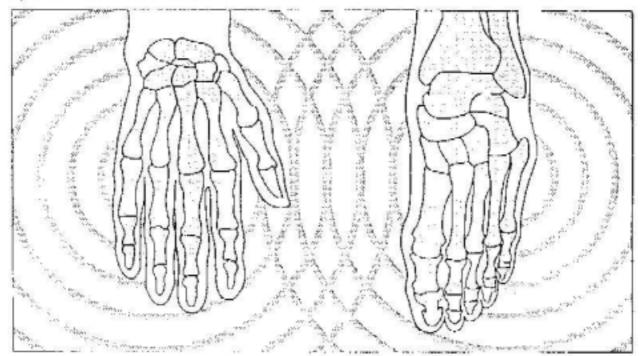
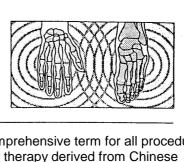
Electroacupuncture According to Voll

by W. John Diamond, M.B., B.Ch.

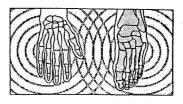




Biologische Heiknittel Heel GmbH Baden-Baden, Gutnany



History of electroacupuncture	Electroacupuncture is a comprehensive term for all procedures based on measurements or therapy derived from Chinese acupuncture, using modern electronics. The word "Electroacupuncture" was first coined by the French acupuncturist Dr. Roger de a Fuye in Paris. He combined an electrical device (Diathermopuncture) with stimulation of inserted needles for 1/8 to 2 seconds. Acupuncture analgesia is also referred to as electroacupuncture.
	In 1953 Werner and Voll developed an instrument for applying electroacupuncture to the skin without the use of needles. The instrument (named the K + F Diatherapuncteur) was able to locate acupuncture points, render them electrically measurable, enable electric treatment to be given and then to be remeasured electrically to assess the effect of the treatment.
	Voll adopted the system of meridians as described in classical acupuncture. These are energy tines accessed by superficial points on the body's surface, each one relating to the energy status of an internal organ.
Electrical measurement at acupuncture points	Topographically, the measuring points are located with reference to certain fixed osseous points - usually the transition zone between the shaft and capitulum. or in the interstices of myofascial planes at the margins of muscles and tendons. The actual point is located on the skin 2-3mm deep in the corium or subcutaneous tissue. The resistance of the corium and subcutaneous tissue is quite high (1 megohm) and in order to measure the points with some degree of reproducibility a state of maximum artifact is induced by applying sufficient pressure over the point. The instrument also has to have s high internal resistance, since the bio-electric energy o~ the acupuncture point is very small. The instrument also has to yield a current polarized in the opposite electrical charge to that of the body. The current must be small or it will distort the measurement and change the physiology of the body. The positive pole is usually the stylus. EAV diagnosis is based on sending direct current obtained from a current source with a high internal resistance through the body using an exactly calibrated value derived from a phantom. Measurement carried out by this direct current as stimulus is a functional test lust strong enough to provoke a healthy body to bring about an equilibrium between the stimulus and its own reaction (stable value).



Clinical measurements

The hand electrode is placed in the patient's hand, while the stylus is stroked to the point of maximum deflection and then a measurement pressure is put on the acupoint. The indicator will travel upwards to a maximum which it either holds or will tend to drop back towards a lower value. This dropping back to a lower value is called an Indicator Drop (ID) and represents the most important reading in EAV. All IDs must be treated first.

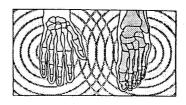
Four consecutive points on each meridian must be measured in order not to miss an ID on a single point. Individual points may be used to diagnose problems with specific parts of specific organs. The hand electrode should be held on the same side that is being measured when measuring the feet. A number of different results can be obtained from a reading of the acupoint:

- A balanced reading: Input voltage is matched by organ output voltage to give a reading of 50 (scale 0-100).
- An irritated or inflamed reading:
 > 60 due to excess discharge from the irritated organ, the intruding current exceeds the capacitance of the organ.
- A degenerative reading: Initial normal or inflamed reading followed by an indicator drop (ID). that shows degeneration of the involved organ. An indicator drop below 30 is serious.
- A low reading or no reading: Due to organ inability to respond at all, a case of advanced de gene ration.
- e. The rate of rise of the indicator gives an indication of the state of energy within the organ. The slower the rate of rise the less energy can be mobilized to balance the intruding current the organ is fatigued.

Precautions during measuring

Measurement of the entire conductance value (hand to hand) have to be made to assess the patient's suitability to ~e measured. If the patient has a reading >85, then he or she has too much sympathetic tone to be measured and needs to be discharged below 82. It the reading is <78, then the patient must be charged up to at least 80. Measurements of foot to foot and diagonal reading may delineate a single quadrant of the body that is the focus of a problem. This whole area may be addressed by

using a SEG or DFM. Balancing of the 4 quadrants can be used



as a basic treatment in many patients.

Measurement parameters

All measurements at acupuncture points should be at 50 with no indicator drops. The readings should only be taken if the hand to hand reading is 82, otherwise all indicator drops will not be seen. Measurement of the hypothalamus point at the root of the ear will indicate the presence of disturbing foci in the head, if the values are discrepant from side to side or if >82 on any side. These focal disturbances usually occur in the teeth, jaws, tonsils, paranasal sinuses or mastoid. These foci will involve the central switchboard functioning of the CNS, the hypothalamus.

Searching for foci

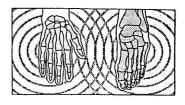
When the hypothalamus point indicates a focus, then the lymph meridian has to be measured to assess whether it is in the teeth and jaws, the sinuses or tonsil. The teeth can be. further assessed as to focal disease, galvanic currents or metal sensitivity. This can be done using nosodes and by electrically stressing each tooth and remeasuring Lymph 2 point. A raised reading after stimulation identifies that tooth as having a focus.

Medication testingMedication testing by Voll was a serendipitous finding. The
patient has all his or her acupoints measured and all abnormal
readings and indicator drops measured. Suitable medication
(homoeopathic. biological or herbal) in ampoules or bottle are
introduced into the measurement circuit by having the patient
hold the medication or inserting it into a honeycomb connected in
series with the hand electrode. If the medication now causes
disappearance of the ID or causes the reading of an abnormal
point to return to 50, then that is the correct medication. If
nosodes are used, then suitable drainage remedies should be
included to enable the body to excrete toxins.

Precautions for medication testing

Pathological oscillations may be transferred from the physician to the patient and render false readings. In order to mitigate this problem, the physician should undergo continuous treatment towards his or her own health and wear cotton gloves to minimize contact with the patient.

The occurrence of geopathic zones of interfering electromagnetic fields in the testing room will also cause false readings. Similar problems can occur with the 50 or 60 Herz oscillating fields of the electricity supply. One should avoid neon lights and CRTs in the



testing room. The floor should have an insulated covering and static electricity should be minimized.

The use of homotoxicology and EAV Homoeopathic remedies may be used to balance each disturbed point that is obtained in EAV measurement. Certain remedies are more likely to balance certain points than others. These remedy clusters are delineated by homoeopathic history of organotrophic effect (certain homoeopathics have known activity on certain organs) and by clinical experience in the general patient population and in individual patients. Combinations of certain remedies can clear the mesenchyme of the patient, detoxify him or her, and clear the homoeopathic field for constitutional therapy. Nosodes can release toxins from old disease and complete the clearing process.

Lymphatic points:

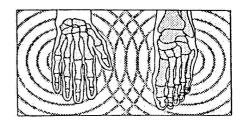
Firstly, we shall consider the organotrophic remedies at each EAV point, taking into consideration the clinical disease pictures characteristic of that point: e.g. use of bronchitis remedies at the lung point. Then we give some general combinations for use on the humoral and cellular phases of the Table of Homotoxicology.

Organotrophic remedies used In EAV Drainage - Lymphomyosot~

Ly 1 (palatine tonsils) Tonsilla compositum Ampullen Streptococcinum Galium-Heel Echinacea compositum Belladonna Homaccord Viral nosode

Lyl. 1 (auricular drainage) Salicyclic acid Mercurius-Heel S Viburcol

Ly2.O (teeth and laws) Osteoheel S Lamioflur Cruroheel S



Traumeel® S Granuloma dentis-Injeel

Ly3.0 (nose/paranasal sinuses) Naso-Heel* S Euphorbium compositum S Sinusitis Nosode-Injeel Mucosa compositum

Drainage - Echinacea compositum

CMP

Drosera-Homaccord® Aconitum-Homaccord® Bryaconeel® Bronchalis-Heel® Carbo compositum Droperteel® Engystol® N Gripp-Heel® Husteel® Tartephedreel® Viral and bacterial nosodes Asthma Nosode-Injeel

Drainage - Nux vomica-Homaccord*

CMP

Anacardium-Homaccord® Atropinum compositum Berberis-Homaccord® Diarrheel® S Hepeel® Heelax® S Podophyllum compositum Spascupreel® Veratrum-Homaccord® Vomitusheel®

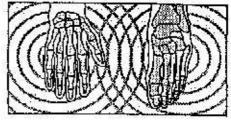
Drainage - Gelsemium-Homaccord®

CMP Chemical toxins Viral nosodes Engystol* N Thalamus compositum Cerebrum compositum

Lung points:

Large intestine:

Nerve degeneration:

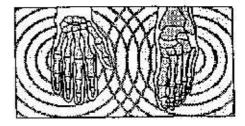


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	Circulation:	Dreinage - Circulo-Injeel*
10		CMP
		Placenta compositum
		Aesculus compositum
		Anglo-Injeel®
		Arteria-Heel [®]
		Barijodeel*
		Caclus compositum
	Allergy:	CMP
	GT 62	Galium Haal
		Allergen
		Liver or kidney remedies
	Organ degeneration:	CMP
	organ degeneration.	Graphites-Homaccord®
		Psorinoheel*
		(Somoneer
	Triple warmer:	TW1 (adrenal/ovary/testis)
		Hormeel [®] S Klimakt-Heel®
	15	Ovarium compositum
		Testis compositum Ampullen
5 2 0		TW2 (thyroid/parathyroids)
		Thyreoidea compositum Strumeel®
		Glonoin-Homaccord*
	Heart:	CMP
		Auðio-aulesi
20 <u>4</u>		Aurumheel* N-Tropfen
9		Cactus compositum
		Cor compositurn
		Cralonin*
		Cralaegus-Heel® S
		Strophantus compositum
	Small intestine:	Drainage - Veratrum-Homaccord®
		CMP
		Nux vomica-Homaccord [®]
		Duodenoheel [®]
		Erigotheel*
		Podophyllum compositum
		Anacardium-Homaccord*



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		50	
		Fatty degeneration:	
		raty degeneration.	Chemicals / pesticides
			Galium-Heel*
			Psorinoheel*
		Gall bladder:	CMP
			Chelidonium-Homaccord*
			Injeel-Chof*
			Berberls-Homaccord*
			Hepar compositum
3		Kidney:	CMP
		a sea service exc	Solidago compositum S
			Berberis-Homaccord®
			Cantharls compositum
			Populus compositum S
			Reneel*
		Bladder:	CMP
			Cantharis compositum S
			Berberis-Homaccord®
			Plantago-Homaccord®
		Bladder 65:	(Prostate/uterus)
			Sabal-Homaccord®
	2		Metro-Adnex-Injee!*
			Lamioflur*
	28 A		Gynäcoheel [»]
		General formulas	
	10	Humoral mixture:	Gripp-Heel*
	.		Traumeel* S
			Engystol* N
			Lymphomyosol*
		Cellular mixture:	Galium-Heel*
			Coenzyme compositum Ampullen
			Ublichinon compositum Ampullen
			Psorinoheel®
		Headaches:	Spigelon*
			Hepar compositum
			Spascupree!"

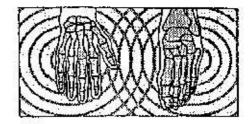


 	
Pancreas/Spleen:	Drainage - Leptandra compositum
	CMP
	Coopethur Hemacoord®
	Syzygium compositum
	Momordica compositum
Liver:	Drainage - Hepeel®
	CMP
	Hepar compositum
	Anacardium-Homaccord®
	Chelidonium-Homaccord*
	Leptandra compositum
	Injeet-Chol*
	Nux vomica-Homaccord®
	Phosphor-Homaccord®
r,	Salmonella/Hepatitis-Nosode-Injeel
Joint:	Drainage - Zeel*
	Traumeel* S
	Rhododendronce!* S
	Rhoumahee!*
	Discus compositum Ampullen
a.	Kalmia compositum
	Neuralgo Rheum-Injeel*
	Coinadul
	Cimiciluga-Homaccord*
	Colocynthis-Homaccord*
Stomach:	Drainage - Gastricumeel*
	Erigotheel*
	Mucosa compositum
	Duodenoheel*
Fibroid degeneration:	CMP
The bid began a and	Graphiles-Homaccord*
· Skin:	Drainage - Schwel-Heel*
	CMP
	Psorinoheel*
	Mezereum-Homaccord*
	Sulfur-Heel*
8	Graphites-Homaccord*
	Cutis compositum
21	Apis-Homaccord*
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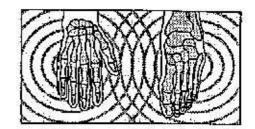
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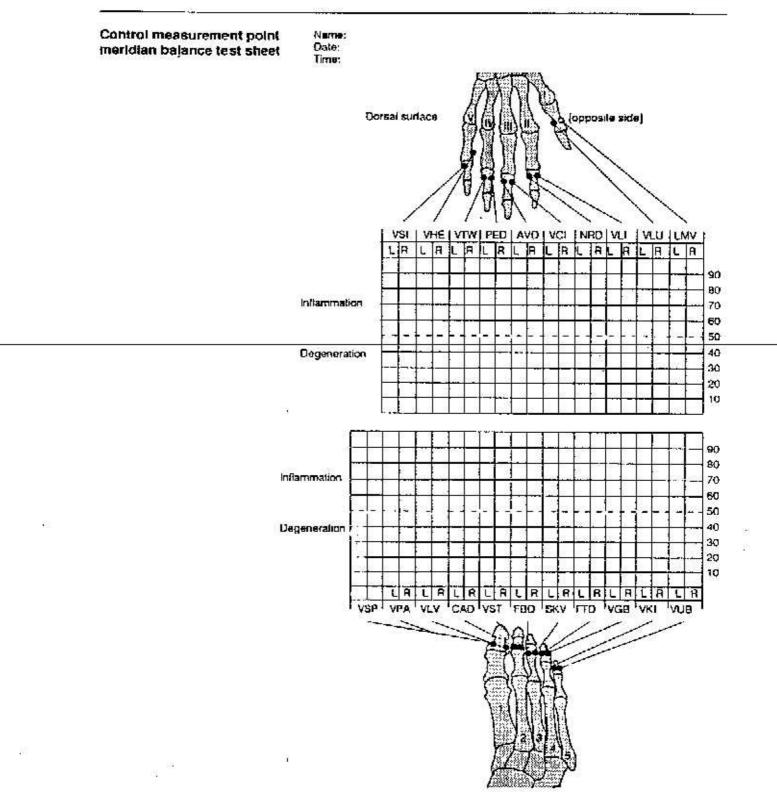
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	Colic:	Spascupreef*
		Nux vomica-Homaccord®
•		Atropinum compositum
	Constipation:	Hepes! ^o
		Heelax*S
	Haemorrhage:	Cinnamomum-Homaccord® N
		Phosphor-Homaccord [®]
		2.
	Laryngilis:	Phosphor-Homaccord®
	2. 1 3 0	20
	Haemorrhoids:	Hamamelis-Salbe-Heel
		Aesculus-Heel [®]
	a.	Paeonia-Heel*
	20	
	Anxiety:	Nervoheel*
	102800000000000000000000000000000000000	Valerianaheel®
	Hypertension:	Melliotus-Homaccord*
		Rauwollia compositum
	Vertigo:	Vertigoheel
	16-54 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 - 1990 -	
	Vomiting:	Viburco!*
		Vomitusheei™
		문제
	Obesity:	Graphites-Homaccord*
		Nux vomica-Homaccord®
		Thyreoidea compositum
	Oedema:	Apis-Horabecord*
14		 Lymphomyosot[®]
		Solidago compositum S
	Sprains/strains:	Traumeel* S
		Amica-Heel*

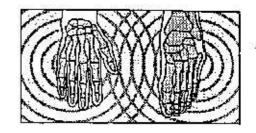
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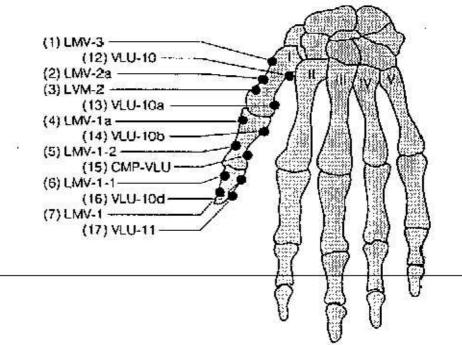


Reference chart 1: Lung meridian and lymph vessel

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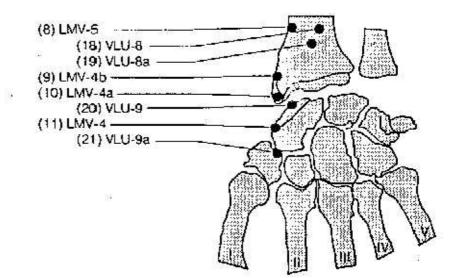
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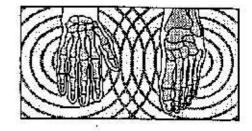


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The paimar aspect.

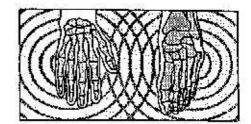


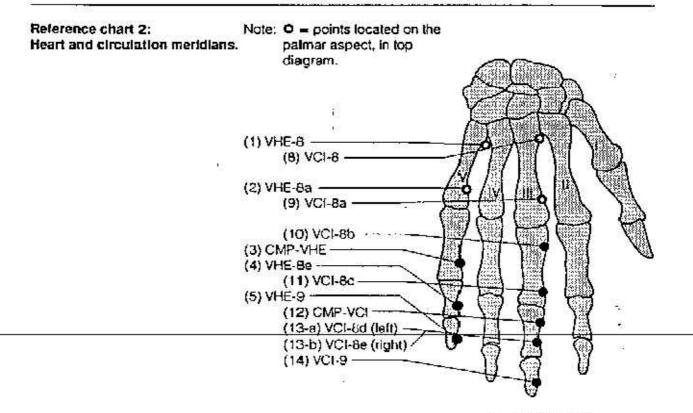
The paimar aspect.



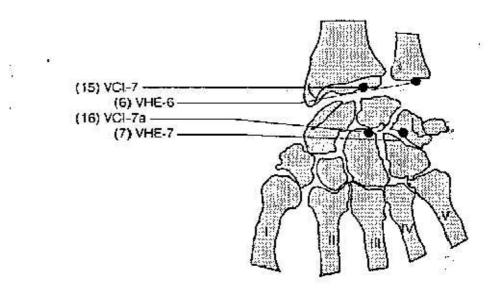
Reference chart 1:	(1) LM	W-3: MP. Lymph drainage of the nose + paranesei
Point names		sínuses.
	(2) LM	IV-2a: MP. Lymph drainage of the eye.
	(3) LM	W-2: MP. Lymph drainage for the upper + lower jaw.
222	(4) LM	W-1a: MP. Tubal tonsil (lateral lymphatic duct).
	(5) LM	IW-1-2: CMP. Five lonsils of the lymphatic pharyngeal ring.
	(6) LM	IV-1-1: MP. Auricular lymph drainage.
	(7) LM	W-1: MP. Palatine tonsil.
	(8) LM	IV-5: MP. Lymph vessels of the heart,
	(9) LM	IV-4b: MP. Lymph drainage of the larynx +
		hypopharync.
	(10) LM	IV-4a: MP. Lymph vessels of the esophagus.
	(11) LM	IV-4: MP. Lymph vessels of the lungs,
	(12) VL	U-10: MP. Bronchi.
	(13) VL	U-10a: MP. Pleura.
	(14) VL	U-10b: MP. Bronchioles.
	(15) CN	AP-VLU: CMP. Lower respiratory passages.
	(16) VL	U-10d; MP. Mediastinal plexus.
	(17) VL	U-11: MP. Alveoli.
	(18) VL	U-8: MP. Veins of the upper extremitiy.
	, (19) VL	
	(20) VL	
	(21) VL	

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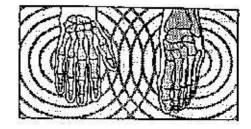




The dorsal aspect.

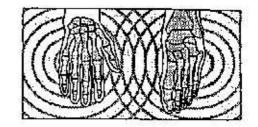


The palmar aspect.



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Reference chart 2:	(1)	VHE-8.	MP. Mitral (bicuspid) and tricuspid valves.
Polnt names	(2)	VHE-8a:	MP. Pericardium.
	(3)	CMP-VHE;	CMP. Heart.
	(4)	VHE-8e:	MP. Cardiac plexus.
	(5)	VHE-9:	MP. Aortic and pulmonary valves.
	(6)	VHE-6:	MP. Myocardium.
	(7)	VHE-7:	MP. Heart conduction system, atrioventricular bundle (bundle of HIS),
	(8)	VCI-8:	SMP. Venous functions,
	(9)	VCI-8a:	MP. thoracic duct.
	(10)	VCI-8b:	MP. Cisterna chyli.
	(22)	VCI-8e:	 a) MP. Abdominal aortic plexus.
			 b) MP. Abdominal aorta.
	(12)	CMP-VCI:	CMP. Arterial, venous, and lymphatic vascular system.
	(13-a)	VCI-8d:	(left side) MP. Thoracic aortic plexus.
	(13-6)	VCI-8e:	(right side) MP. Cardiac ganglia.
	(14)	VCI-9:	SMP. Arterial functions.
<i>x</i>	(15)	VCI-7:	SMP. Coronary vessels.
	(16)	VCE-7a:	MP. Coronay plexus of the heart.



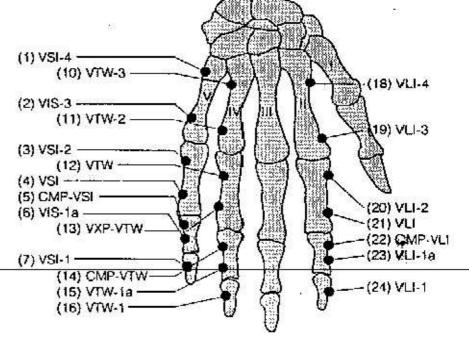
Reference chart 3: Large intestine, small intestine, and triple-warmer (endocrine) meridians.

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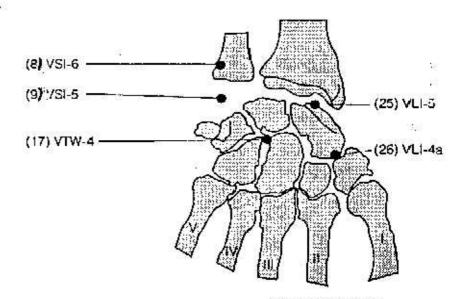
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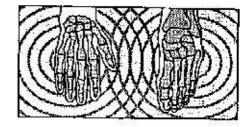
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The dorsal aspect.



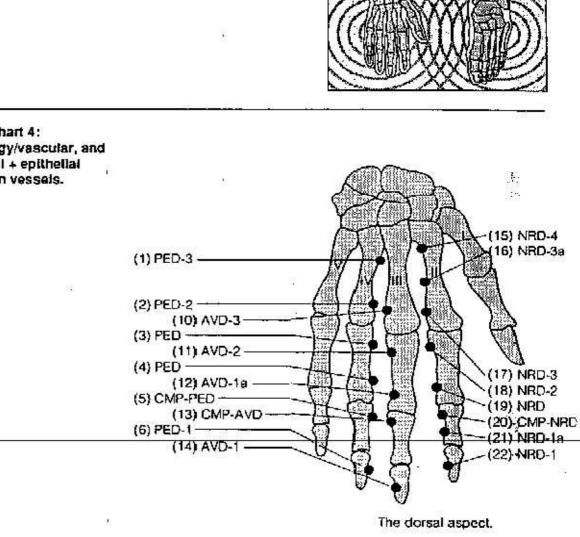
The dorsal aspect.

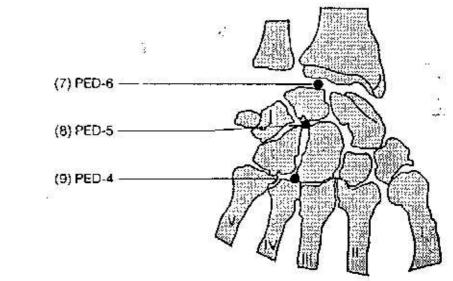


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Reference chart 3:	(1) VSI-4:	MP. Superior (1st) and ascending (4th) parts of
Point names		the duodenum.
	(2) VSI-3:	MP: Descending (2nd) part of the duodoenum
		and duodeojejunal flexure.
	(3) VSI-2.	MP. Horizontal (3rd) part of the duodenum and jejunum.
	(4) VSI:	MP. Peritoneum of the small intestine.
	(5) CMP-VSI:	CMP. Small intestine.
	(6) VSI-1a:	MP. Upper and lower mesenteric plexuses.
	(7) VSI-1:	MP. fleum.
相	(B) VSI-6;	MP. Cervical spine.
)- -	(9) VSI-5:	MP. Ulnar portion of the wrist joint or
		interarticular disc.
	(10) VTW-3:	SMP. Pineal and plluitary glands.
	(11) VTW-2:	SMP. Thymus, thyroid, parathyrold glands.
	(12) VTW:	MP. Mammary gland.
	(13) VXP-VTW,	MP. Incretogenic function (insulin-glucagon) of the pancreas.
	(14) CMP-VTW;	
	(15) VTW-1a;	SMP. Unllateral cervical parts of the
	1963 Fabric 1968 - 96862	Sympathetic nerve.
	(16) VTW-1:	SMP. Gonad and adrenal glands.
	(17) VTW-4:	MP. Distal hand joint.
	(18) VLI-4:	MP. Cecum and transverse colon.
	(19) VLI-3:	MP. Ascending colon and splenic llexure.
	(20) VLI-2:	MP. Descending colon and hepatic llexure.
	(21) VLI:	MP. Periloneum of the large intestine.
	(22) CMP-VLI;	CMP. Large Intestine.
	(23) VLI-1a:	MP. Illac and upper hypogastric plexuses.
	(24) VLI-1:	MP. Transverse colon and sigmoid.
	(25) VLI-5:	MP. Radial portion of the wrist joint,
	(26) VLI-4a:	MP. Appendix and lloogecal (mesenterial)
-e ⁻¹	(lymph nodes.
123		iympi nodes.
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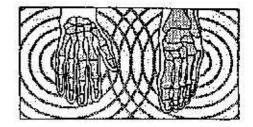


The dorsal aspect.

Heterence chart 4: Nerval, allergy/vascular, and parenchymal + epithellal degeneration vessels.

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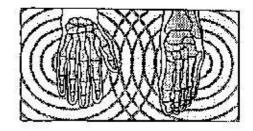


Reference chart 4: Point names,	(1) PED-3:	MP. Parenchymal and epithelial degeneration, organs of the head.
· vin hunch	(2) PEO-2:	MP. Parenchymal and epithelial degeneration, organs of the chest and neck.
	(3) PED:	MP. Degenerative processes in the entire pleura.
	(4) PED:	MP. Degenerative processes in the entire peritoneum.
	(5) CMP-PEC	승규는 것이 같아요. 이번 것이 같아요. 이번 것이 같아요.
	(6) PED-1:	MP. Parenchymal and epithelial degeneration. organs of the abdomen and pelvis.
	(7) PED-6:	MP. Parenchymal and epithelial degeneration, organs of the head.
	(8) PED-5:	MP. Parenchymal and epithelial degeneration, organs of the chest and neck.
	(9) PED-4:	MP. Parenchymal and epithelial degeneration, organs of the abdomen and pelvis.
	(10) AVD-3:	MP. Allergy/vascular degeneration, head region.
	(11) AVD-2:	MP. Allergy/vascular degeneration, upper portions of the body.
	(12) AVD-1a:	MP. Vascular sclerosis.
	(13) CMP-AVE	
	(14) AVD-1:	MP. Allergy/vascular degeneration, lower portions of the body.
	(15) NRD-4:	CMP. Cranial nerves.
	(16) NRD-3a:	CMP. Parasympathelic cranial ganglia.
	(17) NRD-3:	MP. Nerval degeneration of the brain stem and cerebrum.
	(18) NRD-2;	MP. Nerval degeneration of the cervical and thoracic spinal marrow.
	(19) NRD:	MP. Meninges and spinal marrow.
	(an) ON IT KID!	
	а, (20) Сме-ынс	system.
	(21) NRD-1a:	SMP. Enlire autonomic nervous system.
	(22) NRD-1:	MP. Nerval degeneration of the lumbar and sacral spinal marrow.

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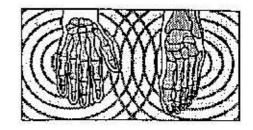


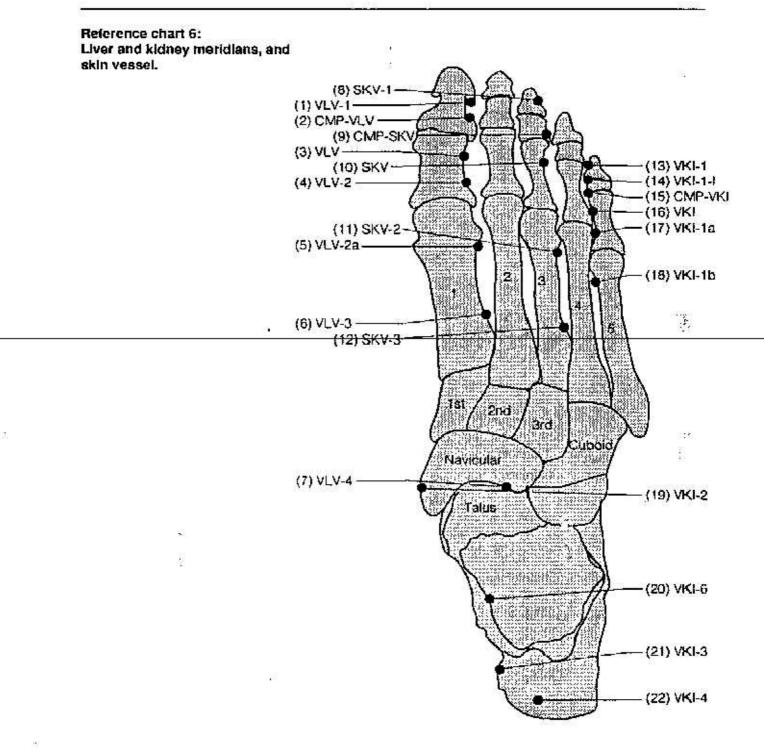


Reference chart 5:	(1a) VSP-1:	MP. White pulp of the spleen.	
Point names.	(1b) VPA-1:	MP. Protein metabolism.	
	(2a) CMP-VSP:	CMP, Spleen.	
52	(2b) CMP-VPA:	CMP. Pancreas,	
	(3a) VSP:	MP. Periloneum of the spleen.	
	(3b) VPA:	MP. Periloneum of the pancreas.	
	(4a) VSP-2:	MP. While pulp of the spleen.	
	(4b) VPA-2:	MP. Nucleoprotein metabolism.	
	(5a) VSP-3:	MP. Red pulp of the spleen.	
	(5b) VPA-3:	MP. Carbohydrate metabolism.	
	(6a) VSP-4.	MP. Reticuloendothelial system and splenic reticulum.	
	(6b) VPA-4;	MP. Lipid metabolism.	
	(7) VSP/PA-5:	MP. Medial portion of the talocrural joint.	
	(8) VUB-67:	MP. Corpus of the urinary bladder.	
	(9) VUB-66c:	MP. Vesical plexus.	
96	(10) CMP-VUB:	CMP. Urinary bladder and genitourinary organs.	
	(11) VUB:	MP. Peritoneum of the urinary bladder.	
	(12) VUB-66:	MP. Fundus, cervix, and sphincter of the	
		urinary bladder.	
	(13) VUB-65:	SMP. Prostate, seminal vesicle, seminal hitlock (colliculus), penis, and urethra in males or uterus, broad ligament, parametrium, vagina, and urethra in females.	
	(14) VUB-64:	SMP. Spermatic cord and epididymis in males or uterine (fallopian) tube in females.	
8	(15) VUB-63:	SMP. Lower hypogastric (pelvic) plexus.	

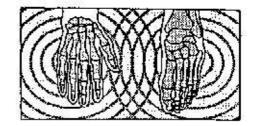
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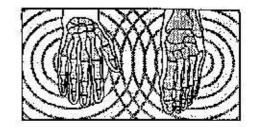
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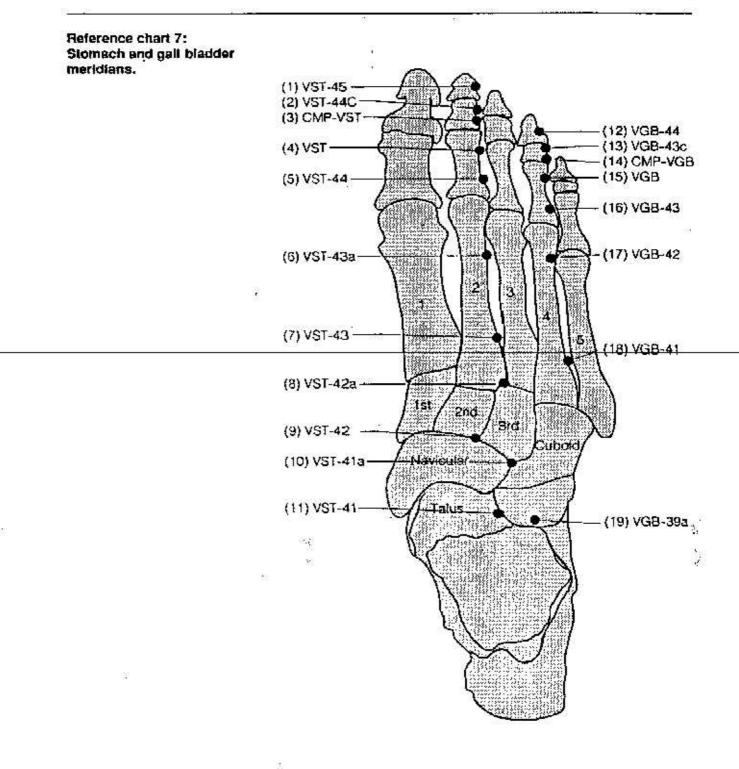


Reference chart 6:	(1) VLV-1:	MP. Central venous system of the liver.
Point names.	(2) CMP-VLV:	CMP, Liver,
	(3) VLV:	MP. Peritoneum of the liver.
	(4) VLV-2:	MP. Liver cell and lobular system.
	(5) VLV-2a:	MP. Interlobular ducts of the liver.
	(6) VLV-3:	MP. Perivascular system of the liver.
	(7) VLV-4:	MP. Talocalcaneonavicular joint.
	(8) SKV-1:	MP. Skin (lower portions of the body).
	(9) CMP-SKV:	CMP. Skin and scars of the skin.
	(10) SKV: .	Scars of the skin.
	(11) SKV-2:	MP. Skin (upper portions of the body).
	(12) SKV-3:	MP. Skin of the head.
	(13) VKI-1;	MP, Renal pelvis.
	(14) VKI-1-I:	MP. Renal plexus.
	(15) CMP-VKI:	CMP, Kidney and ureter.
	(16) VKI:	MP. Peritoneum of the kidneys.
	(17) VKI-1a:	MP. Ureter.
	(18) VKI-1b:	MP. Suprerenal plexus.
	(19) VKI-2;	MP. Pyelorenal boundary layer.
	(20) VKI-6:	MP. Rectum.
	(21) VKI-3:	MP. Renal Cortex.
	(22) VKI-4:	MP. Rectal (middle hemorrhoidal) plexu

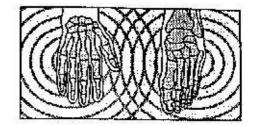
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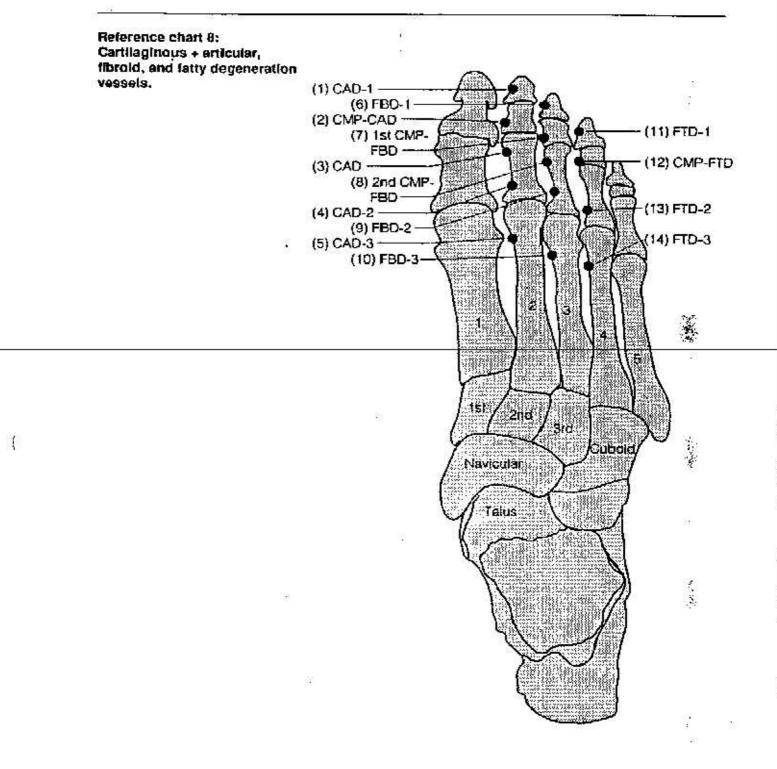
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Reference chart 7:	(1) VST-45:	MP. Corpus and pylorus/pyloric canal.
Point names.	(2) VST-44c:	SMP, Celiac (solar) plexus.
	(3) CMP-VST:	CMP. Slomach.
	(4) VST:	MP. Peritoneum of the Stomach.
	(5) VST-44:	MP. Fundus (vault) and pyloric antrum.
	(6) VST-43a:	MP. Gestric tract.
	(7) VST-43:	MP. Cardia and corpus of the stomach.
	(8) VST-42a:	MP, Esophagus (lower portion).
	(9) VST-42:	MP. Esophagus (upper portion).
	(10) VST-41a:	MP. Mammary gland.
	(11) VST-41:	MP. Lateral portion of the talocrural joint.
	(12) VGB-44:	MP. Common hepatic and bile ducts.
	(13) VGB-43c:	MP. Hepalic plexus.
	(14) CMP-VGB:	CMP. Biliary ducts and gall bladder.
	(15) VGB:	MP. Peritoneum of the gall bladder.
	(16) VGB-43:	MP. Right hepatic and cystic ducts.
	(17) VGB-42:	MP. Gall bladder (corpus) and left hepatic duct.
	(18) VGB-41:	MP, Biliary duct.
	(19) VGB-39a:	MP. Lower portion of the talocrural joint.

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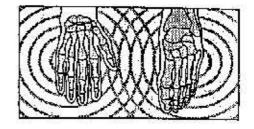
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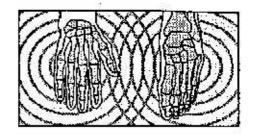


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Reference chart 8: (1) CAD-1: SMP. Joints of the pelvic girdle and lower Point names. extremity. (2) CMP-CAD: CMP. All joints including the spine. (3) CAD: MP. Synovial membranes for all joints. (4) CAD-2: SMP. Joints of the shoulder girdle and upper extremity. (5) CAD-3: SMP. Joints of the atlas, and]aw (temporomandibular). (6) FBD-1: MP. Fibroid degeneration, organs of the abdomen and minor pelvis. (7) 1st CMP-FBD:CMP. Fibroid degeneration of the entire body. (8) 2nd CMP-FBD: CMP. Fibroid degeneration of the mucous membranes. (9) FBD-2: MP. Fibroid degeneration, organs of the chest and neck. (10) FBD-3: MP. Fibrold degeneration, organs of the head. MP. Fatty degeneration, organs of the (11) FTD-1: abdomen. CMP. Fatty degeneration in the entire body. (12) CMP-FTD: MP. Fatty degeneration, organs of the chest. (13) FTD-2: (14) FTD-3: MP. Fally degeneration, organs and vessels of the head. 2



Reference chart 9: Urinary bladder meridian. Medius (1) VUB-27 -(2).VUB-32-(3) VUB-49a Gluteus (4) VUB-33 maximus (5) VUB-29 (6) VUB-34 -(7) VUB-35 · Gracilis (8) VUB-49b (9) VUB-49c-(10) VUB-49d Adductor (11) VUB-50 magnus (12) VUB-50-1 (13) VUB-50-2 (14) VUB-50alliotibiat band Ģ (15) VU8-50b --(8) (16) VUB-50c ---Semitendinosus Ż (17) VUB-51-Long head Biceps femo y. (18) VUB-51a Short head (19) VUB-52- ;-Semimembranosus 4 (20) VUB-54 Plantaris Medial head Gastroc-Inemius 3 Lateral head **Popliteal space**

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