



A first prospective randomized controlled trial to decrease bacterial load on chronic wounds in patients using a cold atmospheric plasma device

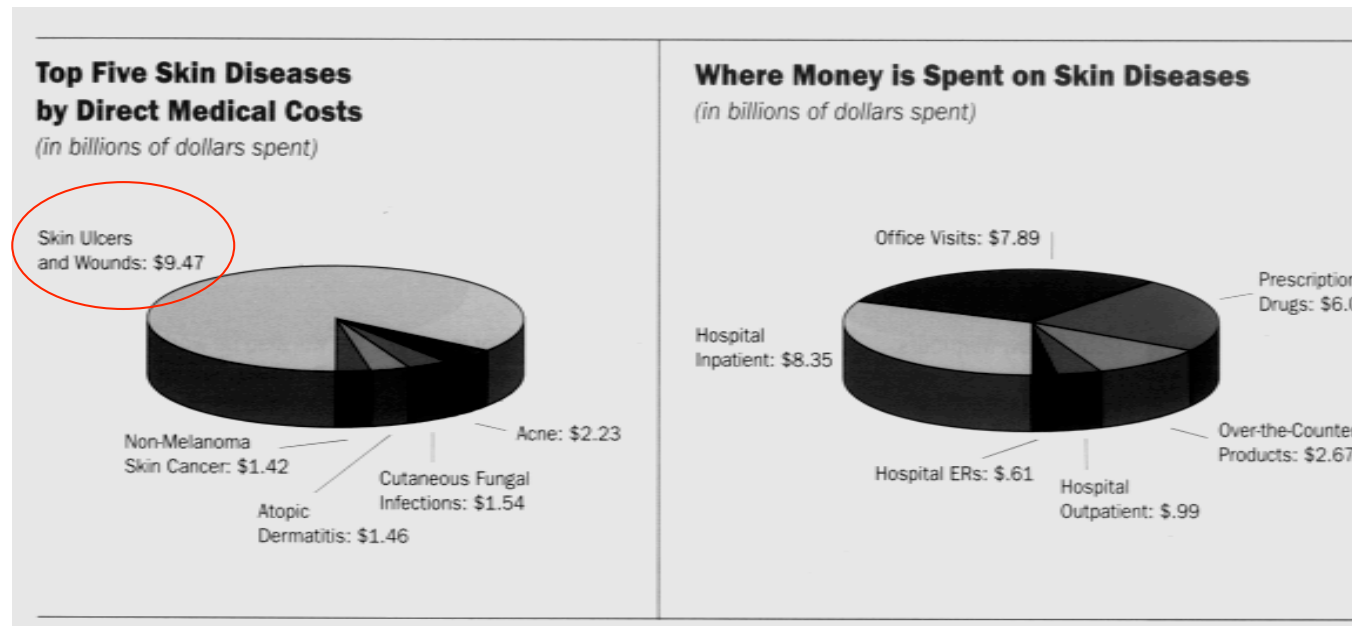
Dr. Georg Isbary

Plasma Project – From medical point of view

- **Importance of the plasma project**
- In vitro proof of principle experiments
- Phase II study – results

Chronic wounds are a major burden for the health system

- Prevalence ~ 1-2 % in German Population (> 1.000.000 patients)
- High costs for the community 1-2 % of annual health care budget*
- Venous ulcers require an average of 24 weeks to heal, 15% never heal, recurrence is found once or multiple times in 15-71% of cases** ***



American Academy of Dermatology Report 2005

*Etufugh CN, Phillips TJ. Venous ulcers. *Clin Dermatol* 2007; **25**: 121-30.

**Kurz et al. VEINES Task Force Report, *Int Angiol.* 1999;18(2):83-102.

***Heit et al. Venous thromboembolism epidemiology *Semin Thromb Hemost.* 2002;28(suppl 2):3-13

Big Issue – Increasing rate of resistance/ multiresistance

- „Bacteria can become resistant to antibiotics“ warned Alexander Fleming, when he landed the Nobel prize in Medicine in 1945.
- European Antimicrobial Resistance Surveillance System (EARSS) 2007: Resistance is becoming a larger problem year after year (especially for *Streptococcus pneumoniae*, *Staphylococcus aureus*, *Enterococcus faecalis*, *Enterococcus faecium*, *Escherichia Coli*, *Klebsiella pneumoniae* and *Pseudomonas aeruginosa*)
- Global Health Care Associations consider multiresistant germs like MRSA as a global threat*
- 19,5 % of all Staph. aureus detected in German hospitals are MRSA, in France 24.5% (EARSS 2008)
- Worrying is the raising resistance against so called reserve drugs within the last 6 years – e.g. Vancomycin (EARSS 2007)
- „Gold standard“ Vancomycin with failure rates of 23-52%!
- November 2008 launch of DART (Deutschen Antibiotika-Resistenzstrategie)

*Grundmann H, Aires-de-Sousa M, Boyce J et al. Emergence and resurgence of methicillin-resistant *Staphylococcus aureus* as a public-health threat. *Lancet* 2006; **368**: 874-85.

Big Issue – Increasing rate of resistance/ multiresistance

- 1999-2005 MRSA-related hospitalizations more than doubled (119%, respectively ~14% per year)*
- Infections with MRSA kill ~19000 hospitalized patients in the U.S. annually (similar to the number of deaths caused by AIDS, tuberculosis and viral hepatitis combined!)**
- In 2006, on any given day >23k pts were hospitalized in the US with MRSA infection***
- 40.000 deaths in 2006 due to infections in Germany (14% Increase 2002-2006)****
- Antimicrobial drug-resistant infections do increase death, illness, and direct costs by 30-100%*****

*Klein E, Smith DL, Laxminarayan R. Hospitalizations and deaths caused by methicillin-resistant *Staphylococcus aureus*, United States, 1999-2005. *Emerg Infect Dis* 2007; **13**: 1840-6

Klevens RM, Morrison MA, Nadle J et al. Invasive methicillin-resistant *Staphylococcus aureus* infections in the United States. *Jama* 2007; **298: 1763-71

***Jarvis, CID Jan 15, 2010

**** Report Deutsche Antibiotika-Resistenzstrategie

*****Cosgrove SE, Carmeli Y. The impact of antimicrobial resistance on health and economic outcomes. *Clin Infect Dis* 2003; **36**: 1433-7.

Side effects of antibiotics

- ~10% of hospitalized patients present an allergy against penicillin (but only 10% of those actually have allergic reactions during treatment)*
- Problematic is the cross-reactivity, which averts the use of many other antibiotics, e.g. cephalosporins*

- Antibiotic associated diarrhea occurs in about 5-30% during therapy or even two month after ending the treatment**, ***

*Greenberger PA. Drug allergy. Part B: Allergic reactions to individual drugs: low molecular weight. *Patterson's Allergic Diseases* 2002: 335-59

McFarland LV. Epidemiology, risk factors and treatments for antibiotic-associated diarrhea. *Dig Dis* 1998; **16: 292-307

***Wistrom J, Norrby SR, Myhre EB et al. Frequency of antibiotic-associated diarrhoea in 2462 antibiotic-treated hospitalized patients: a prospective study. *J Antimicrob Chemother* 2001; **47**: 43-50

New antibiotic drugs

- „Effective antibiotic treatment becomes as precious as clean drinking water“
- Genomic derived or target based antibiotics need a lot of time to brought to the market:
 - for gram + strains ~ 2012*
 - for gram – strains ~ 2016 - 2021*

*Payne DJ, Gwynn MN, Holmes DJ et al. Drugs for bad bugs: confronting the challenges of antibacterial discovery. *Nat Rev Drug Discov* 2007; **6**: 29-40

The New York Times

ON THE WEB

Deadly Germs Largely Ignored By Drug Firms

By ANDREW POLLACK
Published: February 26, 2010

Gram-negative bacteria are practically built to withstand drugs, which is one reason few drug makers have rushed to pursue treatments.

Related

[Rising Threat of Infections Unfazed by Antibiotics](#)
(February 27, 2010)

The bacteria have a double cell membrane to shield them, compared with Gram-positive organisms, which have a single membrane. They can make various enzymes that break down antibiotics. And some,

particularly *Pseudomonas aeruginosa*, have powerful pumps that can expel the drugs.

The bacteria also readily exchange genes, even across different species, that confer drug resistance.

It is likely to be several years before new drugs to treat Gram-negative infections are available. A report last September by European health authorities found only six novel drugs in clinical trials that might work against at least one Gram-negative organism, compared with 13 for Gram-positive bacteria.

A separate study released about a year ago by the Infectious Diseases Society of America found no drugs in middle- or late-stage clinical trials directed specifically at Gram-negative organisms. There were eight drugs in those trials that developers hoped might work against both Gram-negative and Gram-positive microbes.

The difficulty of killing Gram-negative germs is not the only reason for the dearth of new

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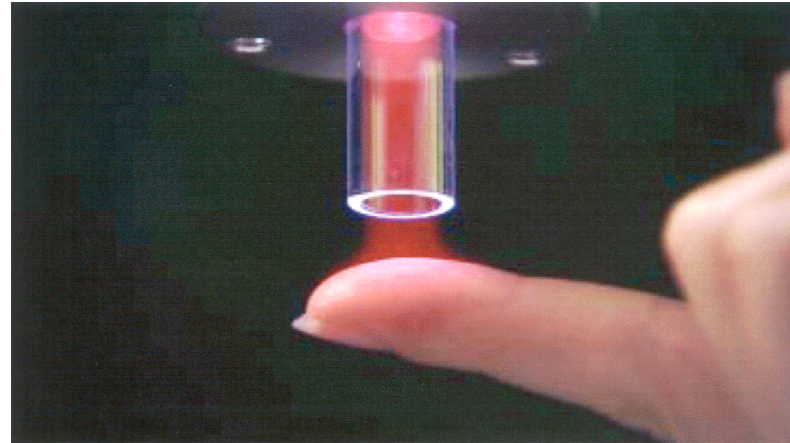


New antibiotic drugs

- „Effective antibiotic treatment becomes as precious as clean drinking water“
- Genomic derived or target based antibiotics need a lot of time to brought to the market:
 - for gram + strains ~ 2012*
 - for gram – strains ~ 2016 - 2021*
- New antibiotic drugs face same problems like usual ones (resistance, allergic reactions and other side effects)

*Payne DJ, Gwynn MN, Holmes DJ et al. Drugs for bad bugs: confronting the challenges of antibacterial discovery. *Nat Rev Drug Discov* 2007; **6**: 29-40

Cold atmospheric plasmas are ideal antibiotics



reactive species (O_3 , NO, NO_2 , N, O, OH,.....)

charged particles (electrons, positive/negative ions)

light (UV, visible and IR)

electric field

heat

Benefits of our indirect low temperature Argon plasma

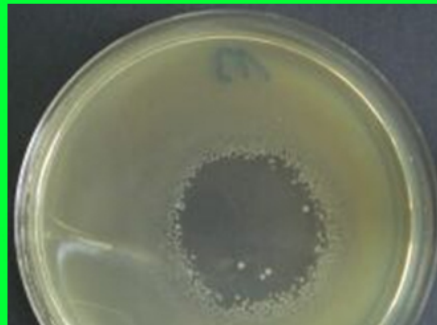
Low temperature argon plasma:

- Allows in-vivo application, without damaging tissue
- Medical cocktail – can be tuned for different purposes
- Contact free application, reaches “rough” surfaces down to micrometer scale
- Bactericidal (fungicidal and virucidal)
- Physical-therapy → Resistance and allergic reactions are less feasible
- Enhanced wound healing

Plasma Project – From medical point of view

- Importance of the plasma project
- **In vitro proof of principle experiments**
- Phase II study – results

Efficiency of 2 min plasma treatment against different germs relevant to wound healing



Escherichia coli

present on
healthy persons



Enterococcus faecalis

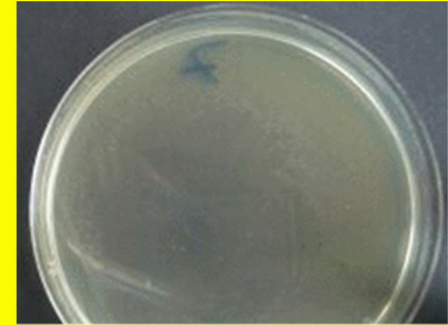


Group A streptococcus

facultative pathogenic, occasional resistance



*methicillin-resistant
Staphylococcus aureus*



*vancomycin-resistant
Enterococcus faecium*

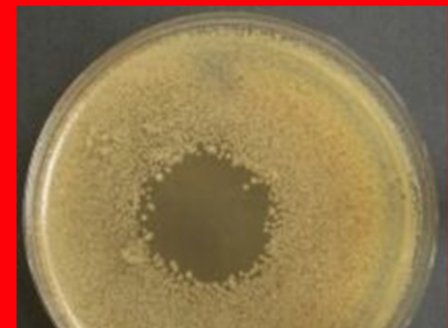
facultative pathogenic, seldom present on healthy skin



*Pseudomonas
aeruginosa*



Burkholderia cepacia



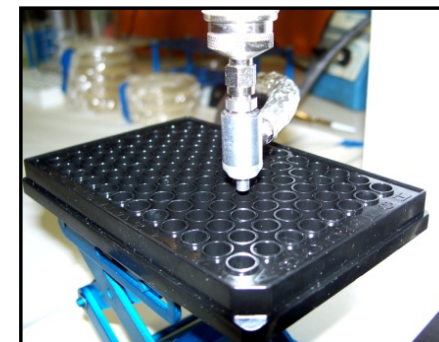
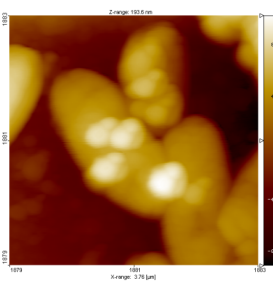
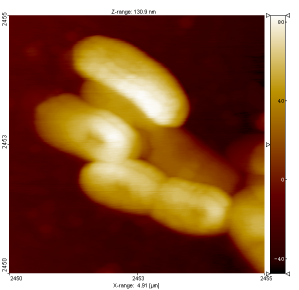
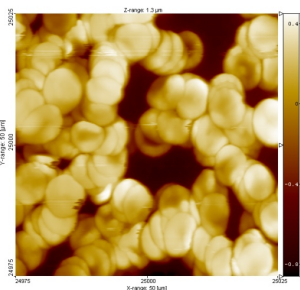
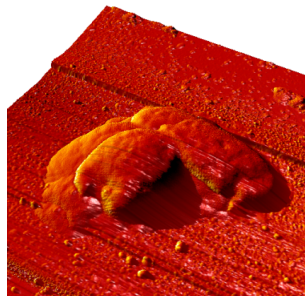
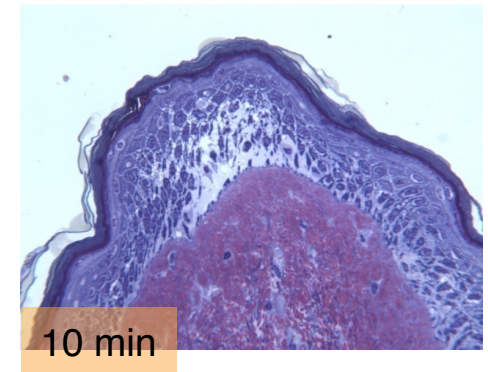
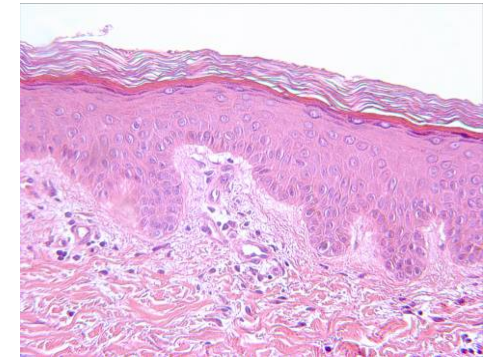
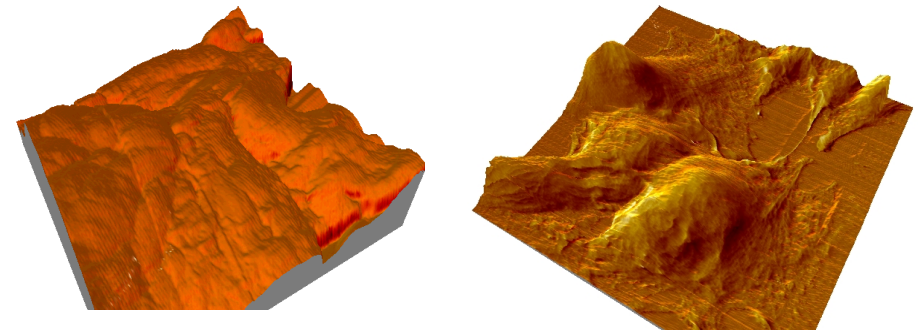
Bacillus cereus

Phase I study

Numerous tests to find dosages and to check harmlessness of the plasma treatment:

e.g. histologies, bloodtests, microscopic images, AFM, cell essays...

Further investigations with fibroblasts, keratinocytes, cell cultures, essays to check toxicity, mutagenicity, and antibodies



Plasma Project – From medical point of view

- Importance of the plasma project
- In vitro proof of principle experiments
- **Phase II study – results**

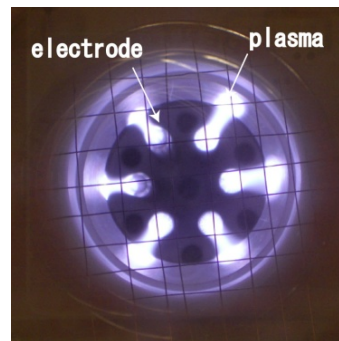
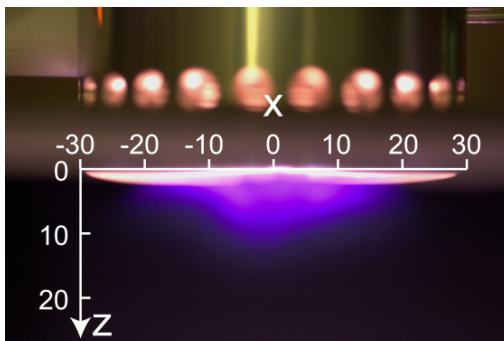
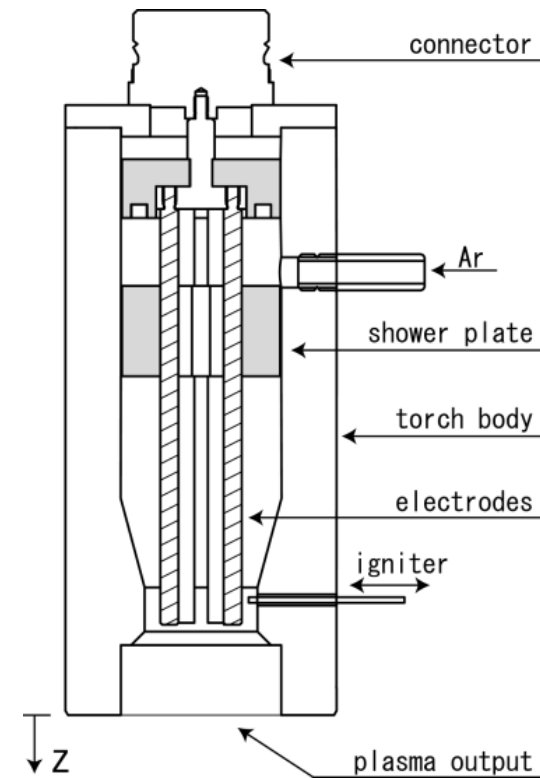
> Klinikum Schwabing

Phase II study: MicroPlaSter (ADTEC Plasma Technology Co. Ltd., Hiroshima/London)

MaryMcGovern@adtec.eu.com



The new device - MicroPlaSter β



- Used gas: argon
- Voltage = 50 - 100 V
- Frequency = 2,3 GHz
- Power = 100 W

⇒ Plasma is generated by microwave-technology

Shimizu et al. 2008

Chronic wounds in dermatology



Venous diseases



Arterial diseases



Infections



Diabetes mellitus



Carcinoma



Pyoderma gangraenosum

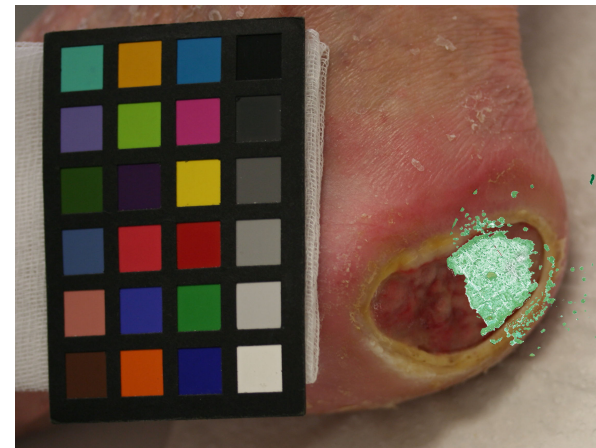
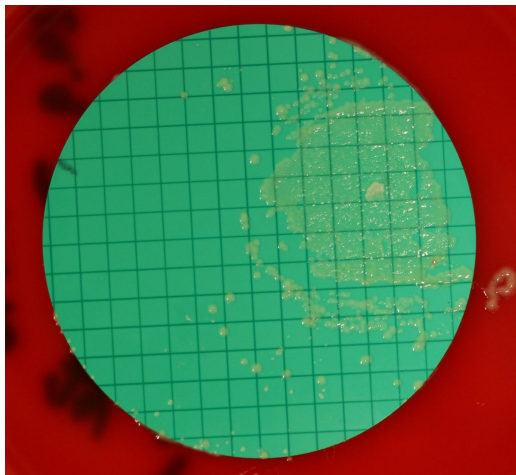
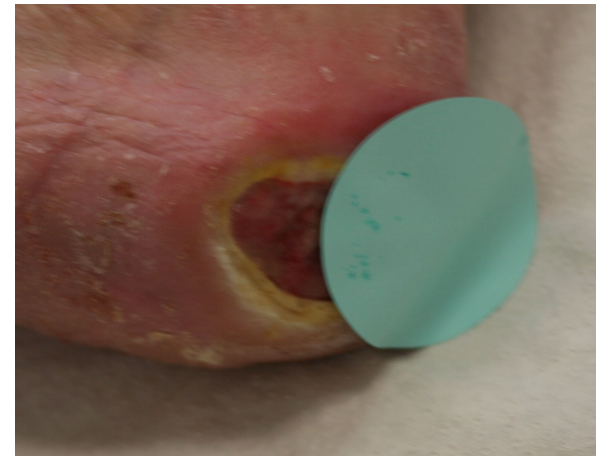
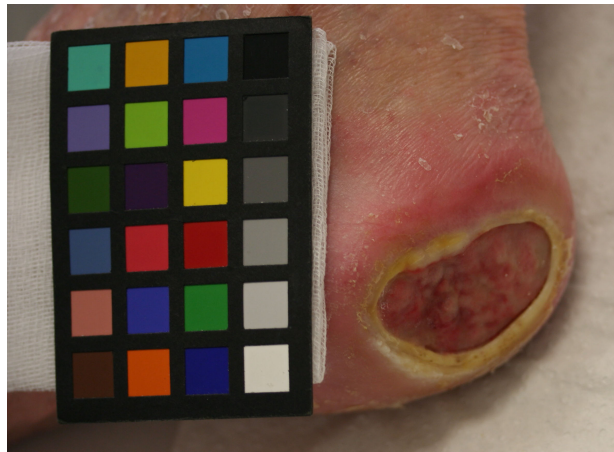
Manual necrolysis or treatment with a high pressure water jet Debritom® (medaxis, Switzerland) to homogenize wound surface



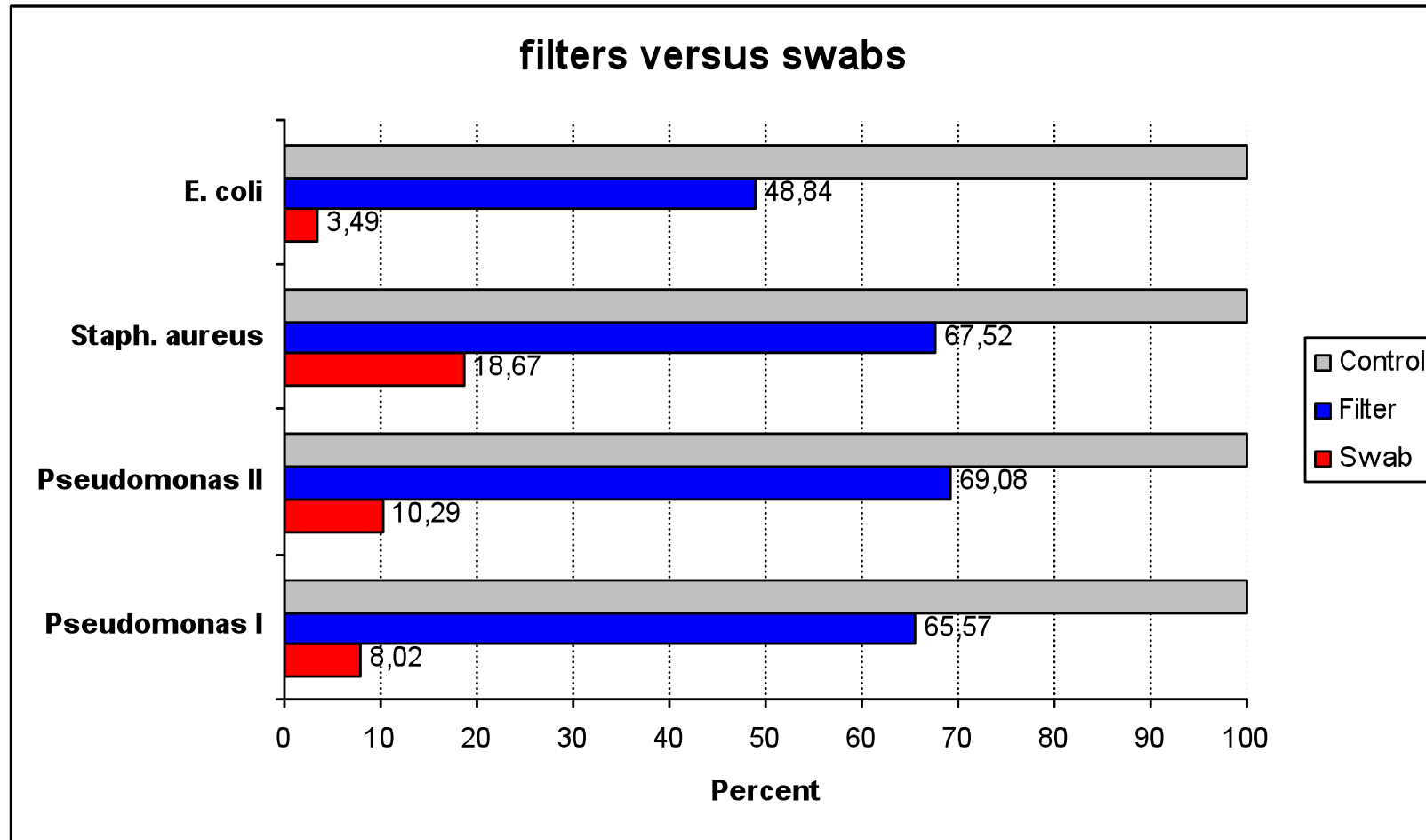
Common swab techniques failed in accuracy and reproducibility of bacterial loads



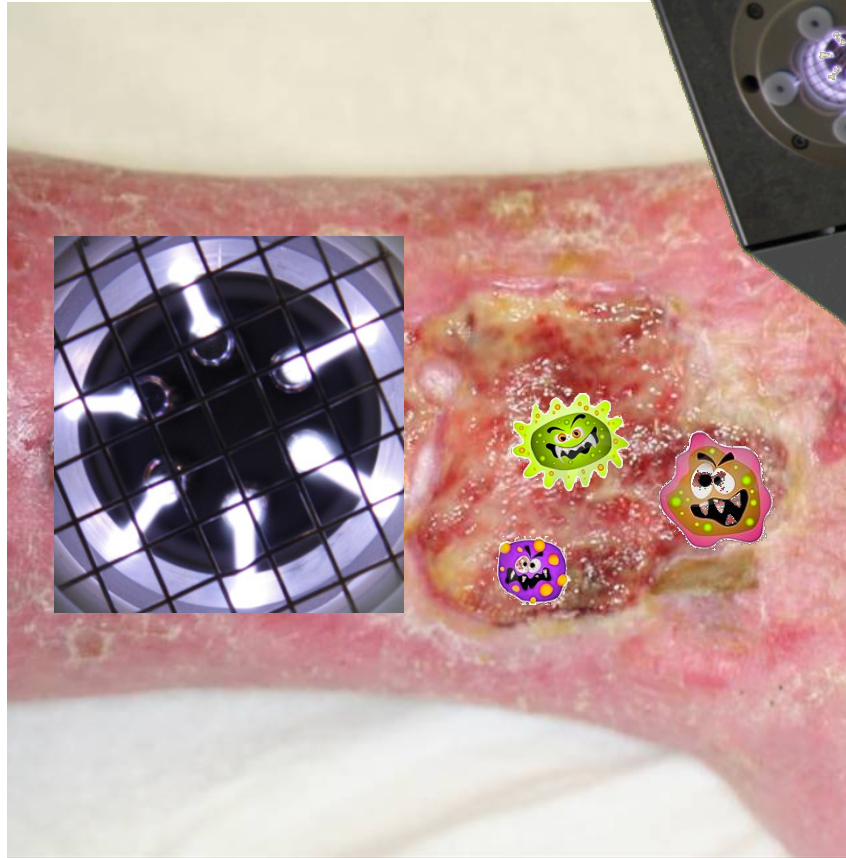
Nitrocellulose filters revealed a higher accuracy and reproducibility



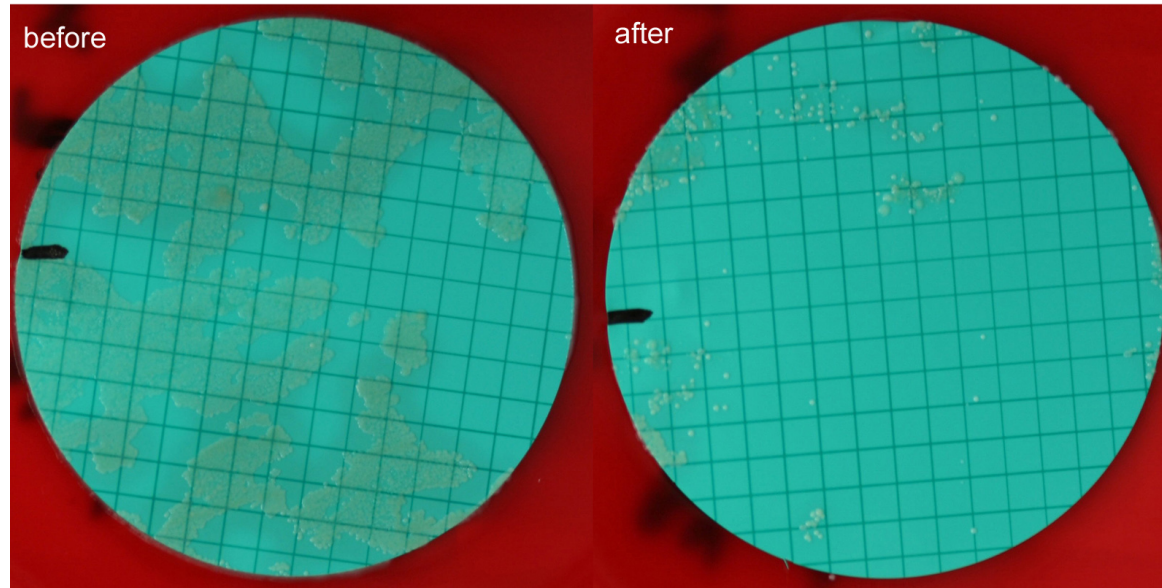
Evaluation of accuracy and reproducibility of swabs vs. nitrocellulose filters



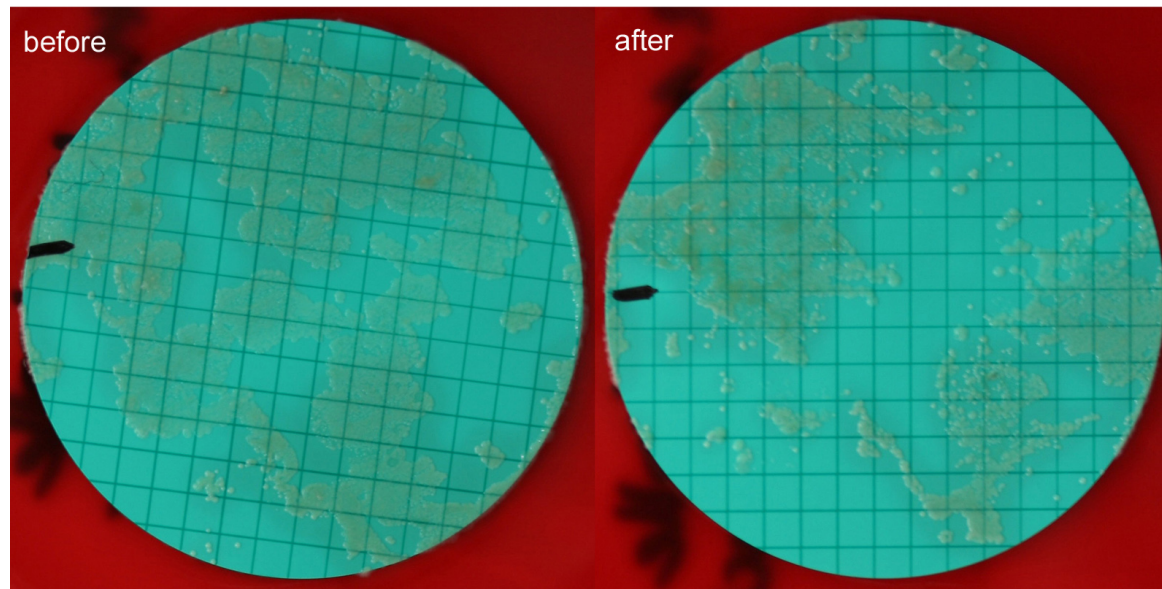
2 min



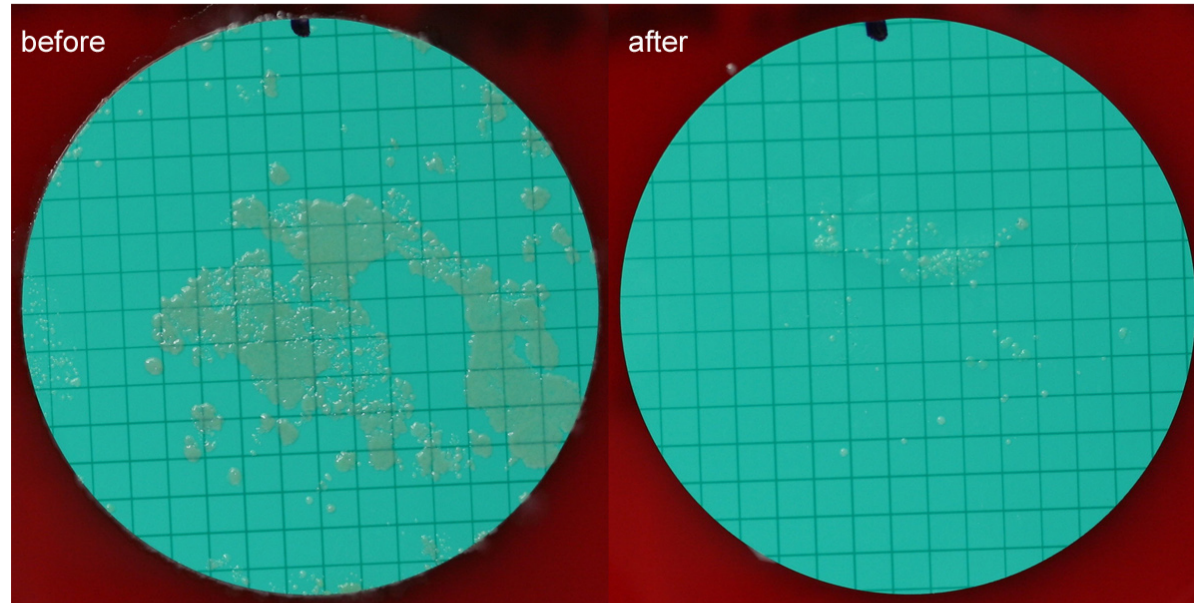
MRSA before and after plasma treatment



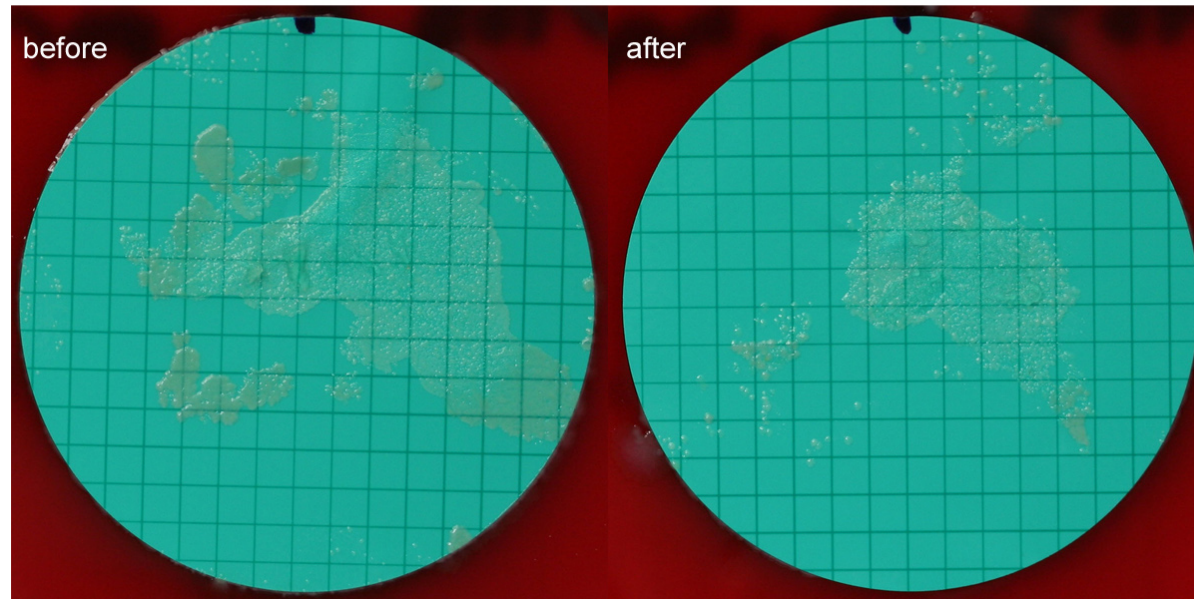
MRSA before and after control



PSAE changes before and after plasma treatment



PSAE changes before and after control



Phase II study up to now – MicroPlaSter alpha

- 1600 treatments (1 to 169, in average 9,1 per patient)
- 166 patients

- diagnosis: mostly infected ulcers of the lower leg

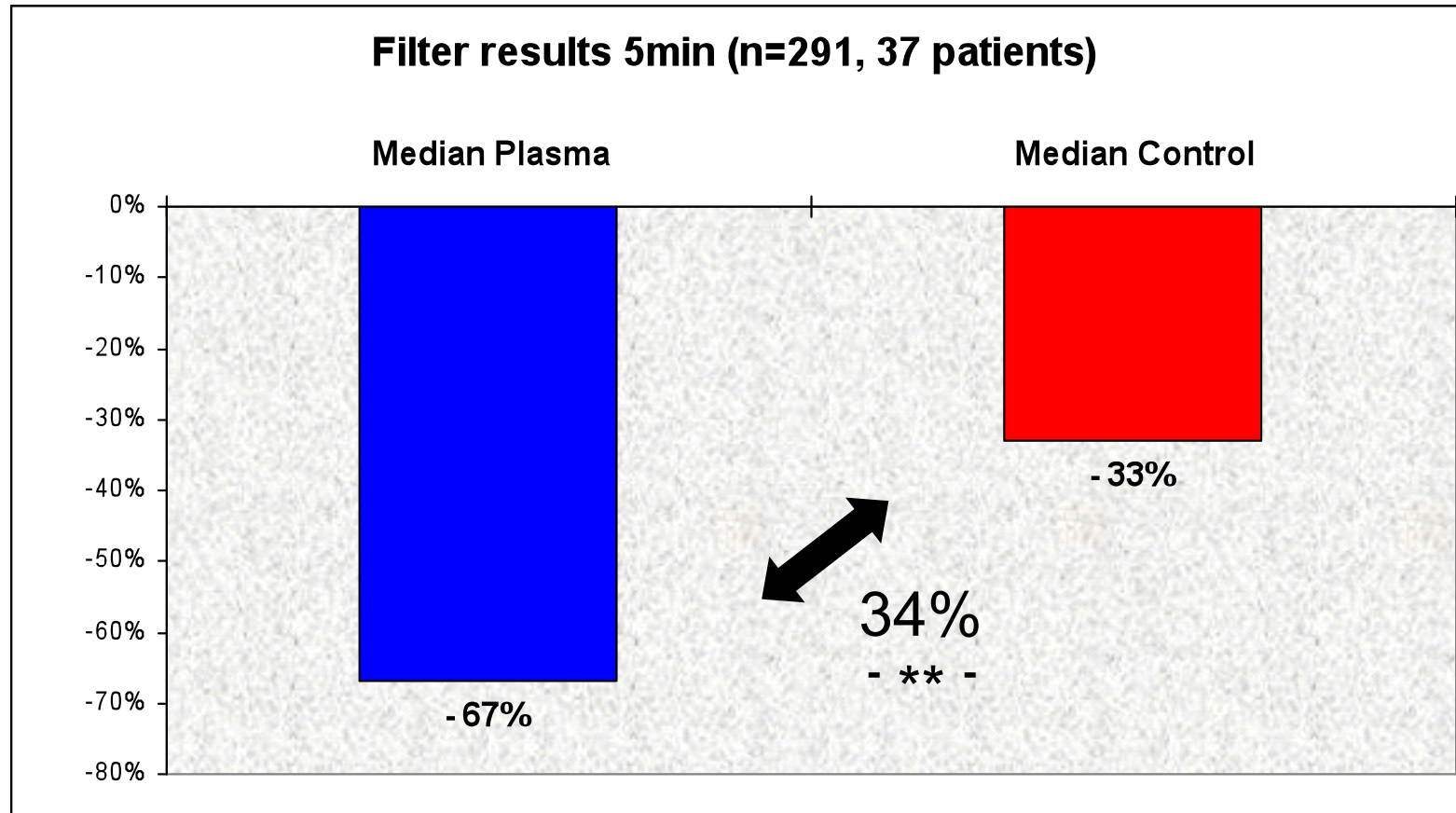
Interim analysis (efficacy of plasma treatment)

- 36 patients
- 291 treatments
- 5 min treatment time

- Primary aetiology of wounds: venous ulcers (47%)

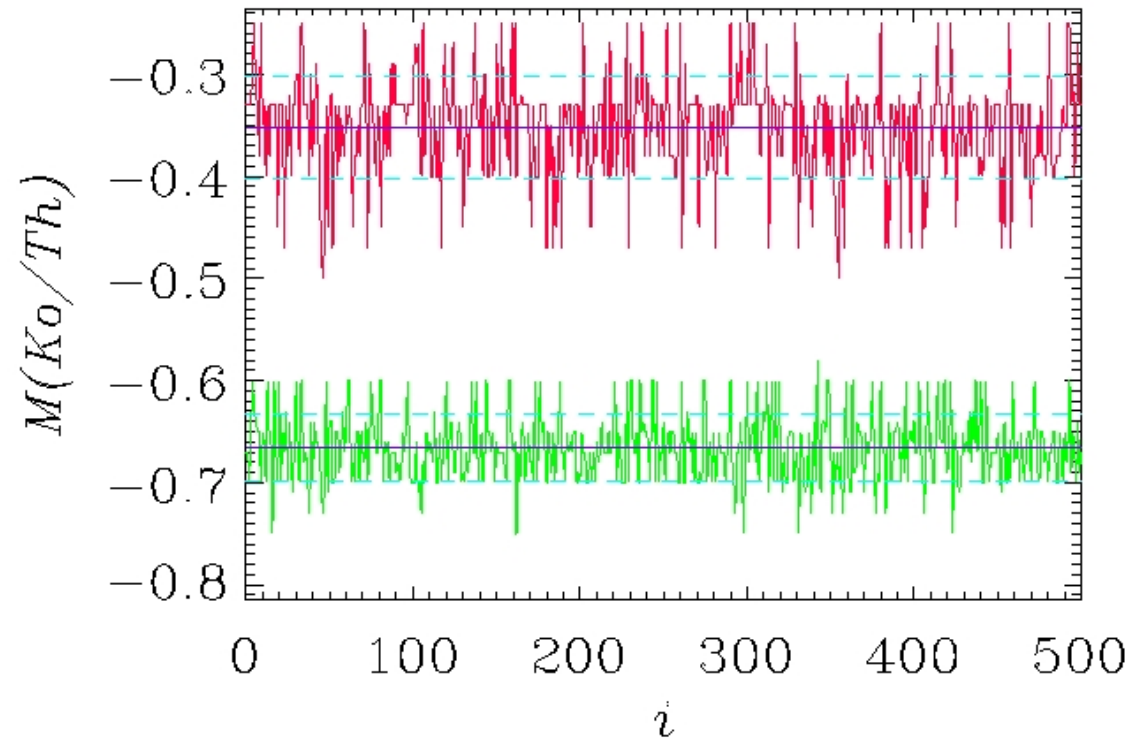
- Filter taken before and after treatment

Results: 5 min treatment time




Highly significant ($p < 10^{-6}$) higher germ reduction (34%) in plasma treated area


Summary of Phase II - Results 5min of treatment time




Results from the corresponding bootstrap-test



British Journal of Dermatology





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A first prospective randomized controlled trial to decrease bacterial load using cold atmospheric argon plasma on chronic wounds in patients

G. Isbary^a, G. Morfill^b, H-U. Schmidt^c, M Georgi^a, K Ramrath^a, J. Heinlin^d, S. Karrer^d, M. Landthaler^d, T. Shimizu^b, B. Steffes^b, W. Bunk^b, R. Monetti^b, J. L. Zimmermann^b, R. Pompl^b, W. Stolz^a

^a Department of Dermatology, Allergology and Environmental Medicine, Hospital Munich Schwabing, Germany
^b Max Planck Institute for Extraterrestrial Physics, Garching, Germany
^c Department of Microbiology, Hospital Munich Schwabing, Germany
^d Department of Dermatology, University of Regensburg, Germany

Correspondence to Georg Isbary, MD
Department of Dermatology, Allergology and Environmental Medicine
Hospital Munich Schwabing
Kölner Platz 1
D-80804 Munich, Germany
Phone: +49(0)89-3068-2904
Fax: +49(0)89-3068-7522
E-mail: oki23@gmx.de

KEYWORDS
plasma medicine • cold atmospheric plasma • argon plasma • infection • chronic wounds • MRSA

ABSTRACT
Background: Bacterial colonization of chronic wounds slows healing. Cold atmospheric plasma has been shown in vitro to kill a wide range of pathogenic bacteria.
Objectives: The safety and efficiency of cold atmospheric argon plasma to decrease bacterial load as a new medical treatment for chronic wounds.

PMID: 20222930

Interim analysis (efficacy of plasma treatment)

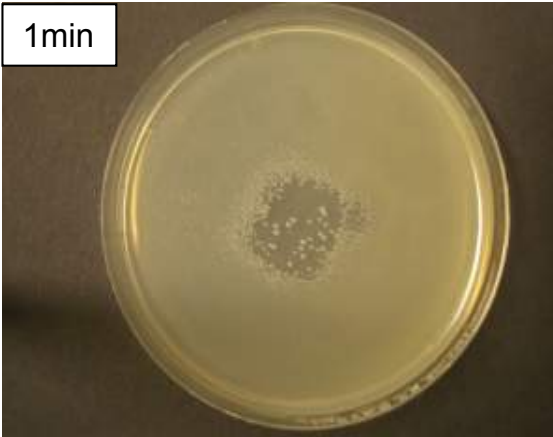
- 14 patients
- 70 treatments
- 2 min treatment time

- Filter taken before and after treatment

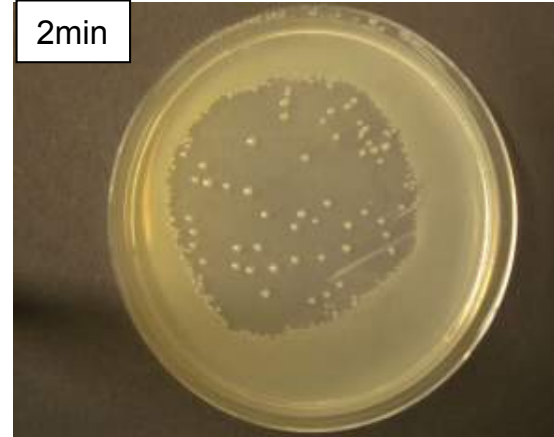
UV effect on bacteria (E. coli)

without quartz glass

1min

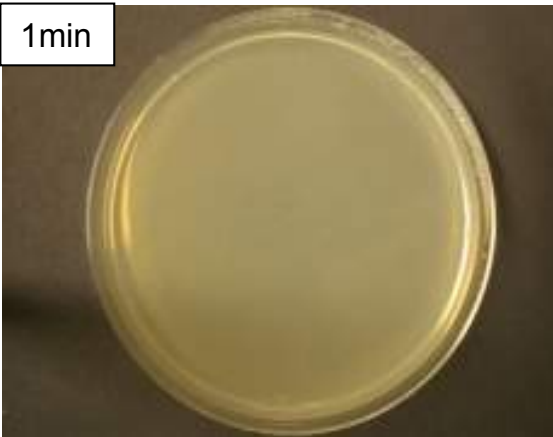


2min

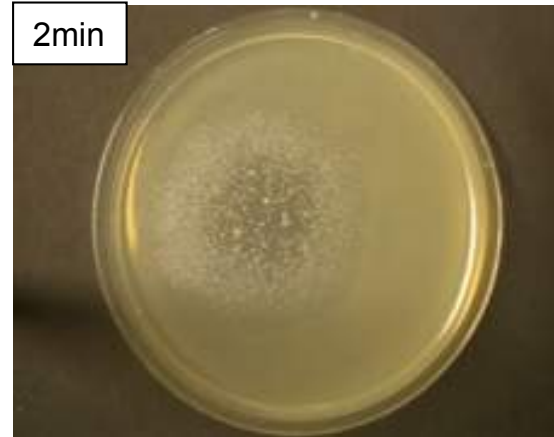


with quartz glass

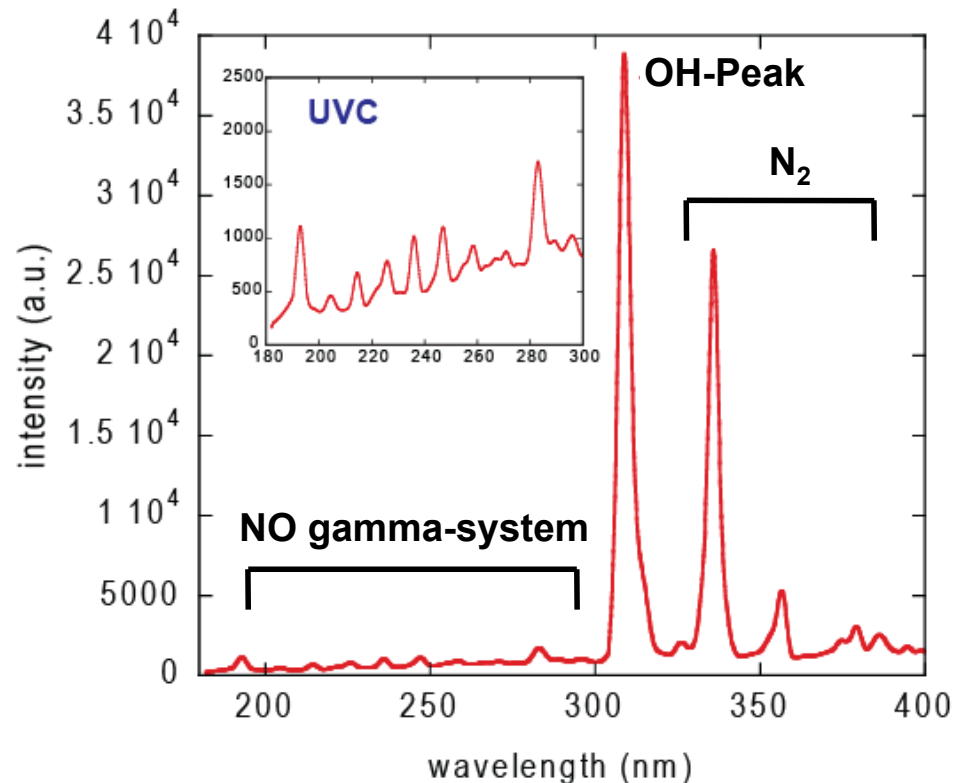
1min



2min



UV-measurements of MicroPlaSter



UV-spectrum of the argon plasma. Control of reactive species and UV radiation:
UVA : N₂ molecules
UVB: OH
UVC: NO

Measurements of reactive species in air revealed:
NO: < 1 ppm
NO₂: ~ 6 ppm at max
O₃: ~ 900 ppb

- The total integrated erythemal-weighted irradiance is:
$$\sum P_{\text{eff}}(\lambda) \times \Delta\lambda = 9.3 \mu\text{W}/\text{cm}^2 = 0.09 \text{ W}/\text{m}^2$$
- Maximum allowed dose = $0.30 \text{ W}/\text{m}^2$
(WHO guidelines – ICNIRP)

Recommendations for open wounds or unprotected skin (SCCP {European Commission} Report 0949/05)

- For **open wounds or unprotected skin** we used a modified erythema action spectrum to calculate the total erythema weighted irradiance:
- $\Sigma P_{\text{eff}}(\lambda) \times \Delta\lambda = 21.1 \mu\text{W}/\text{cm}^2 = 0.21 \text{ W}/\text{m}^2 < 0.3 \text{ W}/\text{m}^2$

Optical emission spectra of UV radiation produced by the MicroPlaSter and the sun

UV Power ($\mu\text{W}/\text{cm}^2$)

	UVC	UVB	UVA
Sun	1-2.5	30-50	~600
MicroPlaSter	10-16	40-60	<100

microwave power 60W, main (Ar) gas flow rate 1300sccm, z 20mm

1 min of MicroPlaSter treatment gives the same UVC dose as 5 min sunlight. For UVB 1 min of treatment is equivalent to 1 min solar exposure. For UVA 1 min of treatment corresponds to 10 s of sun exposure.
(Steffes B., Shimizu T. et al. 2008, 2009)

Background of treatment time reduction: UV-measurements of argon plasma

- There are no regulations and studies about long-term effects of plasma treatment
- We do produce UV, and to some parts UVC as well, which is known to be carcinogenic

To have a „safe“ distance to the aforementioned limits/ recommendations we decided to reduce treatment time to 2 min

FDA approved UVC devices (254nm)

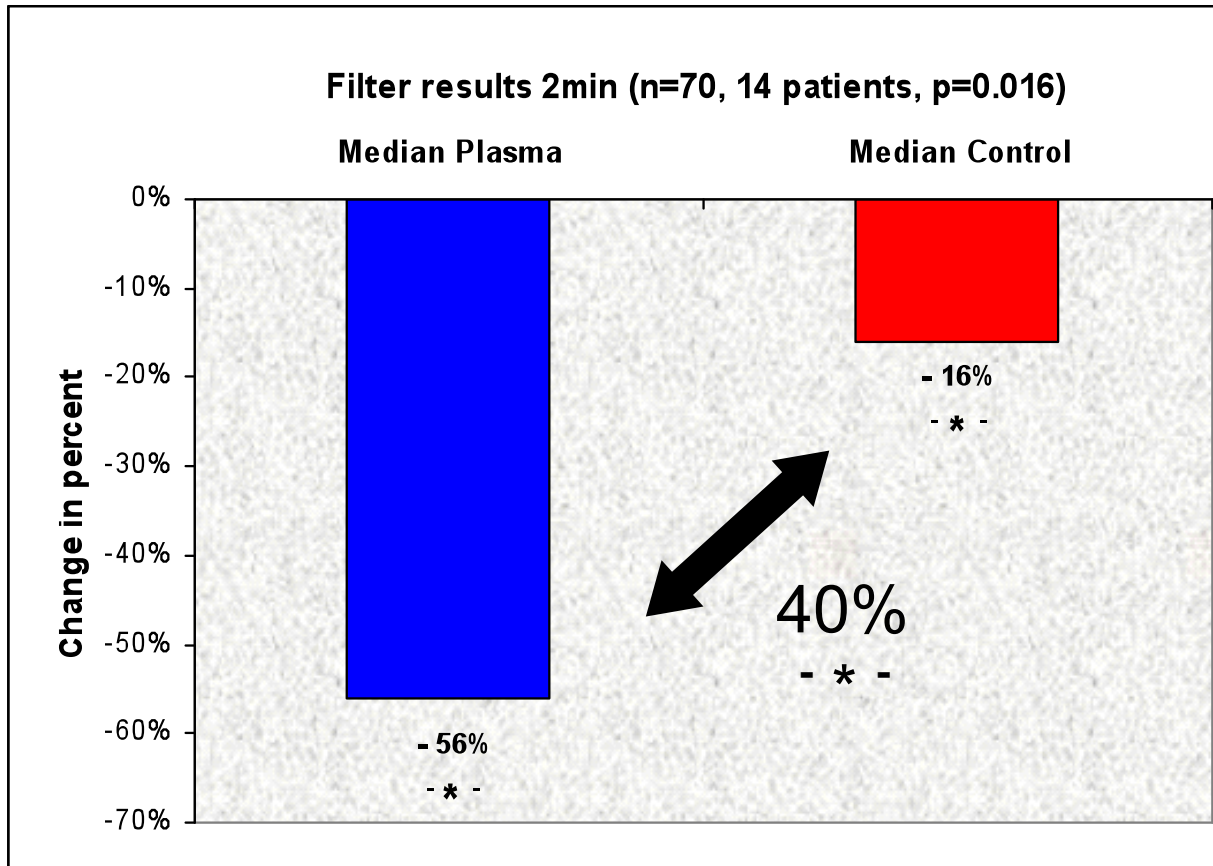


MedFaux



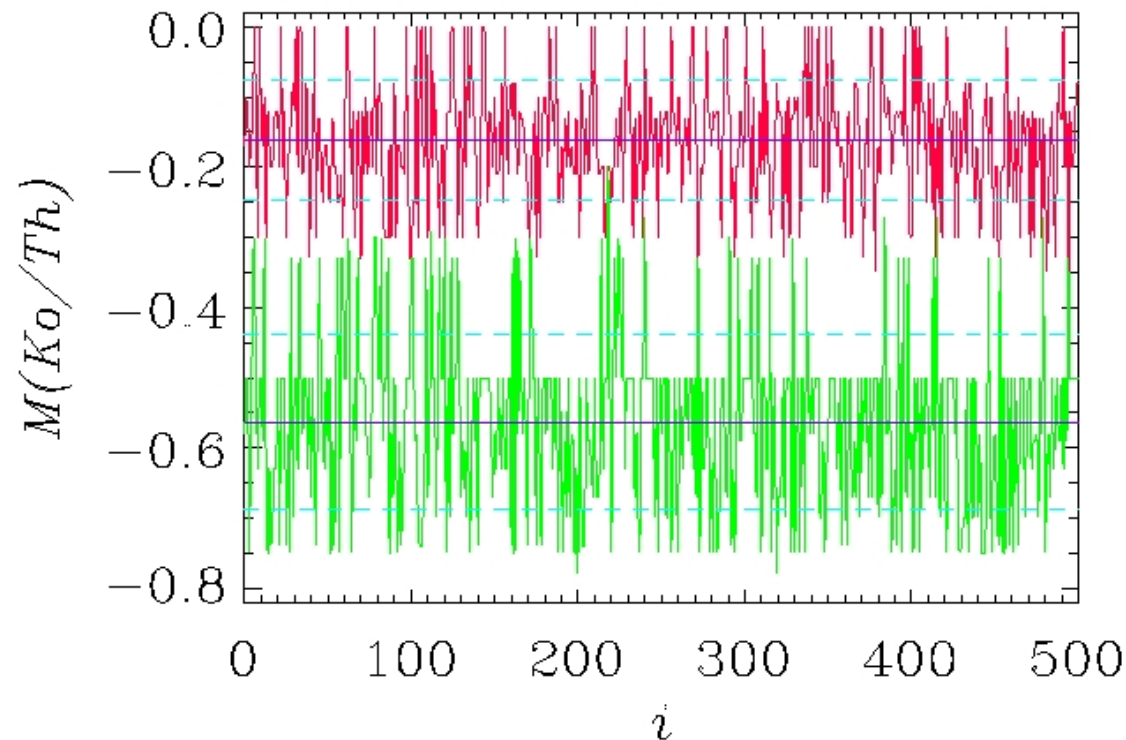
National Biological

Results: 2 min treatment time



Significant ($p < 0.016$) higher germ reduction (40%)
in plasma treated area

Summary of Phase II - Results 2min of treatment time



Results from the corresponding bootstrap-test

Faster wound healing due to plasma therapy?

Very difficult part to measure/evaluate the wound size and changes

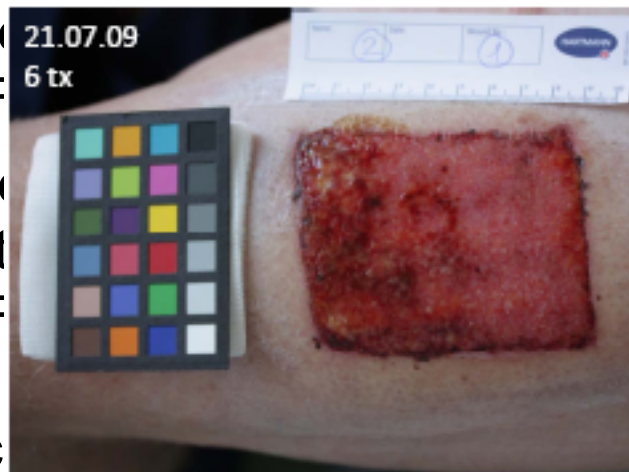
Data in progress, BUT:

Possible faster wound healing due to first „impressions“ of an interim analysis with mesh grafts

Keratinocytes: induce VEGF

Fibroblasts: induce inhibit induces VEGF

reduc



FGF, GM-CSF, 4

(treatment time)
(treatment time)
MMP1,

Results

- A highly significant (34%, $p < 10^{-6}$) higher germ reduction in 5 min plasma treated area vs. control area
- A significant (40%, $p = 0.016$) higher germ reduction in 2 min plasma treated area vs. control area
- No side effects occurred until now, and the treatment is well tolerated in almost all cases
- The use of nitrocellulosis filters revealed a higher accuracy and reproducibility than common swab techniques

www.mpe.mpg.de/theory/plasma-med/index.html

 **PLASMA HEALTH CARE**
A project initiated by the Max Planck Institute for Extraterrestrial Physics

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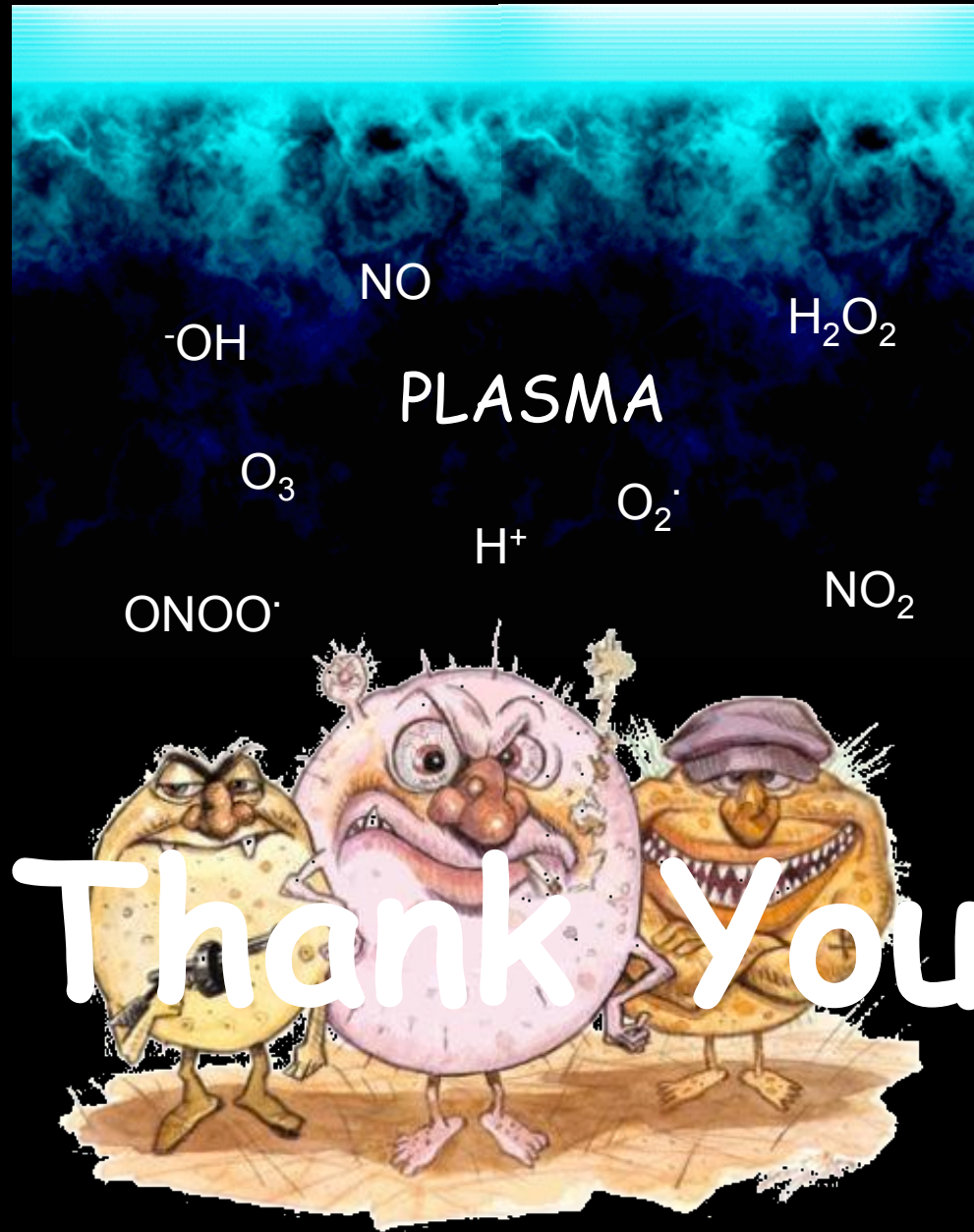
Plasma Health Care in the Media:

German Television:

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- 17-12-2009, ZDF "heute journal": [View](#) (proceed to 16:42 min:sec) or watch clip [here](#).
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- Pforzheimer Zeitung (16-01-2010)
- Physik Journal and 3sat nano (01-2010)
- TAZ (13-02-2010)
- The New York Times (14-02-2010)



We hope that cold atmospheric argon plasma will be an established method to decrease bacterial loads of chronic infected wounds

> Klinikum Schwabing



Gregor Morfill
Tetsuji Shimizu
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Michael Landthaler
Sigrid Karrer
Julia Heinlin
Tim Maisch



Shuitsu Fujii
Mary McGovern
Takuya Urayama

Thank you

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Wilhelm Stolz
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NOVELTIES

Hospital-Clean Hands, Without All the Scrubbing

By ANNE EISENBERG
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HOSPITAL workers often have to wash their hands dozens of times a day — and may need a minute or more to do the process right, by scrubbing with soap and water. But new devices could reduce the task to just four seconds, cleaning even hard-to-reach areas under fingernails.



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Phil Wilson

A prototype hand sanitizer, left, designed by Gregor Morfill.

Instead of scrubbing, the workers would put their hands into a small box that bathes them with plasma — the same sort of luminous gas found in neon signs, fluorescent tubes and TV displays. This plasma, though, is at room temperature and pressure, and is engineered to zap germs, including the drug-resistant supergerm [MRSA](#).



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The technology is being developed in several laboratories. Gregor Morfill, who created several prototypes using the technology at the Max Planck Institute for Extraterrestrial Physics in Garching, Germany, says the plasma quickly

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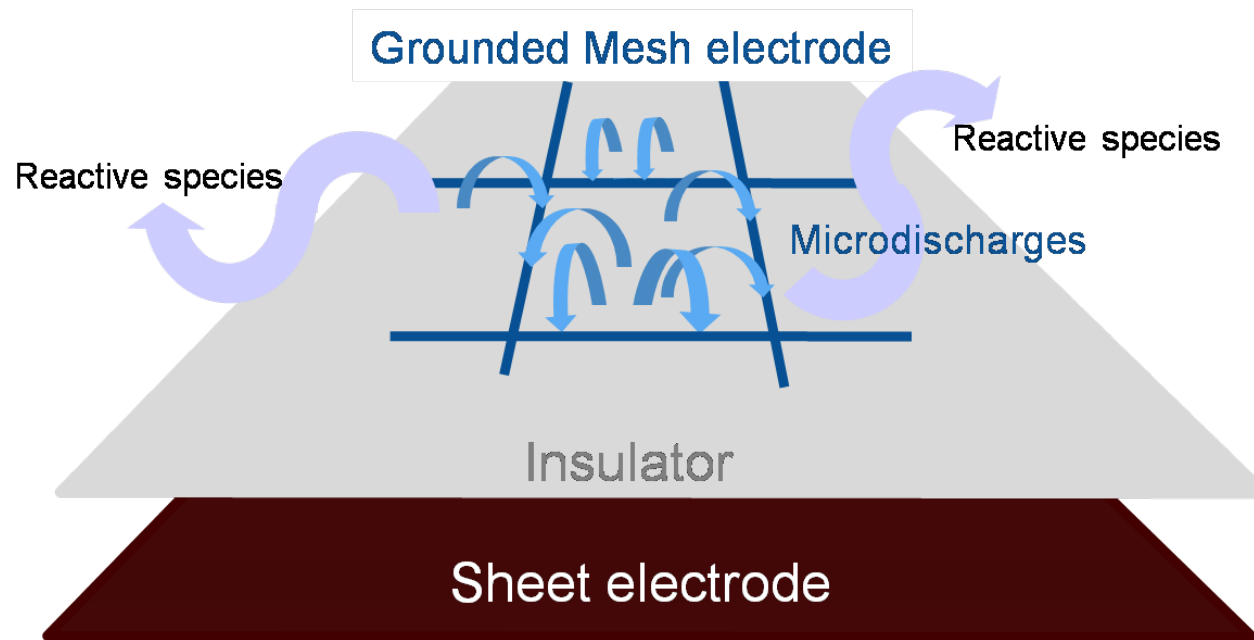
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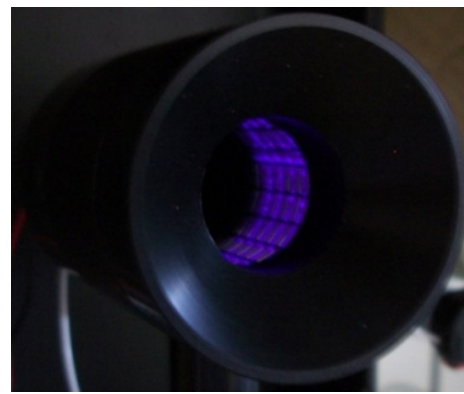
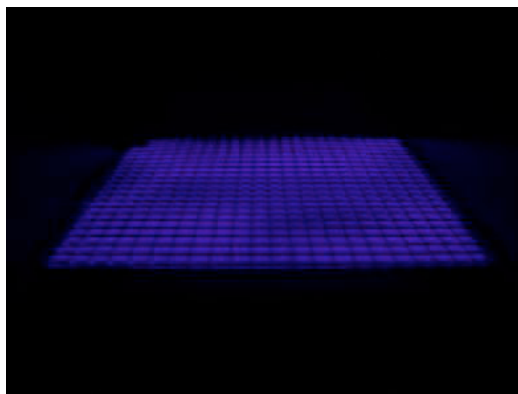
> Klinikum München



Barrier Corona Discharge



- Used gas: air
- Voltage = 18 kV
- Frequency = 12.5 kHz
- Power = 0.5 W/cm²



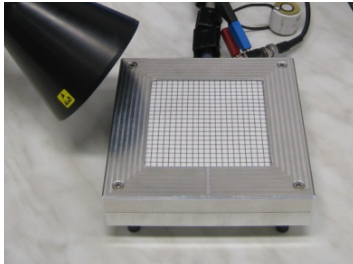
⇒ Plasma is produced by many nano- and microdischarges

Morfill et al. 2009

Possible applications



Handdisinfection (HandPlaSter)



Athlete's foot (FootPlaSter)

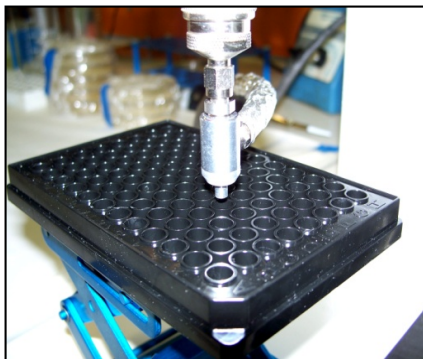


Oral hygiene (OralPlaSter)



Personal hygiene
(DeoPlaSter)

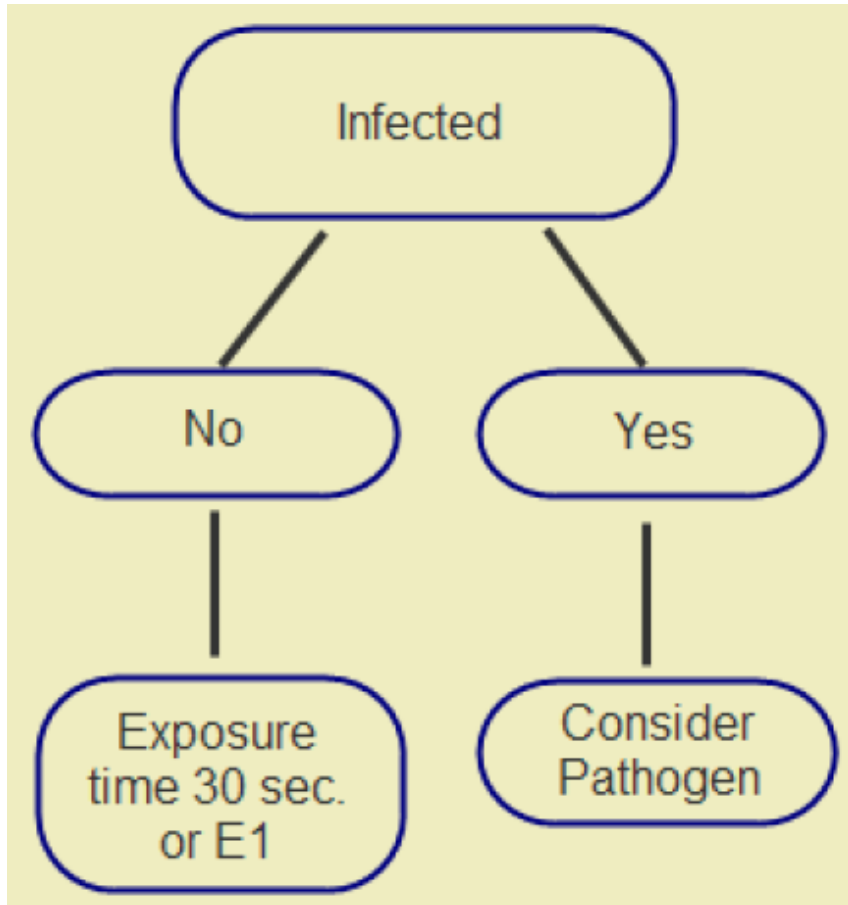
Applications in medicine



- Wound Care
- Treatment of skin diseases (Itching diseases)
- Parodontosis prophylaxy
- Scar prevention
- Treatment of cuts

R&D Network:

- **Plasma physics** (MPE, Eindhoven, Loughborough)
- **Plasma Diagnostics** (MPE, Eindhoven)
- **Plasma Chemistry** (MPE, Berkeley)
- **Plasma Engineering** (MPE, ADTEC)
- **Plasma Biology** (MPE, TUM, Regensburg)
- **Plasma Microbiology** (Schwabing, Regensburg)
- **Plasma Medicine** (Schwabing, Regensburg)
- Also there is a cooperation in all fields with six Research Institutes from the Russian Academy of Science and the Russian Academy of Medical Science
- **Technology Transfer** (Max-Planck Innovation GmbH)



UVC treatment algorithm (Conner-Kerr)

