

HEMATOLOGY • PLATELET DISORDERS • NGN NCLEX 2026

Thrombocytopenia vs Thrombocytosis vs Thrombocythemia

Three "thromb-o-" platelet disorders — same prefix, different problems. Master the differences for safe clinical judgment.

 Normal Platelets: 150,000–400,000/mm³

 Bleeding vs. Clotting Risk

 NCJMM Integrated

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Definition / Quick Compare

Decode the suffix → understand the diagnosis

Thrombocyto-PENIA

"-penia" = deficiency (too few)

< **150,000/mm³**

 **LOW platelets** → **BLEEDING** risk. Caused by destruction (ITP, HIT, DIC) or decreased production (chemo, leukemia, marrow failure).

Thrombocyt-OSIS

"-osis" = condition/state (reactive)

> **450,000/mm³**

 **HIGH platelets** — **REACTIVE / SECONDARY**. Body's response to infection, inflammation, iron deficiency, surgery, splenectomy, or malignancy. Usually *transient*.

Thrombocyt-HEMIA

"-hemia" = blood disorder (primary)

> **450,000/mm³ (often >1 million)**

 **HIGH platelets** — **ESSENTIAL / PRIMARY**. A myeloproliferative neoplasm (often JAK2 mutation). Platelets are *dysfunctional* → BOTH clotting AND bleeding risk.

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Pathophysiology — Side by Side

Why each condition behaves the way it does

Thrombocytopenia

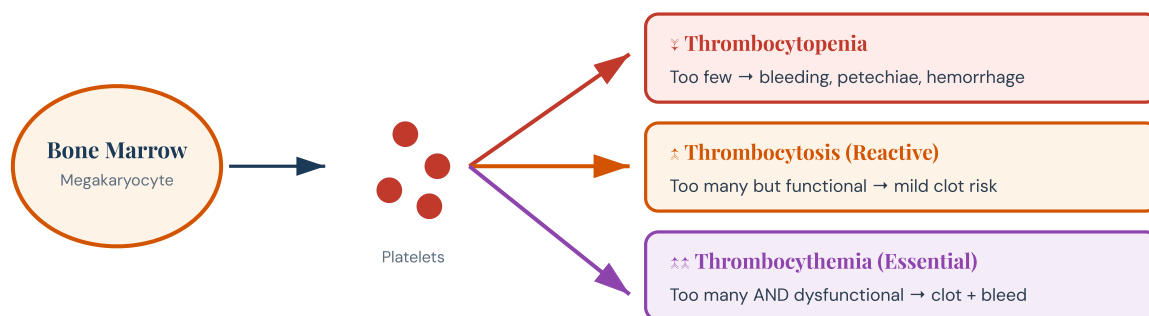
- ▶ Bone marrow makes too few OR spleen/immune system destroys platelets
- ▶ Platelet count drops $< 150,000$
- ▶ Inadequate clot formation at injured vessel
- ▶ → Petechiae, purpura, mucosal bleeding
- ▶ → Risk of intracranial / GI hemorrhage if $< 20,000$

Thrombocytosis (Reactive)

- ▶ Trigger: infection, surgery, inflammation, iron deficiency, splenectomy
- ▶ Cytokines (IL-6) stimulate megakaryocytes
- ▶ Platelet production temporarily $\uparrow$
- ▶ → Platelets are *structurally normal*
- ▶ → Resolves when underlying cause is treated

Essential Thrombocythemia

- ▶ Clonal mutation in megakaryocyte stem cell (often JAK2 V617F)
- ▶ Uncontrolled, autonomous platelet overproduction
- ▶ Platelets are *abnormal & dysfunctional*
- ▶ → Paradoxical THROMBOSIS (clotting) + BLEEDING
- ▶ → Risk: TIA, stroke, MI, DVT, GI bleed



Platelet pathway: one origin (megakaryocyte) → three very different outcomes

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Platelet Count Danger Zones

Memorize these numbers — they drive priority interventions

150K – 400K

NORMAL. Routine monitoring; no special precautions.

50K – 150K

MILD thrombocytopenia. Increased bleeding with trauma/surgery; bleeding precautions begin.

20K – 50K

MODERATE thrombocytopenia. Bleeding with minor trauma; full bleeding precautions; avoid invasive procedures.

< 20,000

SEVERE — SPONTANEOUS BLEEDING RISK. Platelet transfusion indicated; risk of intracranial / GI hemorrhage.

< 10,000

CRITICAL. Life-threatening hemorrhage risk — transfuse immediately; strict bleeding precautions.

> 450,000

THROMBOCYTOSIS / THROMBOCYTHEMIA. Investigate cause; clot risk rises with count, especially > 1 million.

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Signs & Symptoms

Bleeding cues vs. clotting cues — recognize the pattern

Thrombocytopenia (BLEED)

- **Petechiae** (pinpoint red spots)
- **Purpura** & ecchymosis (bruises)
- Bleeding gums, epistaxis
- Hematuria, melena, hematemesis
- Heavy menstrual bleeding
- Prolonged bleeding from puncture sites
- Fatigue (if hemorrhage → anemia)

 **RED FLAG: HA, confusion, ↓LOC, vision changes = possible intracranial hemorrhage**

Reactive Thrombocytosis

- Often **asymptomatic**
- Symptoms of underlying cause: fever, infection, post-op pain, iron deficiency fatigue
- Mild headache (rare)
- Resolves as trigger resolves (days–weeks)
- Thrombosis risk is *low* when platelets are normal-functioning

 **RED FLAG: Sudden splenectomy or surgical patient with new dyspnea or leg swelling = rule out DVT/PE**

Essential Thrombocythemia

- **Erythromelalgia** — burning, red, painful hands/feet
- Headache, dizziness, visual changes
- TIA / stroke symptoms
- Chest pain (MI), DVT, PE
- Splenomegaly
- Paradoxical bleeding: epistaxis, easy bruising, GI bleed (when count is very high)
- Numbness/tingling of extremities

 **RED FLAG: Both clot AND bleed at the same time — highest risk if platelets > 1.5 million**

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Priority Nursing Interventions (ABCs + Safety)

Action verbs: assess, monitor, administer, protect

Bleeding Precautions (Thrombocytopenia)

- ✓ **Assess** for petechiae, ecchymosis, bleeding gums, melena, hematuria every shift
- ✓ **Monitor** platelet count, H&H, neuro status (intracranial bleed)
- ✓ **Use** electric razor — NO straight razors
- ✓ **Use** soft-bristled toothbrush; no flossing if < 50K
- ✓ **Avoid** IM injections, rectal temps, suppositories, enemas
- ✓ **Hold** aspirin, NSAIDs, anticoagulants
- ✓ **Apply** pressure ≥ 5–10 min after venipuncture
- ✓ **Pad** bed rails; remove fall hazards; non-skid footwear
- ✓ **Administer** platelet transfusion if < 20K or active bleed
- ✓ **Avoid** contact sports, straining (stool softeners PRN)

Clot Prevention (Thrombocythemia / Thrombocytosis)

- ✓ **Monitor** for TIA/stroke symptoms (FAST), chest pain, DVT (Homans, leg swelling), PE (sudden SOB, chest pain)
- ✓ **Assess** peripheral pulses, skin color/temp, capillary refill
- ✓ **Encourage** ambulation and hydration
- ✓ **Apply** SCDs / TED hose as ordered
- ✓ **Administer** low-dose aspirin (ET) — paradoxically helps
- ✓ **Administer** hydroxyurea / anagrelide as prescribed
- ✓ **Treat** underlying cause (reactive thrombocytosis): infection, iron deficiency, post-op
- ✓ **Monitor** for paradoxical bleeding when platelets > 1.5 million
- ✓ **Educate** on smoking cessation, lifestyle clot risk reduction
- ✓ **Avoid** dehydration → ↑ blood viscosity

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Pharmacology

Key medications by condition

Drug / Class	Used For	Mechanism (simple)	Key Side Effects / Nursing Considerations
Platelet transfusion	Severe thrombocytopenia (< 20K) or active bleeding	Replaces deficient platelets directly	2 RNs verify; expect ↑ ~10K per unit; monitor for febrile/allergic reactions
Corticosteroids (prednisone)	ITP (immune thrombocytopenia)	Suppresses immune destruction of platelets	↑ glucose, infection risk, GI upset, mood changes, give with food, taper — never stop abruptly
IVIG	ITP (rapid platelet ↑)	Blocks splenic destruction of antibody-coated platelets	Headache, hypersensitivity; pre-medicate with acetaminophen/diphenhydramine; monitor renal function
Romiplostim / Eltrombopag (TPO receptor agonists)	Chronic ITP	Stimulates platelet production	Monitor CBC weekly; thrombosis risk; LFTs (eltrombopag)
Argatroban / Fondaparinux	HIT (heparin-induced thrombocytopenia)	Non-heparin anticoagulants	STOP all heparin! Including flushes. Monitor aPTT (argatroban)
Hydroxyurea	Essential thrombocythemia (1st line)	Suppresses bone marrow platelet production	Myelosuppression, GI upset, skin ulcers; monitor CBC; teratogenic — avoid pregnancy
Anagrelide	Essential thrombocythemia (2nd line)	Inhibits megakaryocyte maturation	Headache, palpitations, fluid retention; cardiac monitoring; avoid in heart failure
Low-dose aspirin (81 mg)	ET — clot prevention	Inhibits platelet aggregation	Hold if platelets > 1.5 million (bleeding risk); GI upset; teach signs of bleeding
Ruxolitinib (JAK inhibitor)	High-risk MPN (incl. some ET cases)	Blocks JAK2 pathway	Anemia, infection risk; monitor CBC; do not stop abruptly



! NCLEX Test Traps

Where students lose points — and how to avoid it

Trap #1: Confusing "-cytosis" with "-cythemia"

✗ THINK

They're the same thing — both = high platelets, no big deal.

✓ KNOW

Cytosis = reactive (functional, transient). Cythemia = primary clonal disease (dysfunctional platelets, lifelong, clot AND bleed risk).

Trap #2: Giving aspirin to a patient with thrombocytopenia

✗ WRONG

"Aspirin is fine — it's just for pain."

✓ RIGHT

Aspirin permanently inhibits platelet function — never give to thrombocytopenic patients. Use acetaminophen instead.

Trap #3: Heparin in a patient developing HIT

✗ WRONG

Continue heparin or switch to LMWH.

✓ RIGHT

STOP all heparin (including flushes & LMWH). Switch to argatroban or fondaparinux.

Trap #4: Treating thrombocythemia like simple thrombocytosis

✗ WRONG

"Just hydrate and treat the underlying infection."

✓ RIGHT

ET is a chronic myeloproliferative disorder — needs cytoreductive therapy (hydroxyurea), aspirin, lifelong monitoring.

Trap #5: Priority — patient with platelets of 8,000 vs. patient with platelets of 800,000

✗ WRONG

"The 800,000 is more abnormal — see them first."

✓ RIGHT

Platelets < 10,000 = imminent spontaneous hemorrhage risk → priority. The high-count patient needs management, but the low-count patient is at acute risk.

Trap #6: Choosing IM injection in a thrombocytopenic patient

✗ WRONG

"Give the IM antibiotic as ordered."

✓ RIGHT

Hold; clarify with provider — IM injections cause hematoma in thrombocytopenia. Use IV or PO route.

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Patient & Family Education

Empower the patient to live safely with their platelet disorder

Daily Self-Care (low platelets)

Use a soft toothbrush, electric razor, and avoid flossing. Blow nose gently; no Q-tips in ears.

Medication Safety

Avoid aspirin, NSAIDs (ibuprofen, naproxen), ginkgo, garlic, fish oil. Read OTC labels — many cold meds contain aspirin.

Activity (low platelets)

Avoid contact sports, lifting > 10 lbs, straining at stool. Wear shoes, prevent falls; no heating pads.

When to Call 911 (low platelets)

Severe headache, vision changes, confusion, unstoppable bleeding, blood in urine/stool, vomiting blood.

Lifestyle (high platelets)

Stay hydrated, no smoking, walk daily, manage cholesterol/BP — all reduce clot risk on top of meds.

When to Call 911 (high platelets)

Sudden weakness/numbness on one side, slurred speech, chest pain, leg pain/swelling, sudden SOB → stroke/MI/DVT/PE.

Wear Medical ID

Bracelet stating diagnosis (ITP, ET, HIT) and current medications. Inform all providers (dental, surgical) before any procedure.

Pregnancy & Hydroxyurea

Hydroxyurea is teratogenic — use reliable contraception. Discuss family planning with hematologist.

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Memory Boosters & Mnemonics

Lock these patterns into long-term memory

B - L - E - E - D

Thrombocytopenia bleeding precautions

- B** – Brush soft (toothbrush)
- L** – Limit injections (no IM, no rectal)
- E** – Electric razor
- E** – Eliminate aspirin/NSAIDs
- D** – Don't strain (stool softeners)

SUFFIX RULE

Decode "thromb-o-____"

-

PENIA "poor" / few → **BLEED**

-

CYTOSIS "condition" → reactive ↑

-

CYTHEMIA "blood disease" → primary, dysfunctional

"-osis" responds; "-hemia" is hereditary-like clonal disease.

P - E - T - S

Spot thrombocytopenia bleeding

- P** – Petechiae (pinpoint)
- E** – Ecchymosis (bruises)
- T** – Tarry stools (melena)
- S** – Saturated bandages / persistent bleeding

"4 T's"

HIT (heparin-induced thrombocytopenia) clinical score

- T** – Thrombocytopenia (drop > 50%)
- T** – Timing (5–10 days after heparin)
- T** – Thrombosis (paradoxical clotting!)
- T** – oTher causes excluded

NUMBERS

Platelet count cliff notes

- 150K** lower limit of normal
- 50K**– bleed with trauma/surgery
- 20K**– spontaneous bleed risk → transfuse
- 10K** – critical hemorrhage risk
- 450K** upper limit of normal

F - A - S - T

Recognize stroke in thrombocythemia

- F** – Face drooping
- A** – Arm weakness
- S** – Speech difficulty
- T** – Time to call 911

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Rapid-Fire: Commonly Confused

Last-minute clarifiers

Question	Quick Answer
ITP vs. TTP?	ITP = isolated low platelets, immune-mediated. TTP = pentad: fever, low platelets, hemolytic anemia, neuro changes, renal failure → emergency, plasma exchange.
HIT vs. ITP?	HIT = thrombocytopenia + clotting from heparin exposure. ITP = autoimmune, no clotting (just bleeding).
Reactive thrombocytosis vs. ET?	Reactive resolves with the trigger. ET is clonal, lifelong, JAK2 mutation common, requires cytoreduction.
Why does ET cause BOTH bleeding and clotting?	Platelets are too many AND dysfunctional — clot in some places, fail to clot in others. Worse risk if > 1.5 million.
Goal platelet count when transfusing for < 20K?	Generally > 50K for active bleeding; > 100K for neurosurgery / intracranial procedures.
Splenectomy & platelets?	Reactive thrombocytosis is expected post-splenectomy (loss of platelet pooling/destruction site) — usually transient.



NCJMM Clinical Judgment Scenario

Apply the 6-step Clinical Judgment Measurement Model

Scenario: A 62-year-old client admitted 7 days ago for a DVT on heparin infusion is found this morning with a platelet count drop from 240,000 to 88,000. The client now reports a swollen, painful right leg and a new dusky toe. Vital signs: BP 138/82, HR 102, RR 20, SpO₂ 96% on RA, T 37.6°C.

STEP 1 — RECOGNIZE CUES

What stands out?

Platelets dropped > 50% within 5–10 days of heparin start; new thrombosis (dusky toe, swollen leg) despite anticoagulation; mild tachycardia.

STEP 2 — ANALYZE CUES

What's connecting?

Pattern fits the "4 T's" of HIT (Heparin-Induced Thrombocytopenia) — thrombocytopenia WITH paradoxical thrombosis, not just bleeding.

STEP 3 — PRIORITIZE HYPOTHESES

Most likely?

HIT (Type II) is the leading hypothesis. Less likely: post-op reactive change, sepsis-induced consumption (DIC), drug-induced thrombocytopenia from another agent.

STEP 4 — GENERATE SOLUTIONS

What needs to happen?

STOP all heparin sources (IV drip, line flushes, LMWH). Notify provider. Anticipate non-heparin anticoagulant (argatroban or fondaparinux). Send HIT antibody panel (PF4/heparin ELISA).

STEP 5 — TAKE ACTION

First action?

 **Discontinue the heparin drip immediately** and remove heparin flushes from the unit's care plan. Place "no heparin" alert on chart. Contact provider for argatroban order. Apply continuous cardiac/SpO₂ monitoring.

STEP 6 — EVALUATE OUTCOMES

Did it work?

Platelets begin rising within 2–3 days off heparin; toe perfusion improves; no new clots; HIT antibody confirms; client educated to wear medical ID stating "HIT — no heparin lifetime."

Diane's NGN NCLEX 2026 Study Series

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