

EFFECT ON THE CCRF-CEM CELLS EXPOSED TO RF/MW

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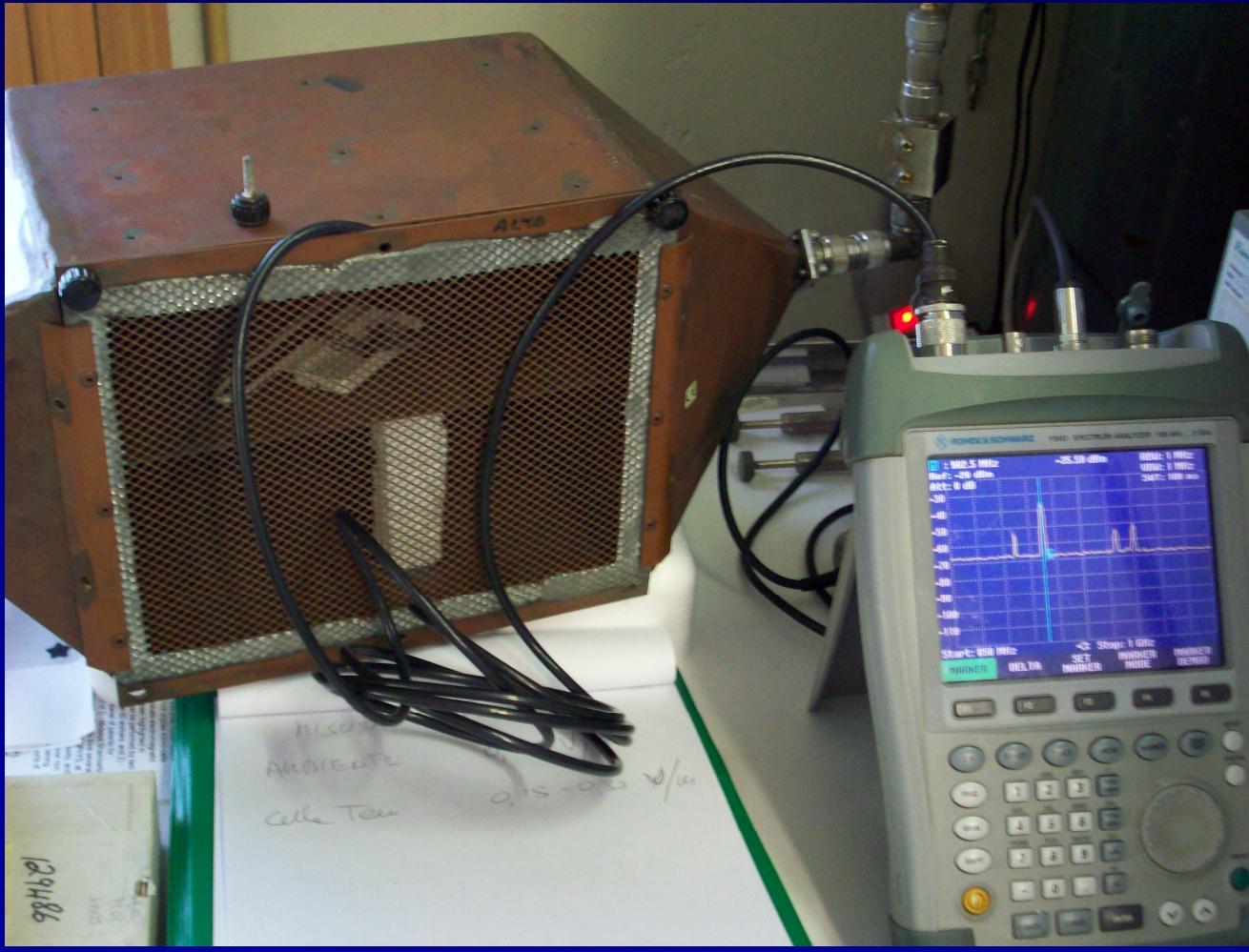
PROTOCOLLO SPERIMENTALE TEST MOBILE BADGE

- EXPOSURE IN TEM CELL of the cells CCRF-CEM by electromagnetic field of 900 MHz in presence and without MOBILE BADGE..
- 900 MHz telephone connected to CMD 55
- PROLIFERATION TEST
- WB analysis CASPASI Bcl2
- FACS analysis

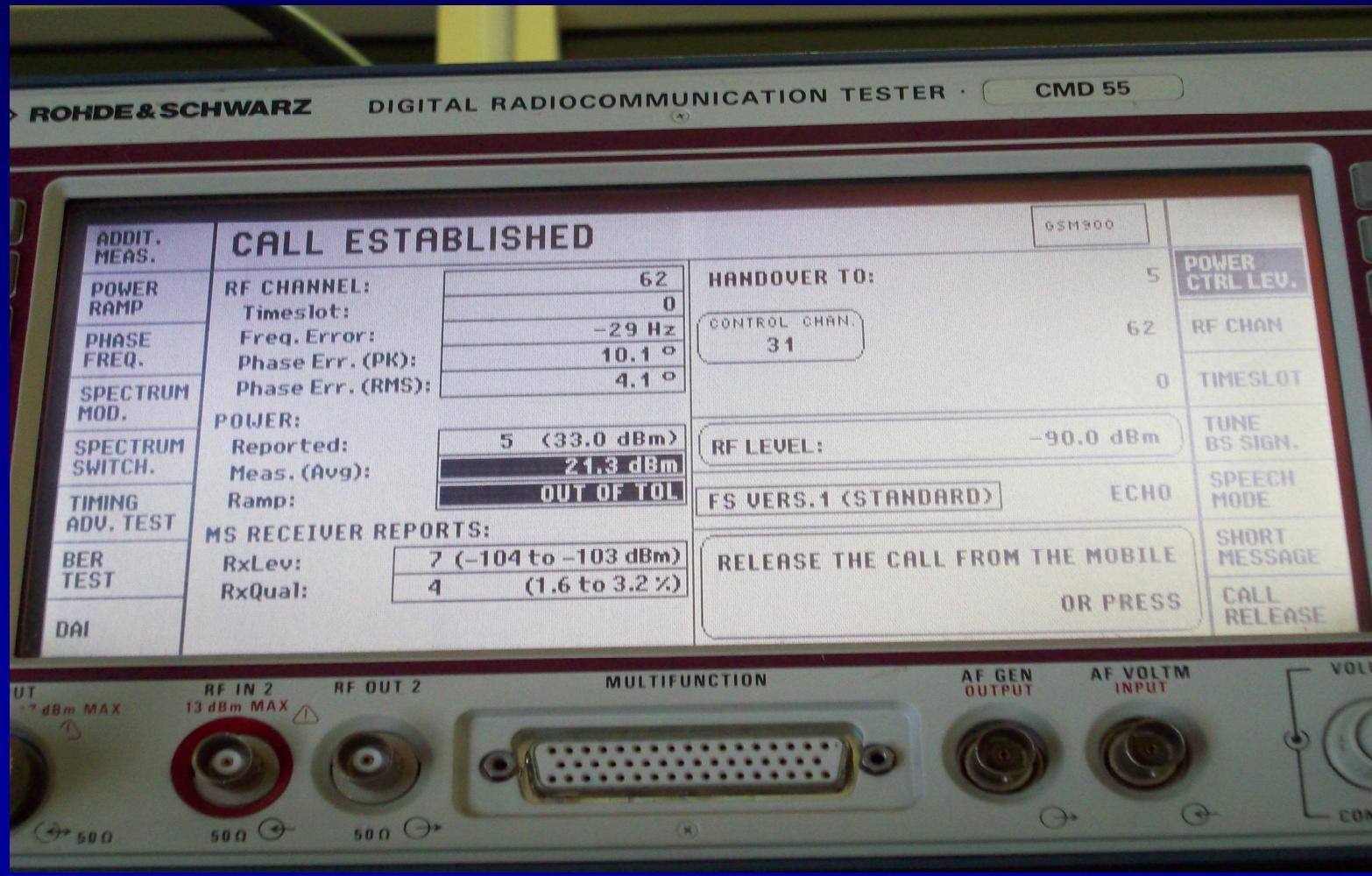
GSM PER ESPOSIZIONI



SEGNALE GSM 900 MHz IN CELLA TEM

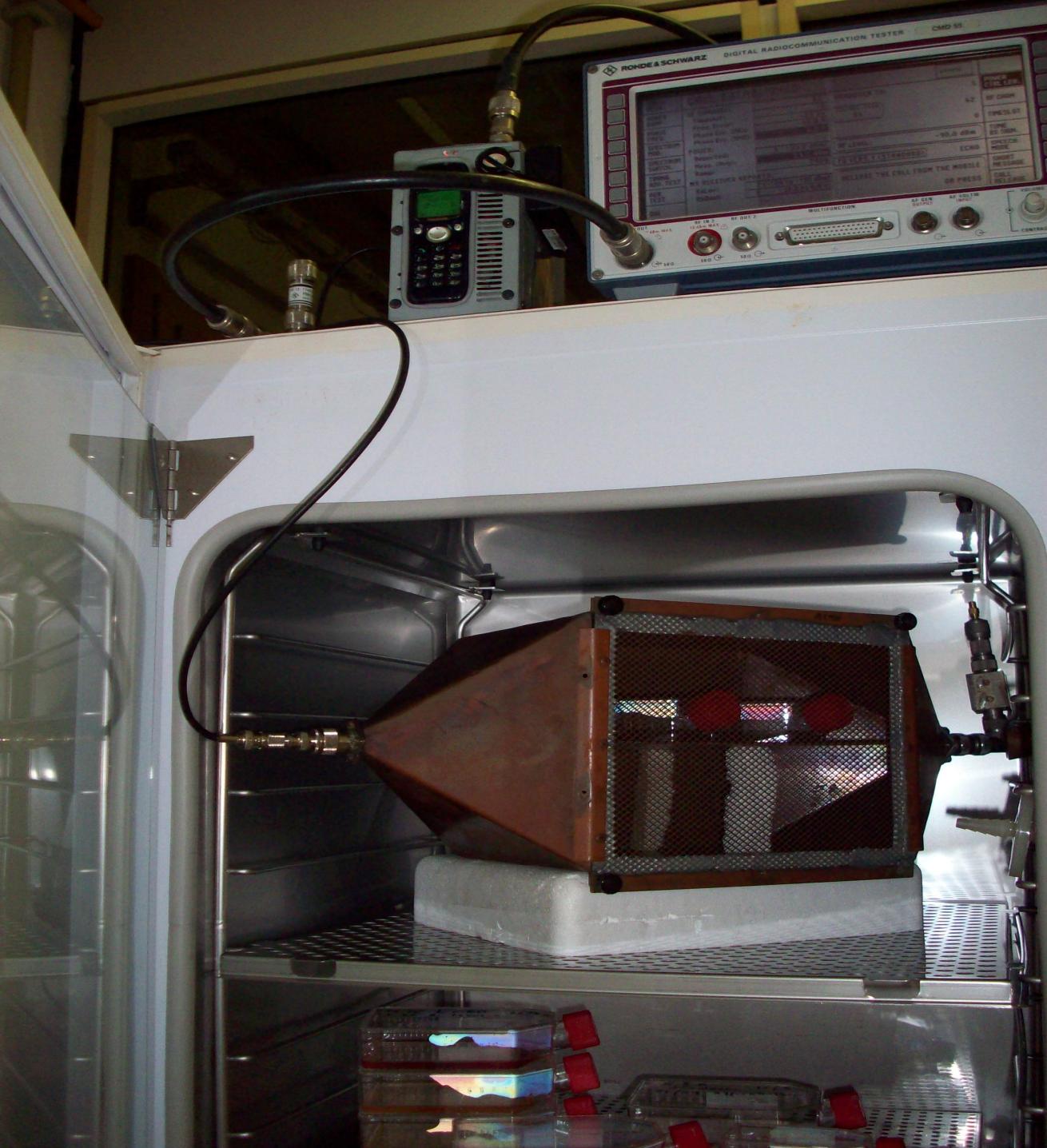


CARATTERIZZAZIONE DEL SEGNALE GSM



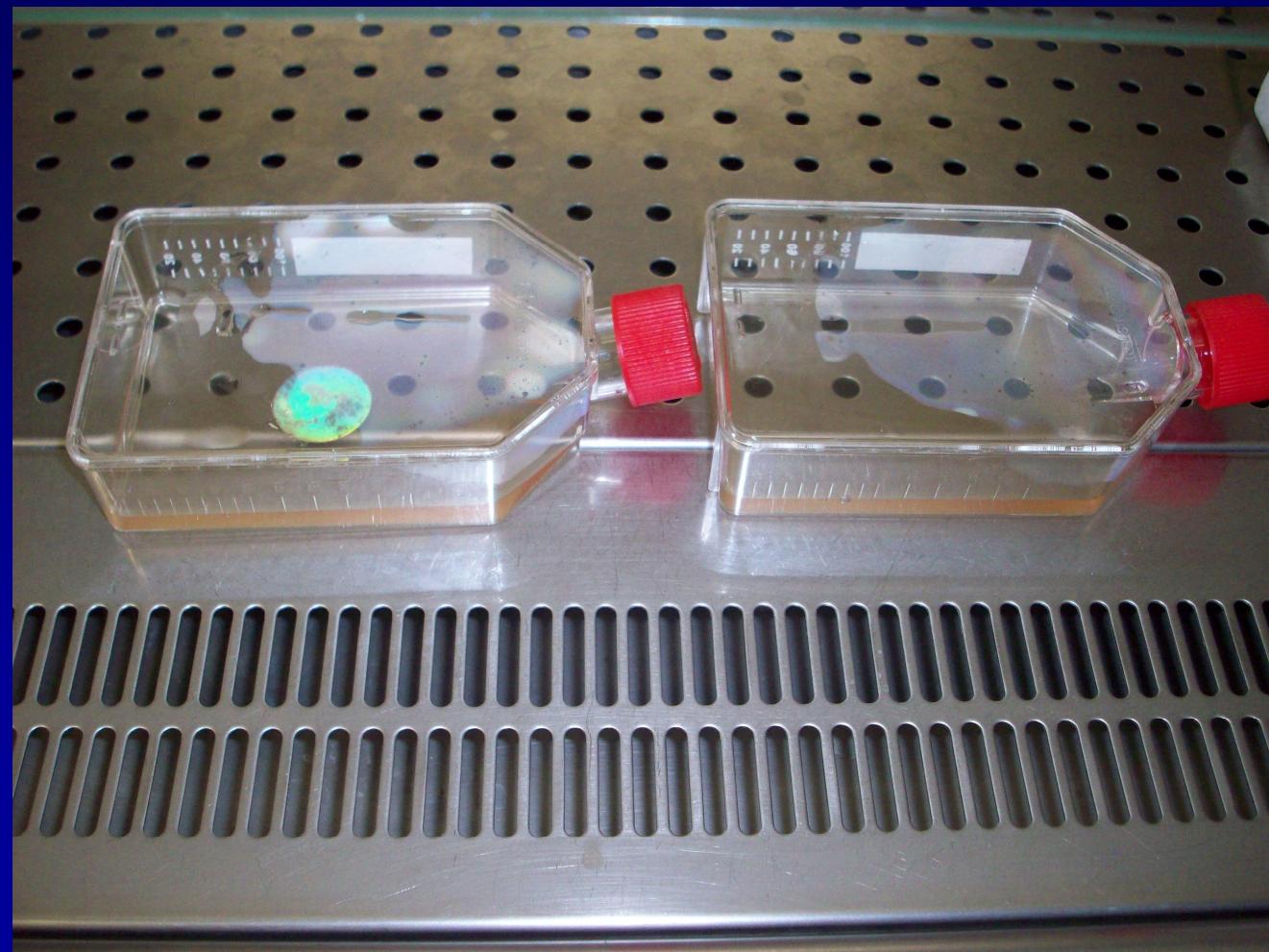
GSM -27.4 dBm





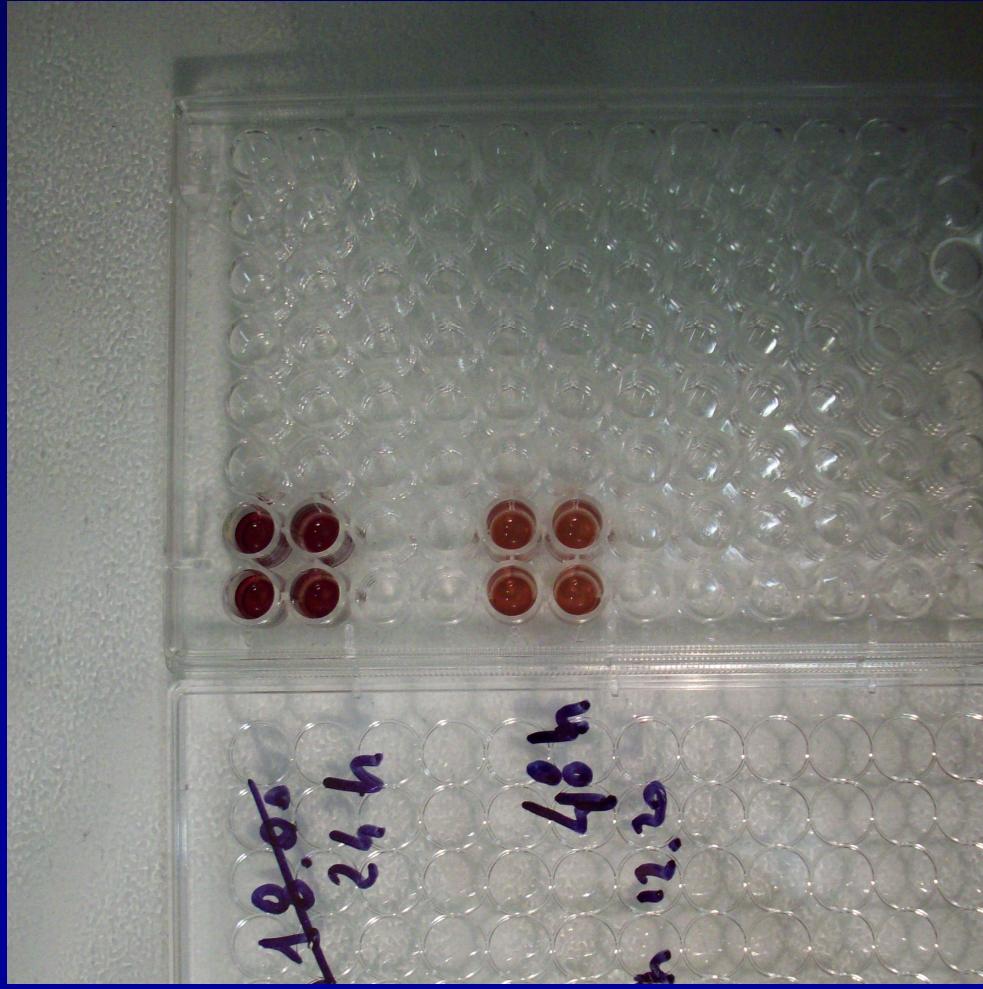








PROLIFERAZIONE CELLULARE



PRELIMINARY RESULTS WB

Caspase-3

- 1- Ctrl 24h Out
- 2- E + B Exp 24h In
- 3- Exp 24h In
- 4- Ctrl 24h
- 5- Exp 24h



Risultati I° Esperimento WB

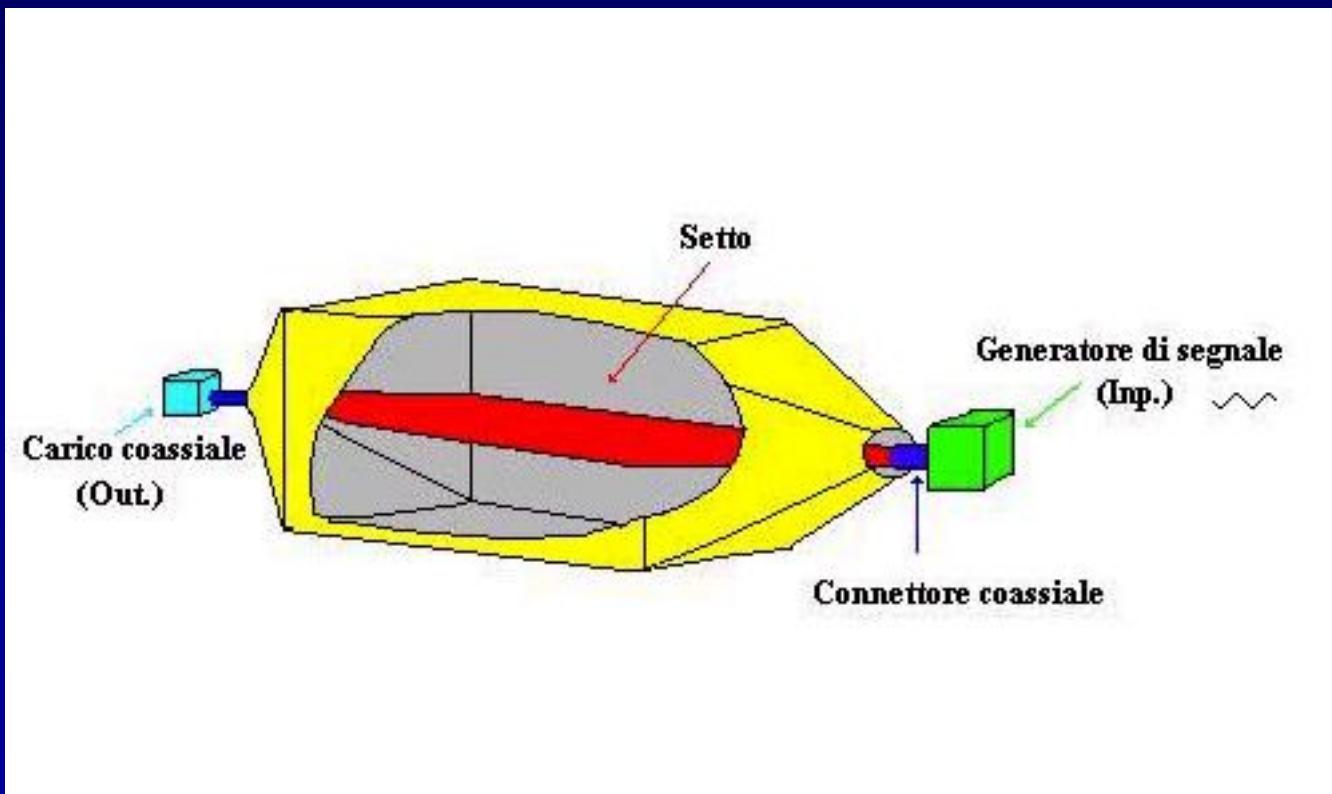
- Effetto del Badge relativo alla espressione genica della Caspasi3 che induce apoptosi nelle cellule Esposte a Segnale GSM
- .
- Da testare l'antagonista BC12
- Quantificare le proteine espresse.







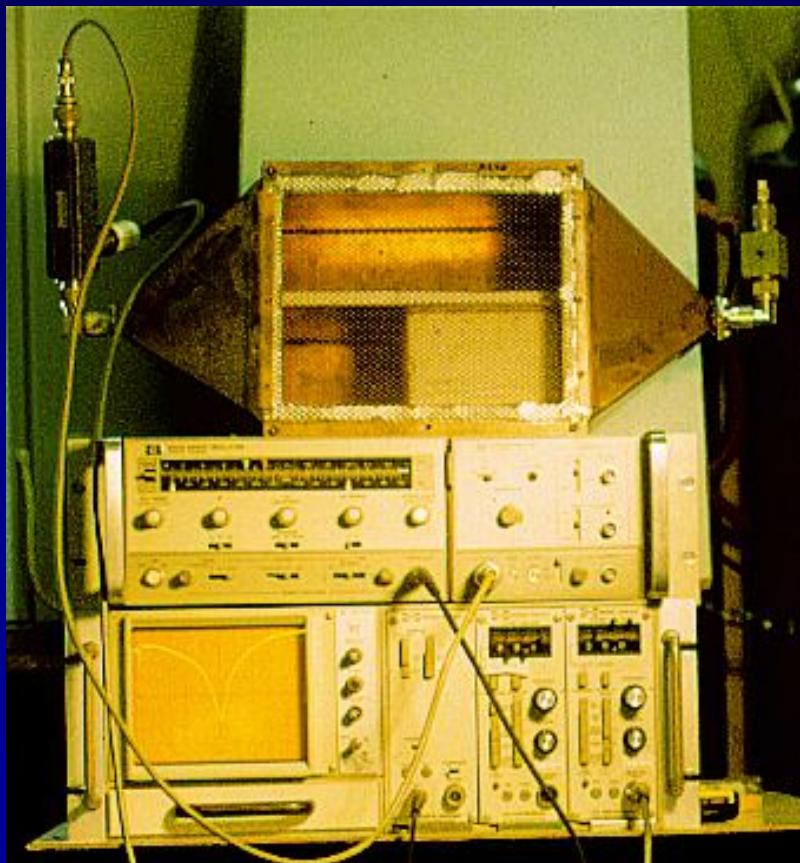
EXPOSURE DEVICE TEM (transverse EM) CELL



TEM CELL 900 MHz



800-1000 MHz EXPOSURE SYSTEM



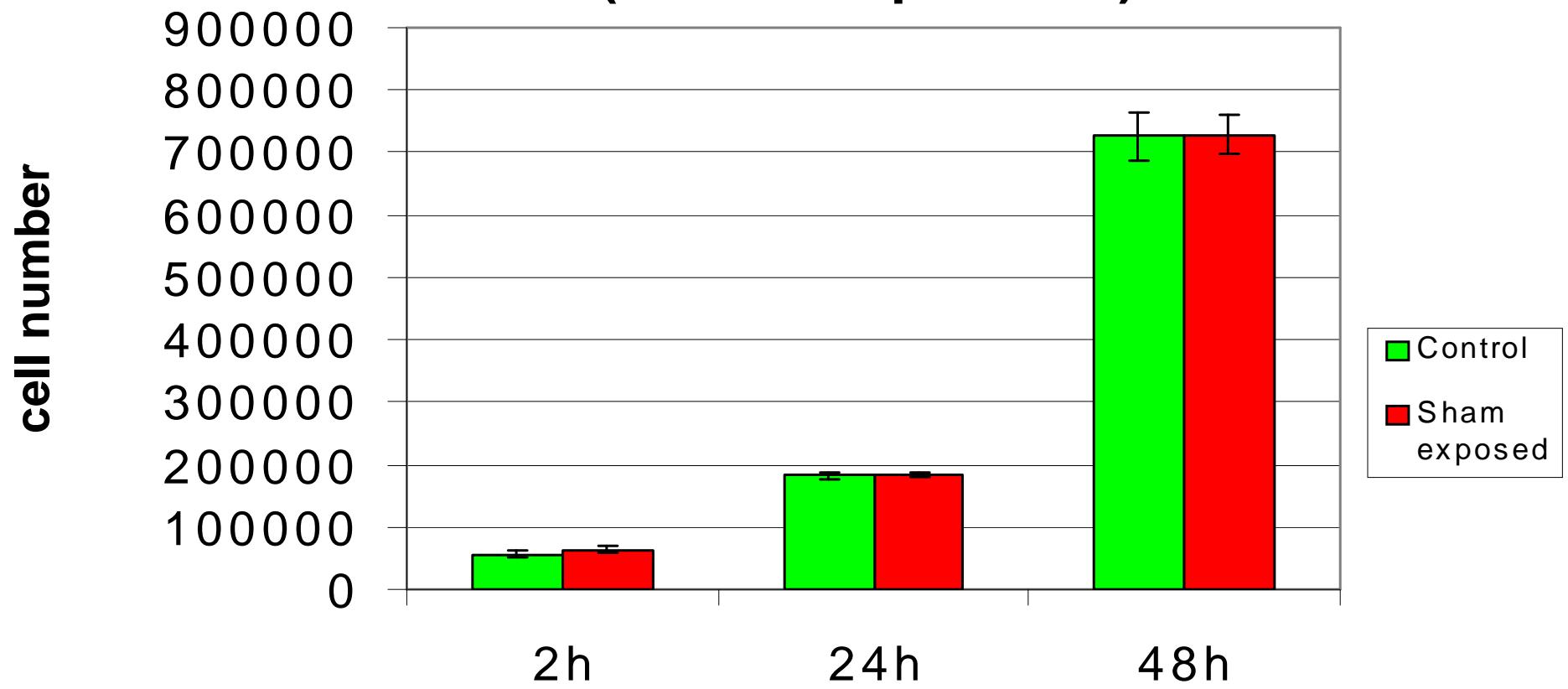
- Cella TEM
- Sweep Oscillator HP 8620 C
- Amplificatore HP 86222B RF 10 - 2400 MHz
- Accoppiatore direzionale HP 796D
- Testina bolometrica PM 10-0328
- Rivelatore PM 1038
- Universal counter HP 5316A
- Power meter HP 431 A

TEMPERATURE CONTROL

hours	T1 (CONTROL FLASK)	T2 (EXPOSED FLASK)	T3 (INCUBATOR)
2	37.01	37.01	37.00
24	37.00	37.00	37.00
48	37.00	37.00	37.00

SHAM EXPOSURE

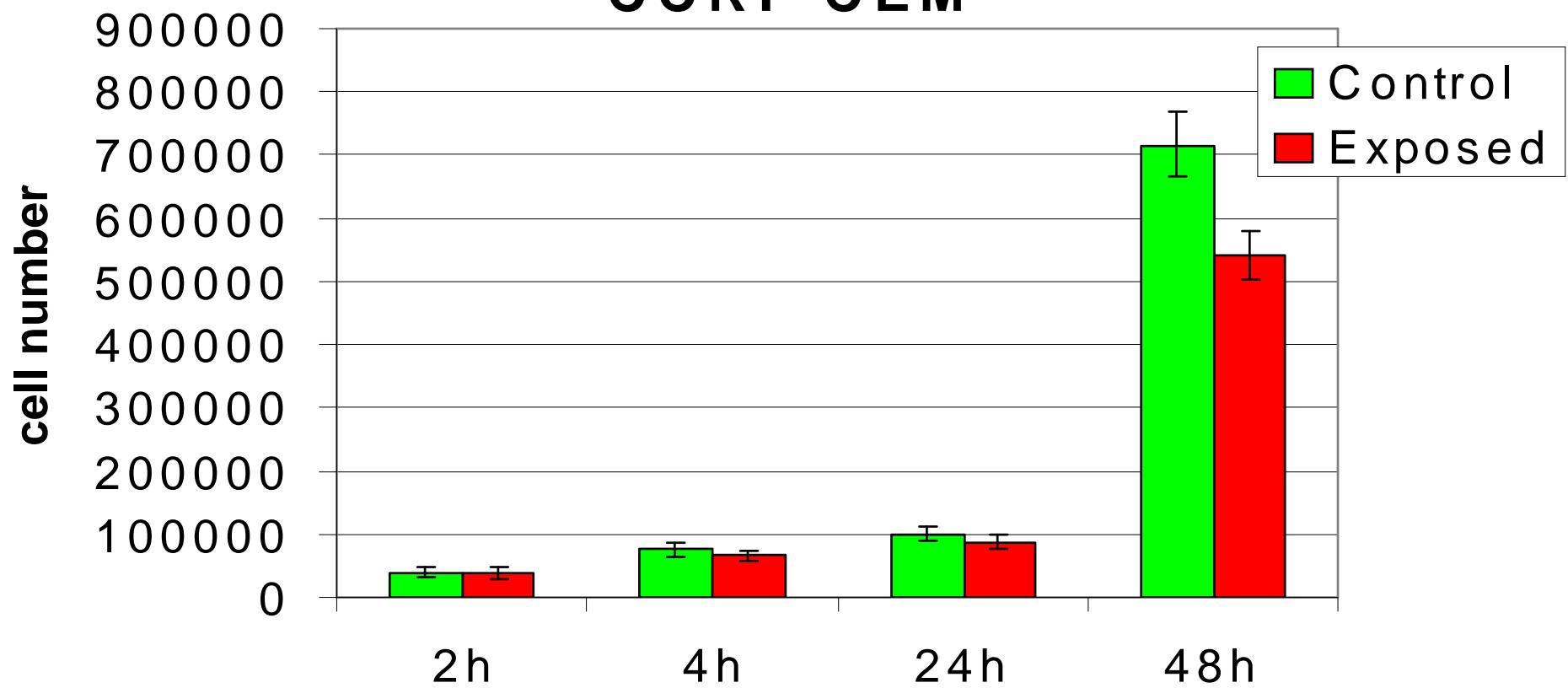
Test di proliferazione in cellule CCRF-
CEM (sham exposure)



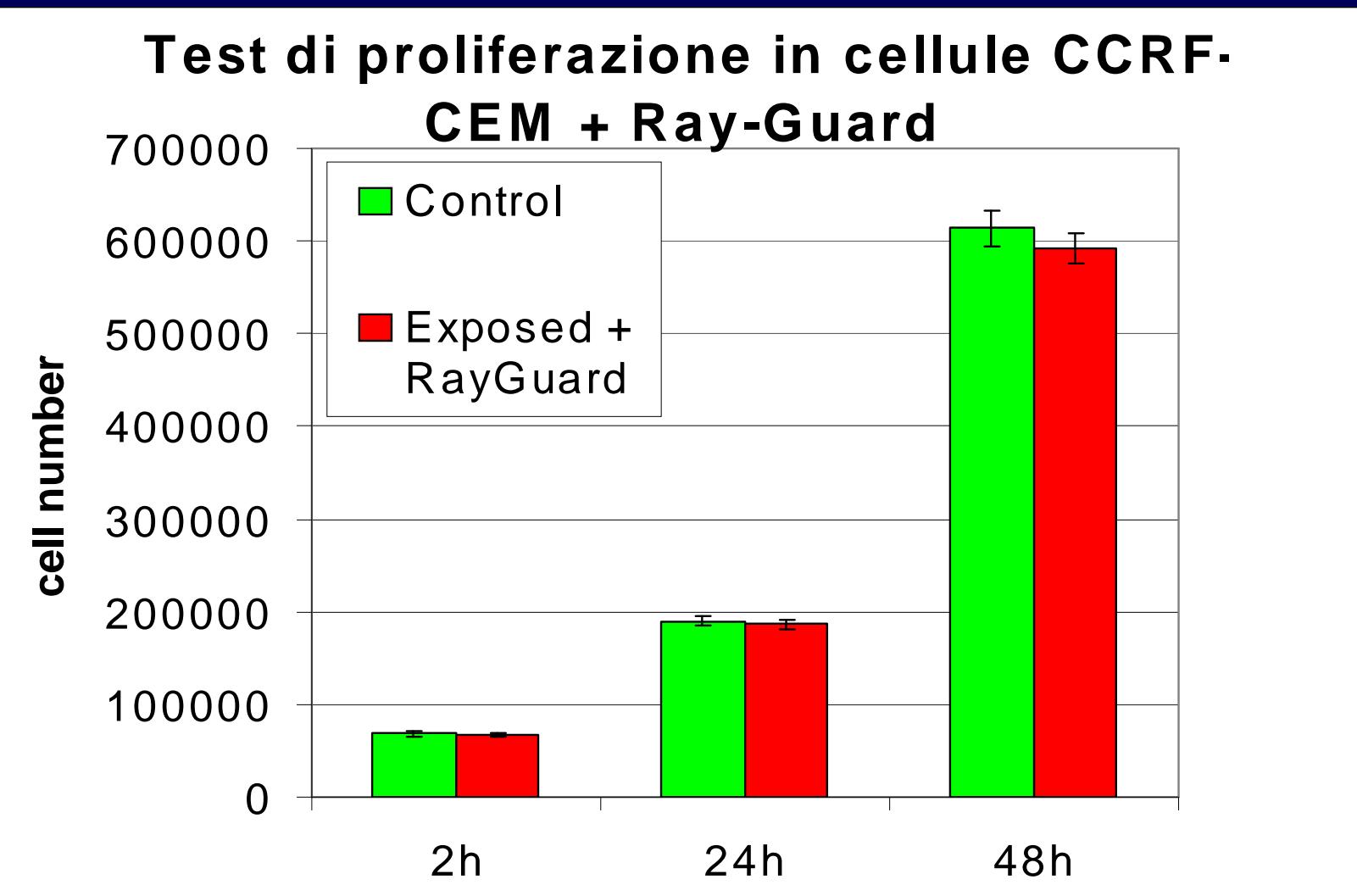
BIOLOGICAL EFFECTS

PROLIFERATION TEST

Test di proliferazione in cellule
CCRF-CEM

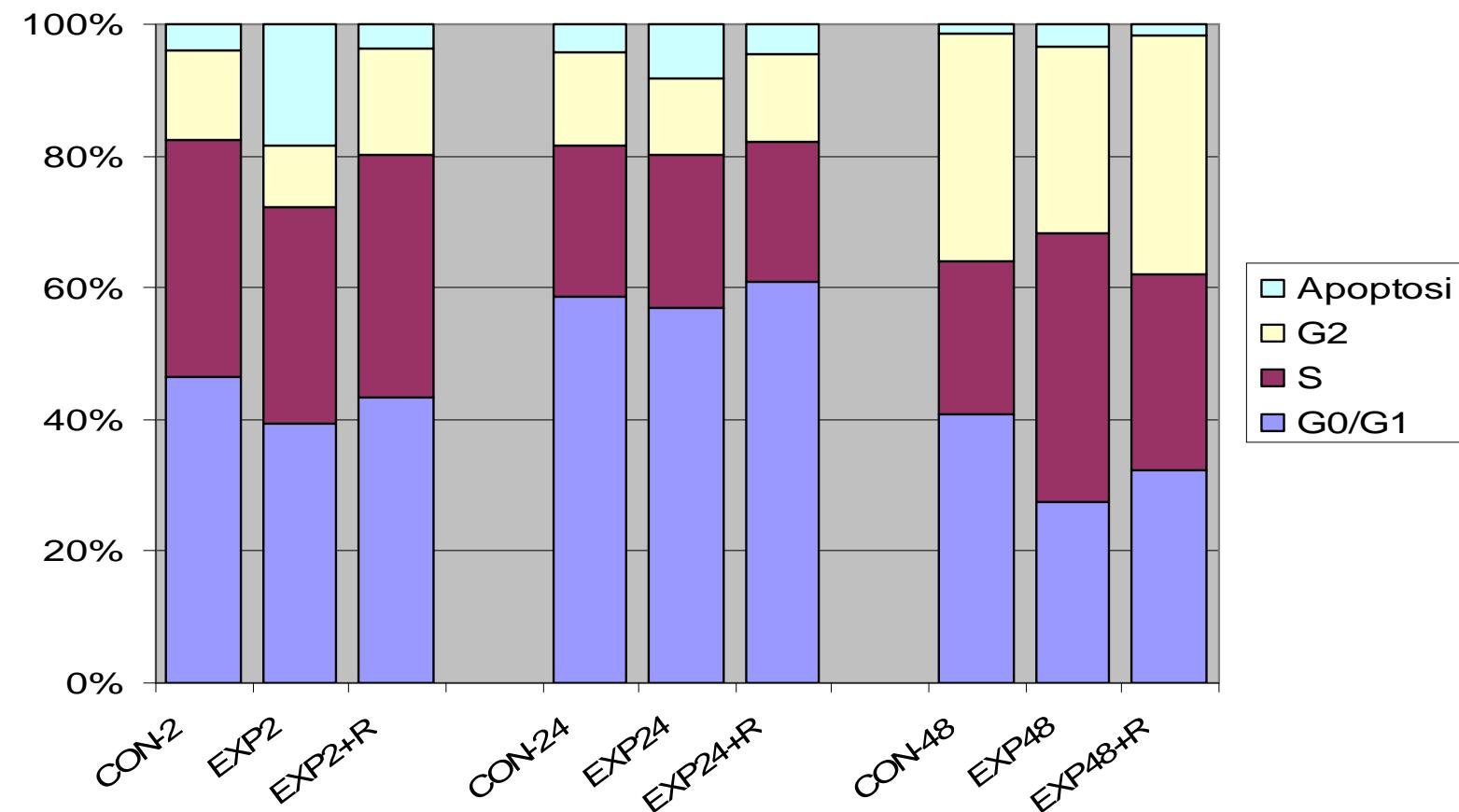


EMF and RAY-GUARD



% OF CELL IN G0/G1, S, G2/M and APOPTOSIS IN CELL CYCLE PHASES

CCRF-CEM FACS Analysis



2 HOURS 900 MHz EXPOSURE

	G0/G1 (Quiescence)	S (DNA synthesis)	G2/M (mythosis)	APOPTOSIS
CONTROL	45.75	35.55	13.42	3.89
NIR EXPOSED	38.5	32.38	9.13	18.07
NIR EXPOSED + RAY-GUARD	42.43	36.2	15.83	3.75

24 HOURS 900 MHz EXPOSURE

	G0/G1 (Quiescence)	S (DNA synthesis)	G2/M (mitosis)	APOPTOSIS
CONTROL	56.31	21.95	13.79	4.3
NIR EXPOSED	54.82	22.15	11.09	7.89
NIR EXPOSED + RAY-GUARD	58.31	20.15	12.98	4.22

48 HOURS 900 MHz EXPOSURE

	G0/G1 (Quiescence)	S (DNA synthesis)	G2/M (mitosis)	APOPTOSIS
CONTROL	40.05	22.60	33.97	1.37
NIR EXPOSED	26.68	39.83	27.62	3.38
NIR EXPOSED + RAY-GUARD	31.74	29.35	35.39	1.7

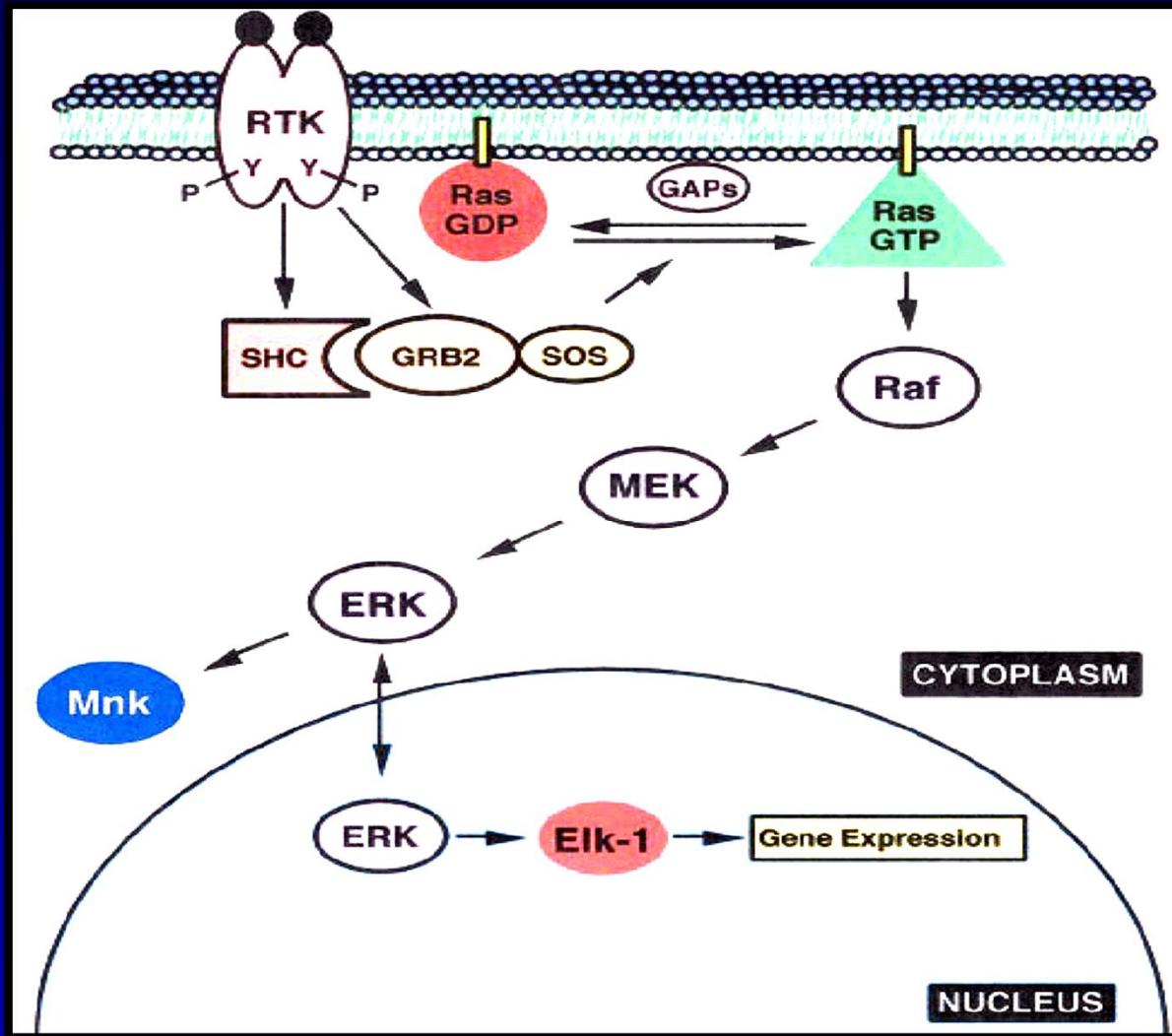
CONCLUSIONS 1

1. Short exposure time (2 and 24 hours) at 900 MHz and 4.8 V/m electromagnetic fields induce apoptotic response in CEM cells. This effect has been strongly reduced by the presence of the RAY-GUARD device inside the TEM cell during exposure.
2. Longer exposure time (48 hours) induces selection of tumorigenic clones, which show an higher level of DNA synthesis and a reduction of resting cells. The RAY-GUARD reduce the cycling cells and increase the resting quiescent cells. Moreover the apoptotic effect of NIR exposure is neutralized by the RAY-GUARD device

CONCLUSIONS 2

- 3. The biological effects induced by electromagnetic field are not related to the thermal effect.**
- 4. The electromagnetic fields can activate the cell cycle genes (i.e. RAS proto-oncogene) involved in the DNA synthesis control.**
- 5. Should their expression be controlled by RAY-GUARD?**

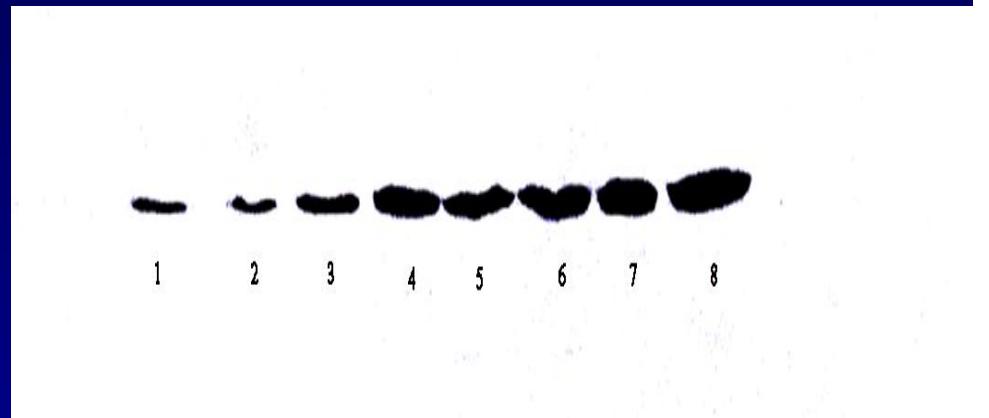
RAS SIGNALING



from Vojtek AB and Der CJ,
JBC 273 n.32:19925-19928

WB ANALYSIS H-RAS EXPRESSION

- **1,3,5,7 SAMPLES= CONTROL CELLS**
2,4,24,48 hs



- **2,4,6,8 SAMPLES= EXPOSED CELLS**
2,4,24,48 hs

Power Density	Reported Biological Effects	References
0.168 - 1.053 $\mu\text{W}/\text{cm}^2$	Irreversible infertility in mice after 5 generations of exposure to RFR from "antenna park"	Magras & Xenos, 1997
0.16 $\mu\text{W}/\text{cm}^2$	Motor function, memory and attention of school children affected (Latvia)	Kolodynki, 1996
0.2 - 8 $\mu\text{W}/\text{cm}^2$	Two-fold increase in childhood leukemia / RFR exposure to AM/FM towers	Hocking, 1996
1.0 $\mu\text{W}/\text{cm}^2$	Whole body microwave irradiation of male mice caused a significant effect on the immune system	Fesenko, 1999
1.0 $\mu\text{W}/\text{cm}^2$	Irradiation (5 hours) with low-power microwaves stimulates the immune potential of macrophages and T cells	Novoselova, 1999
1.3 - 5.7 $\mu\text{W}/\text{cm}^2$	Two-fold increase in leukemia in adults from AM RF exposure	Dolk, 1997
~2-4 $\mu\text{W}/\text{cm}^2$	Direct effect of RFR on ion channels in cells/opening of acetylcholine channels	D'Inzeo, 1988
4-10 $\mu\text{W}/\text{cm}^2$	Visual reaction time in children is slowed//lower memory function in tests	Chiang, 1989
5 - 10 $\mu\text{W}/\text{cm}^2$	Impaired nervous system activity	Dumansky, 1974
10 $\mu\text{W}/\text{cm}^2$ (0.0027 W/Kg SAR)	Changes in active avoidance conditioned reflex (behavioral change) after 0.5 hour exposure	Navakatikian, 1994
10-20 $\mu\text{W}/\text{cm}^2$	Increase in micronuclei (aberrant DNA form) found in workers chronically exposed to microwaves at 1250-1350 MHz.	Garaj-Vrhovac, 1995
10 - 25 $\mu\text{W}/\text{cm}^2$	Changes in the hippocampus of the brain	Belokrinitsky, 1982
30 $\mu\text{W}/\text{cm}^2$ (0.015 W/Kg SAR)	Immune system effects - elevation of PFC count (antibody-producing cells)	Veyret, 1991
50 $\mu\text{W}/\text{cm}^2$	An 18% reduction in REM sleep (important to memory and learning functions)	Mann, 1996
100 $\mu\text{W}/\text{cm}^2$	Changes in immune system function	Elekes, 1996
100 $\mu\text{W}/\text{cm}^2$ (0.027 W/Kg SAR)	A 24% drop in testosterone after 6 hours exposure	Navakatikian, 1994

THE RESEARCH GROUP

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REMARKS

- It is hardly possible to transfer these results automatically to humans.
- We do not know if the Ray-Guard can protect efficiently the organisms.
 - We suppose it! As we obtained preliminary results on human lymphocytes

New experimental aims on normal lymphocytes

- 1.Exposure of healthy donor lymphocytes to RF/MW 900 MHz in presence of Ray-Guard device.
- 2.Analysis of the electromagnetic field effect on cell cycle in presence of Ray-Guard device.

New proposal research

- Ray-Guard effect on modulation of cell cycle genes expression in CCRF-CEM cells exposed to NIR, such as oncogenes (i.e.Ras) and onco-suppressor genes.