Article Title/ Author/Year	Article Type	Subjects (Number and Criteria)	Outcome Measures and Timeframes	Interventions or Methods	Results or Key Points	Conclusions
Description of Dry Needling In Clinical Practice: An Educational Resource Paper 2013	Educational Resource Paper	N/A	N/A	N/A	Definition of DN: skilled intervention that uses thin filiform needles to penetrate skin and stimulate myofascial trigger points, muscle, and connective tissue to improve pain and movement impairments to improve activity and participation  Definition of trigger points: hyperirritable spots within taut band of contractured skeletal muscle fibers that produce local and/or referred pain when stimulated;	N/A

characterized by local ischemia and hypoxia, low ph, chemical differences, pain, and altered muscle activation patterns; generate motor endplate noise and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful, latent trigger points are only painful when stimulated  Theories of physiological effects:	
and hypoxia, low pH, chemical differences, pain, and altered muscle activation patterns; generate motor endplate noise and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful; latent trigger poinful are only painful when stimulated	
pH, chemical differences, pain, and altered muscle activation patterns; generate motor endplate noise and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	
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muscle activation patterns; generate motor endplate noise and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	
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generate motor endplate noise and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	muscle activation
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and excessive release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	generate motor
release of acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	endplate noise
acetylcholine; peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	and excessive
peripheral sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	release of
sources of constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	acetylcholine;
constant nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	peripheral
nociceptive input that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	sources of
that can contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	constant
contribute to peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	nociceptive input
peripheral and central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	that can
central sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	contribute to
sensitization; active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	peripheral and
active trigger points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	central
points are spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	sensitization;
spontaneously painful; latent trigger points are only painful when stimulated  Theories of physiological	active trigger
painful; latent trigger points are only painful when stimulated  Theories of physiological	points are
trigger points are only painful when stimulated  Theories of physiological	spontaneously
trigger points are only painful when stimulated  Theories of physiological	painful; latent
only painful when stimulated  Theories of physiological	
Theories of physiological	
physiological	stimulated
physiological	
	Theories of
	physiological
	effects:
mechanotransduc	mechanotransduc

tion (body
converting
mechanical
loading by needle
into cellular
response)
achieved by
needle rotation
that generates
fibroblast
activation and
collagen
reorganization.
Types of DN:
deep DN with
penetration of
trigger point
which causes
local twitch
response (spinal
cord reflex that
causes
involuntary
contraction of
contractured taut
band) which is
associated with
alleviation of
motor endplate
noise, chemical
imbalance
(nociceptive,
inflammatory,

immune), and
ischemia, as well
as fiber
relaxation,
decreased pain,
and improved
ROM. Superficial
DN just into
muscle within
proximity to
trigger point
which activates
mechanoreceptor
s, resulting in
decreased pain
and improved
ROM. No local
twitch response
produced.
Indications for
DN: presence of
trigger points,
ROM restrictions
due to
contractured
muscle
fibers/taut bands
Safety: requires
knowledge, skills,
and attributes to
perform; adhere
to OSHA Blood

Borne Pathogens standard (gloves)  Contraindications /precautions: needle phobia, significant anxiety, children 12 years of age or younger, significant cognitive impairment, unable to communicate directly or via interpreter, patient unwilling to receive treatment, patient unable to give consent, local skin lesions, local or systemic infections, local lymphedema, severe hypperalgesia, metal allergy, abnormal bleeding (on anticoagulants or with hypophocytopen)		
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anxiety, children 12 years of age or younger, significant cognitive impairment, unable to communicate directly or via interpreter, patient unwilling to receive treatment, patient unable to give consent, local skin lesions, local or systemic infections, local lymphedema, severe hyperalgesia, metal allergy, abnormal bleeding (on anticoagulants or with		significant
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severe hyperalgesia, metal allergy, abnormal bleeding (on anticoagulants or with		
metal allergy, abnormal bleeding (on anticoagulants or with		
metal allergy, abnormal bleeding (on anticoagulants or with		hyperalgesia,
abnormal bleeding (on anticoagulants or with		
bleeding (on anticoagulants or with		
anticoagulants or with		
with		
		thrombocytopeni

		a), compromised
		immune system,
		pregnancy,
		vascular disease,
		post-surgery of
		open joint
		capsule
		Adverse events:
		minor bleeding
		most common
		Application: DN is
		rarely performed
		alone and should
		be included in a
		broader physical
		therapy
		approach,
		including manual
		therapy,
		therapeutic
		exercise,
		neuromuscular
		re-education, and
		functional
		training
		Reimbursement:
		physical
		therapists should
		check with
		insurance payers

					to determine	
					billing policies	
Dry Needling:	APTA Magazine	N/A	N/A	N/A	Definition of DN: N/A	
Getting to the	web article				instrument-	
Point					assisted manual	
					therapy, based on	
Ries					western	
					neuroanatomy	
2015					and modern	
					science	
					"single tool in the	
					PT's toolbox"	
					Acupuncture: DN	
					and acupuncture	
					differ in terms of	
					historical,	
					philosophical,	
					indicative, and	
					practice context	
					Theories of	
					physiological	
					effects: largely	
					unclear	
					Personal scope of	
					practice/PT	
					education: basic	
					anatomical,	
					physiological, and	
					biomechanical	
					knowledge for DN	
					taught as part of	

	core PT
	education; DPT
	programs are
	teaching students
	about DN,
	movement to
	incorporate into
	residencies
	Professional
	scope of practice:
	included in APTA
	Board of Directors
	policy Guidelines:
	Physical Therapist
	Scope of Practice
	and listed as
	manual therapy
	technique in
	Guide to Physical
	Therapist Practice
	3.0 based on
	2011 review of
	evidence that
	found DN has
	mid-range
	research support
	Reimbursement:
	not CPT code,
	some payers do
	not reimburse ,
	APTA defines DN
	as a manual

therapy
technique but
does not say it
should be coded
or billed as such,
should check with
payers to
determine if DN is
covered and
which code to
use, many
provide DN on a
cash basis to
avoid this
Historical
perspective:
David Simons and
Janet Travell
authored
textbook
Myosfasical Pain
and Dysfunction:
The Trigger Point
Manual
Application: DN
should not be a
standalone
procedure
Adverse events:
most serious

					being	
					pneumothorax	
FSBPT Analysis of	Practice analysis	N/A	N/A	Purpose of	86% of	N/A
Competencies for	web report			document:	knowledge	
Dry Needling by				determine	requirements	
Physical				measurable or	(evaluation,	
Therapists				observable	assessment,	
				knowledge, skills,	diagnoses, plan of	
Caramagno et al.				and/or abilities	care,	
				(competencies) a	documentation,	
2015				PT must possess	safety,	
				to	professional	
				perform DN	responsibilities)	
				competently	to be competent	
					in DN is acquired	
				Methods to	during PT entry	
				determining	level education	
				competencies:		
				background	14% of	
				literature review,	knowledge	
				practitioner	requirements	
				survey, task force	(patient selection,	
				meeting of 7 DN	needle placement	
				experts to define	and manipulation,	
				DN and standards	identification of	
				for competence,	contraindications)	
				review DN tasks	must be acquired	
				and knowledge	through post-	
				requirements,	graduate or	
				and identify DN	specialized	
				skills and abilities	education	
					The only skills	
					that entry level	

_	1	1	1	T	T	
					PT's do not have	
					that is required of	
					DN is ability to	
					handle needles	
					and palpate	
					tissues, which	
					requires	
					specialized	
					training	
					Historical	
					perspectives:	
					pioneered by	
					Travell and	
					Simons (MDs)	
					Definition DN:	
					based on	
					philosophical and	
					theoretical	
					framework	
					supported by	
					modern science,	
					use of needle	
					without injectate	
Considering	APTA Magazine	N/A	N/A	N/A	Professional	N/A
Providing Dry	web article				scope of practice:	
Needling					defined by	
Services?					education,	
Understand your					research, and	
professional,					APTA positions;	
legal, and					recognized by	
personal scopes					APTA as within	
of practice					scope of practice	

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		and listed in
Markels		Guide to Physical
		Therapist Practice
2021		3.0 as manual
		therapy
		technique; APTA
		House of
		Delegates
		position
		"Interventions
		Performed
		Exclusively by
		Physical
		Therapists" says
		DN should not be
		delegated
		Legal scope of
		practice: depends
		on state licensure
		law/practice
		act/state
		licensure board
		positions
		Personal scope of
		practice: activities
		that a PT is
		educated,
		trained, and
		competent to
		perform,
		individual states
		often have

					different educational requirements for DN  Reimbursement: CPT codes 20560 (needle insertion without injection, 1-2 muscles) and 20561 (needle insertion without injection, 3 or more muscles)	
State Laws and Regulations Governing Dry Needling Performed by Physical Therapists in the U.S.	N/A	N/A	N/A	N/A	DN by PTs permitted by law in 36 states plus Washington D.C. (including NC)  DN by PTs prohibited by law in 6 states  DN by PTs ambiguous in law in 8 states	N/A
Scope of Practice 2022	NC PT Board website	N/A	N/A	N/A	Legal and student scope of practice: Board previously determined that DN is "advanced" skills that requires advanced training	N/A

(beyond entry-
level); "Students
who are in the
process of
didactic and
clinical training do
not meet the
definition of
'advanced'"
meaning that
they cannot
perform DN;
there are no
specific
requirements for
the education and
training required
for performing
dry needling by a
physical therapist
licensee;
certification is not
currently required
by the Board; "it
is very useful to
keep (certification
if it is obtained)
on file as part of
documentation of
competence
related to the
personal practice
of DN), DN
courses currently

					approved per Continuing Competence rules 12 NCAC 48G .01050112.; "it is incumbent upon the licensee to obtain the appropriate training, education and be competent to perform DN"	
ADVERSE EVENTS ASSOCIATED WITH THERAPEUTIC DRY NEEDLING  Boyce et al. 2020	Prospective questionnaire	420 physical therapists included	Survey collected information on minor (bleeding, bruising, pain, feeling faint, nausea, headache, drowsiness) and major (pneumothorax, punctured organ, broken or forgotten needle, excessive bleeding, nerve injury, infection, prolonged pain, fainting, convulsion, vomiting, skin reactions)	N/A	Information from a total of 20,464 DN sessions was used.  7,531 minor events reported; 36.7% of treatments resulted in minor events including bleeding (16.04%), bruising (7.71%), pain during treatment (5.93%), and others (<3%).  20 major events reported; <0.1% of treatments	Minor adverse events are relatively common as a result of DN; however, major adverse events are extremely rare. Because the risk of major adverse event is very small, DN can safely be used among patients.

			adverse events		resulted in major	
			during DN over a		events including	
			6-week period.		symptom	
			o week periou.		aggravation (6	
					participants),	
					fainting (4	
					participants),	
					forgotten needles	
					(3 participants),	
					flu like symptoms	
					(2 participants),	
					infection (2	
					,	
					participants),	
					lower extremity	
					weakness (1	
					participant),	
					excessive	
					bleeding (1	
					participant),	
					upper extremity	
					numbness (1	
		0.55   1   1	6 11 . 1	21/2	participant).	
A survey of	Cross sectional	865 physical	Survey collected	N/A	95.3% of	Minor adverse
American physical	observational	therapists	information on		participants	events are
therapists'	survey	included	general		report using a	relatively
current practice			information		deep DN	common as a
of dry needling:			about DN, DN		technique and	result of DN.
Practice patterns			practice patterns		54% report using	While major
and adverse			and training,		superficial DN.	adverse events
events			adverse events		Many participants	can occur, they
			occurrence		also reported	are far less
Gattie et al.			(major and		almost always	common. Adverse
			minor), and		using the	events resulting
2020			demographics.		pistioning (in and	from DN are

Information	out) technique lower or
regarding adverse	(37.4%). comparable to
events is most	other treatments
relevant to this	97.4% of (thrust joint
capstone project.	participants manipulations or
	report typically medications) for using DN in musculoskeletal
	using DN in musculoskeletal combination with conditions.
	other treatments.
	Of the 412
	Of the 413
	participants who
	performed DN,
	minor events
	such as pain
	during (39.56%)
	and after
	(23.38%)
	treatment were
	the most
	common. Other
	minor events
	included bleeding
	(17.11%), bruising
	(11.81%), fatigue,
	emotional,
	headache,
	drowsiness,
	shaky, itching,
	numbness, and
	claustrophobia
	(all less than 5%).

Of the 413 participants who performed DN, major events such as prolonged symptom aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE		1	I		1		1
PERTINENT DRY NEEDLING COMSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE						Of the 413	
major events such as prolonged symptom aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING COMSIDERATIONS FOR MINIMIZING COMSIDERATIONS FOR COMSIDERATIONS FO						participants who	
as prolonged symptom aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY REEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE  ADVERSE EFFECTS — PART ONE						performed DN,	
symptom aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE  Symptom aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A  Theories of physiological effects: peripheral and central pain modulation including the gate						major events such	
aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE  ADVERSE EFFECTS PART ONE  Aggravation (15%) and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A  N/A  N/A  N/A  Theories of physiological effects: peripheral and central pain modulation including the gate						as prolonged	
and fainting (15%) were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE  ANA  N/A  N/A  N/A  N/A  N/A  N/A  N						symptom	
were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING COMMENTARY  PERTINENT DRY Commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS  PART ONE  Were the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  Prefixed the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  Prefixed the most commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PROTINENT DRY  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						aggravation (15%)	
Commonly reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING COMMENTARY COMMENTARY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE  REPART ONE  RE						and fainting (15%)	
reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE  Reported. Other major events included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						were the most	
PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE  Minimizing ADVERSE EFFECTS PART ONE  Minimizing ADVERSE EFFECTS PART ONE  Minimizing ADVENSE Minimizing ADVENSE EFFECTS PART ONE  Minimizing ADVENSE MINIMIZING AD						commonly	
included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE  Included forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A N/A Theories of physiological effects: peripheral and central pain modulation including the gate						reported. Other	
PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS – PART ONE    Forgotten needle (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).    Major adverse events were rare.						major events	
PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE    (6.3%), and vomiting, subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).   Major adverse events were rare.						included	
PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS PART ONE						forgotten needle	
Subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE  Subdural hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						(6.3%), and	
hematoma, pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING COMMENTARY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE  N/A NA N/A N/A N/A N/A Theories of physiological effects: peripheral and central pain modulation including the gate						vomiting,	
pneumothorax, nerve injury, infection, or broken needle (all less than 5%).  PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE  Pineumothorax, nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						subdural	
nerve injury, infection, or broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS – PART ONE  N/A N/A N/A N/A N/A Theories of physiological effects: peripheral and central pain modulation including the gate						hematoma,	
PERTINENT DRY Clinical commentary  PERTINENT DRY N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/A						pneumothorax,	
broken needle (all less than 5%).  Major adverse events were rare.  PERTINENT DRY NEEDLING commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS — PART ONE  Droken needle (all less than 5%).  Major adverse events were rare.  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						nerve injury,	
PERTINENT DRY NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE    Less than 5%).   Major adverse events were rare.   Major adverse events were rare.   Minimizer   N/A						infection, or	
PERTINENT DRY Clinical commentary  CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS – PART ONE  Major adverse events were rare.  N/A  N/A  N/A  N/A  N/A  N/A  N/A  N/						broken needle (all	
PERTINENT DRY Clinical N/A N/A N/A Theories of physiological effects: FOR MINIMIZING ADVERSE EFFECTS – PART ONE						less than 5%).	
PERTINENT DRY Clinical N/A N/A N/A Theories of physiological effects: FOR MINIMIZING ADVERSE EFFECTS – PART ONE							
PERTINENT DRY NEEDLING COMMENTARY CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE  Clinical N/A N/A N/A N/A N/A N/A N/A Theories of physiological effects: peripheral and central pain modulation including the gate						Major adverse	
NEEDLING CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE  physiological effects: peripheral and central pain modulation including the gate						events were rare.	
CONSIDERATIONS FOR MINIMIZING ADVERSE EFFECTS - PART ONE  effects: peripheral and central pain modulation including the gate	PERTINENT DRY	Clinical	N/A	N/A	N/A	Theories of	N/A
FOR MINIMIZING ADVERSE EFFECTS - PART ONE  peripheral and central pain modulation including the gate	NEEDLING	commentary				physiological	
MINIMIZING ADVERSE EFFECTS - PART ONE  Central pain modulation including the gate	CONSIDERATIONS					effects:	
ADVERSE EFFECTS  - PART ONE  modulation including the gate	FOR					peripheral and	
- PART ONE including the gate	MINIMIZING					central pain	
	ADVERSE EFFECTS					modulation	
control theory	– PART ONE					including the gate	
Control theory						control theory	

Halle and Halle	and endogenous
Traile and traile	opioid system,
2016	disruption of
2010	hyperalgesia and
	central
	sensitization,
	disruption of
	trigger points
	with local
	ischemia/hypoxia
	related to access
	acetylcholine
	release and
	endplate noise,
	increases in blood
	flow and oxygen
	saturation levels,
	increased
	fibroblastic
	activity,
	endocrine/neurol
	ogic activity that
	decrease
	activation of
	limbic system
	Safety concerns:
	pneumothorax,
	penetration of
	pericardial
	sac/cardiac
	tamponade,
	hematoma, CNS
	injury when

	1	1	1	1		
					needling in and	
					around thorax	
					(shoulder, neck,	
					etc); can be	
					mitigated with	
					knowledge of	
					anatomy and	
					training in needle	
					application	
PERTINENT DRY	Clinical	N/A	N/A	N/A	Safety concerns:	N/A
NEEDLING	commentary				puncture of	
CONSIDERATIONS					peritoneal cavity	
FOR					or internal organs	
MINIMIZING					when needling in	
ADVERSE EFFECTS					and around the	
– PART TWO					abdomen, pelvis	
					and back;	
Halle and Halle					vasovagal	
					responses leading	
2016					to	
					lightheadedness	
					or syncope;	
					importance of	
					conveying	
					possible adverse	
					events when	
					obtaining	
					informed	
					consent; use of	
					universal	
					precautions	
					(glove wear to	
					protect patient	
					and therapist);	

Physiologic	Review article	N/A	Review of basic	N/A	can be mitigated with knowledge of anatomy, training in needle application, and training in response to adverse events Peripheral pain	The proposed
Effects of Dry Needling Cagnie et al.	TREVIEW dition		and clinical published research		modulation is achieved by chemical changes in local tissues including nociceptive and	effects of DN are not definitively supported by research, but are complex and involve peripheral
2013					inflammatory substances.  Central pain modulation is achieved by the gate control theory, endogenous opioid system, and other neural or endocrine pathways.  Trigger point	and central pain modulating systems.
					formation: development of taut band by excessive	

T
acetylcholine
release and end
plate noise,
sustained
contracture of
sarcomere results
in ischemia and
hypoxia, this
results in release
of nociceptive
chemicals, which
can cause
peripheral and
central
sensitization
Theories of
physiologic
effects: disruption
of motor
endplate noise
and acetylcholine
release by
mechanical
stimulation,
increase blood
flow and
oxygenation by
microinjury and
release of
vasodilators, gate
control theory by
mechanical
stimulation,

			endogenous opioid release or neurologic/endoc	
			rine response by	
			microinjury	

Abbreviations: DN (dry needling), ROM (range of motion), PT (physical therapy or physical therapist), CPT (Current Procedural Terminology), APTA (American Physical Therapy Association), CNS (central nervous system)

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