THERAPEUTICAL MANAGEMENT

According to the European consensus conference « European approach for the medical management of mass radiation exposure » updated in October 2017

Beyond the first 48 hrs, a second patient scoring is done by organs (Haematopoietic, Gastrointestinal, Cutaneous, Neurovascular) according to the METREPOL document for therapeutical management and Multiple Organ Failure (MOF) prediction.

Cytokines

Score I: Monitoring. No cytokine

- Outpatient clinical monitoring.
- Blood count day 1 2 and then once a week for 2 months.

Score II: Cytokines (curative)

- G-CSF (Pegylated or not) should be used within 48 hrs or as soon as possible until neutrophil recovery (ANC > 0.5 x 10°/L). EPO and TPO agnoists can be used if needed. Routine marrow failure support with antibiotics, blood products as per routine haemato-oncology care.
- Symptomatic treatment of gastrointestinal damage.
- If severe aplasia → Protected environment.
- Accidental radiation exposure is generally heterogeneous.
 Some under-exposed/protected regions of bone marrow can give rise to endogenous haematopoietic recovery.

Score III: Cytokines (until reappraisal of score)

- Patients to be treated as score II until it is clear that they are score III.
- Palliative and end of Life care to be initiated.
- Re-evaluation of score during the first week based on laboratory or clinical symptoms revealing irreversible organ damage or MOF.

ALL BLOOD PRODUCTS SHOULD BE IRRADIATED.

SEVERE RADIATION SKIN LESIONS HAVE A PECULIAR EVOLUTION. CONSIDER MESENCHYMAL STEM CELLS AT SPECIALIST CENTRES.

References:

- Gorin NC et al Ann Hematol, 85: 671-679, 2006.
- Fliedner TM et al Medical Management of Radiation Accidents Manual of the acute radiation syndrome, published by BIR, 2001.
- Powles R et al Health Phys, 98: 810-814, 2010.

Haematopoietic stem cell (HSC) transplantation

Background

- HSC transplantation is not an emergency.
- It is crucial to avoid GVHD in order not to compromise an endogenous recovery.
- If severe aplasia persists under cytokines for more than 14 days, the possibility of an haematopoietic stem cell (HSC) transplantation is discussed (as below).

Criteria to transplant

- Severe marrow aplasia persisting 14 21 days despite cytokines.
- No residual haematopoiesis on bone marrow biopsy.
- No other irreversible organ damage.
- Treated or controlled infection, if present

Graft

- Type of graft:
 - Bone marrow.
 - Peripheral blood HSC (depleted or not).
 - Cord blood.
- Donor in the following order of priority (as per current transplant criteria):
 - HLA-identical sibling.
 - HLA-identical unrelated donor.
 - Cord blood > 4/6 matched.
 - Haplo-identical.
- Doses of cells to be grafted:

At least:

- 2x10⁶ CD34 cells. kg⁻¹ (peripheral blood).
- 2x108 nucleated cells.kg-1 (bone marrow).
- 3x10⁷ nucleated cells (cord blood).

Conditioning and GVHD prevention (as per current transplant criteria):

- Reduced intensity conditioning.
- No Methotrexate.

EUROPEAN APPROACH

FOR THE MEDICAL MANAGEMENT OF MASS
RADIATION EXPOSURE
THERAPEUTICAL MANAGEMENT



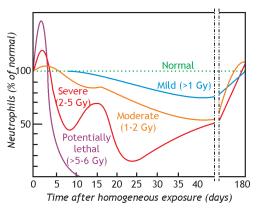




What can the European Blood and Marrow Transplant Group (EBMT) offer?

Advice on:

- 1. Secondary Triage and Treatment
- 2. Ongoing feedback from experts on how the clinical scenario is evolving
- 3. Optimizing care outside of national borders
- 4. Skilled network of 500 BMT Centres
- 5. Generating prospective database of event



The amount of energy absorbed by the organs of the body is measured in Gray (Gy); the effects of this radiation is given by the equivalent dose which is measured in Sievert (Sv).

For x-rays, gamma rays, and beta particles 1 Gray \approx 1 Sievert.

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THE FIRST 48 HOURS

Decontamination after stabilisation.

Life-threatening wounds and burns should be treated first.

Radiation dose review comes later – Irradiation is not contamination – An irradiated person is not a source of radiation.

Acute Radiation Injury

The severity of prodromal clinical features is indicative of probable significant injury.

- Extensive and immediate erythema.
- Early Transient Incapacitation Syndrome (temporary loss of consciousness).
- High fever.
- Hypotension; Early Vomiting.
- Immediate diarrhoea.

Accident Characterisation

- Inquiry: circumstances of the accident (is irradiation +/-contamination present; use contamination monitoring device), source characteristics, source-victim geometry, duration of exposure, shielding, homogeneous / heterogeneous irradiation.
- Labelling and storage of personal belongings and clothes, biological material (hair, nails).

Urgent sampling

- Blood cell counts (+ differentials) every 4-8 hours for the 1st 24 hours, 12-24 h every day after.
- Red cell group typing
- Standard biochemistry + amylasemia.
- Urine and faeces if radionuclide contamination is suspected
- Store serum and cells or DNA for further analyses including HLA typing.
- Chromosome aberrations on blood lymphocytes (biodosimetry) (15 ml + heparin). Seek advice from national / international biodosimetry networks as soon as possible.

Primary scoring

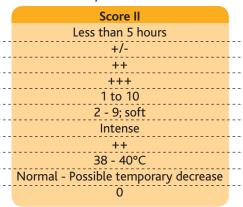
Record all clinical symptoms on a date and hour-stamped chart

1	Score I
Average delay before symptoms appear	Less than 12 hours
Cutaneous erythema	0
Asthenia /Weakness	+
Nausea	+
Vomiting per 24 hrs	Maximum 1
Diarrhea / Number of stools per 20 hrs	Maximum 2 - 3; bulky
Abdominal pain	Minimal
Headaches	0
Temperature	Below 38°C
Blood pressure	Normal
Temporary loss of consciousness	0

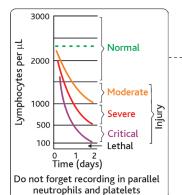
At 24 hours

At 48 hours

Above 1 500 / μL	
Above 1 500 / μL	



Score III
Less than 30 minutes
 +++ ; before 3 nd hour
 +++
 (-)
 Above 10; intractable
 Above 10; watery
 Excruciating
 Excruciating; Signs of intra-cranial HT
 Above 40°C
 Systolic below 80
 + / Coma



Below 1 500 / μL Below 1 500 / μL

Depletion of blood lymphocytes

Below 500 / µL Below 100 / µL

Outpatient monitoring

Hospitalisation for curative treatment

Hospitalisation (MOF predicted) Multiple Organe failure (MOF)

WARNING: the symptoms and values indicated above are reliable only in case the whole body or large parts of the body have been externally exposed to a high radiation dose delivered within few minutes or few hours. Fill and fax EBMT MED A to: (+33)1 71 97 04 88 To download EBMT MED A: www.ebmt.org in Data-Management/Registry structure/data collection forms & manuals