

# Fresh Whole Blood – a possible role in civilian preparedness

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# Back to the future

Whole blood -> Blood components ->  
Whole blood?

# Blood component therapy

- Designed for the haemato-oncology patients
- Designed for the anaemic patients
- Designed for plasma fractionation
- What about the acute severely bleeding patient ?
- Who are the patients needing transfusion in (civilian) disasters?

# Blood component therapy

- The blood bank facility may be damaged
- The blood bank capacity may be too limited for the supply needed
- May be dependent on “import” of blood components

# Back to the future?

- NO!
- Fresh whole blood/whole blood – what is what?

# Disposition

- When the helicopter can not fly
- Regulatory issues
- Extreme situations
- How Fresh Whole Blood/Whole Blood must be improved



## 2.2 REGULATORY CONCERNS

The availability of blood may be the primary concern in a disaster, but the safety of the blood supply is also paramount. Adherence to Food and Drug Administration (FDA) regulations is crucial.



# Regulatory issues

- Register Fresh Whole Blood as a “Blood component” (EU Blood Directive)





# What is Fresh Whole Blood

- Storage of red cells
- Storage of platelets
- Storage of plasma
- Removal of leukocytes?

# Fresh Whole Blood

Experiences from the past:

Blood stored less than 24h

Blood stored less than 48 h

# Storage solution

- Reduce activation during blood donation
- Reduce storage lesion
  
- Combined additive solution for red cells and platelets is possible (Heaton, 1990)
- Reducing citrate concentration may stabilize coagulation factors

# Storage temperature

- Reduce metabolism
- Reduce risk of bacterial contamination (growth)
- Simple in theory, difficult in real life

# Red cell storage lesion

- Temperature 0 -> - 2°C (Strumia, 1954)
- Or 1-2°C (Strumia, 1954-2)
- Or 4-25°C (Strauss, 1974)
- 22°C is good for short-term (Avoy, 1978)
- 37°C is bad! (Jandl, 1958)
- Improving red cell storage solution (Sparrow, 2011)



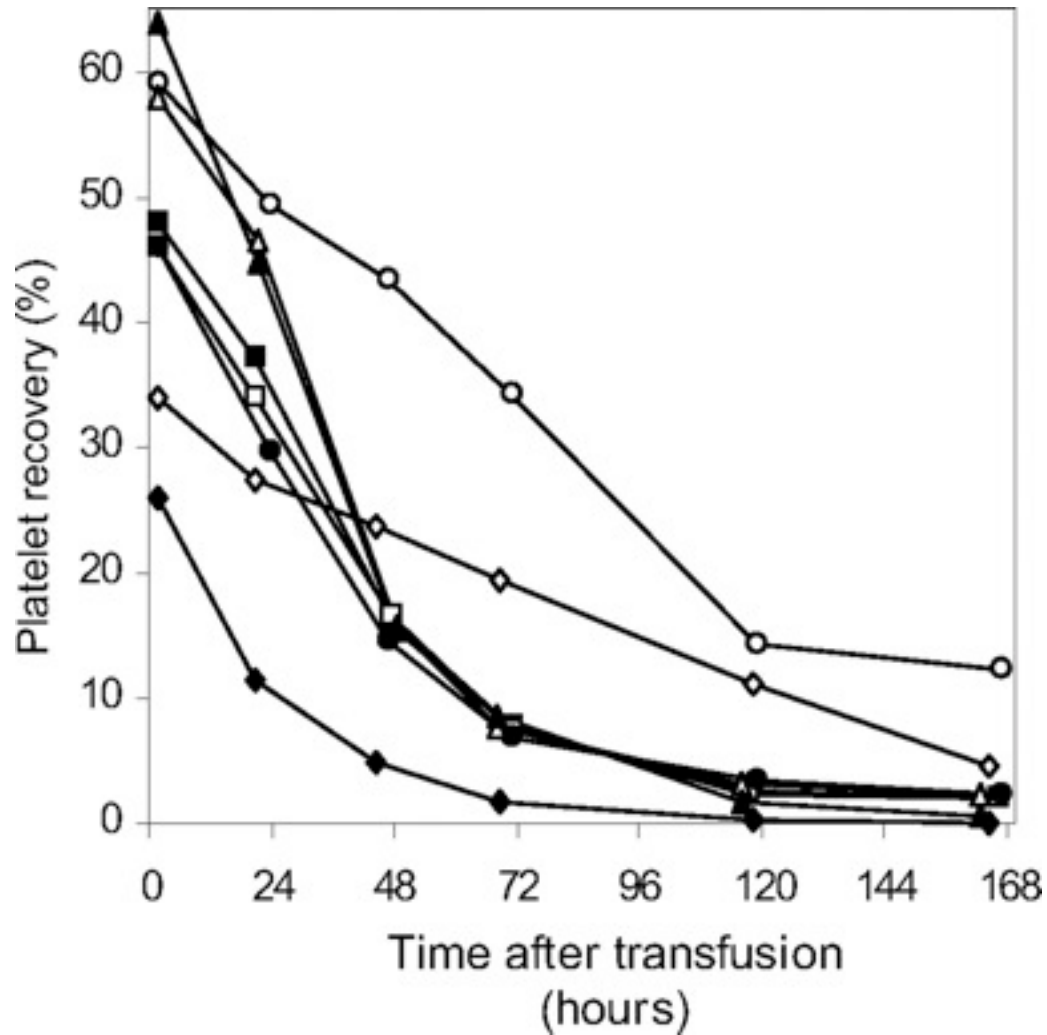
# Platelet storage lesion

Recovery dependent on storage temperature

- 4°C recovery 45%, 2d 15%
- 22°C recovery 63% (Becker, 1973)
- Slichter et Harker, 1973

	4°C	22°C
Recovery	63%	51%
Survival	1.3d	8.2d
The GP Ib alpha effect		

## In vivo autologous radiolabeled human platelet recoveries and survivals.



Wandall H H et al. Blood 2008;111:3249-3256

# Stability of plasma proteins and coagulation factors

- F VIII is reduced to 40% after 24h storage (Slichter, 1976)
- F VIII is reduced by 23% after 24 h storage temperature. Only minor reduction of other factors, including protein S and protein C (Cardigan, 2011)

# Leukocyte depletion of Fresh Whole Blood

- Leukocytes perform phagocytosis
- But, then.....

# Fresh Whole Blood and Graft-versus-Host Disease

- Data from cardiac surgery in Japan
- T-lymphocytes from the donor attack the host's immune system

# Prevention

- Gamma irradiation
- Removal of leukocytes (filtration)
- Keep the platelets alive

# HLA-immunization

- Children receiving whole blood in connection with cardiac surgery form HLA-antibodies (Opheim, submitted 2011)
- Primary immunization can be prevented by filtration

# Conclusions

Fresh Whole Blood will not replace blood components

The component must be improved and better defined

If this will be successful, the clinical usage may be increased – and during disasters it may be THE blood component of choice