Pathophysiology, classification, and complications of common asymptomatic thrombocytosis in newborn infants

Running title: Thrombocytosis in newborn infants

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Abstract

We frequently encounter newborn infants with thrombocytosis in the neonatal intensive care unit. However, neonatal thrombocytosis is not yet fully understood. Thrombocytosis is more frequently identified in newborns and young infants, notably more often in those younger than two years than in older children or adults. The production of megakaryocytes (megakaryopoiesis) and platelets (thrombopoiesis) is mainly regulated by thrombopoietin (TPO). Increased TPO levels during infection or inflammation can stimulate megakaryopoiesis, resulting in thrombopoiesis. TPO concentrations are higher in newborn infants than in adults. Levels increase after birth, peak on the second day after birth, and start decreasing at 1 month of age. Initial platelet counts at birth increase with gestational age. Thus, preterm infants have lower initial platelet counts at birth than late-preterm or term infants. Postnatal thrombocytosis is more frequently observed in preterm infants than in term infants. A high TPO concentration and low TPO receptor expression on platelets leading to elevated plasma-free TPO, increased sensitivity of megakaryocyte precursor cells to TPO, a decreased red blood cell count, and immaturity of platelet regulation are speculated to induce thrombocytosis in preterm infants. Thrombocytosis in newborn infants is considered a reactive process (secondary thrombocytosis) following infection, acute/chronic inflammation, or anemia. Thrombocytosis in newborn infants is benign, resolves spontaneously, and, unlike in adults, is rarely associated with hemorrhagic and thromboembolic complications.

Keywords: infant, newborn; infant, premature; platelets; thrombocytosis; thrombopoietin
Key Message

- Thrombocytosis, common in newborns and infants (<2 years) (3%–13%), is caused by elevated thrombopoietin (TPO) concentrations.
- Serum TPO levels are significantly higher immediately to 1 month postnatal and decrease with age.
- Platelet counts are positively correlated with gestational age at birth and postnatal age.
- Thrombocytosis is more common in preterm than in term infants.
- Thrombocytosis in newborns is reactive and resolves spontaneously without complications.

Graphical abstract

Thrombocytosis is frequently observed in newborn infants. Platelets, non-nucleated fragments arising from megakaryocytes, are activated by endothelial injury as well as infection, acute/chronic inflammation, and anemia. Activated platelets can interact with leukocytes, monocytes, and bacteria through multiple receptors. Reactive (secondary) thrombocytosis in newborn infants is benign and can spontaneously resolve. It is rarely associated with complications, unlike thrombocytosis in adults. Essential (primary) thrombocytosis, a myeloproliferative neoplasm, is rare in newborn infants and characterized by megakaryocytic hyperplasia in the bone marrow caused by mutations in the Janus kinase 2 (JAK2) or myeloproliferative leukemia (MPL) gene.
Introduction

Thrombocytosis is known to increase the risk of hemorrhagic and thromboembolic complications in adults for which anti-platelet or cytoreductive treatment is required.\textsuperscript{1,2)\textsuperscript{1}} Thrombocytosis frequently occurs in newborns and young infants younger than 2 years, with an estimated incidence of 3\%–13\%,\textsuperscript{3)\textsuperscript{1}} although the exact prevalence is unknown. Thrombocytosis in newborn infants is considered a reactive process in most cases, and it is rarely associated with complications.\textsuperscript{4)\textsuperscript{1}} Whether similar risks such as hemorrhagic and thromboembolic complications in adults exist in newborn infants with thrombocytosis remains unclear. Reasons for frequent reactive thrombocytosis in newborn infants include higher levels of thrombopoietin (TPO) and more sensitive megakaryocyte precursor cells to TPO.\textsuperscript{5)\textsuperscript{1}}

Although thrombocytosis is frequently encountered in newborn infants, especially preterm infants admitted to neonatal intensive care units, few studies have evaluated neonatal thrombocytosis. Therefore, this review aimed to evaluate the pathophysiology, classification,
and complications of thrombocytosis in newborn infants. Neonatal thrombocytosis according to gestational age at birth and postnatal age was also evaluated.

**Pathophysiology**

**Megakaryopoiesis and thrombopoiesis**

Little is known about the pathophysiology underlying thrombocytosis in neonates. Platelets are non-nucleated fragments arising from megakaryocytes. Hematopoiesis transitions from the yolk sac to the liver and then to the bone marrow. Megakaryocytes are found in the yolk sac at a gestational age of 5 weeks and in the liver by a gestational age of 10 weeks, while platelets are first found in the circulation at a gestational age of 8–9 weeks. Bone marrow megakaryopoiesis starts at a gestational age of 11 weeks and then dominates at a gestational age of 20 weeks. Thus, bone marrow is the primary site for megakaryopoiesis in newborn infants.

Megakaryopoiesis and platelet production (thrombopoiesis) are mainly regulated by TPO and other cytokines and hematopoietic growth factors such as interleukin-3 (IL-3), IL-6, and IL-11. The increased production of these factors during infection or inflammation can stimulate megakaryocyte production, resulting in elevated platelet counts. Ishiguro et al. reported that elevated serum TPO and IL-6 concentrations precede thrombocytosis during acute infection.

Platelets are activated by endothelial injury, infection, acute/chronic inflammation, and anemia. Following infection, platelets are activated to interact with leukocytes, monocytes, bacteria, and others through multiple receptors. Activated platelets can interact with bacteria through various receptors such as Toll-like receptor (TLR)-4, a low-affinity receptor for the constant fragment (Fc) of immunoglobulin G (FcγRIIa) and glycoprotein (Gp) IIb/IIIa. As a result, platelets assist in clearing bacteria by presenting them to neutrophils, thus enhancing phagocytosis during infection.
Thrombopoietin

The regulation of platelet counts relies on the negative feedback mechanism of TPO, which binds to the TPO receptor on platelets and megakaryocytes, reducing plasma-free TPO levels and decreasing thrombopoiesis. A reduction in platelet counts leads to elevated plasma-free TPO, resulting in increased thrombopoiesis. Platelets are mainly removed from the blood in the spleen and liver.

It is well established that TPO concentrations in newborn infants are higher than those in adults. Levels increase after birth, peak on the 2nd day of life, and start decreasing at 1 month of age, and they are inversely correlated with platelet counts. Nakayama et al. reported similar results that serum TPO levels soon after birth to 1 month of age are significantly higher than those at 2–6 months after birth. Thereafter, TPO levels decrease with age (Fig.1). The low expression of TPO receptor on the platelets of newborn infants until 1 month of age decreases TPO clearance, leading to elevated plasma-free TPO levels and increased thrombopoiesis. Megakaryocyte precursor cells of newborn infants are more sensitive to TPO than those of adults. Platelet counts are typically low in small-for-gestational-age (SGA) newborns at birth. However, TPO concentrations in SGA newborns remain controversial. TPO concentrations were significantly elevated in SGA newborns compared to appropriate-for-gestational-age newborns (135 pg/mL vs. 94 pg/mL, respectively) and healthy term newborns (86 pg/mL vs. 72 pg/mL, respectively). On the other hand, TPO concentrations were not elevated in SGA newborns, especially those with liver problems.

Anemia and erythropoietin

Erythropoietin (EPO), a regulator of erythrocyte production, is structurally similar to TPO with an amino terminal domain containing a sequence homologous to that of EPO. Preterm infants with thrombocytosis show decreased red blood cell counts and elevated endogenous
EPO levels due to anemia, which probably results in elevated platelet counts. Increased production of thrombopoietic factors, cytokines, and other hematopoietic growth factors due to anemia might be related to elevated platelet counts as well. Yadav et al. and Özcan et al. reported that anemia was present in 29% and 9% of pediatric patients with thrombocytosis, respectively. It was also reported that 34% of newborn infants with anemia have thrombocytosis. These findings support a negative feedback mechanism of endogenous EPO in thrombocytosis, which has functional as well as structural similarities to TPO. Recombinant human EPO, a potent thrombopoietic factor, also affects thrombopoiesis and erythropoiesis.

**Infection and inflammation**

Platelets have pro-inflammatory activity in addition to pro-coagulant activity. Infectious pathogens and immune complexes formed by them can activate platelets, which then secrete molecules to activate immune reactions. Thom et al. suggested that rebound vigorous multi-lineage hematopoiesis following myelosuppression due to critical illnesses such as infection, inflammation, or iron deficiency can lead to hyperactive bone marrow and result in thrombocytosis.

**Classification of thrombocytosis (Table 1)**

**Classification by platelet count**

Thrombocytosis in newborn infants has variably been defined as a platelet count of > 400,000, > 600,000, or > 900,000/µL. Thrombocytosis as defined and classified by Sutor is widely used. Thrombocytosis has been defined as a platelet count ≥ 500,000/µL and classified as mild (platelet count 500,000–699,000/µL), moderate (platelet count 700,000–899,000/µL), severe (platelet count 900,000–999,000/µL), or extreme (platelet count ≥ 1,000,000/µL) according to peak platelet count.
Classification by etiology

Neonatal thrombocytosis is also classified as essential (primary) or reactive (secondary). In most cases, this is a reactive process. Essential thrombocytosis, a myeloproliferative neoplasm with an estimated annual incidence rate of approximately 1.00 per 100,000 adults, is extremely rare in newborns.\(^1\) The incidence of essential thrombocytosis in newborn infants is not well known. Essential thrombocytosis is represented by megakaryocytic hyperplasia in the bone marrow with thrombocytosis, which is mainly caused by mutations in Janus kinase 2 (\(JAK2\)) or myeloproliferative leukemia (\(MPL\), a TPO receptor gene).\(^2\) It is known to increase the risk of vascular events such as hemorrhage and thrombosis due to thrombocytosis. Antiplatelet or cytoreductive treatments are needed to reduce these complications. The differential diagnosis of essential thrombocytosis includes reactive (secondary) thrombocytosis, spurious thrombocytosis, or other forms of myeloproliferative neoplasms such as polycythemia vera or primary myelofibrosis. On the other hand, reactive thrombocytosis is considered a benign condition that is more common in newborns than in adults. Increased platelet production as a result of megakaryopoiesis occurs due to several conditions, such as infections, inflammation, tissue injury, and anemia, resulting in reactive thrombocytosis. Iron deficiency, hemorrhage, stress, exposure to epinephrine, vitamin E deficiency, and rebound from thrombocytopenia can also cause thrombocytosis.\(^3\) Denton and Davis\(^4\) reported that all pediatric cases of extreme thrombocytosis (platelet count \(\geq 1,000,000/\mu\text{L}\)) are reactive, requiring no treatment. Wiedmeier et al.\(^5\) reported that all cases of extreme thrombocytosis (platelet count \(\geq 1,000,000/\mu\text{L}\)) in newborns and young infants are reactive. All resolved without any hemorrhagic or thromboembolic complications. They also reported that infection (mostly respiratory tract infections), postoperative status, and anemia could cause thrombocytosis, generally a benign and self-limiting condition.\(^6\) Reactive thrombocytosis rarely increases the
risk of hemorrhagic and thromboembolic complications in newborn infants, and it usually does not require antiplatelet or cytoreductive treatment. Elevated platelet counts have also been reported in infants with severe osteogenesis imperfecta, notably those younger than two years of age. The authors suggested that elevated platelet counts in infants with severe osteogenesis imperfecta can serve as a marker of inflammation. Asplenia and sickle cell disease can also cause thrombocytosis, especially cases of extreme thrombocytosis (platelet count $\geq 1,000,000/\muL$) due to reduced platelet storage and removal.

**Initial platelet count at birth by gestational age**

Platelet counts increase with advancing gestational age. The initial platelet count at birth is correlated with gestational age, which is increased by 2,089/µL as gestational age increases by 1 week. Thus, preterm infants have lower initial platelet counts than late-preterm or term infants (Fig. 2). These findings are speculated to be related to higher TPO concentrations at birth in preterm infants than in term infants. There is an inverse correlation between TPO concentrations and platelet counts due to the negative feedback mechanism of TPO.

**Platelet count by postnatal age**

Platelet counts increase with advancing postnatal day, indicating a positive correlation with postnatal age. According to Wiedmeier et al., platelet counts are as high as 750,000/µL at 2–3 weeks and 6–7 weeks after birth. The cause of thrombocytosis at 2–3 weeks of age was speculated to be a neonatal surge in TPO. The cause of thrombocytosis at 6–7 weeks was not apparent. It has been suggested that infection, hypoxemia, and anemia can temporarily elevate platelet counts. According to Nakayama et al., the initial platelet count in preterm infants at birth was 230,000/µL, and it gradually increased to 579,000/µL 5 weeks of life. This increase continued, reaching a relatively high level at approximately 8 weeks of life.
Thrombocytosis in preterm infants

Thrombocytosis is more frequently observed in preterm infants than in term infants during the first few postnatal weeks to months. In one study, thrombocytosis was found in 38% of low birth weight preterm infants.\(^3\) Thrombocytosis was observed in 32.6% of preterm infants with a gestational age < 32 weeks who had not received recombinant human EPO.\(^4\) Higher TPO concentrations\(^3,32\) and more sensitive megakaryocyte precursor cells\(^3\) in preterm versus term infants are the speculated cause (Fig. 1).\(^3\) Low expression of TPO receptor in preterm infants leads to the accumulation of free TPO, which has been suspected to induce thrombocytosis in preterm infants.\(^12\) Anemia was frequently diagnosed in preterm infants with a gestational age < 32 weeks by Del Rey Hurtado de Mendoza et al., who reported an inverse correlation between decreased red blood cell counts and increased platelet counts in preterm infants.\(^4\)

Thrombocytosis was reportedly more common in preterm infants with a gestational age < 27 weeks than in those with a gestational age ≥ 27 weeks.\(^20\) Thrombocytosis in preterm infants was most common at a postmenstrual age (PMA) of 35 weeks, indicating that postnatal platelet regulation would mature at a PMA of approximately 35 weeks. McPherson and Juul\(^20\) suggested that the maturation of platelet regulation develops with PMA. The overall prevalence of abnormal platelet counts decreases with maturity in preterm infants.\(^20\) Developmental immaturity of platelet regulation might be associated with frequent thrombocytosis in preterm infants.

Complications of thrombocytosis

Thrombocytosis can increase the risk of arterial thrombosis, myocardial infarction, hemorrhage, and microvascular occlusion in adults.\(^1,2\) Moreover, adults with essential thrombocytosis are at an increased risk of leukemic transformation or evolution to myelofibrosis.\(^33\) However, reactive thrombocytosis, notably extreme thrombocytosis, in
newborn infants does not seem to be associated with an increased risk of hemorrhagic or thromboembolic complications.\textsuperscript{24,26,28,29) Preterm infants with a gestational age < 32 weeks and thrombocytosis had a higher incidence of retinopathy of prematurity according to Del Rey Hurtado de Mendoza et al.\textsuperscript{4) Conversely, other studies reported that thrombocytopenia, not thrombocytosis, is a risk factor for retinopathy of prematurity.\textsuperscript{34) Thus, the association between thrombocytosis and retinopathy of prematurity remains controversial.

\textbf{Conclusion}

Thrombocytosis is frequently found in newborns and young infants, especially those younger than 2 years of age. We often encounter preterm infants with thrombocytosis in neonatal intensive care units. Initial platelet counts at birth increase with advancing gestational age. Preterm infants have lower initial platelet counts than late-preterm or term infants. Thrombocytosis during the first few postnatal weeks to months is more frequently observed in preterm than term infants. A high TPO concentration and low TPO receptor expression on platelets leading to elevated plasma-free TPO levels, more sensitive megakaryocyte precursor cells to TPO, decreased red blood cell counts, and immaturity of platelet regulation have been speculated to cause thrombocytosis in preterm infants. Thrombocytosis in newborn infants, a reactive process following infection, acute/chronic inflammation, or anemia, is benign and resolves spontaneously, and unlike in adults, rarely associated with hemorrhagic or thromboembolic complications.

\textbf{Footnotes}

\textbf{Conflict of interest}

No potential conflict of interest relevant to this article was reported.

ORCID
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### Table 1. Classification of thrombocytosis

#### Classification by platelet count

<table>
<thead>
<tr>
<th>Classification</th>
<th>Peak platelet count (/µL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mild</td>
<td>500,000–699,000</td>
</tr>
<tr>
<td>Moderate</td>
<td>700,000–899,000</td>
</tr>
<tr>
<td>Severe</td>
<td>900,000–999,000</td>
</tr>
<tr>
<td>Extreme</td>
<td>≥ 1,000,000</td>
</tr>
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</table>

#### Classification by etiology

<table>
<thead>
<tr>
<th>Incidences</th>
<th>Essential (primary)</th>
<th>Reactive (secondary)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Causes</td>
<td>Mutations in (JAK2) or (MPL) (TPO receptor gene) → megakaryocytic hyperplasia in the bone marrow</td>
<td>Infection</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Acute/chronic inflammation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Tissue injury</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Anemia</td>
</tr>
<tr>
<td>Differential diagnosis</td>
<td>Reactive (secondary) thrombocytosis</td>
<td>Essential (primary) thrombocytosis</td>
</tr>
<tr>
<td></td>
<td>Other forms of myeloproliferative neoplasms (polycythemia vera or primary myelofibrosis)</td>
<td></td>
</tr>
<tr>
<td>Treatments</td>
<td>Anti-platelet or cytoreductive treatment</td>
<td>Requiring no treatment for thrombocytosis</td>
</tr>
<tr>
<td>Complications</td>
<td>Arterial thrombosis, myocardial infarction, hemorrhage, microvascular occlusion in adults</td>
<td>Rare</td>
</tr>
</tbody>
</table>

\(JAK2\), Janus kinase 2; \(MPL\), myeloproliferative leukemia; \(TPO\), thrombopoietin
Figure legends

**Fig. 1.** Corresponding changes in platelet counts and thrombopoietin (TPO) concentration.

TPO concentrations in newborn infants are increased after birth, peak on the 2nd day after birth, and start to decrease at 1 month of age. TPO levels then decrease with age. We found an inverse correlation between TPO concentration and platelet count. Preterm infants had higher initial TPO concentrations and lower initial platelet counts than term infants. Thrombocytosis during the first few postnatal weeks is more common in preterm infants than in term infants. Plotted from the results of Matsubara et al.\(^3\) and Ishiguro et al.\(^13\)

**Fig. 2.** Platelet counts by gestational age.

Platelet counts are shown for neonates with a gestational age of 22–42 weeks. Platelet counts increase with advancing gestational age, which is increased by 2,089/µL as gestational age advanced by 1 week. Thus, preterm infants have lower initial platelet counts than term infants. The mean values and 5th and 95th percentiles are given. Adapted from Wiedmeier et al.\(^11\) with permission from Springer Nature.
Corresponding changes in platelet counts and TPO concentration