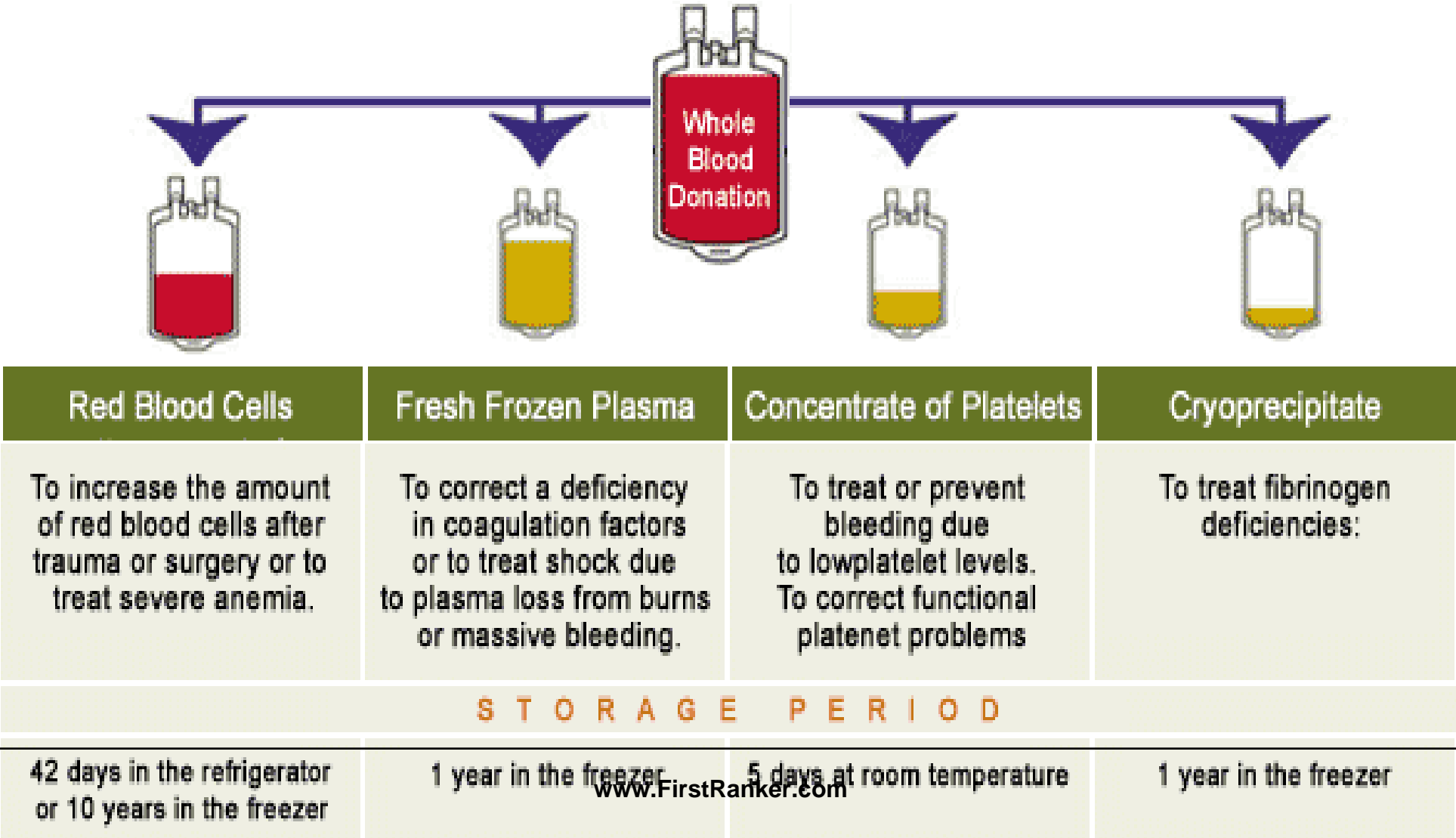


Rational Use of Blood and Blood Components

The potential of HUMAN BLOOD



Best Transfusion is “No Transfusion”

Why Avoid Blood Transfusion?

- **Infection Risk**
 - HIV, Hepatitis
- **Other Complications**
 - Febrile reactions
 - Allergic, urticarial reactions
- **Clerical Errors**
 - ABO mismatch
- **Immunologic Issues**
 - TA-GvHD
 - Immunosuppression
- **Religious Reasons**

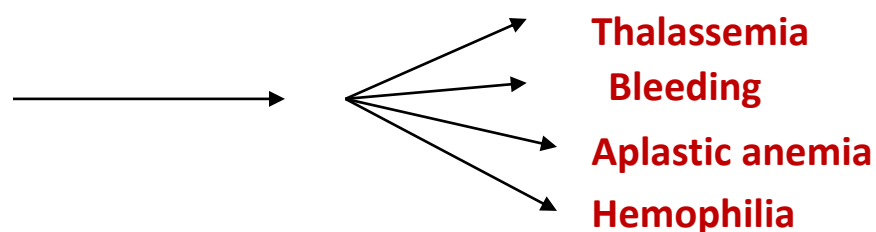
Misconceptions and Myths

- Whole blood
- “Fresh” Blood
- Empirical Transfusion
 - Nutritional Anemia
 - Pre Surgical
 - Wound Healing
 - Enhancement of well being

Why whole blood not rational

■ Maximize blood resource

one unit of whole blood



■ Better patient management

- concentrated dose of required component
- avoid circulatory overload
- minimize reactions

■ Specific storage requirements of components

- Red Blood cells +2-6⁰ C
- Platelets +22⁰ C
- Fresh frozen plasma - 30⁰ C

■ Decrease cost of management

- except for the cost of bag, other expenses remain same

Whole Blood Vs Packed Red Cells

Parameter	Whole blood	Packed red cells
Volume	350 – 450 ml	200 – 240 ml
Increment in Hb	1 -1.5 gm/dl	1 -1.5 gm/dl
Red cell mass /ml	Same as PRBC	Same as WB
Viable platelets	No	No
Labile factors	No	No
Plasma citrate	++++	+
Allergic reactions	++++	+
FNHTR	++++	+
Risk of TTI	++++	+
Waste of components	Yes	No

“Fresh blood” – misconception.

❖ What is “fresh blood”?

- varying definition
- any unit kept at 4°C for 4 hours is no longer “fresh”

❖ Increased disease transmission

- Intracellular pathogens (CMV, HTLV) survive in leukocyte in fresh blood
- Syphilis transmission- tryponema can’t survive > 96 hours in stored blood (JAMA,95)
- Malaria transmission- malaria parasite cannot survive > 72 hours in stored blood (Mollison)

“Fresh blood” – misconception.

❖ Immunological complication due to WBCs in fresh blood

- Transfusion Associated-Graft vs Host Disease – 90% fatality
- TA-immunomodulation
- Alloimmunization- Red cell / platelet

❖ Logistics

- no time for component preparation
- less time for infection screening
- storage lesions in different constituents due to storage temp

Rational Use of Blood

- **Right product**

- **Right dose**

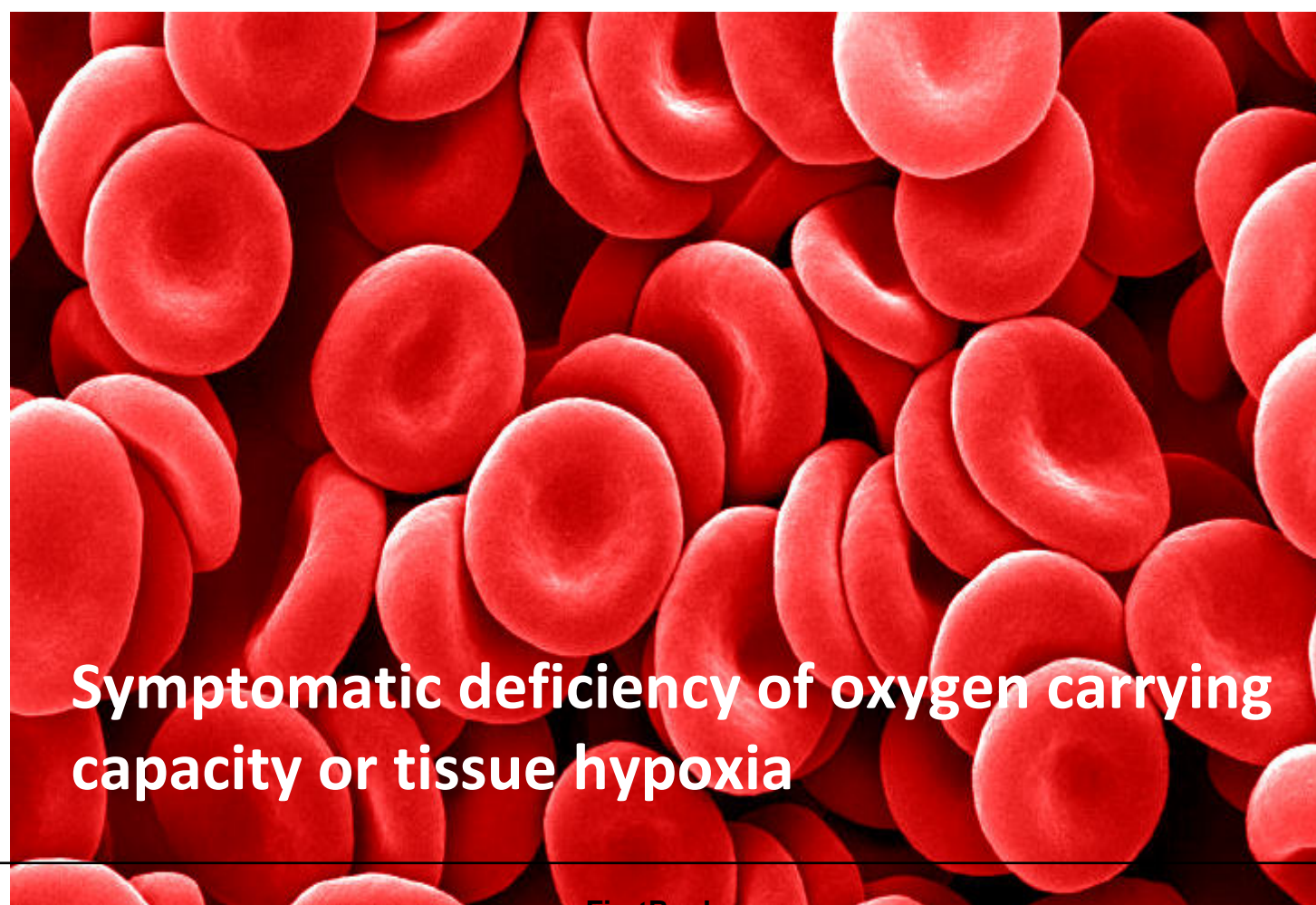
- **Right time**

- **Right reasons**

Answer 4 Qs before transfusion

- **Why to transfuse ?**
 - benefit > risk
 - patients symptoms Vs lab levels
 - prophylactic Vs therapeutic
- **What to transfuse ?**
 - whole blood **NO**
 - components / fractions
- **How much to transfuse ?**
 - Single unit **NO**
- **How to transfuse ?**
 - use of filter
 - rate of transfusion
 - warming

Packed Red Cells (PRBC)



Symptomatic deficiency of oxygen carrying capacity or tissue hypoxia

Appropriate use of Packed red cells

- Should be ABO and Rh compatible
- Clinical judgment- a vital role
- Co-existing conditions – age, general health, cause of anemia, its severity and chronicity
- Not for conditions like Iron/ B12/ Folate deficiency

PRBC - Triggers

- **Preoperative / peri-procedural** : Hb < 6g/dl
Hb 6- 10 g/dl
(bleeding, cardio resp. disease)
- **Symptomatic chronic anemia** : Hb < 6 g/dl
- **Acute blood loss** : > 40% blood loss
> 30% continued
blood loss or on
respiratory support

Neonates

- **Hemoglobin**

- <12g/dl in first 24 hrs
- <12 g/dl with intensive support care
- <11 g/dl with chronic oxygen need
- < 7 g/dl in a stable infant

- **Blood loss**

- Stable infant > 10% loss of estimated volume
- Unstable infant > 5% loss of estimated blood volume

PRBC - Dosing

- One unit of compatible RBC – **1 g/dl or Hct by 3%**
- Neonates
Dose – **10- 15 ml/kg**
Increase Hb - **2-3 g/dl**

Issues in red cell transfusion

One unit of PRBC

- Vol 250 ml
- Hct 65%
- Raise Hb by 1 gm/dl
- 200 mg iron
- 70% post transfusion survival

Age of blood

- concerns regarding K level
- decreased post transfusion survival

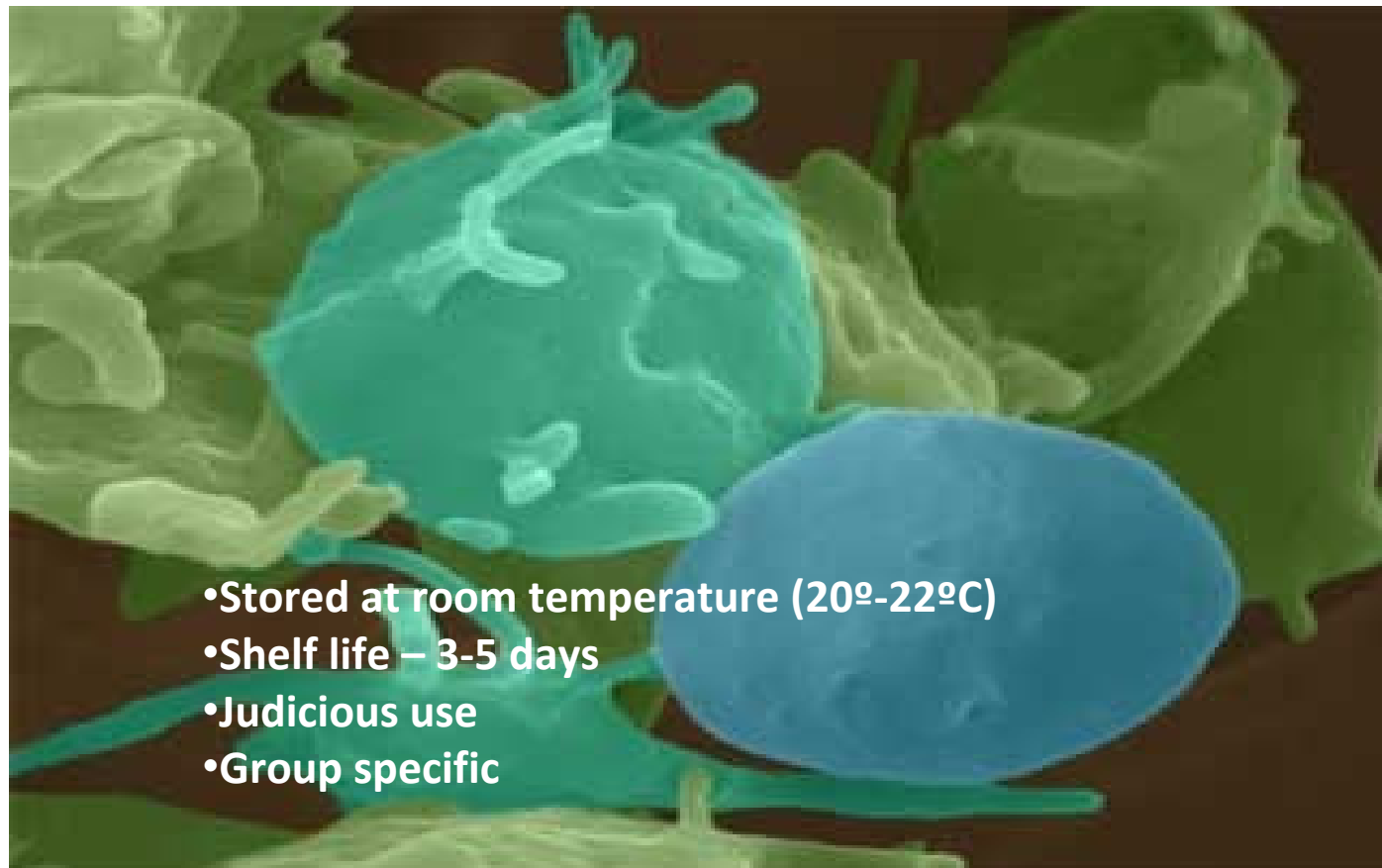
Specific conditions

- intrauterine transfusion < 3 days old
- thalassemics < 5 days
- open heart surgery < 10 days

Cardinal principles in red cell transfusion in chronic anemia

- Evaluate etiology of anemia - **AIHA, IDA**
- Do not transfuse just on the basis of given Hb level
- Try to establish whether Signs / Symptoms are due to anemia
- Determine if Signs / Symptoms of anemia are alleviated by transfusion
- Determine that temporary relief of symptoms warrants continued transfusion

Platelets



Appropriate Transfusion of Platelets

- **Symptomatic platelet problems**
Number related – eg. Aplastic anemia
Function related – eg. Glanzmann's thrombasthenia
- **Do not treat the number in isolation –**
eg Chronic ITP with no bleeds
- **Prophylactic in specific situations**
CNS, eye surgery, other major surgeries, acute leukemia, patients on chemoradiotherapy

Dose: 1 RDP/10 Kg

Platelet- Triggers

Condition	Platelet count
Prophylaxis against bleeding	< 10,000/ μ l
Bedside invasive procedures	< 50,000/ μ l
Neurosurgical procedures, Ophthalmic surgeries	< 100000/ μ l
Massive Transfusion	< 50,000/ μ l

Neonates – Prophylactic Platelet Triggers

Term Neonates

- Clinically stable - 20,000/ μ l
- Clinically sick - 30,000/ μ l

Preterm Neonates

- Clinically stable - 30,000/ μ l
- Clinically sick - 50,000/ μ l

Contraindications

- Thrombotic Thrombocytopenic purpura
- Heparin induced thrombocytopenia
- Immune Thrombocytopenic purpura

Fresh Frozen Plasma



Appropriate Transfusion of FFP

- Replacement of multiple factors: **DIC, liver disease, warfarin reversal, snake bite**
- **PT/ INR** should be determined
- Dose: **10-15 ml/kg**
- **Not** for volume expansion
- **Not** for nutritional support/ hypoproteinemia

Cryoprecipitate

- Out of group can be transfused but preferably ABO compatible
- RhD type need not be considered
- Thawed Cryoprecipitate transfused within 6 hours
- Indicated for bleeding associated with fibrinogen deficiency and factor XIII deficiency

- Hemophilia A or von Willebrand disease when appropriate substitute not available
- Bleeding with fibrinogen levels< 100mg/dl
- Dose - **one unit/10 kg body weight**
- Raises fibrinogen concentration by **50 mg/dl**

Choice for ABO Blood Groups

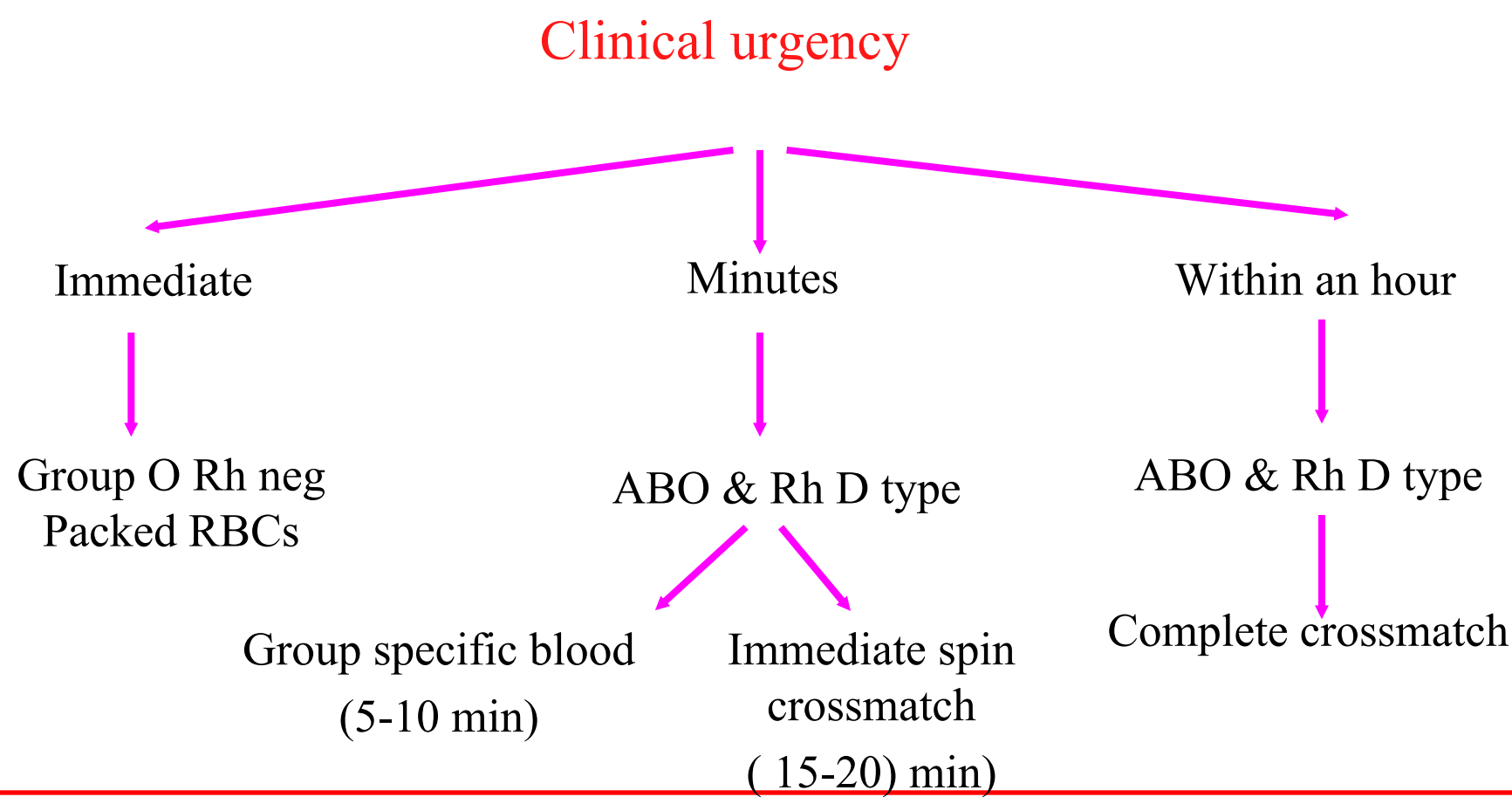
Patient type	Donor PRBC	Donor FFP	Donor PC
O Positive	O	O,B,A, AB	O,B,A, AB
A Positive	A, O	A, AB	A, AB ,O,B
B Positive	B, O	B, AB	B, AB ,O,A
AB Positive	AB,B,A, O	AB	AB ,B,A,O

Choice for Rh Blood group

- Rh (D) negative patient transfused with Rh (D) positive components**

PRBC	Only as a life saving measure and with consent from treating physician & patient's relative
FFP	No anti-D immunoprophylaxis required
PC	Anti D immunoprophylaxis required (300 µg anti-D gives protection for 7 plateletpheresis units or 30 Rh (D) positive platelet concentrates for 6 weeks)





Cross matching: Special Circumstances



If units are issued without X match – written consent of physician to be taken, -complete X match protocols followed after issue

Take Home Messages

- **No place for Whole Blood in clinical medicine**
- **Component preparation and use is the demand of time**
- **Best Transfusion is “No Transfusion”**
- **Promotion of judicious use of blood / components**

-  Audit of transfusion practices
-  CME on use of components
-  Promote autologous use of blood
-  Discourage single unit / fresh blood

Thank You