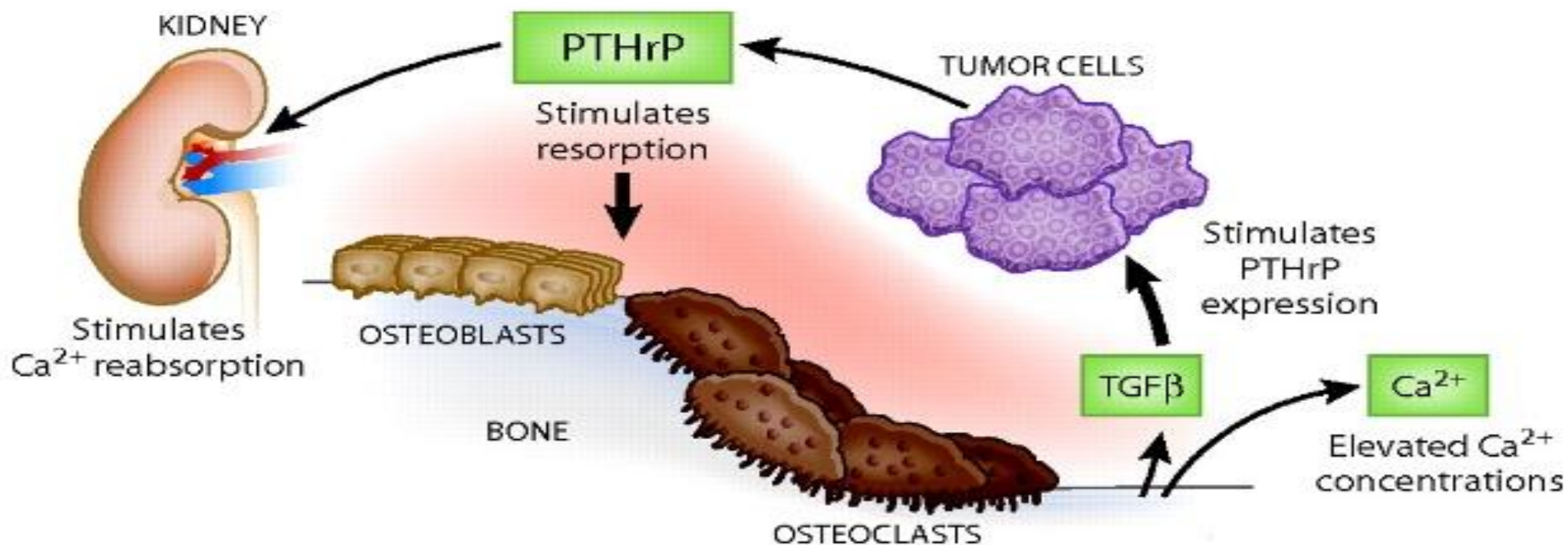
Ασβέστιο: 15.2

Σύγχυση από 2ημέρου

PATHOPHYSIOLOGY



Parathyroid hormone related peptide



Degrees of hypercalcemia



Mild

10.5-11.9mg/dL



Moderate

12-13.9 mg/dL



Severe

≥ 14 mg/dL

Anorexia N/V
Polydipsia
Constipation Polyuria
Weight Loss
Fatigue Bone Pain

Confusion
Lethargy
Coma

Corrected Calcium = Calcium + 0.8 (4-Albumin)

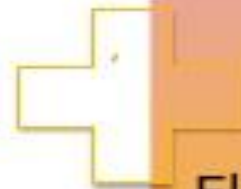
TREATMENT

Stop calcium retaining products



Thiazides
Calcium
Vitamin D
Lithium

Remove calcium from blood



Fluids + Diuretics
Calcitonin
Bisphosphonates
Glucocorticoids

TREATMENT



Hydration (6-12h*)

NS 300-500cc/hr until euvolemic

Caution: HF, Renal failure*



Diuresis (6-12h)

Lasix 20-40 mg IV q12-24h

Avoid Thiazides



Calcitonin (12-24h)

Calcitonin 4-8IU/kg **IM/IV** q12h

Tachyphylaxis often occurs, IN ineffective



Bisphosphonates (48-72h)

Zoledronic Acid 4 mg IV

Pamidronate 60-90 mg IV

Caution: poor dentition (ONJ) or reduced CrCl



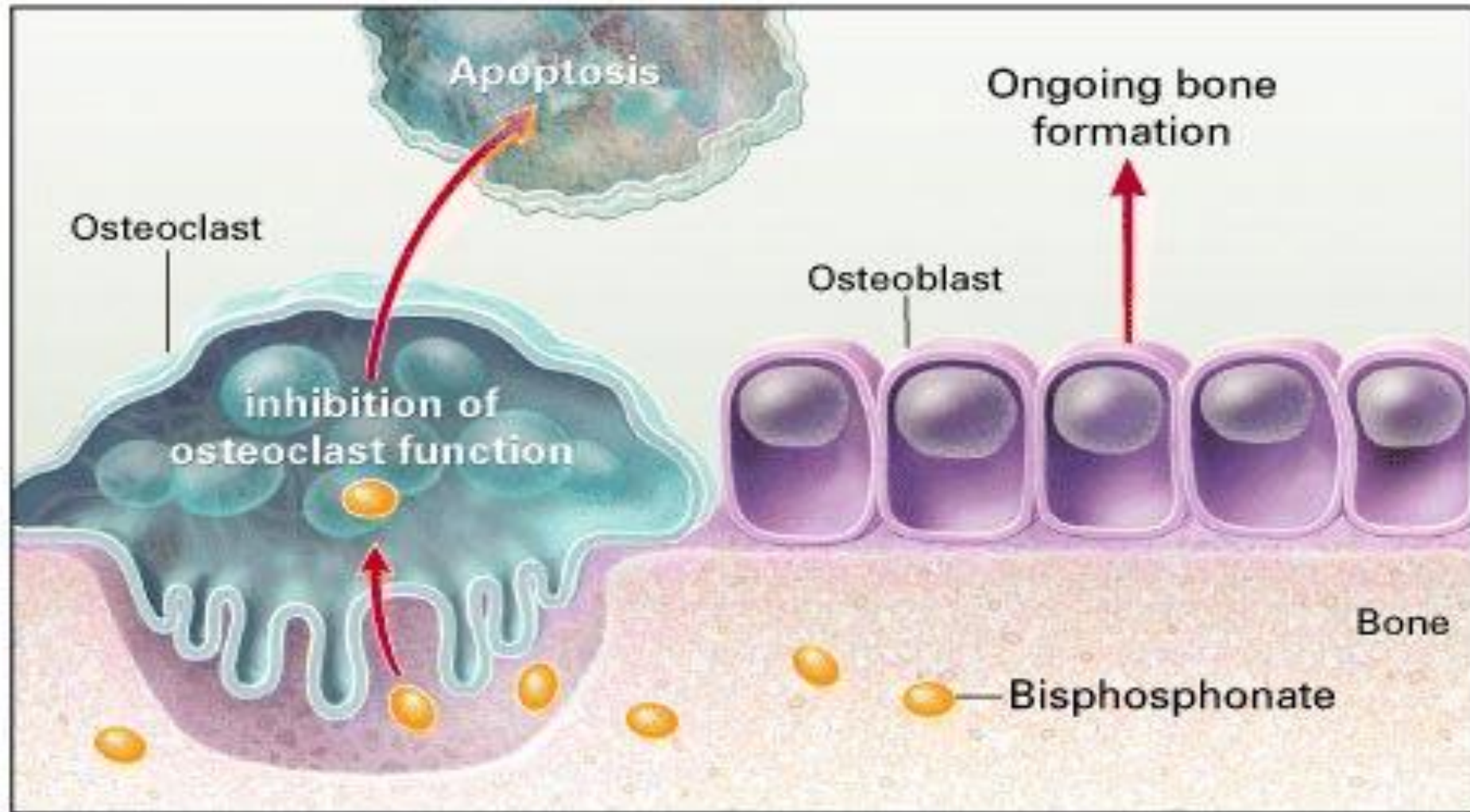
Glucocorticoids (3-5 days)

Hydrocortisone 100 mg IV q6h

Prednisone 60 mg PO daily

Usually limited to lymphomas

Bisphosphonates



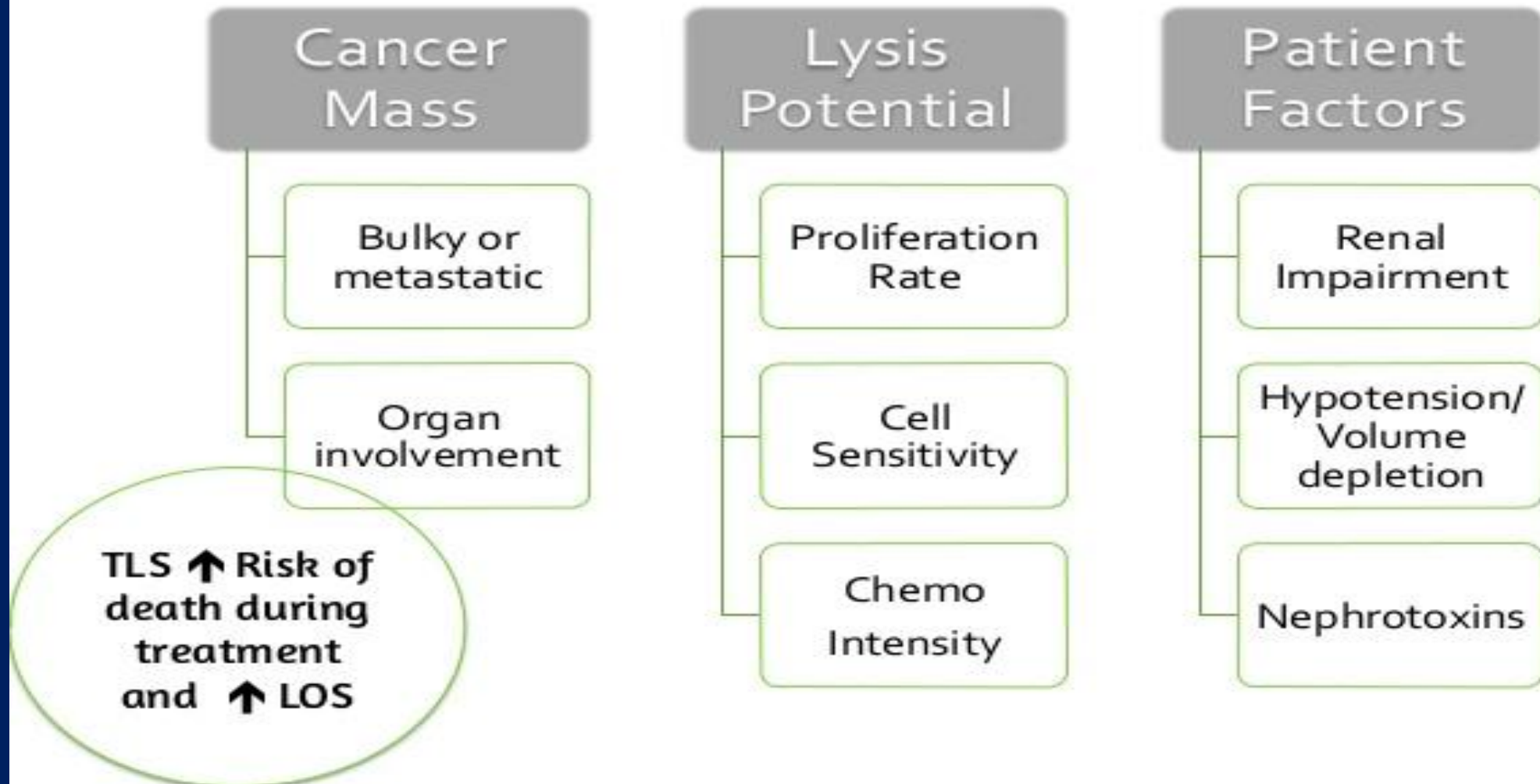
Γυναίκα 61 ετών

Πρόσφατη λήψη χημειοθεραπείας λόγω λευχαιμίας

Κοιλιακό άλγος, ανορεξία, έμετοι

Αδυναμία,ολιγουρία

Epidemiology & Prognosis



Definition

Cairo-Bishop Laboratory TLS

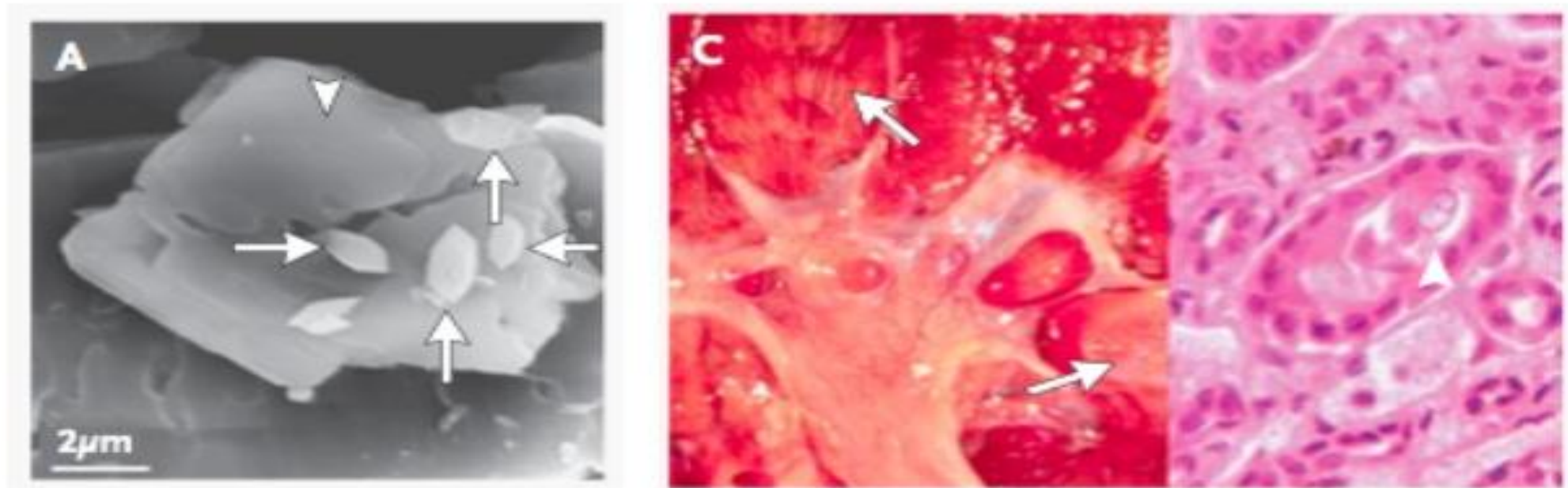
≥ 2

Uric Acid ≥ 8 mg/dL
Potassium ≥ 6mEq/L
Phosphorous ≥ 6.5 mg/dL
Calcium ≤ 7 mg/dL
OR a 25% change from baseline

Clinical TLS

SCr ≥ 1.5 x ULN
Cardiac arrhythmia
Seizure
Sudden death

TLS Induced Renal Failure



Treatment Strategies

Prevention

Evaluate risk factors

- 1) Tumor burden
- 2) Sensitivity
- 3) Underlying dysfunction

Determine **need** and
INTENSITY of prophylaxis

Treatment

Manage lab abnormalities

- Hyperkalemia
- Hyperphosphatemia
- Hyperuricemia

Dose adjust medications
based on **renal** function

Prevention

Hyperhydration

Caution: Heart or Renal failure

- NS IV 2500-3000mL/m²/day
- Goal: UO – 80-100 mL/m²/hr

Allopurinol

No need for renal dose adjust

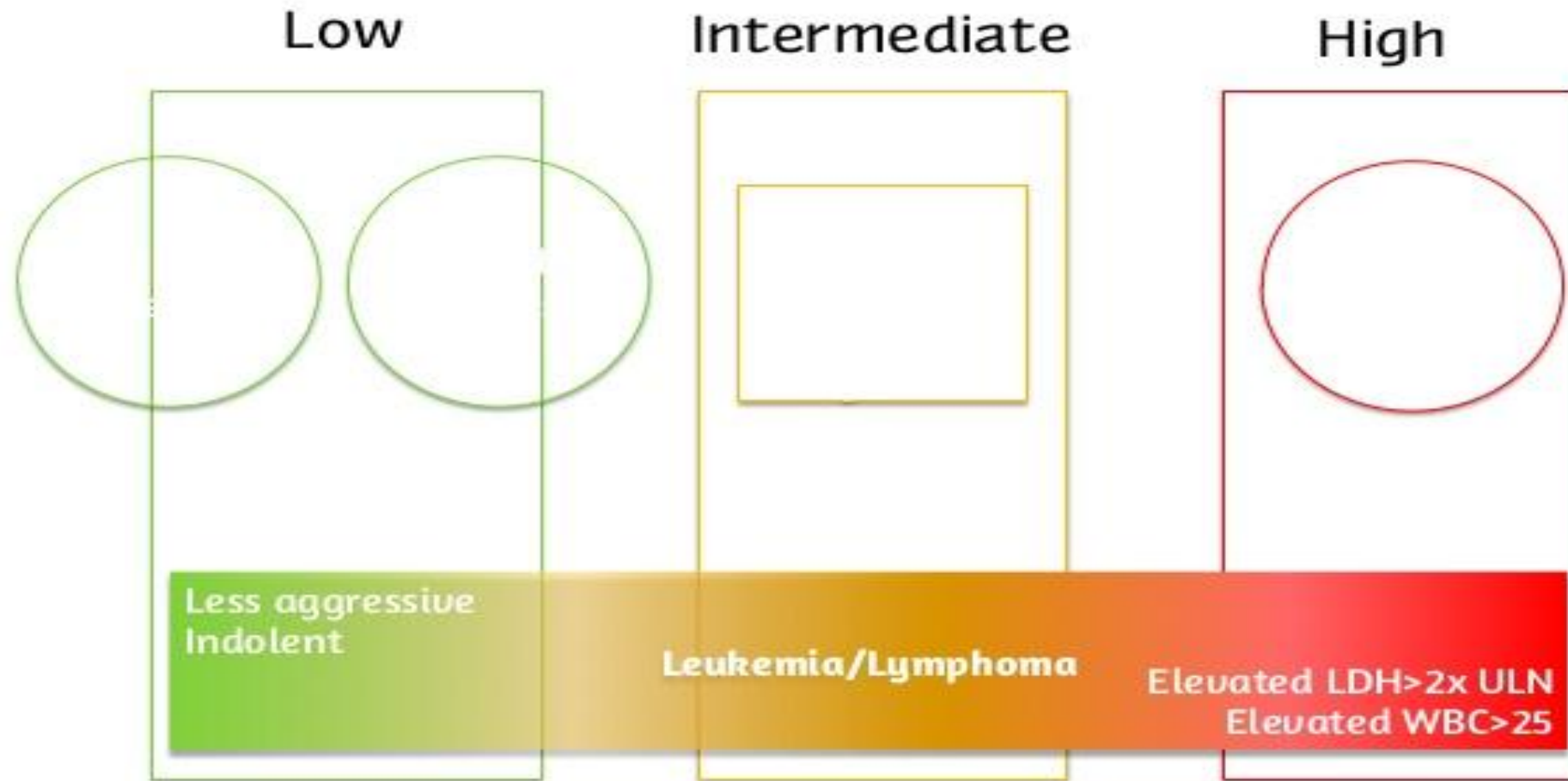
- 300 mg/m²/day starting 48h prior to chemotherapy

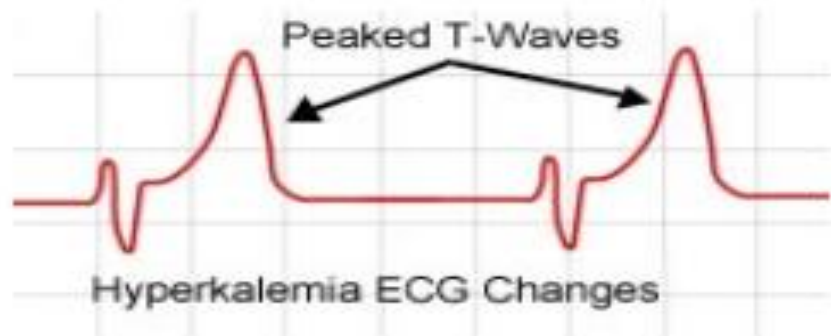
Rasburicase

Ensure proper G6PD function

- 0.15-0.2 mg/kg IV x 1 then reassess

Risk Stratification





Hyperkalemia

Potassium (3.7-5.2 mEq/L)

Mild: 5.5-6.5

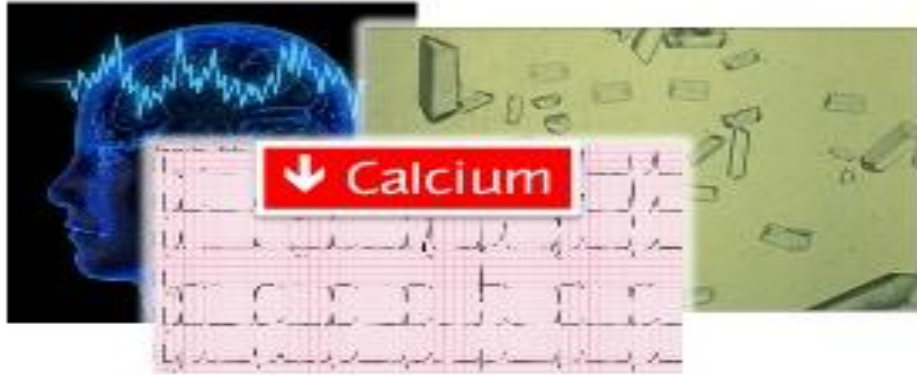
Moderate: >6.5

Severe: >6.5 + ECG changes

- Calcium gluconate 1-2 g IV over 5-10 mins, may repeat

- Insulin 10 units + D50W 50mL IVP
- Albuterol 10-20 mg nebulized over 10 mins
- Sodium bicarbonate 50-100mEq IV over 2-5 mins

- Lasix 20-40 mg IV
- SPS 15-60 gm PO q6h



Hyperphosphatemia

Phosphate (2.4-4.1 mg/dL)
 $\text{Ca} \times \text{Phos} > 70$

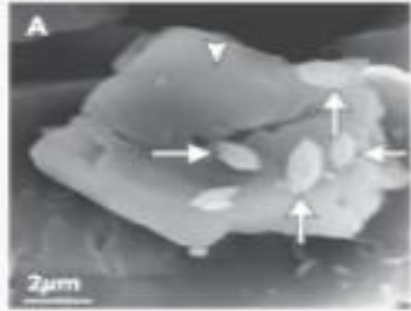
Phosphate Binders
(prevent absorption via GI tract)

↓ Renal
fxn

↑ Ca

Calcium Acetate 1337 mg PO TID AC

Sevelamer 800-1600mg PO TID AC



Hyperuricemia

Uric Acid

Low <4

Intermediate: 4-8

High >8

- NS 2500-3000mL/m²/day
- Goal: UO – 80-100mL/m²/hr

- Rasburicase 0.2 mg/kg IV x 1, redose based upon UA levels

Άνδρας 70 ετών

Ca πνεύμονα σταδίου IV

Λήψη χημειοθεραπείας πρό 10 ημερών

Εμπύρετο έως 38.6, αδυναμία-καταβολή

Febrile Neutropenia

■ Definition:

□ Fever:

- Oral temperature ≥ 38.3 without signs of noninfectious causes of increased temperature
OR
- temperature of ≥ 38.0 C twice, lasting for at least 1 h or measured twice within 12 h

□ Neutropenia

- Neutrophil count $<500/\text{mm}^3$
OR
- Neutrophil count $<1000/\text{mm}^3$ with predicted decline to 500/ml within the next 2 days

Febrile Neutropenia

- Investigations
 - cultures
 - CBC
 - BUN/Cr
 - transaminase measurements
- If respiratory symptoms are present → CXR
 - may be clear due to lack of inflammatory response
- Can consider outpatient treatment of low risk patients >16 years old

Febrile Neutropenia

Table 4. Scoring index for identification of low-risk febrile neutropenic patients at time of presentation with fever.

Characteristic	Score
Extent of illness ^a	
No symptoms	5
Mild symptoms	5
Moderate symptoms	3
No hypotension	5
No chronic obstructive pulmonary disease	4
Solid tumor or no fungal infection	4
No dehydration	3
Outpatient at onset of fever	3
Age <60 years ^b	2

NOTE. Highest theoretical score is 26. A risk index score of ≥ 21 indicates that the patient is likely to be at low risk for complications and morbidity. The scoring system is derived from [50].

^a Choose 1 item only.

^b Does not apply to patients ≤ 16 years of age. Initial monocyte count of ≥ 100 cells/mm³, no comorbidity, and normal chest radiograph findings indicate children at low risk for significant bacterial infections [46].

Febrile Neutropenia

■ Initial treatment algorithm

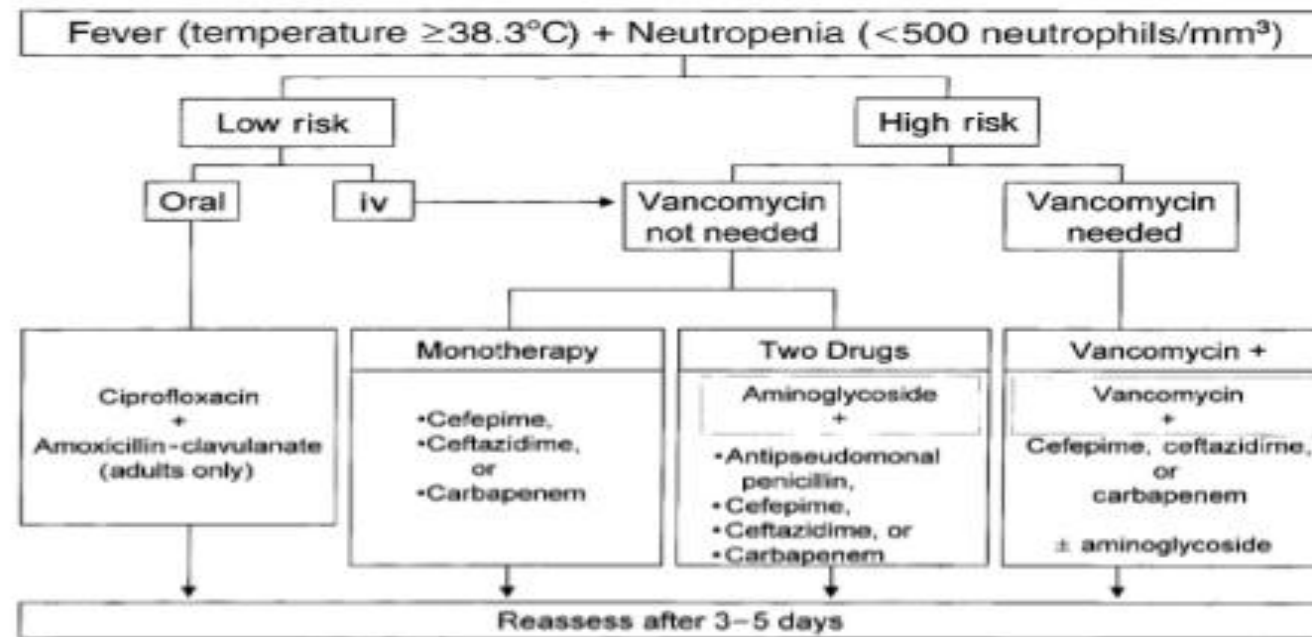
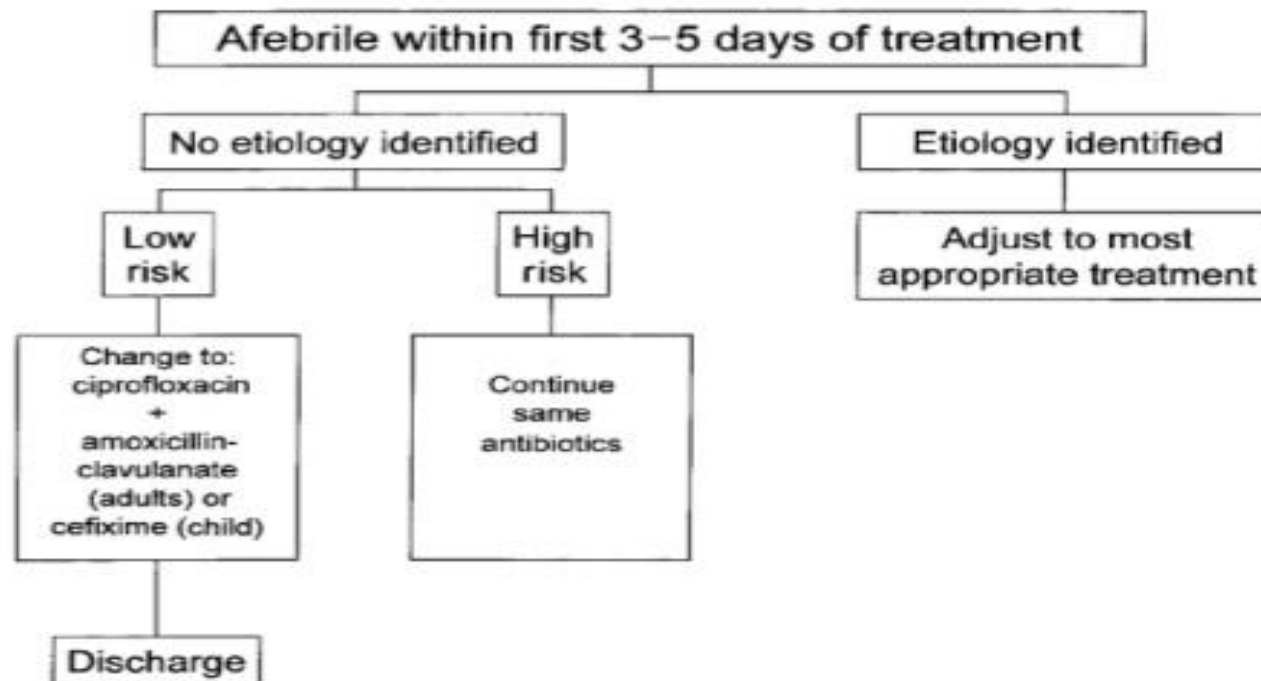


Figure 1. Algorithm for initial management of febrile neutropenic patients. See tables 3 and 4 for rating system for patients at low risk. Carbapenem, imipenem or meropenem.

Febrile Neutropenia

- Continuing treatment algorithm



Febrile Neutropenia

■ Continuing treatment algorithm

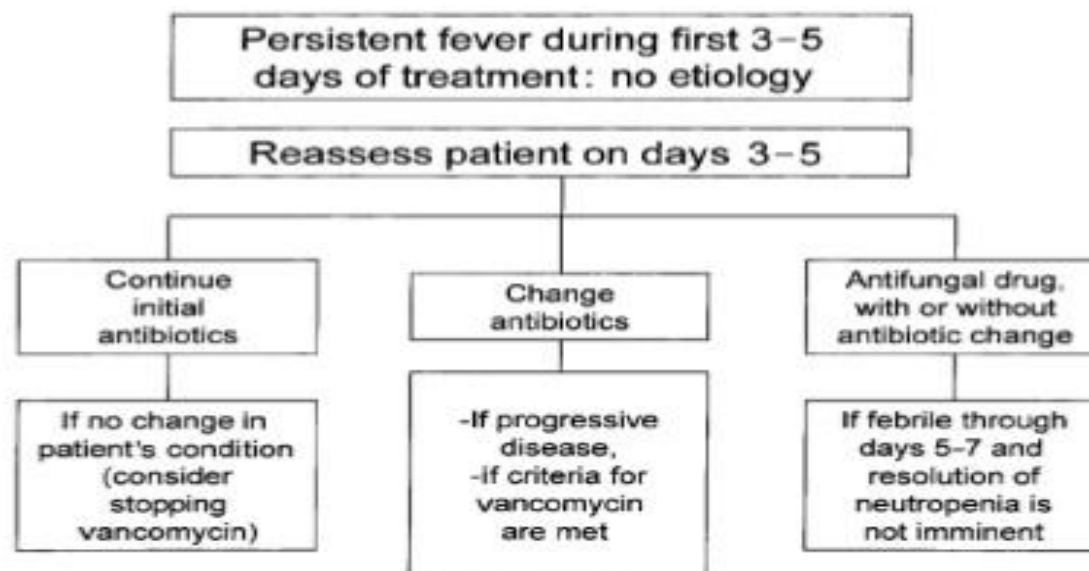


Figure 3. Guide to treatment of patients who have persistent fever after 3-5 days of treatment and for whom the cause of the fever is not found.