



World Health
Organization

Session 3

Radiation effects and health risks from radiation exposure at the workplace

Contributed papers

Rapporteur: Dr María del Rosario Pérez

Department of Public Health, Environmental and Social Determinants of Health

Summary of contributed papers

- Two contributed papers from **Iran** and **Egypt**
 - 1. Occupational Radiation Exposure, DNA Damage and Genetic Polymorphisms in DNA Repair Genes**
(F. Zakeri et al)
 - 2. Endothelial Progenitor Cells in Peripheral Blood of Cardiac Catheterization Personnel**
(S. Korraa et al.)

These papers have two common features

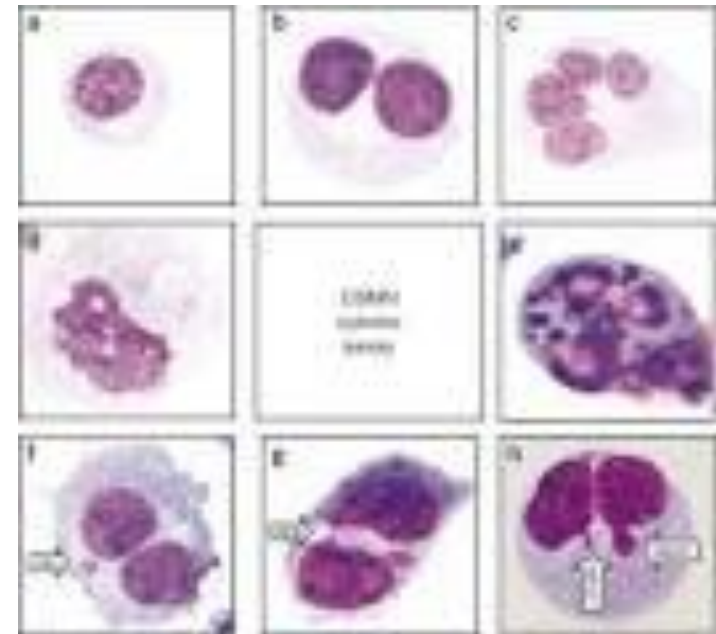
Study population:

- *health workers exposed during fluoroscopy procedures*



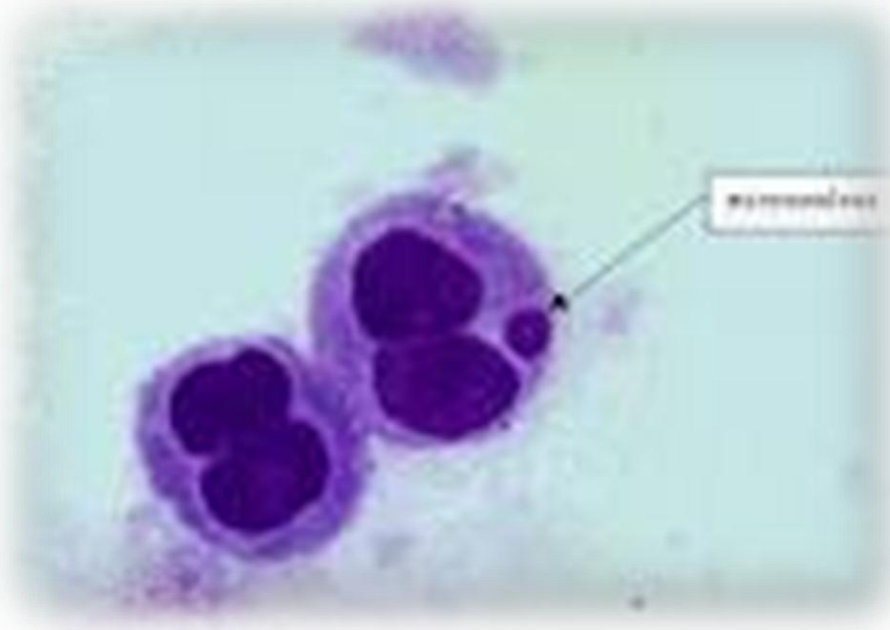
One of the end points:

- *frequency of micronucleus (MN) in peripheral lymphocytes (cytokinesis-block micronucleus assay)*



Frequency of MN in peripheral lymphocytes

- The cytokinesis-block MN assay in peripheral blood lymphocytes is a validated technique of biological dosimetry (e.g. accidental exposures). It has been also used to evaluate levels of DNA damage in workers occupationally exposed to radiation.



- *In addition to radiation, other genotoxic agents (e.g. tobacco) can increase the MN frequency.*
- *Baseline MN frequency depends strongly on age and gender.*

Paper N° 131

Occupational Radiation Exposure, DNA Damage and Genetic Polymorphisms in DNA Repair Genes

F. Zakeri^{a,b}, MR Farshidpour^b, MR Rajabpour^b, MJ Ahmadpour^b, F. Mianji^{a,b}

- a) Nuclear Science and Technology Research Institute, Tehran, Iran*
- b) Iran Nuclear Regulatory Authority, Tehran, Iran*

Objective: to determine the relationship between genetic polymorphisms in genes coding DNA repair enzymes and the levels of DNA damage in interventional cardiology staff



Study population

| Characteristic | Exposed group ^(a) | Control group |
|----------------------------|------------------------------|---------------|
| Males | 59 | 38 |
| Females | 31 | 22 |
| Total | 90 | 60 |
| Mean age (years +/- SD) | 41.5 +/- 7.6 | 41.4 +/- 9.1 |
| Mean last year exposure | 3.5 +/- 2.7 mSv | - |
| Mean last 5 years exposure | 11.2 +/- 10.5 mSv | - |
| Mean years of employment | 9.5 +/- 6.7 | - |

^(a) *interventional cardiologists, technicians and nurses*

End points in peripheral blood lymphocytes

- **Frequency of micronucleus**

by cytokinesis-block micronucleus test in binucleated cells



- **Single nucleotide polymorphisms (SNPs)**

by polymerase chain reaction combined with restriction fragment length polymorphism (PCR-RFLP genotyping assay)

In genes coding DNA repair enzymes:

1. XRCC1
2. OGG1
3. APE1
4. XRCC3
5. XPG

Results

- MN frequency significantly higher in:
 - *exposed group vs. control group*
 - *within exposed group $>3\text{mSv/y}$ vs. $\leq 3\text{mSv/y}$*
 - *within exposed group >10 years vs. ≤ 10 years of exposure*
 - *exposed group carrying SNPS in the genes XRCC3 and XPG*
 - *control group carrying SNPs in the gene OGG1*

Authors' conclusion

- Occupational exposure to IR in interventional cardiologists, technicians and nurses is associated to increased DNA damage (expressed as higher MN frequency).
- DNA damage was higher in individuals carrying genetic polymorphisms in DNA repair enzymes (SNPs), suggesting that this might represent a particularly vulnerable population (mutagenic and cancer risk).
- The relationship between MN and SNPs in genes involved in DNA repair may contribute to evaluate susceptibility to ionizing radiation in individuals occupationally exposed.



Paper N° 168

Endothelial Progenitor Cells in Peripheral Blood of Cardiac Catheterization Personnel

S. Korraa^a, MS Tawfik^a, A Zaher^b, M Maher^c

- a) *Department of Radiation Health, National Centre for Radiation Research and Technology, Cairo, Egypt*
- b) *National Health Institute, Cairo, Egypt*
- c) *Faculty of Science, Suez Canal University, Cairo, Egypt*

Objective: to evaluate the level of circulating endothelial progenitor cells and its correlation with DNA damage in staff exposed during fluoroscopy cardiac procedures.

Study population

| Characteristic | Exposed group ^(a) | Control group |
|-------------------------|------------------------------|---------------|
| n Total | 70 | 40 |
| Smokers | 34 | - |
| Non-smokers | 46 | 40 |
| Mean age (years +/- SD) | 42.8 +/- 5.2 | 42 +/- 4.8 |
| Annual dose (range) | 2.16 – 8.44 mSv/year | - |

(a) Staff involved in fluoroscopy-guided cardiac catheterization in 3 hospitals in Cairo, Egypt

Biological end points

- **Frequency of micronucleus**

by cytokinesis-block micronucleus test in binucleated cells

- **Number of Endothelial Progenitor Cells (EPCs) in peripheral blood (PB)**

by flow cytometry with monoclonal antibodies for CD133, CD34 and kinase domain receptors (KDR)

- **Plasma levels of stromal growth factor (SDF-1)**

using ELISA assay

Results

- The staff involved in fluoroscopy-guided cardiac catheterization (CC) presented:
 - *Significantly higher micronucleus (MN) frequency*
 - *Significantly higher number of Endothelial Progenitor Cells (EPCs) in peripheral blood*
 - *Significantly higher plasma levels of stromal growth factor (SDF-1)*
- Smoker CC staff exhibited higher MN frequency and SDF-1 and lower levels of EPCs than non-smokers

Authors' conclusion

- Staff involved in fluoroscopy-guided cardiac catheterization present and increased MN frequency, which is higher in smokers vs. non-smokers.
- Circulating EPCs numbers and SDF-1 plasma levels, which are markers of endothelial activation/damage, are significantly increased in the radiation exposed group.
- The dual effect of IR and smoking (i.e. additive effect on inducing SDF-1 expression as other DNA damaging agents, with an opposite effect on the number of EPCs in peripheral blood) is interpreted as a regenerative process decreased by smoking.
- Further studies are needed to elucidate the role of EPCs as a potential marker of radiation exposure.

