IFVBESA

Information is crucial

P50 2.1 BESA-Detailed Project

various chemical stress factors

Human Protect SD1

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Detailed- Project P50 2.1
to BESA expert opinion

Bioenergy informative system analysis within the framework of the BESA seal of approval about the effectiveness
of the product „Human Protect SD1“
Made in Germany / SN-213509-NH
of the company IPC Europe UG / D 77866 Rheinau
for various ingredients of a mRNA vaccine
in the project also called "test object“

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Client:
IPC Europe UG
Norbert Heuser
Schwarzwaldstraße 48
D 77866 Rheinau

Project participants:

Project leader:	Wolfgang Hans Albrecht, president and scientific director of the IFVBESA
Testing person:	Eva Krankl, Vice president and deputy scientific director of the IVFBESA
Respondent:	1 anonymou test person with different health conditions and age or gender. Subject aged 80 years, 2x Corona mRNA inoculated, was tested for the effect of the test object against spike proteins, virus fragments and graphene oxide.
other participants:	none

Project location: location of IFVBESA (international professional association for bio energie information system analysis), Hauptstraße 1, A-4861 Kammer/Schörfling am Attersee

Date: 27.10.2021 until 16.11.2021

Project duration: 4 Days

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Important notes

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BESA- legend for interpretation of BESA-measurement result

The measured value of 50 on the tested meridian represents an optimal energetic state in this organ or its subordinate and superordinate levels. Also measured values in the range of 50 to max. 70 still count as a neutral and balanced energy status. The organism is able to regulate irritations of the system (wrong environmental signals) very well.

Readings above 70 to 100 represent the inflammatory range or a so-called energy surplus in response to the irritation of the system by environmental signals corresponding to it. After reaching the maximum values, the energy state tilts into the degenerative (blue) range.

Readings from below 50 to near 0 represent the so-called degenerative measurement range or a lack of energy in response to the stimulation of the system by environmental signals corresponding to it.

Measured values represented by a so-called pointer drop of more than 3 scale lines indicate a total deregulation. The influence of certain environmental signals then leads to such strong system overloads, which can only be harmonized by corresponding new signals.

BESA Indicators:

up 0,79	very deep energetic regulation disorder (SSD) energy deficiency
0,8 to 1,19	severe energy regulation disorder (SD) degeneration/energy deficiency
1,2 to 1,59	energetic regulation disorder (D) degeneration/energy deficiency
1,6 to 1,99	degenerative transition area (DÜ)
2,0 to 2,39	optimale regulation (OR)
2,4 to 2,79	in regulation (R)
2,8 to 3,19	partial inflammation = regional energy excess (PE)
from 3,2	total inflammation = strong general energy surplus (TE)

Basics of the research project creation P50 2.1

The international professional association for bioenergetic system analysis was commissioned by the company IPC Europe UG to test the effect of the test object "Human Protect SD1" by means of bioenergetic system analysis (BESA) or to prove its effect. The testing took place independently of the subjective feeling of the test person.

Description of the „Human Protect SD1“ by the client:

The name of the test object already says for which field of application it is designed:
"For people and protection"!
On the front of the test object are the areas: "Body, mind, psyche".
On the back are the "elements of holistic physiology: "was, is, I am, will".
These elements are supposed to be an interpretation about the scope of the test object's mode of action and an indication of its holistic, i.e. holistic, orientation. In order to meet this requirement, a specific program was developed by the client and the test object was programmed with the software and hardware developed by the client.

The test object forms a virtual sphere with an effective radius of 5 meters and a diameter of 10 meters. The effect is created by means of interactions between the so-called hyperspace and the reference space of the test object and biological objects."

So, the concept of this test object should be to harmonize, neutralize disturbances, problems, blockades, disharmonies in biological objects and thus replace negative states with positive states. This has been done in the projects

Furthermore, the client assumes that side effects triggered by the mRNA vaccination on the body or in the brain can be reversed by the test object. The client also assumes that these effects should also occur with other similar vaccinations or vector-like vaccinations.

The test object should also protect unvaccinated people from infection by shedding.

Due to its mode of operation, there is a wide range of applications for the test object. In the case of this project it is about the application of the "Human Protect SD1" with various modern, so-called mRNA or vector-based vaccines and its effect on the human body.

Research Support Services of the IFVBESA - BESA Reference Tests

Project P50 2.1 specifically addresses the proof of concept of "Human Protect SD1" against a subject vaccinated 2x Corona mRNA.
About the pre-existing conditions:
The subject suffers from advanced Alzheimer's dementia disease as well as mild hypertension. He takes appropriate medication such as donepezil (cholinesterase inhibitor for dementia) as well as anticoagulants (blood thinners) and ACE inhibitors against the increased blood pressure. In addition, the test person is almost permanently in the environment of Corona

vaccinated people. Thus, it can be assumed that at least the transmission of spike proteins can occur within these people or groups of people.

In this detailed project, special attention is paid to spike proteins as well as virus fragments and graphene oxide, which, according to scientific studies (see list of sources on pages 35 to 37), suppress the repair mechanism of DNA and thus irritate the immune system or cause autoimmune reactions. Spike proteins, virus fragments as well as graphene oxide are digitized and their effect simulated accordingly in the BESA test.

The "Human Protect SD1" is tested according to the client's request within the framework of the applicable conditions of the IFVBESA for the award of quality seals. In principle, seals of approval are awarded in three categories depending on the significance of the test results, taking into account all the tests of a project. For the "Human Protect SD1" it is to be determined whether by its application stresses over already mentioned stress factors like the administered mRNA vaccine and as a result of it in the energy system of the test person (biological system) arising or existing disturbances, problems, blockades, disharmonies can be harmonized, neutralized and thus negative pathological conditions can be replaced by positive conditions. This is questioned in the following commissioned tests of this project

Research project description

The reason for the test is to prove the functionality of the "Human Protect SD1" by test results obtained by confronting the test person with the digitized spike proteins, the virus fragments as well as the graphene oxide in order to significantly prove and compare the reactions without the "test object" and with the "test object". In a BEFORE measurement, the subject is exposed to the listed stress factors and tested. In the AFTER measurement, in addition to the stress factors mentioned, the test person is brought into contact with the test object, the "Human Protect SD1", and tested.

- The BEFORE measurements are performed without the „Human Protect SD1“
- The AFTER measurements are performed with the „Human Protect SD1“

The question for each AFTER measurement is: "Is the "test object" suitable and able to harmonize or neutralize the so perceived negative effects of the mentioned stress factors on the organism"?

The correspondingly designed tests should provide information about this by comparing the pre-measurement without the "Human Protect SD1" with the test results of the post-measurements to be carried out using the "Human Protect SD1".

The client's concern is to determine whether the test object, the "Human Protect SD1", as noted in the product description, is suitable for harmonizing the disturbances, problems, blockages, disharmonies in the meridian system of the test person (biological object) resulting from the above-mentioned stress factors.

General information transfer of the test object

The information transfer takes place from the hyperspace of the test object to the hyperspace of biolo-gical objects (humans, animals, plants). From there the information reaches the reference space or the energy space via so-called interaction channels. This is a union of, among other things, all organs and energy forms in the biological object. There, the information of the program can dynamically materialize and thus change current states. The changes can appear in the form of neutralization or harmonization of disturbances, dissolution of problems, blockages and disharmonies..

Conditions:

The BESA tests are performed on the premises of the IFVBESA under laboratory conditions, at a room temperature of 20°Celsius, on a natural wooden floor. In principle, the test persons are deswitcht (made testable) before the BESA tests or the test possibility is questioned with the test persons.

Pos.1 BESA 1 testing Basic (bioenergetic status) on the test person

Pos.2 BESA 2 testing Basic-BREVIS (extended bioenergetic status) on the test person

Pos.3 BESA 3 testing when the test person is confronted with the digitized spike proteins and virus fragments.

Pos.4 BESA 4 testing when the subject is confronted with the digitized spike proteins, the virus fragments and the graphene oxide.

Pos.5 BESA 5 testing when the test person is confronted with the already mentioned stress factors and the test object, the "Human Protect SD1".

Pos.6 Evaluation of the results in the project as well as summary in a corresponding expert opinion according to the sample.

Procedure and specifications for the implementation

1. **BESA-basic measurement of the test person** at all previously determined measurement points (TING points) serves to determine the actual condition. The results are determined exactly according to the BESA specifications and documented via the BESA graphs.
2. **Basic BESA measurement BREVIS of the test person** at all previously determined measuring points (BREVIS points) serve to determine the actual condition. BREVIS refers to an extended matrix of acupuncture points to be measured. These points are applied or tested in order to get a deeper insight into the energy-informative events of the functional circuits of the test person. The results are again determined exactly according to the BESA specifications and documented via the BESA graphs.
3. **The test person** is brought into **contact with the listed stress factors** depending on the project, whereby the sequence discussed with the developer is considered as a

specification and is adhered to accordingly. In order to be able to determine the current energy state, the measuring points mentioned under point 1 are measured in the same sequence and for the same duration for each test of the load factors. The results are determined exactly according to the BESA specifications and documented via the BESA graphs.

4. **Activation of the „Human Protect SD1“ test object**

   4.1. When **activating the "Human Protect SD1"**, it is brought into the measurement area according to the client's instructions.

   4.2. The test person is brought into contact with the load factors. The measuring points mentioned under point 1 are measured in the same order and for the same duration in order to determine the current energy state. The results are determined exactly according to the BESA specifications and documented via the BESA graphs.

Test procedure

BESA 1 testing BASIC BEFORE as status

In the first step, a basic bioenergetic test (bioenergetic status) is performed on the meridian end points (TING points) of the test person.

BESA 2 testing BASIC BREVIS BEFORE as status

In the next step, a basic bioenergetic BREVIS testing (bioenergetic status) is performed on the meridian end points (TING points) of the subject.

BESA 3 testing BEFORE with the mentioned stress factors like spike-proteins and virus-fragments.

In the further BESA testing process, the subject is contacted and tested with listed digitized spike proteins and the virus fragments. The questions are: How does the meridian system react? How does the energetic status change when confronted with the stress factors? What are the differences between BESA 1 and 2 BASIC testing BEFORE?

BESA 4 testing BEFORE with the mentioned stress factors from BESA 3 testing BEFORE and the graphene oxide.

In the further course of BESA testing, the test person is brought into contact with the digitalized stress factors and tested. The questions are: How does the meridian system react? How does the energetic status change when confronted with the stress factors? What are the differences compared to the BESA 1,2 and 3 tests BEFORE?

BESA 5 testing AFTER with the mentioned stress factors as well as the "Human Protect SD1".

In the last BESA test, all the load factors already tested above and the "Human Protect SD1" are introduced into the measurement circuit. Now the question is: How does the meridian

system of the test person react within the effective range of the "Human Protect SD1", if all the already tested stress factors are activated at the same time?

BESA 1 Testing BASIC BEFORE

BESA 1 testing BASIC BEFORE as status

Eva Krankl performed a BESA basic measurement on the test person. All BESA tests were performed at the TING points (40 nail fold points on the fingers and toes).

Goal: The creation of a basic test (status) for the representation of the energetic initial situation for all further BESA tests.

BESA-test evaluation P50 2.1

from **29-11-2021 at 10:35 to 10:39** (4 minutes) page 11 and 12

result: The measurement result indicated partly strong energetic stresses at the meridian end points and subsequently on the subordinate metabolic situation of the test person.

97 % in the blue area

2 % in the red area

1 % in the yellow transition area

Conclution: As the graphs show, all measuring points are in the degenerative blue range (energy deficiency) although 2% of them are even in the red measuring range. The red measured values interpret a current total deregulation in this control circuit (organ large intestine). The comparisons of the BESA graphs confirm the stressful influences on the energy-informative events in the meridian system of the test person.

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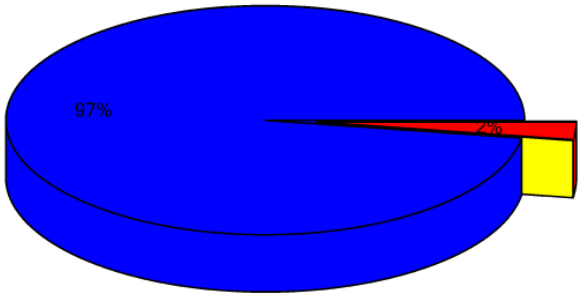
Overview of BESA measuring



Red: values with indicator decline
Decline greater than 2 (2%)

Yellow: high values without indicator decline
Values over 70 ()

Blue: low values
Values lower 50 (97%)

Green: standard values
Values between 50 and 70 ()

BESA basic test

+++: Indicator decline > 15 Skt.
++: Indicator decline 6-15 Skt.
+: Indicator decline 3-5 Skt.

T: Total inflammation (>89 Skt.)
P: Partial inflammation (70-89 Skt.)

D: Degeneration (< 50 Skt.)

Standard values: (50-70 Skt.)

Element: lu - sk - li - ct

	Right	Left
Lung		
Lu 1 (11.) Parenchym	24/1	27/0
Skin		
Sk 1 (1.) Lower Body	31/1	19/0
Large intestine		
Li 1 (1.) Colon transv./sigm.	32/3 +	31/0
Connective tissue		
Ct 1 (1.) Abdomen	23/1	32/0

Element: st - nd - ps - od

	Right	Left
Stomach		
St 1 (45.) Pylorus/body	32/1	18/0
Nerves deg.		
Nd 1 (1.) lumb./sakral.	25/2	18/0
Pancreas-spleen		
Ps 1 (1.) Eiw./w.Pulpa	35/0	38/1
Organ deg.		
Od 1 (1.) Abdominal region	10/0	10/0

Element: bl - ly - ki - al

	Right	Left
Bladder		
BL 1 (67.) body	30/1	29/0

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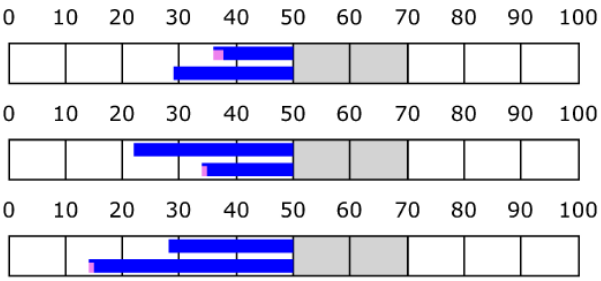
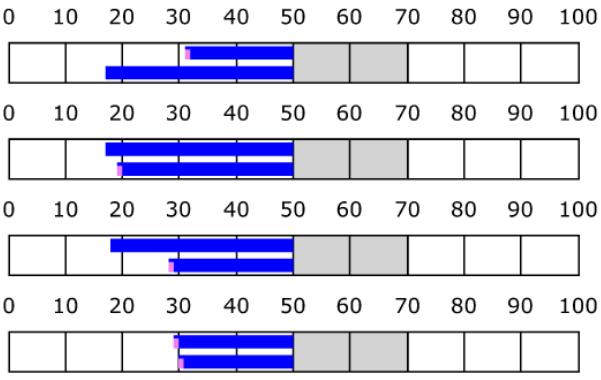
Standard values: (50-70 Skt.)

Element: bl - ly - ki - al



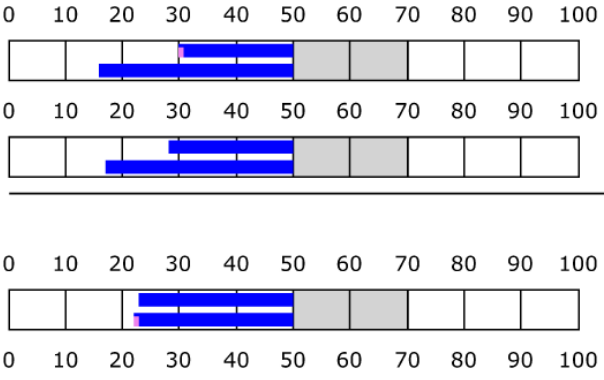
	Right	Left
Lymph		
Ly 1 (1.) Tons.Palat.	36/2	29/0
Kidney		
Ki 1 (11.) pelvis	22/0	34/1
Allergy		
Al 1 (1.) Lower body	28/0	14/1

Element: ga - jd - li - gd



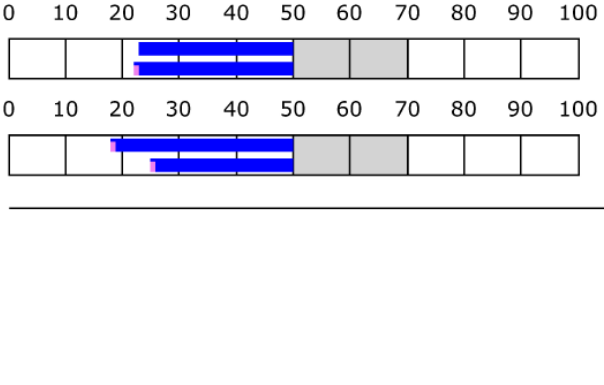
	Right	Left
Gall		
Ga 1 (44.) Duct.choled./hep.	31/1	17/0
Joint deg.		
Jd 1 (1.) lower extremities	17/0	19/1
Liver		
Li 1 (1.) Central veins	18/0	28/1
Greasy deg.		
Gd 1 (1.) Abdomen	29/1	30/1

Element: he - si



	Right	Left
Heart		
He 1 (9.) Pulm.sm./Aortic valves	30/1	16/0
Small intestine		
Si 1 (1.) Ileum	28/0	17/0

Element: bc - 3e



	Right	Left
Blood circulation		
Bc 1 (9.) SMP Arteries	23/0	22/1
Endocrine		
3e 1 (1.) Gonads/NNI	18/1	25/1

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BESA 2 testing BEFORE

BESA 2 testing BASIC BREVIS BEFORE as extended status

Eva Krankl performed a second, extended basic BESA measurement on the test person. All BESA tests are now performed at the BREVIS points (78 measuring points on the fingers and toes including the base joints). In this BESA testing design, in addition to the acute points on the toes (40measuring points), other acupuncture points on the fingers and toe joints are tested.

Goal: The creation of an extended basic test (status) for the presentation of more profound questions about the respective organs. This BESA testing is intended to provide a more comprehensive and deeper understanding of the subject and his or her energy-informative initial situation.

BESA-test evaluation P50 2.1

from **29-11-2021 at 11:04 to 11:15** (11 minutes) Page 14 to 16

result: The measurement result also indicated again strong energetic stresses at the tested meridian points and in further consequence on the subordinate metabolic situation of the proband.

100 % in the blue area

Conclution: As the graphics show, all measuring points are also in the degenerative blue area (energy deficiency). This BREVIS test variant was selected in order to gain a deeper insight into the energetic events in the meridian system of the test person. As can be seen in the BESA graphs, this matrix allows a larger spectrum of organ details and thus also shows the chronified events in the energy-informative area in contrast to the TING points (acute events). The deeper measurements at the extended acupuncture points also show similarly deep, degenerative readings. The BESA graph shows the chronic course of the stresses. In contrast to the "acute events" from the BESA 1 test BEFORE, the red measured values are missing, but this test is about the deeper, chronic progressions in the energy-informative system. But also in chronic course the measuring point of the "peritoneum" (abdominal membrane) as part of the large intestine with 22/2 (see page 15) also just scratches a red measured value. The comparisons of the BESA graphs thus confirm that, in addition to the acute measuring points, all chronic measuring points also lie deep in the degenerative range and thus confirm the stressful influences on the energy-informative events in the meridian system of the test person.

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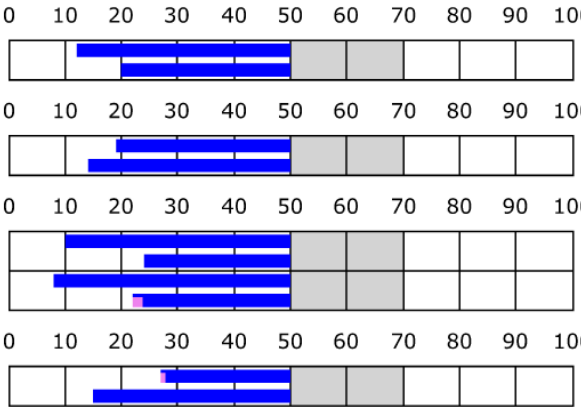
D: Degeneration (< 50 Skt.)

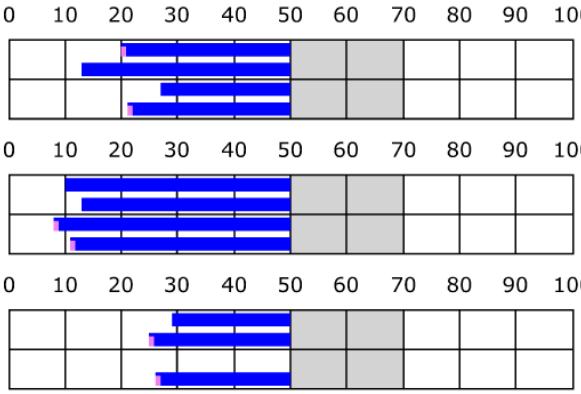
Standard values: (50-70 Skt.)

Element: lu - sk - li - ct



	Right	Left
Lung		
Lu 1 (11.) Parenchym	12/0	20/0
Skin		
Sk 1 (1.) Lower Body	19/0	14/0
Large intestine		
Li 1 Checkpoint	10/0	24/0
Li 1 (1.) Colon transv./sigm.	8/0	22/2
Connective tissue		
Ct 1 (1.) Abdomen	27/1	15/0

Element: st - nd - ps - od



	Right	Left
Stomach		
St 1 (45.) Pylorus/body	20/1	13/0
St Checkpoint	27/0	21/1
Nerves deg.		
Nd 1 (1.) lumb./sakral.	10/0	13/0
Nd Checkpoint	8/1	11/1
Pancreas-spleen		
Ps Checkpoint	29/0	25/1
Ps 1 (1.) Eiw./w.Pulpa		26/1

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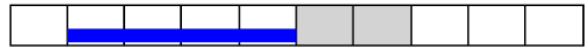
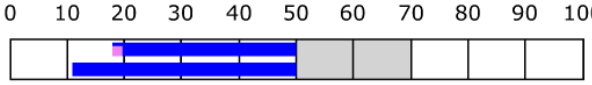
Element: st - nd - ps - od

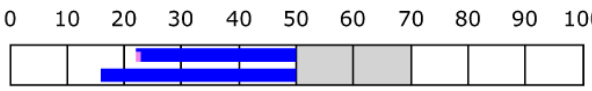


Pancreas-spleen	Right	Left
Ps 1 (1.) Eiw./w.Pulpa		10/0

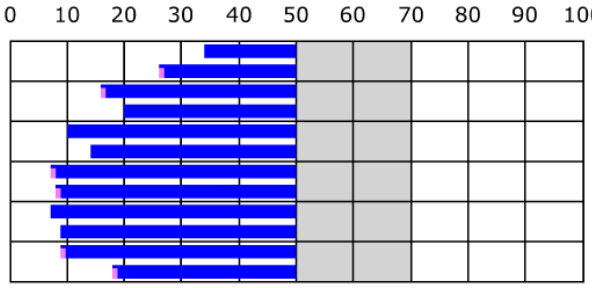


Organ deg.	Right	Left
Od 1 (1.) Abdominal region	18/2	11/0

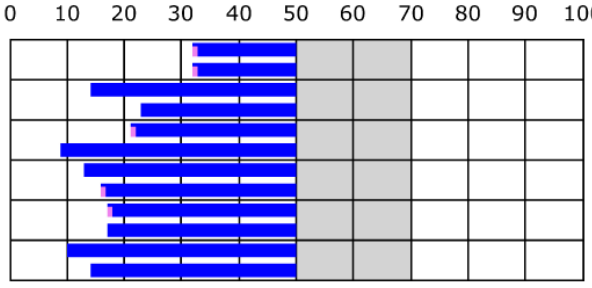
Element: bl - ly - ki - al



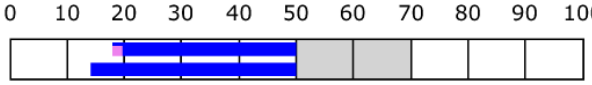
Bladder	Right	Left
BL 1 (67.) body	22/1	16/0



Lymph	Right	Left
Ly 1 (1.) Tons.Palat.	34/0	26/1
Ly Checkpoint 1	16/1	20/0
Ly Checkpoint 2	10/0	14/0
Ly Checkpoint 3	7/1	8/1
Ly Checkpoint 4	7/0	9/0
Ly Checkpoint 5	9/1	18/1

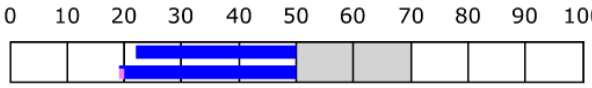


Kidney	Right	Left
Ki 1 (11.) pelvis	32/1	32/1
Ki Checkpoint 1	14/0	23/0
Ki Checkpoint 2	21/1	9/0
Ki Checkpoint 3	13/0	16/1
Ki Checkpoint 4	17/1	17/0
Ki Checkpoint 5	10/0	14/0

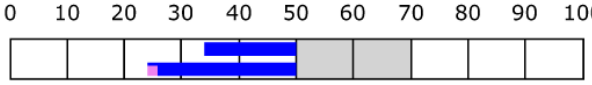


Allergy	Right	Left
Al 1 (1.) Lower body	18/2	14/0

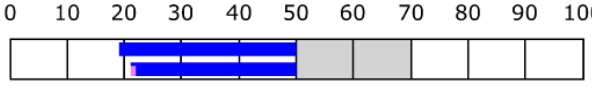
Element: ga - jd - li - gd



Gall	Right	Left
Ga 1 (44.) Duct.choled./hep.	22/0	19/1



Joint deg.	Right	Left
Jd 1 (1.) lower extremities	34/0	24/2



Liver	Right	Left
Li 1 (1.) Central veins	19/0	21/1

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BESA basic test

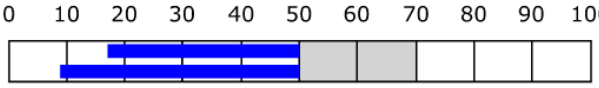
+++: Indicator decline > 15 Skt.
++: Indicator decline 6-15 Skt.
+: Indicator decline 3-5 Skt.

T: Total inflammation (>89 Skt.)
P: Partial inflammation (70-89 Skt.)

D: Degeneration (< 50 Skt.)

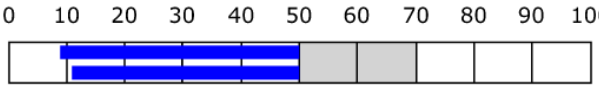
Standard values: (50-70 Skt.)

Element: ga - jd - li - gd

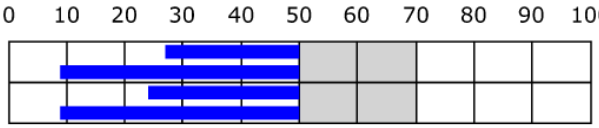


Greasy deg.	Right	Left
Gd 1 (1.) Abdomen	17/0	9/0

Element: he - si

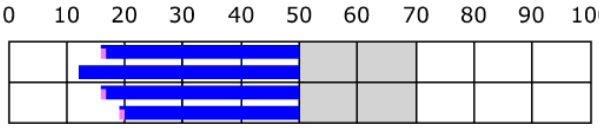


Heart	Right	Left
He 1 (9.) Pulm.sm./Aortic valves	9/0	11/0

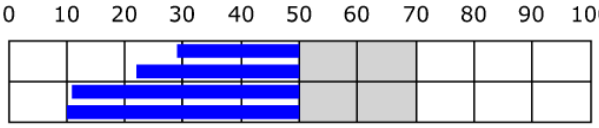


Small intestine	Right	Left
Si 1 (1.) Ileum	27/0	9/0
Si Checkpoint	24/0	9/0

Element: bc - 3e

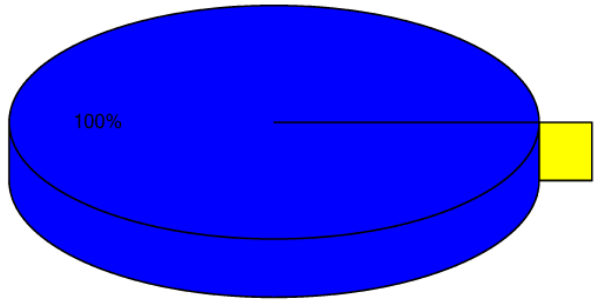


Blood circulation	Right	Left
Bc 1 (9.) SMP Arteries	16/1	12/0
Bc Checkpoint	16/1	19/1



Endocrine	Right	Left
3e 1 (1.) Gonads/NNI	29/0	22/0
3e Checkpoint	11/0	10/0

Overview of BESA measuring



Red: values with indicator decline
Decline greater than 2 ()

Yellow: high values without indicator decline
Values over 70 ()

Blue: low values
Values lower 50 (100%)

Green: standard values
Values between 50 and 70 ()

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BESA 3 testing BEFORE

BESA 3 testing BEFORE; testing for the presence of spike proteins and virus fragments - as a result of the modified spike proteins of the mRNA vaccine

In the further BESA testing procedure, the subject was tested "for the presence of spike proteins" and "viral fragments" in the organism. For this purpose, the digitized loading factors were introduced into the measurement circuit and tested at the specific measurement points. All BESA tests were again performed at the TING points (40 nail fold points on the fingers and toes) as in the BESA 1 test BASIC BEFORE.

Goal: Determination of the reaction in the meridian system of the test person to the question in which control circuit (measuring points of the respective organs) spike proteins and virus fragments can be detected bioenergy-informatively.

ATTENTION: this BESA test is not a status measurement like BESA 1 and 2 test BASIC. Unlike the **"BESA status test"**, this BESA test is about measuring a reaction to the nosodes (digitized spike proteins and virus fragments) in the measuring circuit. We refer to this as **"BESA resonance testing"**. **Question**: Do the digitized nosodes in the measuring circuit show a resonance at the acupuncture points indicated? It is important to understand that in the case of the presence of spike proteins at the respective measurement points, the "measurement response" must indicate a green measurement value (50-70Skt). The possible green measurement results confirm, so to speak, the resonance with the respective nosode from the measurement circuit. "Yes, these substances are detectable in the respective organs tested"!

BESA-test evaluation P50 2.1
from **29-11-2021 at 10:40 to 10:45** (5 minutes) Page 18 to 22

result: ATTENTION, the bioenergy-informative measuring result from the "resonance testing" shows a resonance and thus the load of spike proteins and virus fragments within these tested organs via the representation of the green measuring values.
ATTENTION: Only the green measured values are relevant in this BESA test. The blue measured values play only a subordinate role in this case.

22 % in the blue area

77 % in the green area

1 % in the yellow transition area

Conclutio: As the BESA graphs show, after introduction of the nosodes with the digitized spike proteins and the virus fragments into the measuring circuit of the test person, 77% of the measured values are still in the green range.

ATTENTION: The "green readings" are "positive" because they confirm the resonance and thus the presence of the tested substances in these organs. This is response of BESA resonance test and thus confirms the presence of spike proteins and virus fragments within these organs.

It is also interesting to see in this subject how quickly the ingredients have spread, or are expected to spread, incriminatingly to almost the entire body of the subject. According to the bioenergy-informative measurement results (green area), these listed and tested ingredients (resonance of the digitalized nosodes) are located in the following organs:

The lung- left, skin- right, large intestine- right and left, connective tissue- left, stomach, nerve degeneration, pancreas/spleen as well as organ degeneration right and left, bladder, lymph, as well as kidney right and left, veins (allergy point) left, heart, small intestine right and left as well as circulation and endocrine right and left.

The representation of the green BESA graphs confirms the existing influences on the subject's organs.

ATTENTION: Please note the correspondingly marked measurement results of the organs from the BESA resonance test in the green area!

It is difficult to establish a relationship to the symptoms of the test person (especially Alzheimer's dementia) because the subject matter is too complex. However, a closer look at these measured values, also in comparison to other test persons, raises the question of what influence the medications taken by the test person have in correlation with the ingredients of the vaccine.

For simple explanation:

What are spike proteins (S-proteins)?

An S-protein (S-glycoprotein) is an outwardly protruding viral envelope that is studded with so-called spikes. This S-protein consists of viral membrane proteins, which is anchored in a lipid membrane. The outwardly projecting spikes or spines are glycosylated by sugar residues (saccharification of e.g. proteins). These spines enable the S protein to bind to the surface receptors of the target cell, thus enabling the so-called virus to enter the cell.

About the virus: *Scientifically, a so-called virus could never be identified as what it is assumed to be, as dangerous for humans! Let us leave for the time being the designations in such a way, as they are quoted currently pseudo-scientifically.*

Via the so-called "spikes", the virus and its function or property (for us, a specific piece of information) interact with the host's antibodies when it binds to the host cell.

Never has anything attracted so much attention in the past months/years since Corona as the "spike protein". It is important to understand that in addition to the typical S protein found in influenza disease, there is another. This is the one that is produced by genetically engineered vaccines such as the mRNA vaccine or the vector vaccines. In principle, the natural S protein

is already a challenge for the human immune system, but the modified S protein that the body produces in response to the vaccine represents a real challenge for the human organism.

In an article by Stephanie Seneff and Greg Nigh titled "Worse Than The Disease: Reviewing Some Possible Unintended Consequences of mRNA Vaccines Against COVID-19: Reviewing Some Possible Unintended Consequences of mRNA Vaccines Against COVID-19), published in the International Journal of Vaccine Theory, Practice and Research explains that a major part of the problem is that while natural spike protein is bad, spike protein produced by the body in response to the vaccine is even worse.

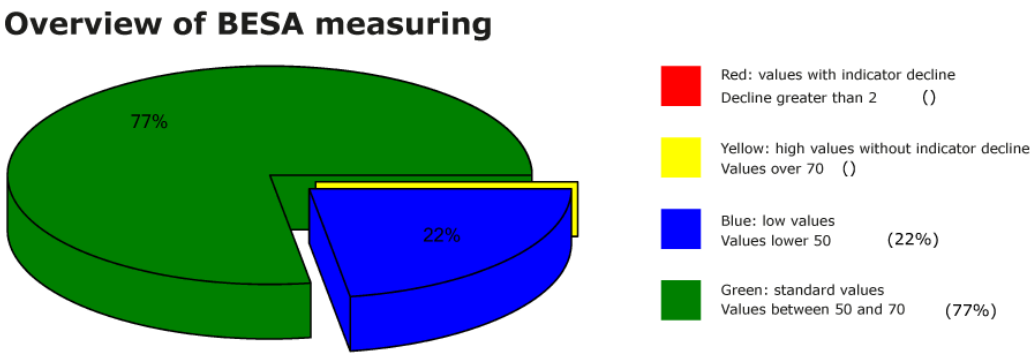
What are modified spike proteins: For this, it is necessary to first return to modified RNA (mRNA). RNA stands for ribonucleic acid. This means that RNA or RNA is a chain of so-called nucleotides. They are the basic building blocks of DNA and RNA and also have regulatory functions in cells. This synthetic RNA (mRNA) has been manipulated to produce an "artificial spike protein". The difference to the natural S-protein is that it does not collapse as soon as it binds to an ACE2 receptor, but it remains open and attached to the ACE2 receptor. This overrides the receptor and allows for appropriate immunological reactions, leading to challenges such as cardiopulmonary and autoimmune reactions.

ACE2 (Angiotensin Converting Enzyme 2) is a protein compound that is mainly produced at the vascular endothelial cells of the heart, kidneys as well as respiratory epithelia and the gastrointestinal tract. It thus also plays an important role in the regulation of blood pressure as well as in anti-inflammatory and lung-protective effects.
Exactly these or similar symptoms could be detected in this and the following subjects with BESA.

What are viral spikes:
Here is the further explanation of what happens after the ACE2 receptors are inhibited by the spike proteins. These enter the cell and nucleus through the injected mRNA vaccine, suppressing the human body's DNA repair mechanism and triggering an explosion of immune deficiency,autoimmune,or other severe complications.
New research published in Viruses, part of MDPI's SARS-CoV-2 Host Cell Interactions Edition (open access journals), shows that vaccine spike proteins enter cell nuclei and destroy the cells' DNA repair mechanism by suppressing DNA repair by up to 90%.
The research paper is titled "SARS-CoV-2 Spike Impairs DNA Damage Repair and Inhibits V(D)J Recombination In Vitro" and was authored by Hui Jiang and Ya-Fang Mei at the Department of Molecular Biosciences, The Wenner-Gren Institute, Stockholm University, SE-10691 Stockholm, Sweden, and the Department of Clinical Microbiology, Virology, Umeå University, SE-90185 Umeå, Sweden, respectively.

Mechanistically, the researchers found that the spike protein is localized in the nucleus and inhibits DNA damage repair by interfering with the recruitment of the important DNA repair proteins BRCA1 and 53BP1 to the damage site.

This means that the spike protein, which is formed in the cell's ribosomes after cells are hijacked by mRNA vaccines, does not always leave the cell and enter the bloodstream, as proponents of mRNA vaccines tell us.
In some cases, the spike protein enters the nucleus. There it interferes with the DNA repair mechanism, as described in this article. It confirms that such vaccines do indeed destroy genetic integrity and have side effects that were not predicted or described by proponents of mRNA vaccines. These SARS-CoV-2 viral fragments are referred to as nonstructural protein or "Nsp1, Nsp5, Nsp13, and Nsp14." Overexpression of these viral fragments and spike proteins reduces DNA repair efficiency (NHEJ repair) according to this study.



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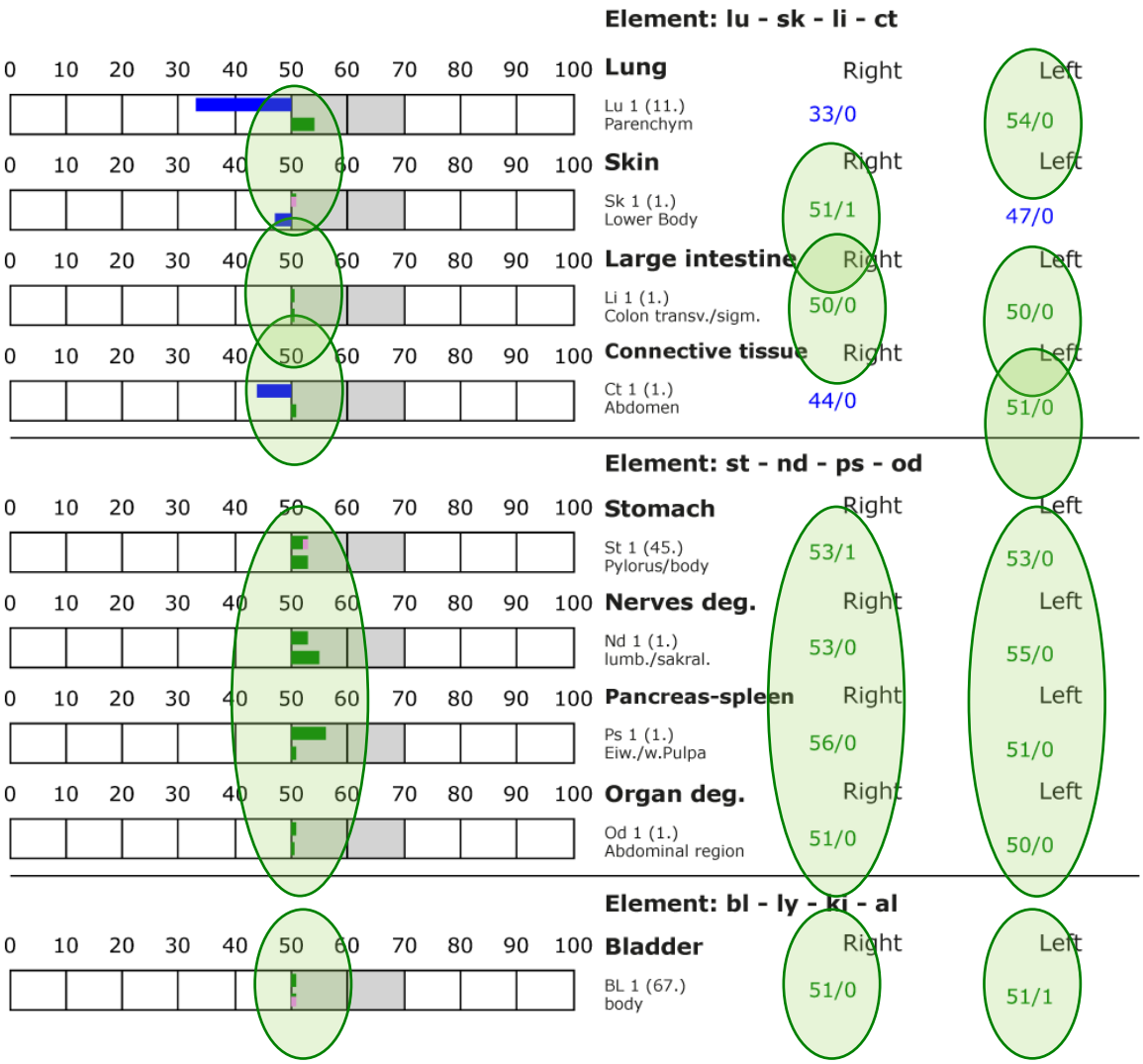
D: Degeneration (< 50 Skt.)

Standard values: (50-70 Skt.)



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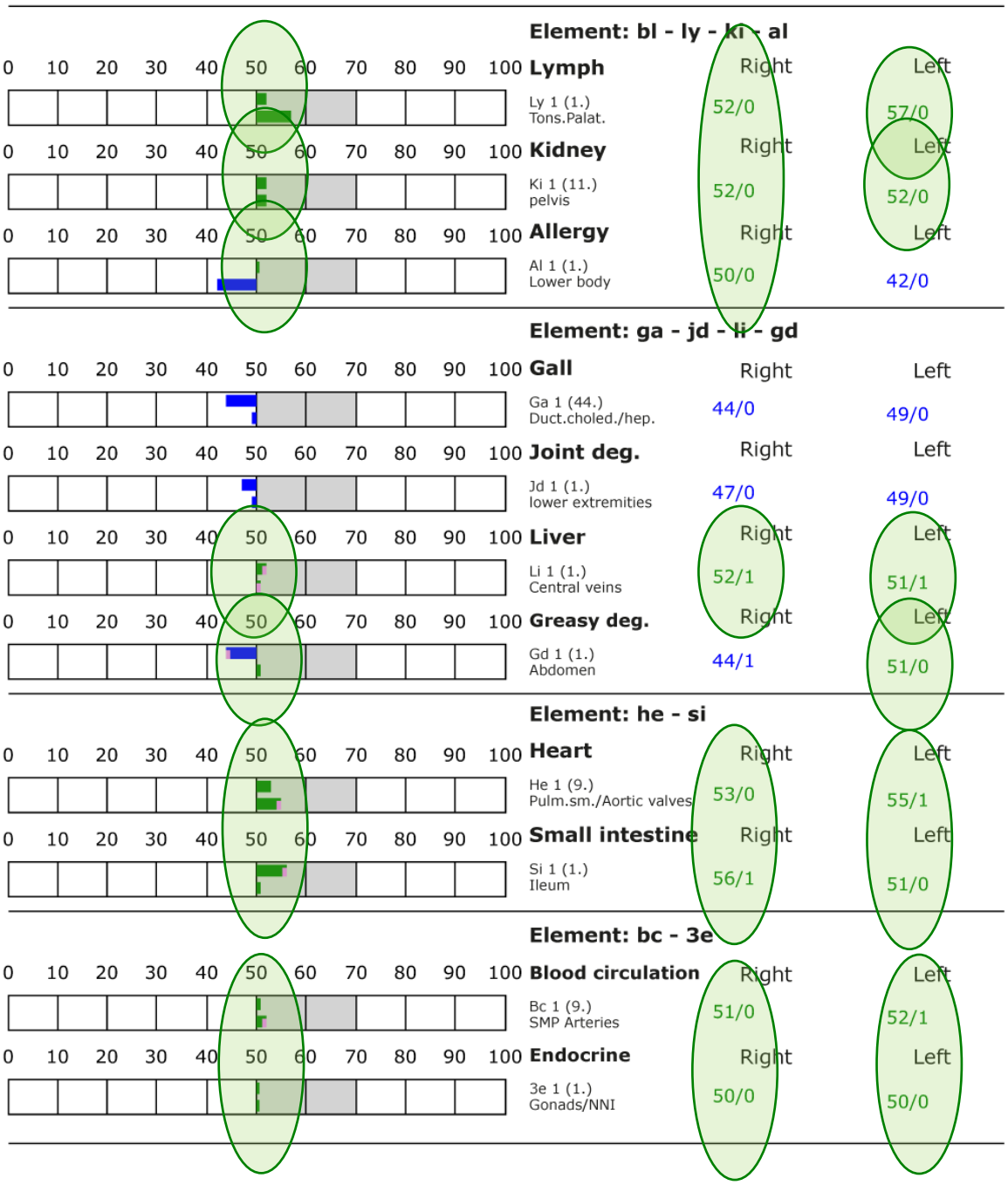

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BESA 4 testing BEFORE

BESA 4 testing BEFORE; testing for the presence of stress factors from the BESA 3 testing BEFORE as well as graphene oxide

In the further course of the BESA test, the subject was tested with the nosode (digitized graphene oxide) for the "presence of graphene oxide" in connection with the stress factors from the BESA 3 test BEFORE in the respective control circuits of the organism (measuring points). For this purpose, all the mentioned digitized loading factors from BESA 3 testing BEFORE were introduced together into the measuring circuit and tested at the specific measuring points. All BESA tests were again performed as in BESA 1, and 3 testing BEFORE at the TING points (40 nail fold points on the fingers and toes).

Goal: Determination of the reaction in the meridian system of the test person to the question, in which control circuit (measuring points of the respective organs) the mentioned stress factors can be tested energy-informatively.

ATTENTION: this BESA test, like the BESA 3 test BEFORE, does not represent a status measurement like BESA 1 and 2 test BASIC. Unlike the **"BESA status test"**, this BESA test is about "measuring a reaction" to the nosodes (digitized graphene oxide) in the measuring circuit. We speak of a **"BESA resonance test"**.
One could also equate this with a question: Do the digitized nosodes in the measuring circuit show resonance at the acupuncture points indicated? It is important to understand that in the "case of the presence of graphene oxide" at the respective measuring points, the "measuring answer" must show a green measuring value (50-70Skt). The possible green measurement results confirm, so to speak, the resonance with the respective nosode from the measurement circuit. So according to the motto: "Yes, these substances are detectable in the respective organs tested".

BESA-test evaluation P50 2.1
from **29-11-2021 at 10:47 to 10:52** (5 minutes) page 24 to 27

result: ATTENTION, the bioenergy-informative measuring result from the "resonance testing" shows "over the representation of the green measuring values" a resonance behavior and thus the load of graphene oxide, spike proteins and virus fragments" within these tested organs.

ATTENTION: In this BESA test "only the green measured values" are decisive. The blue measured values play only a subordinate role in this case. The absence of the blue measured values or, in other words, the presence of exclusively green measured values shows that all bioenergy-informatively tested organs of the test person are loaded with these tested nosodes!

100 % in the green area

Conclution:

ATTENTION: "100% in the green range" means - 100% resonance behavior to the nosodes (digitized loading factors) located in the measuring circuit!

As the graphics show, after the introduction of the mentioned nosodes (digitized stress factors such as spike proteins, virus fragments and graphene oxide) into the measuring circuit of the test person, "all measured values are in the green range".

ATTENTION: The "green readings" are "positive" because they confirm the resonance with the nosodes and thus the presence of the tested substances. This is response of BESA resonance test (as resonance behavior) and thus confirms the presence of graphene oxide, spike proteins and virus fragments within these organs.

It may sound or seem paradoxical, but the organism of the subject is, as the BESA graphs show, permeated with the tested substances.

ATTENTION: Because of this, we refrain from a separate representation (delimitation) of the measurement results with green circles. The representation of the BESA graphs in the green area confirms the presence of the indicated stress factors on the tested energy-informative control circuits of the test person.

For a detailed description of the spike proteins and virus fragments, please refer to pages 19 to 20.

For a simple explanation:

What is graphe oxid?

Graphene oxide is obtained from graphite by the action of strong oxidants. Graphene oxide is potentially used in biomedical applications to deliver drugs to target organs and cells via the bloodstream. Graphene oxide and cationic lipids (cationic lipids are used as lipid shells for drug delivery to the mRNA particles of COVID vaccines) have toxic effects.

Graphene is 200 times stronger than steel, 1 million times lighter than paper and transparent, thus information about laser light or similar beams can be transported optimally and mega fast. Graphene conducts electricity and heat better than a diamond. It conducts 250 times better than silicon, so it is the new superchip.

Under an electron microscope, saliva can be used to see exactly how tiny white or silvery spots or particles move and crosslink with each other after the mRNA injection. Subsequently, they crystallize and branch into rectangular antenna-like structures.

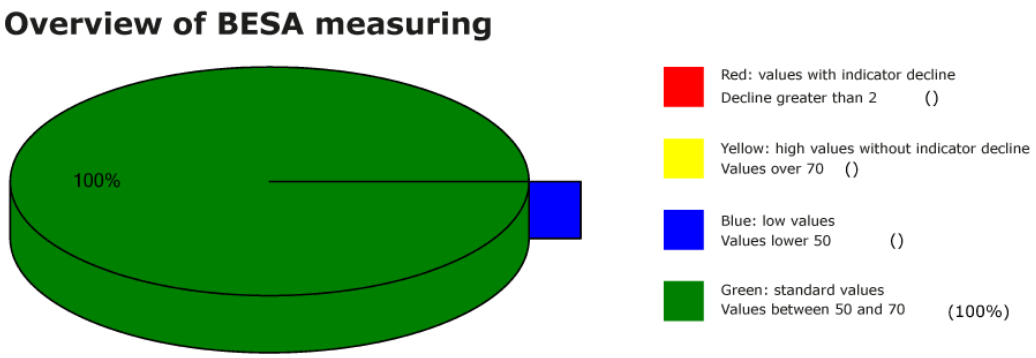
Several scientific works show that graphene oxide is used in gene therapy as a platform for delivering biomolecules such as mRNA into cells. The background lies in its high electrical conductivity and ability to penetrate cell membranes.

The crystalline and rectangular networks that form in the body fluid after mRNA vaccination and in the vaccine itself look like high-voltage antennas.

The "National Graphene Institute" is located in Manchester, where extensive research is being conducted. It was at this university that the existence of graphene antennas was first proven. It has been shown that graphene liquid crystals form spontaneously in the presence of an external magnetic field. Further scientific results showed that electric fields can change the crystal structure of graphene.

Graphene is considered a foundational technology for 5G by graphene manufacturers such as Grolltex, who are working on the future of graphene and 5G. Graphene has also been successfully linked to neurons and graphene-based neurotechnologies have been and continue to be the subject of intense research.

More details or scientific background on graphene can be found at the end of this detailed project.



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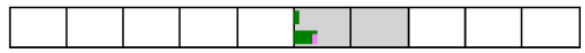
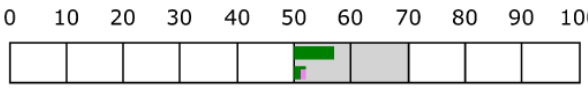
Standard values: (50-70 Skt.)

Element: lu - sk - li - ct



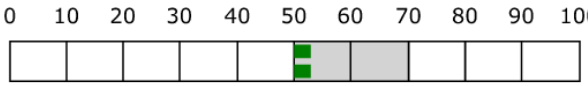
Lung — Right — Left

Lu 1 (11.) Parenchym — 51/0 — 54/1



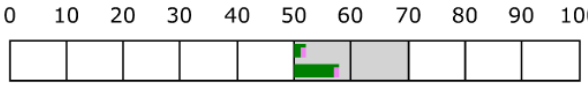
Skin — Right — Left

Sk 1 (1.) Lower Body — 57/0 — 52/1



Large intestine — Right — Left

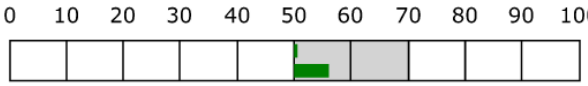
Li 1 (1.) Colon transv./sigm. — 53/0 — 53/0



Connective tissue — Right — Left

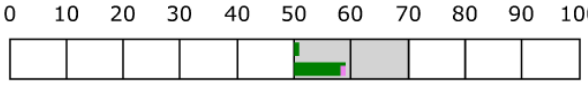
Ct 1 (1.) Abdomen — 52/1 — 58/1

Element: st - nd - ps - od



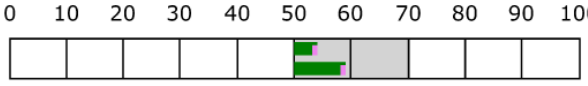
Stomach — Right — Left

St 1 (45.) Pylorus/body — 50/0 — 56/0



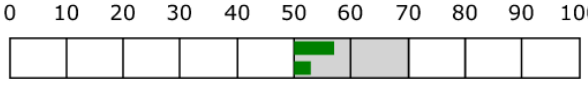
Nerves deg. — Right — Left

Nd 1 (1.) lumb./sakral. — 51/0 — 59/1



Pancreas-spleen — Right — Left

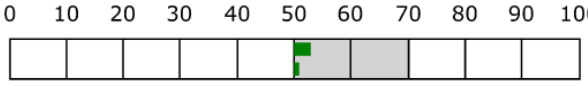
Ps 1 (1.) Eiw./w.Pulpa — 54/1 — 59/1



Organ deg. — Right — Left

Od 1 (1.) Abdominal region — 57/0 — 53/0

Element: bl - ly - ki - al



Bladder — Right — Left

BL 1 (67.) body — 53/0 — 51/0

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D: Degeneration (< 50 Skt.)

Standard values: (50-70 Skt.)

Element: bl - ly - ki - al

0 10 20 30 40 50 60 70 80 90 100		Right	Left
	Lymph		
	Ly 1 (1.) Tons.Palat.	60/1	55/0
	Kidney		
	Ki 1 (11.) pelvis	56/0	51/1
	Allergy		
	Al 1 (1.) Lower body	53/0	52/0

Element: ga - jd - li - gd

0 10 20 30 40 50 60 70 80 90 100		Right	Left
	Gall		
	Ga 1 (44.) Duct.choled./hep.	51/0	51/0
	Joint deg.		
	Jd 1 (1.) lower extremities	51/0	53/1
	Liver		
	Li 1 (1.) Central veins	52/0	58/1
	Greasy deg.		
	Gd 1 (1.) Abdomen	52/0	51/0

Element: he - si

0 10 20 30 40 50 60 70 80 90 100		Right	Left
	Heart		
	He 1 (9.) Pulm.sm./Aortic valves	57/0	52/0
	Small intestine		
	Si 1 (1.) Ileum	54/0	53/0

Element: bc - 3e

0 10 20 30 40 50 60 70 80 90 100		Right	Left
	Blood circulation		
	Bc 1 (9.) SMP Arteries	50/0	51/1
	Endocrine		
	3e 1 (1.) Gonads/NNI	54/1	52/1

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BESA 5 testing AFTER

BESA 5 testing AFTER with „Human Protect SD1“

In the further BESA test procedure, the test person was brought into contact with the test object, the "Human Protect SD1". This means that the test object was in close proximity to the subject.
ATTENTION: the nosodes with the digitized loading factors such as modified spike proteins, virus fragments and the graphene oxide are now no longer in the measurement circuit. In the BESA 3 and 4 tests BEFORE, the presence of the stress factors in the control circuit of the test person was already proven via the "BESA resonance tests!

All BESA tests were again performed as in the BESA 1, 3 and 4 tests at the TING points (40 nail fold points on the fingers and toes).

Goal: to determine the reaction of the subject's meridian system in connection with the test object. Can the "Human Protect SD1" neutralize the enormous influence of the tested stress factors on the bioenergy informative system of the test person? Determine the differences of BESA 5 testing AFTER compared to BESA 1, 2, 3 and 4 testing BEFORE.
ATTENTION: this BESA test is again a status measurement like the BESA 1 test BEFORE. In contrast to the **"BESA Resonance Test"**, this BESA test alone is only about "measuring the reaction" of the energy-informative system of the test person to the "Human Protect SD1"!

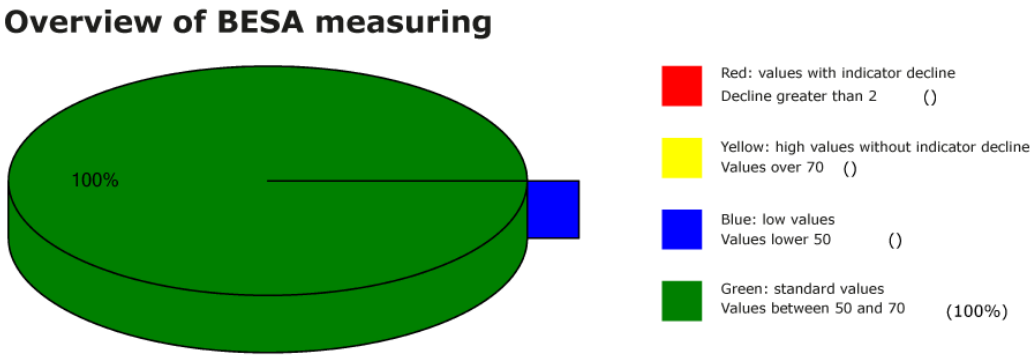
BESA-Testauswertung P50 2.1
vom **29-11-2021 um 10:53 bis 10:58** (5 Minuten) Seite 29 bis 31

result: The measurement result shows significant improvements at the meridian end points or at the energetic state of the test person.

100 % in the green area

Conclution: As the graphics show, 2 minutes after inserting the "Human Protect SD1" into the measuring circuit, all measuring points are in the green, optimal and harmonized range (balanced energy system). The BESA test results in a significant improvement of the energy situation in the meridian system of the test person compared to the BESA 1 and 2 tests BEFORE. All readings were at or just above 50 sct (50 to 54 sct). It can be seen that the "Human Protect SD1" is able in a very short time to give the necessary impulse for a harmonization (neutralization) into the life-promoting range to the severe stresses determined in the BESA 1 and 2 tests BEFORE by the stress factors tested, such as spike proteins, virus fragments and graphene oxide (see the green measured values as an expression of the positive measurement response). The comparisons of the BESA graphs confirm the change and harmonization of the stress factors on the meridian system of the test person.

These BESA 5 tests show that the test object, the "Human Protect SD1" is basically able and suitable to produce a harmonization of the stressing information (of the spike proteins, virus fragments and graphene oxide tested in the organism of the test person) up to the deeper structures of the cell.



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BESA basic test

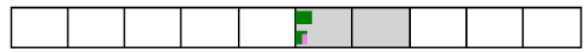
+++: Indicator decline > 15 Skt.
++: Indicator decline 6-15 Skt.
+: Indicator decline 3-5 Skt.

T: Total inflammation (>89 Skt.)
P: Partial inflammation (70-89 Skt.)

D: Degeneration (< 50 Skt.)

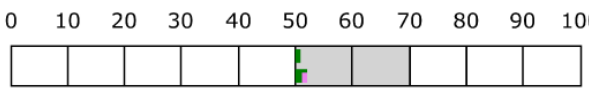
Standard values: (50-70 Skt.)

Element: lu - sk - li - ct



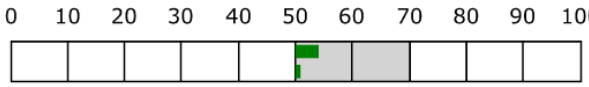
Lung — Right — Left

Lu 1 (11.) Parenchym — 53/0 — 52/1



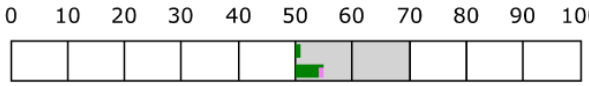
Skin — Right — Left

Sk 1 (1.) Lower Body — 51/0 — 52/1



Large intestine — Right — Left

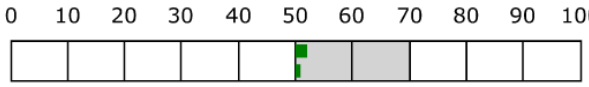
Li 1 (1.) Colon transv./sigm. — 54/0 — 51/0



Connective tissue — Right — Left

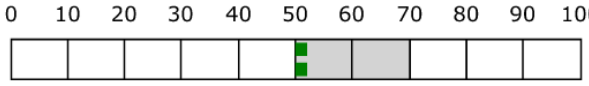
Ct 1 (1.) Abdomen — 51/0 — 55/1

Element: st - nd - ps - od



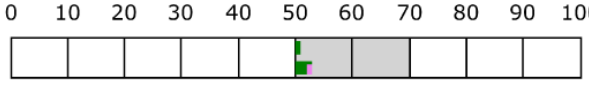
Stomach — Right — Left

St 1 (45.) Pylorus/body — 52/0 — 51/0



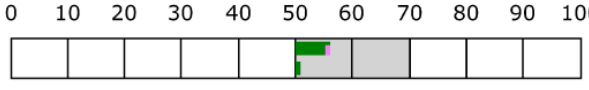
Nerves deg. — Right — Left

Nd 1 (1.) lumb./sakral. — 52/0 — 52/0



Pancreas-spleen — Right — Left

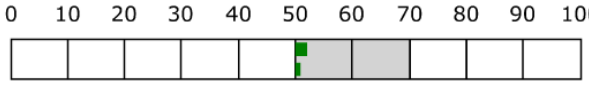
Ps 1 (1.) Eiw./w.Pulpa — 51/0 — 53/1



Organ deg. — Right — Left

Od 1 (1.) Abdominal region — 56/1 — 51/0

Element: bl - ly - ki - al



Bladder — Right — Left

BL 1 (67.) body — 52/0 — 51/0

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Standard values: (50-70 Skt.)

Element: bl - ly - ki - al

	Right	Left
Lymph		
Ly 1 (1.) Tons.Palat.	53/0	51/0
Kidney		
Ki 1 (11.) pelvis	53/1	51/0
Allergy		
Al 1 (1.) Lower body	53/1	53/0

Element: ga - jd - li - gd

	Right	Left
Gall		
Ga 1 (44.) Duct.choled./hep.	51/0	52/0
Joint deg.		
Jd 1 (1.) lower extremities	51/1	54/1
Liver		
Li 1 (1.) Central veins	52/0	51/0
Greasy deg.		
Gd 1 (1.) Abdomen	52/1	50/0

Element: he - si

	Right	Left
Heart		
He 1 (9.) Pulm.sm./Aortic valves	52/1	54/1
Small intestine		
Si 1 (1.) Ileum	53/1	52/0

Element: bc - 3e

	Right	Left
Blood circulation		
Bc 1 (9.) SMP Arteries	52/0	52/0
Endocrine		
3e 1 (1.) Gonads/NNI	51/0	54/0

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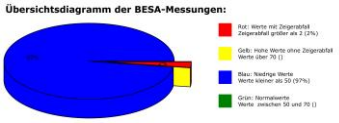
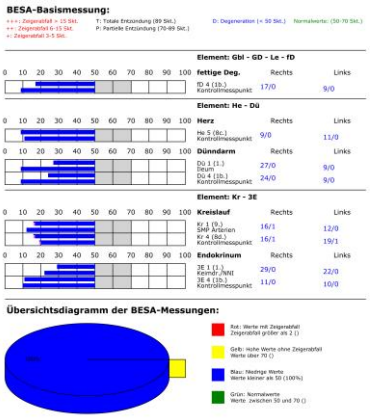
The results of the BESA tests at a glance

Respondent

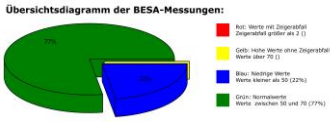
BESA 1 testing BASIC BEFORE as energetic status



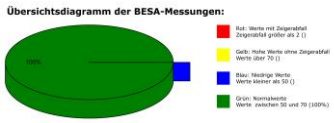
BESA 2 testing BASIC BREVIS BEVORE as 2nd energetic status



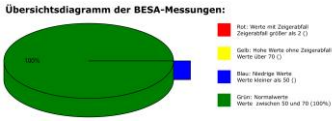
BESA 3 testing BEFORE with spike-proteins and virus fragments



BESA 4 testing BASIC BEFORE with spike-proteins, virus-fragments and graphen oxid



BESA 5 testing BASIC BEFORE with the test object, the „Human Protect SD1“



Please note the respective explanations of the BESA graphs following the BESA tests BEFORE and AFTER to avoid misunderstandings in the interpretation of the measurements.

General information about the test result

Man represents a kind of receiving antenna for environmental information. This is because human life depends fundamentally and exclusively on environmental information. Our organism is biologically very sensitive where natural information (fields) are located or where this natural information is subject to interactions and fluctuations. The situation is even more dangerous when such fields of building structures are introduced (injected) into the body as nanoparticles via a so-called mRNA vaccine. **For this reason, detected informative electromagnetic interference fields are biologically highly relevant. Any reduction or transformation of these interference fields (ideally to 100 percent) is biologically very important, in the case of these tested vaccines possibly even life-decisive.** These information loads from the vaccine as well as from our artificial environment are only compatible with life if they can be readjusted to a natural fluctuation tolerance. Disturbances, problems,

blockades, disharmonies in the biological control circuit of man find their causes in such disturbing electromagnetic information influences.

Neutralizing or harmonizing effects could be proven in this project P50 2.1 to determine the effect of the test object, the "Human Protect SD1". The "Human Protect SD1" was able to neutralize the biologically detrimental effects and effects of the stress factors tested on the test subject.

Due to the ingredients in the Corona mRNA and vector-based vaccines, a completely new situation and challenge has been created for the human immune system. It will occupy the prevailing science for decades to come to find solutions to the resulting questions. Many solutions and answers already exist. They offer an important approach in understanding what is going on at the human level right now.

But the biggest challenge is facing those people who already have pre-existing health conditions and are taking medications. This is because no one can imagine what effects these drugs will produce when combined with mRNA and/or vector-based vaccines. Many question marks have already arisen within this detailed project. It will take more projects to resolve them.

But critical to this detail project is the definitive ability of the test subject, the "Human Protect SD1" to neutralize and harmonize the severe strain factors tested in this P50 2.1 project. The transformation of the tested stress factors into bioenergy information with biological and life-promoting goodness is proven with this project.

Authorized summary

The BESA tests carried out by IFVBESA on the energetic and physical effectiveness of the test object, the "Human Protect SD1", have clearly shown that this test object is able to neutralize or harmonize biologically significant stress factors such as spike proteins, virus fragments and graphene oxide at the acupuncture points of the test person. Via the bioenergy-informative system analysis, the effect of the above-mentioned stress factors on the test person, his meridian system and his energetic-biological control circuits was questioned and systemically tested on the energy-informative level. The BESA tests BEFORE - AFTER show significant changes at the tested acupuncture points on the meridian system of the test person. The measurement data as well as their key figures impressively confirm on the one hand the stresses that the tested factors cause on the human organism, and on the other hand clarify how the deregulating energies are transformed into body-immanent and biocompatible energies after application of the test object, the "Human Protect SD1".

From a holistic point of view, it can be assumed that the positive effect on the test subjects also occurs in other people. That the positive influence by the "Human Protect SD1" is actually possible with high precision is clearly shown by this test through the BESA-PRE-AFTER-AFTER comparison. All measured values improved significantly from the mostly 100% blue measuring

range to the green, mostly 50 Skt. range (Skt = scale value), i.e. the range of optimal measured values. This means that an optimal regulation dynamic has taken place. Here, in the sense of the IFVBESA, one can clearly speak of an optimal, significant improvement of the body's own energy situation.

Result

The test person was brought into contact with the mentioned, digitalized ingredients of an mRNA vaccine during each BESA BEFORE test in order to prove its presence in the organism. In contrast to the BESA BEFORE tests, in which the test object, the "Human Protect SD1", was not used, consistently positive measurement results were found, which indicate that neutralization or harmonization has taken place. The regulation dynamics developed into an optimal effective range.

By demonstrating the energy informative effectiveness of the test object "Human Protect SD1" in this project P50 2.1, the requirements for obtaining a BESA seal of approval by the International Professional Association for BESA were met.

Appendix - List of sources

at spike-proteins and virus-fragments:
Zombies and Mutants. HORRIFYING new research reveals how vaccines suppress DNA repair mechanism in your cells
https://www.brighteon.com/cb351cd3-6c94-4f2b-a05d-bbda757d4472

https://www.naturalnews.com/2021-11-02-science-horror-vaccine-spike-protein-enters-cell-nuclei-suppresses-dna-repair-engine-of-the-human-body-cancer-aging.html#

https://www.naturalnews.com/files/viruses-13-02056-v2.pdf

Non-Homologous End Joining
NHEJ is an error-prone repair pathway that can occur throughout all cell cycle phases, whereas HR is an error-free pathway that predominantly occurs in late S and G2 phases.

https://www.sciencedirect.com/topics/biochemistry-genetics-and-molecular-biology/non-homologous-end-joining

at graphen oxid:
https://www.extremetech.com/extreme/150121-magnetogenetics-a-new-technique-to-control-the-inner-workings-of-human-cells-and-build-neural-circuits
https://www.theguardian.com/science/neurophilosophy/2016/mar/24/magneto-remotely-controls-brain-and-behaviour

"Graphene-based "neuromodulation" technology is REAL: Brain-controlling biocircuits with AI-driven graphene" July 21, 2021 In "General"

DNA as data storage: new Microsoft technology is part of Covid-19 vaccines June 9, 2021 In "General"

In Biontech-Pfizer's mRNA vaccine, 94-96% graphene oxide was found to be an ingredient.
https://www.businessinsider.com/military-sensor-skin-covid-darpa-19-2021-4

Definitions of COVID vaccines by Dr. Sherri Tenpenny; Cleveland, - 2021. www.drTenpenny.com

REF: Classen JB. COVID-19 RNA-based vaccines and the risk of prion disease. Microbiol Infect Dis. 2021; 5(1):
https://scivisionpub.com/pdfs/covid19-rna-based-vaccines-and-the-risk-of-prion-disease-1503.pdf

Graphene toxicity
https://www.sciencedirect.com/topics/pharmacology-toxicology-and-pharmaceutical-science/graphene

Graphene in drug research
https://www.sciencedirect.com/science/article/abs/pii/S1359644617301733?via%3Dihub

(2) Reveal related to graphene in Pfizer mRNA vaccine.
https://everydayconcerned.net/2021/08/02/evidence-of-nano-graphene-oxide-go-poisoning-body-brain-in-covid-flu-vaccines-chem-trails-rainwater-saline-plus-pfizer-whistleblower-karen-kingston-confirms-go-in-pegylated-lipid-nano-in/

DARPA-funded microchip technology optimizes convalescent plasma therapy for COVID-19 patients | School of Medicine | University of California, Irvine
https://www.som.uci.edu/news_releases/Convalescent-plasma-emerging-pathogens.asp

Science-Fiction instead of test: Pentagon chip detects coronavirus
https://www.computerbild.de/artikel/cb-News-Internet-Pentagon-Chip-erkennt-Coronavirus-Infektion-30060881.html

PCR test
https://arrangement-group.de/horror-was-der-covid-pcr-test-in-ihrem-koerper-anrichtet-aktuelle-studie-in-krankenhaeusern/

graphen
https://odysee.com/@FreieMedien:d/Graphenoxid-und-Lungentoxizit%C3%A4t:f

Darpa and COVID 19
https://www.darpa.mil/work-with-us/covid-19

PIL Genozid - by PCR-Tests and FFP2-Masks (using Darpa-Hydrogel) - PIL Genocide study confirms the - StuDocu
https://www.studocu.com/de-at/document/universitat-salzburg/public-international-law/pil-genozid-by-pcr-tests-and-ffp2-masks-using-darpa-hydrogel/15354125