





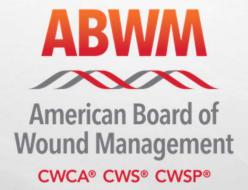






CERTIFICATION ELIGIBILITY

Separate Certifications for Different Levels of Providers & Others Examination for Certification CME Requirement Nationally Recognized



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CERTIFIED WOUND CARE ASSOCIATE® (CWCA®)

- HEALTHCARE PROFESSIONAL WITH AT LEAST 3 YEARS OF WOUND CARE EXPERIENCE
- CANDIDATES WITH A PROFESSIONAL LICENSE APPLYING FOR THE CWCA® MUST POSSESS A
 <u>FULL AND UNRESTRICTED PROFESSIONAL LICENSE</u> IN AT LEAST ONE STATE AND IN ALL STATES IN
 WHICH THE CANDIDATE CURRENTLY PRACTICES
 - EXAMPLES: LPNS, LVNS, PTAS, ALL CERTIFIED HEALTHCARE ASSISTANTS (CNAS, CMAS), HEALTHCARE
 ADMINISTRATORS, DIETITIANS, SALES AND MARKETING PROFESSIONALS, AND ACADEMIC RESEARCHERS

CERTIFIED WOUND SPECIALIST® (CWS®)

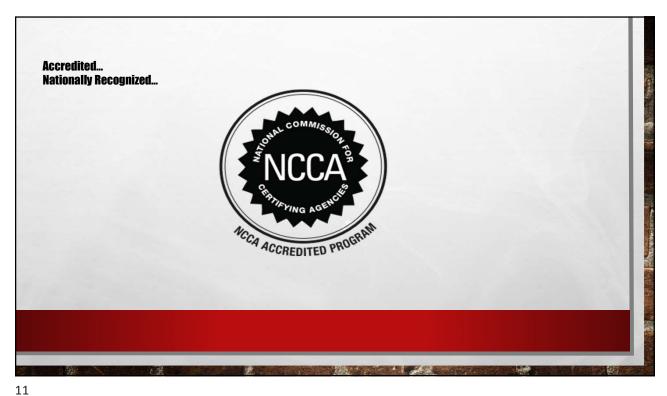
- ullet A Licensed Healthcare Professional with a Bachelor's. Master's, or doctoral degree and ullet or more years of clinical wound care experience
- A LICENSED RN WITH AN ASSOCIATE'S DEGREE AND 3 MORE YEARS OF CLINICAL WOUND CARE
 EXPERIENCE
- CANDIDATES MUST POSSESS A <u>FULL AND UNRESTRICTED PROFESSIONAL LICENSE</u> IN AT LEAST ONE STATE
 AND IN ALL STATES IN WHICH THE CANDIDATE CURRENTLY PRACTICES
 - EXAMPLES: REGISTERED NURSES, NURSE PRACTITIONERS, VETERINARY PHYSICIANS, PHYSICIAN ASSOCIATE, PHYSICAL THERAPISTS, OCCUPATIONAL THERAPISTS, PHARMACISTS, AND REGISTERED DIETITIANS

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CERTIFIED WOUND SPECIALIST PHYSICIAN® (CWSP®)

- MDS, DOS, AND DPMS WITH THREE OR MORE YEARS OF CLINICAL WOUND CARE EXPERIENCE
- CANDIDATES MUST POSSESS A <u>FULL AND UNRESTRICTED PROFESSIONAL LICENSE</u> IN AT LEAST ONE STATE
 AND IN ALL STATES IN WHICH THE CANDIDATE CURRENTLY PRACTICES

EXAMPLE: PHYSICIANS AND PODIATRIST



CWSP NATIONAL BOARD CERTIFICATION EXAMINATION

- EXAMINATIONS ARE DELIVERED BY COMPUTER AT 300+ PEARSON VUE TEST CENTERS THROUGHOUT AMERICA (APPOINTMENT ONLY)
- 180 COMPUTER-BASED, MULTIPLE-CHOICE OUESTIONS, GIVEN OVER A PERIOD OF THREE AND A HALF HOURS
- 84.4% PASS RATE
- SPECIFIC METHODOLOGIES FOR ESTABLISHING THE PASSING SCORE FOR EXAMS. AND ENSURING TEST FORMS ARE EQUIVALENT. WILL BE SELECTED BY A QUALIFIED PSYCHOMETRICIAN BASED ON COMPLIANCE WITH NATIONAL ACCREDITATION STANDARDS

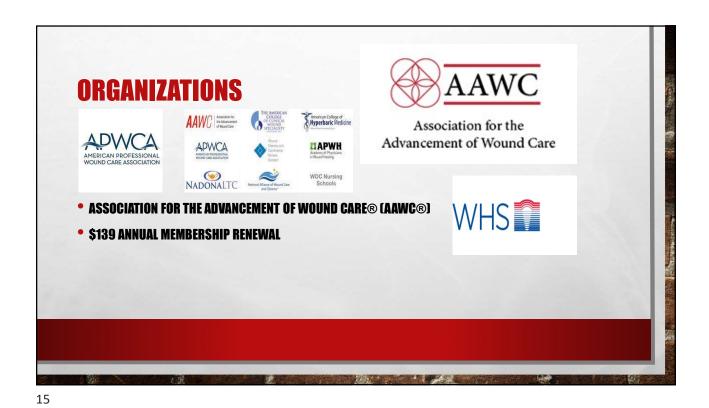
- \$150 ANNUAL RENEWAL FEE
- 10 YEAR RECERTIFICATION



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CONTINUING EDUCATION HOURS

- A MINIMUM OF SIX (6) HOURS OF CONTINUING EDUCATION IN THE FIELD OF WOUND CARE MANAGEMENT ARE REQUIRED EACH YEAR TO STAY CERTIFIED
- HTTPS://ABWMCERTIFIED.ORG









WOUND ASSESSMENT AND DOCUMENTATION

- TYPE OF WOUND:
 - ARTERIAL ULCERS & PERIPHERAL ARTERIAL DISEASE (PAD)
 - ADVANCED VENOUS INSUFFICIENCY & ULCERS
 - LYMPHEDEMA
 - DIABETES AND WOUNDS
 - RIIRNS
 - PRESSURE ULCERS
 - ATYPICAL
 - PRE-MALIGNANT AND MALIGNANT SKIN TUMORS
 - SURGICAL/TRAUMATIC

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WOUND SPECIFIC GRADE/ STAGING SYSTEMS

- NPUAP/ EPUAP STAGING/ GRADING OF PRESSURE ULCERS
- WAGNER SCALE FOR DIABETIC FOOT ULCERS (DFUS)
- UT DIABETIC WOUND CLASSIFICATION- DFUS
- CEAP FOR VENOUS LEG ULCERS
- TISSUE LOSS (PARTIAL/ FULL)- UNIVERSAL
- PAYNE-MARTIN & STAR FOR SKIN TEARS

WAGNER ULCER GRADE CLASSIFICATION DIABETIC ULCERS

Grade	Characteristics
0	Pre-ulcerative lesions, healed ulcers, neuropathy, presence of bony deformity
1	Superficial ulcer without subcutaneous tissue involvement, not infected
2	Penetration through the subcutaneous tissue; may expose bone, tendon, ligament, or joint capsule
3	Extensive ulceration with exposed bone and/or deep infection (i.e., osteomyelitis or abscess)
4	Gangrene of toes or forefoot
5	Gangrene of the whole foot requiring disarticulation

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THERMAL INJURY CLASSIFICATION

Classification According to Depth		
Superficial	1 st Degree	Epidermal Damage
Partial-Thickness Superfical Deep	2 nd Degree	Epidermis and Superficial (Papillary) Dermis Damage Damage Extends into the Reticular Dermis
Full-Thickness	3 rd Degree 4 th Degree	Complete Loss of Dermis Damage to Underlying Fascia, Muscle, Bone

PRESSURE ULCER CLASSIFICATION

Classification	Description
Stage I	Intact skin with non-blanchable redness of a localized area usually over a bony prominence
Stage II	Partial-thickness loss of dermis presenting as a shallow open ulcer with a red/ pink wound bed, without slough. May also present as an intact or open/ ruptured serum filled blister.
Stage III	Full-thickness tissue loss. Subcutaneous fat may be visible but bone, tendon, or muscle are not exposed.
Stage IV	Full-thickness tissue loss with exposed bone, tendon, or muscle.
Unstageable	Full thickness tissue loss in which actual depth of the ulcer is completely obscured by slough and/or eschar in the wound bed.

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VENOUS ULCER CLASSIFICATION (CEAP) CLINICAL PRESENTATION, ETIOLOGY, ANATOMIC DISTRIBUTION, PATHOPHYSIOLOGIC RASIS

Class	Description
0	No signs of venous disease
1	Telangiectasias
2	Varicose veins
3	Edema
4	Skin changes - C4a: Pigmentation, Eczema - C4b: Lipodermatosclerosis
5	Healed ulcer
6	Active ulceration

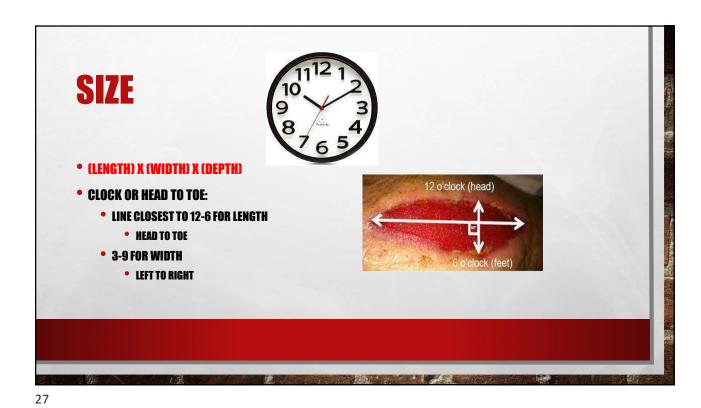
LOCATION

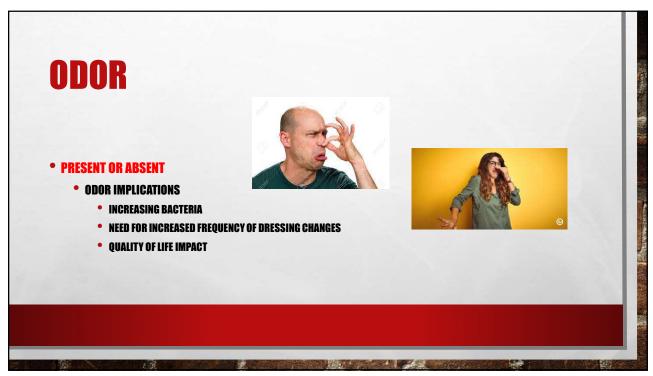
- DOCUMENT IN REFERENCE TO HEAD, FRONT OR BACK
- REFERENCE NEAREST BONE STRUCTURE
- COMMONLY USED TERMS
 - PROXIMAL, DISTAL
 - SUPERIOR, INFERIOR
 - MEDIAL, LATERAL
 - ANTERIOR, POSTERIOR
 - DORSAL, PLANTAR

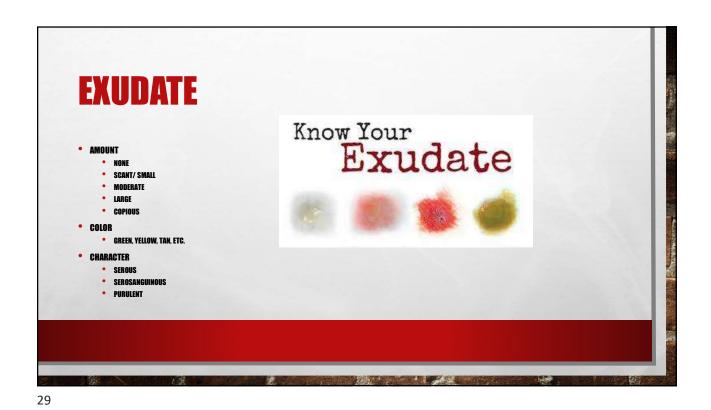
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DEGREE OF TISSUE DESTRUCTION

- PARTIAL-THICKNESS
 - LOSS OF EPIDERMIS AND DOWN INTO BUT NOT THROUGH THE DERMIS
 - ABRASIONS, SKIN TEARS, BLISTERS, SKIN GRAFT DONOR SITES (SPLIT-THICKNESS)
- FULL-THICKNESS
 - THROUGH THE DERMIS EXTENDING DOWN TO SUBCUTANEOUS TISSUE, MUSCLE, MAY HAVE EXPOSED STRUCTURE

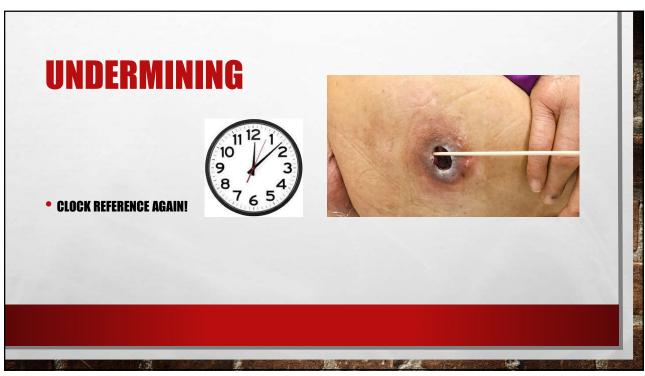
















DEBRIDEMENT - AUTOLYTIC - ENDOGENOUS ENTYMES - ENZYMATIC - COLLAGENASE - MECHANICAL - WOUND SCRUBBING, WET-TO-DRY DRESSINGS, HYDROTHERAPY, ULTRASONIC - SHARP - SURGICAL - STERILE MAGGOTS - LOW FREQUENCY ULTRASOUND - CHEMICAL

MOIST WOUND HEALING & AUTOLYTIC DEBRIDEMENT

- MOIST WOUND HEALING GOAL:
 - PROMOTE PHYSIOLOGICAL HEALING
- AUTOLYTIC DEBRIDEMENT GOAL:
 - THE PROMOTION OF ENDOGENOUS ENZYMES TO DEBRIDE NECROTIC TISSUE

AUTOLYSIS

- AUTOLYSIS: THE NATURAL DEGRADATION OF DEVITALIZED TISSUE USING ENDOGENOUS ENZYMES:
 - PROTEINASES: PROTEOLYTIC ENZYMES AND WBC IN EXUDATE TO BREAKDOWN AND LIQUEFY NECROTIC TISSUE
- MOISTURE RETENTIVE OR MOISTURE DONATING DRESSINGS USED TO PROMOTE MOIST WOUND
 HEALING AND PREVENT EVAPORATION OF FLUIDS FROM THE WOUND SURFACE

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AUTOLYTIC DEBRIDEMENT

- MOISTURE-RETENTIVE OR MOISTURE DONATING DRESSING IS APPLIED COVERING WOUND, RETAINING ENDOGENOUS ENZYMES
- ESCHAR AND SLOUGH ARE LIQUEFIED BY REHYDRATION AND ACTIVITY OF LYSOSOMES NORMALLY PRESENT IN WOUND FLUIDS

AUTOLYTIC DEBRIDEMENT Indications All wounds with necrotic tissue Beneficial with wounds covered by dry eschar Cross-hatching eschar will facilitate autolysis Cross-hatching eschar will facilitate autolysis

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ADVANTAGES OF AUTOLYTIC DEBRIDEMENT

- NON INVASIVE- SAFEST
- DOES NOT DISRUPT HEALTHY TISSUE
- MAY BE USED IN CONJUNCTION WITH OTHER METHOD OF DEBRIDEMENT
- PAINLESS
- EASY TO PERFORM
- SELECTIVE
- REQUIRES MINIMAL PROFESSIONAL EXPERTISE/ MONITORING
- **CAREGIVER FRIENDLY**

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AUTOLYTIC DEBRIDEMENT DRESSING DECISION TREE

- ASSESS WOUND DRAINAGE
 - MINIMAL
 - MODERATE
 - COPIOUS

DRAINAGE DRIVES DRESSING CHOICES

- AMOUNT OF DRAINAGE
- CONDITION OF PERIWOUND SKIN AT DRESSING CHANGE
- FREQUENCY OF DRESSING CHANGES
- QUESTION: WHAT CHOICE OF DRESSING/ PRODUCTS ALLOWS THE WOUND TO BE DISTURBED THE LEAST?
- DRESSING MAY HAVE TO BE CHANGED MORE OFTEN DURING DEBRIDEMENT

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DRESSING CATEGORIES THAT PROMOTE AUTOLYTIC DEBRIDEMENT

- MAINTAIN MOIST ENVIRONMENT
 - TRANSPARENT FILMS (THIN FILMS)
 - HYDROCOLLOIDS
 - HYDROGELS
 - ALCINATES
 - SEMIPERMEABLE FOAMS
 - ABSORBENT FILLER
 - ABSORBENT PAD
 - IMPREGNATED DRESSING
 - GRANULAR EXUDATE ABSORBERS

ENZYMATIC DEBRIDEMENT

- "ENZYMATIC DEBRIDEMENT IS ACCOMPLISHED BY APPLYING TOPICAL DEBRIDING AGENTS TO DEVITALIZED TISSUES ON THE WOUND SURFACE.
- THIS OPTION SHOULD BE CONSIDERED WHEN INDIVIDUALS CANNOT TOLERATE SURGERY OR ARE IN LONG-TERM CARE FACILITIES OR RECEIVING CARE AT HOME AND WHEN THE ULCER DOES NOT APPEAR INFECTED
- IF INFECTION SPREADS BEYOND THE ULCER (E.G., ADVANCING CELLULITIS, SEPSIS), THERE IS URGENT NEED FOR SHARP DEBRIDEMENT"

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ENZYMATIC DEBRIDEMENT

- DEFINITION- USE OF TOPICALLY APPLIED CHEMICAL AGENT TO STIMULATE THE BREAKDOWN OF NECROTIC TISSUE
- ENZYMATIC AGENTS
 - COLLAGENASE
 - PAPAIN-UREA PRODUCTS NO LONGER AVAILABLE IN U.S.

COLLAGENASE

- COLLAGENASE (AN ENZYME) DIGESTS NECROTIC COLLAGEN
- SPECIFIC FOR DENATURED COLLAGEN
- MEDICATION HYDROPHILIC
- COLLAGENASE IS MORE LIKELY TO BE USEFUL FOR A VERY PROLONGED PERIOD OF TIME IN THE WOUND IN WHAT IS CALLED MAINTENANCE DEBRIDEMENT

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EXOGENOUS ENZYMES

- ENZYMES DERIVED FROM OTHER SOURCE OUTSIDE THE BODY
- BIOLOGICALLY ENGINEERED



OTHER CONSIDERATIONS WHEN USING ENZYME MEDICATIONS

- SALTS OF HEAVY METALS SUCH AS SILVER, LEAD, MERCURY, INACTIVATE THE ENZYMES
- HYDROGEN PEROXIDE SOLUTION MAY INACTIVATE PAPAIN

MECHANICAL WOUND DEBRIDEMENT

- APPLICATION OF SOME OUTSIDE "FORCE" OR ENERGY TO DISLODGE NECROTIC TISSUE
- METHODS
 - WOUND SCRUBBING
 - WET-TO-DRY DRESSINGS
 - HYDROTHERAPY
 - ULTRASONIC

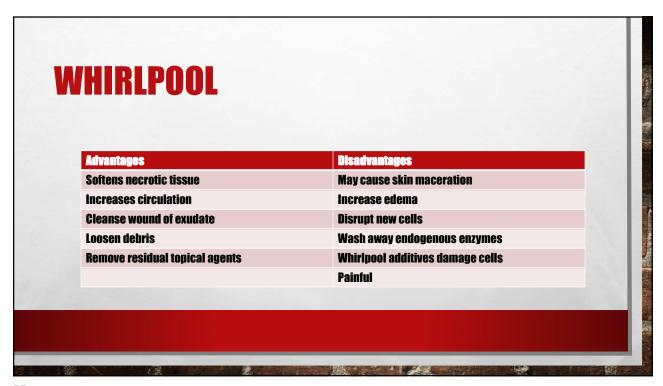
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WOUND SCRUBBING

- USE FOR SOFT SLOUGH DEBRIS
- REMOVES SURFACE DEBRIS
- SCRUB FROM CENTER OF WOUND OUTWARD
- USE GAUZE OR SPONGE
- SCRUB 2-3 MINUTES
- AVOID SCRUBBING HEALTHY GRANULATION TISSUE



HYDROTHERAPY WHIRLPOOL WOUND IRRIGATION PULSE LAVAGE WITH SUCTION (PLWS)



WOUND IRRIGATION • USE 35-50 CC SYRINGE • 19' GAUGE ANGIOCATHETER • 4-8 PSI RECOMMENDED • AVOID > 15 PSI

PULSED LAVAGE WITH SUCTION (PLWS)

- PLWS PROVIDES CLEANSING AND DEBRIDEMENT WITH PULSED IRRIGATION COMBINED WITH SUCTION
- PROVIDES NEGATIVE PRESSURE TO REMOVE THE IRRIGANT AND DEBRIS TO HELP REDUCE INFECTION AND TO ENHANCE GRANULATION TISSUE FORMATION

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PLWS

- INITIALLY USED BY SURGEONS IN OR
- IRRIGATION IN SURGICAL PROCEDURES
- CLEANSE WOUNDS OF DEBRIS
- IRRIGATION AND DEBRIDEMENT TO CLEANSE AND ENHANCE HEALING OF SOFT TISSUE WOUNDS



- THEORY AND SCIENCE OF THERAPY
 - CLEANSING VIA GENTLE PULSATILE LAVAGE TO STRONGER IRRIGATION AND DEBRIDEMENT

- REDUCES BACTERIA AND INFECTION
- PROMOTES ANGIOGENESIS- GRANULATION AND EPITHELIALIZATION
- THEORY: NEGATIVE PRESSURE OF SUCTION STIMULATES CELLS AND GRANULATION

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PLWS

- ELIMINATED NEED FOR WHIRLPOOLS EXCEPT IN LIMITED CIRCUMSTANCES
- NOTE: WP CONTRAINDICATED FOR DFUS

PULSATILE LAVAGE

- GOOD FOR LARGE OR MULTIPLE WOUND SITES
- VARIABLE PRESSURE THAT IS CONTROLLABLE
- 4-8 PSI RECOMMENDED
- > 15 PSI AVOID
- RETURN SUCTION ASSISTS WITH DEBRIDEMENT
- SELECTIVE APPLICATION TIPS

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PULSATILE LAVAGE WITH SUCTION

- STANDARD PRECAUTIONS
 - PRIVATE ROOM ONLY- NO FAMILY/VISITORS
 - AEROSOLIZATION STUDY (LOEHNE 1998)
 - MICROORGANISMS-INFECTION POTENTIAL
 - TESTED 2' & 8' FROM WOUND
 - 30 MINUTES BEFORE, DURING, AFTER RX
 - INCREASED MICROORGANISMS DETECTABLE DURING AND AFTER RX
 - HARRIETT LOEHNE, DPT, CWS

LOW FREQUENCY ULTRASOUND

- EMERGING AS AN ALTERNATIVE METHOD FOR WOUND BED PREPARATION AND WOUND HEALING
- Ultrasound vibration applied to the wound bed via saline creating deep tissue penetration
- CONTACT ULTRASOUND IS AN EFFICIENT DEBRIDEMENT TOOL FOR WOUNDS WITH DENSE FIBRIN AND SLOUGH
- DISRUPTS BIOFILM COLONIES
- CAUSES BACTERIAL DESTRUCTION
- **CONTACT AND NON-CONTACT LFU**

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INDICATIONS FOR LOW FREQUENCY ULTRASOUND

- LOCALLY INFECTED WOUNDS
- WOUNDS WITH IMPAIRED CIRCULATION
- WOUNDS WITH THE NEED FOR DEBRIDEMENT, IRRIGATION, AND TOPICAL TREATMENT
- PRESSURE, DIABETIC, ARTERIAL AND VENOUS ULCERS, POSTTRAUMATIC, AND SURGICAL

PROPOSED MECHANISMS OF ACTION AT CELLULAR LEVEL

Cavitation	Acoustic Microstreaming
Ultrasound induced pressure changes in the tissue fluids, which create gas-filled bubbles that expand and compress	Unidirectional movement of fluids along cell membranes
Stable macro cavitation may change cell membrane permeability	Results from mechanical pressure changes within the ultrasound field
	Acoustic microstreaming may change cell membrane permeability

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CONTACT LOW FREQUENCY US

- WOUND IRRIGATION FLUID DIRECTED THROUGH AN OPENING IN A PROBE'S TIP
- ADMINISTERS FLUID DIRECTLY TO WOUND SURFACE
- FLUID SERVES AS COUPLING MEDIUM, COOLANT, WOUND LAVAGE, OR FLUSH
- TOPICALLY TREAT THE WOUND BASE

OUTCOMES OF LOW FREQUENCY ULTRASOUND

- DEBRIDEMENT:
 - LOOSENS THE NECROTIC TISSUES FROM THE WOUND AREA SO IT CAN BE WASHED AWAY
- BIOBURDEN CONTROL
 - SIGNIFICANTLY DECREASES THE BACTERIA CONTENT OF THE WOUND AREA
- INCREASES THE MOVEMENT OF FIBROBLAST CELLS INTO THE WOUND AREA
- STIMULATES THE GROWTH OF FIBROBLAST CELLS
- IMPROVES THE GROWTH OF CAPILLARY BLOOD VESSELS
 - IMPROVE CIRCULATION IN THE AREA SURROUNDING THE WOUND
- DEDONTO OF REORESCED WOUND CITE DAW

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BIOSURGERY



- MAGGOTS HAVE BEEN USED SUCCESSFULLY TO DEBRIDE:
 - PRESSURE ULCERS
 - VENOUS INSUFFICIENCY ULCERS
 - DIABETIC FOOT ULCERS
 - SOFT TISSUE WOUNDS RESULTING FROM NECROTIZING SOFT TISSUE INFECTIONS

MAGGOT THERAPY • STUDIES IN SUPPORT OF DEBRIDEMENT • BAER- 1929 • HOBSON-1931 • MASERITZ- 1934 • ZIFFREN ET AL- 1953 • WATERHOUSE & IRZYKIEWICZ- 1957 • FRASER ET AL; BROOKES- 1961 • PENDOLA & GREENBERG- 1975 • VISTNES ET AL- 1981 CASU ET AL- 1994

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MAGGOTS PHAENICIA (LUCILIA) SERICATA LARVA COLORIZED SCANNING ELECTRON MICROGRAPH



** OF THE STATE OF

MAGGOT THERAPY

- JANUARY 12, 2004- FDA APPROVES THE PRODUCTION AND MARKETING OF THE BRAND: MEDICAL MAGGOTS AS A MEDICAL DEVICE (510K#K033391) FOR THE FOLLOWING INDICATIONS
 - "... DEBRIDING NON-HEALING NECROTIC SKIN AND SOFT TISSUE WOUNDS, INCLUDING PRESSURE ULCERS, VENOUS STASIS ULCERS, NEUROPATHIC FOOT ULCERS, AND NON-HEALING TRAUMATIC OR POST SURGICAL WOUNDS."

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CHEMICAL DEBRIDEMENT

- CHEMICAL DEBRIDEMENT
- AKA CHEMICAL CAUTERIZATION
- SILVER NITRATE STICKS

INDICATIONS AND CHEMICAL MAKEUP

- SKIN OR MUCOUS MEMBRANE CAUTERIZATION
- REMOVING WARTS
- HYPER-GRANULATED TISSUE
- 75% SILVER NITRATE AND 25% POTASSIUM NITRATE

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APPLICATION CONSIDERATIONS

- RIGID WOODEN APPLICATOR FOR PRECISION PLACEMENT
- SILVER NITRATE TIPS ARE SOLUBLE IN ALL SECRETIONS
- CONTROL STRENGTH BY DIPPING IN STERILE WATER
- INCREASE IN WOUND DRAINAGE- INFLAMMATORY RESPONSE
- EDUCATE PATIENT/ FAMILY REGARDING BLACK/ GRAY STAINING
- PROTECT PERIWOUND AREA WITH PETROLATUM- WILL INJURE HEALTHY TISSUE
- PHYSICIAN ORDER NEEDED

SELECTIVE SHARP DEBRIDEMENT





- SHARP DEBRIDEMENT IS THE MOST RAPID METHOD
- SELECTIVE FOR NON-VIABLE TISSUE ONLY
- SIGNS OF ADVANCING CELLULITIS OR SEPSIS REQUIRES RAPID EMERGENT DEBRIDEMENT

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INDICATIONS FOR SHARP DEBRIDEMENT

- EXTENSIVE DEVITALIZED TISSUE
- SIGNS OF ADVANCING CELLULITIS OR SEPSIS
- PRESENCE OF THICK ADHERENT ESCHAR- INDICATES FULL-THICKNESS WOUND
- AS AN ADJUNCT IN COMBINATION WITH OTHER METHODS
- CALLOUS FORMATION- INSENSATE FOOT

PRECAUTIONS AND CONTRAINDICATIONS FOR SHARP DEBRIDEMENT

- ARTERIAL INSUFFICIENCY
 - GANGRENE
 - LOW ABI, LOW TCP02
- PAIN CONTROL CANNOT BE ASSURED
- EXCESSIVE BLEEDING MIGHT RESULT
- DIAGNOSIS OF THE LESION MAY BE UNCERTAIN
- PATIENT IMMUNOCOMPROMISED
- STABLE VERSES UNSTABLE HEEL ULCERS
- UNIDENTIFIABLE STRUCTURES
- TERMINALLY ILL- DEPENDS ON GOALS FOR CARE
- NOTE CHARD CHROIGH DERRIDEMENT MAY CAUSE GROANISMS TO BE CHOWERED INTO BLOODSTREAM RESULTING IN PACTEREMIA

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REASONS TO STOP SHARP DEBRIDEMENT

- CLINICIAN FATIGUE/ PATIENT FATIGUE
- BLEEDING
- PAIN
- TO VIABLE TISSUE
- LOCATION OF A FASCIAL PLANE
- LOCATION OF A NAMED STRUCTURE
- HIGH ANXIETY LEVEL (CLINICIAN OR PATIENT)
- ACHIEVED SET TIME LIMIT (15-30 MINUTES)

HEMOSTASIS

- PRESSURE WILL STOP MOST BLEEDING
- ELEVATION ABOVE HEART IF EXTREMITIES
- NITRATE STICKS
- CALCIUM ALGINATE
- LIDOCAINE JELLY
- ARTICAINE AND EPINEPHRINE ANESTHETIC
- HEMOSTATIC AGENTS
 - TOPICAL THROMBIN
- COLLAGEN
- QR POWDER

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MAINTENANCE DEBRIDEMENT

 REPEATED ONGOING DEBRIDEMENT OF CHRONIC WOUNDS TO MAINTAIN THE WOUND BED IN A STATE OF READINESS TO HEAL

SURGICAL DEBRIDEMENT

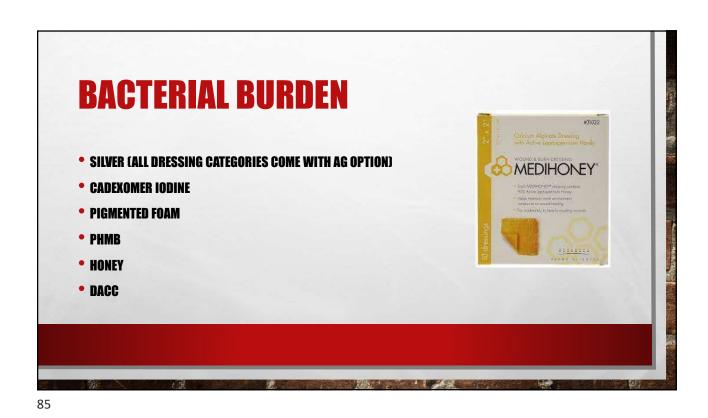
- SHARP SURGICAL DEBRIDEMENT
- LASER DEBRIDEMENT
- FASTEST METHODS OF DEBRIDEMENT
- CAN GET ALL NECROTIC OR UNHEALTHY WOUND TISSUE IN ONE PROCEDURE
- SELECTIVE IF NONVIABLE TISSUE ONLY
- NON-SELECTIVE IF VIABLE TISSUE INCLUDED
- SURGICAL DEBRIDEMENT CAN BE PERFORMED IN THE OPERATING ROOM OR AT BEDSIDE

CLINICIAN MUST BE CREDENTIALED AS MD/DO, DPM

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SURGICAL DEBRIDEMENT

- ADVANTAGES
 - GOOD FOR WOUNDS WITH A LARGE AMOUNT OF NECROTIC TISSUE
 - APPROPRIATE FOR REMOVAL OF INFECTED TISSUE
 - FAST
- DISADVANTAGES
 - PAINFUL TO PATIENT
 - COSTLY, ESPECIALLY IF OPERATING ROOM IS REQUIRED



DRESSING FUNCTION Films Hydrogels Alginates/ **FOAMS HYDROCOLLOIDS Hydrofibers** COVER/ PROTECT X X X X **HYDRATE** X **MAINTAINS** X X **MOISTURE/ AUTOLYTIC SUPPORT ADDS MOISTURE** X **ABSORPTION** X **FILL SPACE** X X

WOUND FLUID MANAGEMENT

LOW	MODERATE	HIGH
FILMS	CALCIUM ALGINATES Hydrofibers	CALCIUM ALGINATES Hydrofibers
FOAMS	FOAMS	FOAMS
HYDROCOLLOIDS	HYDROCOLLOIDS	
HYDROGELS		

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TISSUE ENGINEERED PRODUCTS

- CAN BE CLASSIFIED AS CELLULAR OR ACELLULAR
 - ACELLULAR ARE TISSUE MATRICES THAT CONTAIN NO CELLS
 - CONSIST OF A MATRIX THAT FUNCTIONS BY BINDING TO THE HOST, ALLOWING MATRIX-CELL INTERACTIONS
 - THE MATRIX ALLOWS HOST CELLS TO INFILTRATE
 - CELLULAR DEVICES USUALLY KERATINOCYTES AND FIBROBLASTS
 - SINGLE LAYER KERATINOCYTES GROWN ON POLYGLACTIN MESH
 - BILAYER KERATINOCYTES AND FIBROBLASTS ON COLLAGEN MATRIX

FUNCTION OF CELLULAR SKIN SUBSTITUTES

- GOAL OF USE IS TO RESTORE SKIN BARRIER
- SECRETE EXTRACELLULAR MATRIX (COLLAGEN)
- PRODUCE GROWTH FACTORS NEEDED BY THE WOUND AT THE NEEDED TIME AND NEEDED AMOUNTS
- PROVIDE TEMPORARY WOUND COVERAGE
- PRODUCTS WITH DIFFERENTIATED KERATINOCYTE LAYER PROVIDE PROTECTION FROM MOISTURE LOSS, BACTERIAL PROTECTION



CELLULAR SKIN SUBSTITUTES DO NOT "TAKE"

- THEY ARE NOT SKIN GRAFTS
- THERE IS NO PERSISTENCE
- THERE IS NO VASCULARIZATION
- THERE IS NO INTEGRATION

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HUMAN FIBROBLAST-DERIVED DERMAL SUBSTITUTE (DERMOGRAFT)



 A CRYOPRESERVED HUMAN FIBROBLAST DERIVED DERMAL SUBSTITUTE; IT IS COMPOSED OF FIBROBLASTS, EXTRACELLULAR MATRIX, AND A BIOABSORBABLE SCAFFOLD

BI-LAYER BIOENGINEERED SKIN SUBSTITUTE (APLIGRAFT) Organogenesis



- A LIVING, BI-LAYER SKIN SUBSTITUTE:
 - EPIDERMAL LAYER IS FORMED BY HUMAN KERATINOCYTES AND HAS A WELL-DIFFERENTIATED STRATUM CORNEUM
 - CONTAINS KERATINOCYTE STEM CELLS
 - DERMAL LAYER IS COMPOSED OF HUMAN FIBROBLASTS IN A BOVINE TYPE I COLLAGEN LATTICE
 - HUMAN MATRIX PROTEINS AND CYTOKINES FOUND IN HUMAN SKIN ARE PRESENT
 - DOES NOT CONTAIN LANGERHANS CELLS, MELANOCYTES, MACROPHAGES, LYMPHOCYTES, BLOOD VESSELS OR HAIR
 FOLLICLES (NO IMMUNE CELLS)

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ACELLULAR TISSUE ENGINEERED PRODUCTS



- MAY FUNCTION AS A BIOLOGIC MODULATOR
 - MATERIAL OR SUBSTANCE DERIVED FROM BIOLOGIC OR SYNTHETIC SOURCES
 - INFLUENCES BIOLOGICAL PROCESSES SUCH AS WOUND HEALING
- PROVIDE A SCAFFOLD FOR TISSUE REPAIR
 - NEEDS TO REMAIN IN PLACE FOR A SUFFICIENT LENGTH OF TIME
 - PROVIDES A SUPPORTING STRUCTURE INTO WHICH CELLS CAN MIGRATE







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