

## PAININTHEOLDER PERSONAND COGNITIVELYIMPAIRED Nele Van Den Noortgate IUC 21January 2022





### <u>SUMMARY</u>

Influence of ageing and cognitive impairment on perception of

pain

— Is pain an important problem?

- Prevalence of pain
- Consequences
- Management of pain
  - Challenges
  - Evaluation of pain
  - Medical treatment: Specific aspects





# 'NORMAL' AGEING PROCESS OF SOMATOSENSORY PAIN SYSTEM

- Relation age ~ pain is still unclear
  - Reduction in afferent transmission
  - Reduction in endogenous pain inhibitory system
- Little or no change in acute pain perception
- Alterations in pain threshold ~ intensity, area, modality and duration of the stimulus



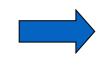
## Reduced ability to detect signals harmful to the body





## 'NORMAL' AGEING PROCESS OF SOMATOSENSORY PAIN SYSTEM

- Longer period of central hyperalgesia
  - For comparable levels of spontaneous pain, thermal hyperalgesia and flare
- Tenderness after injury appears to be prolonged

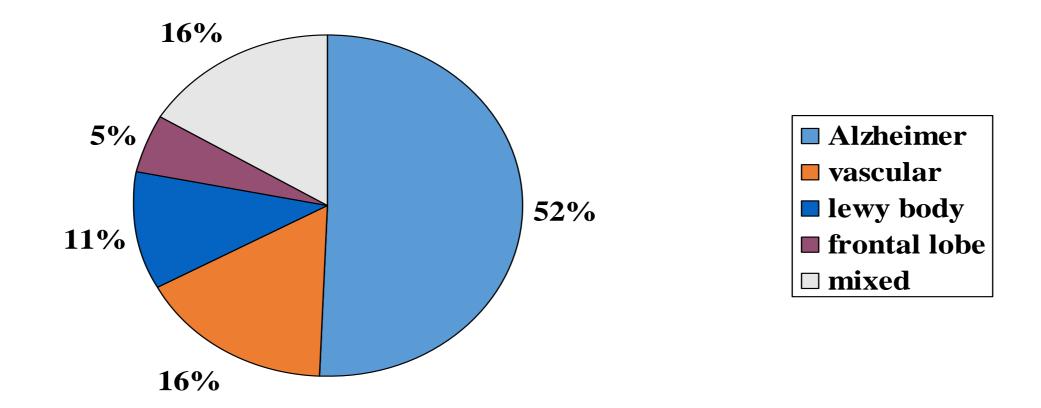


Higher risk of chronic pain and decreased pain tolerance





### PAIN AND COGNITIVE IMPAIRMENT







#### PAIN AND COGNITIVE IMPAIRMENT

#### **ACUTE PAIN**

ALZHEIMER DISEASE

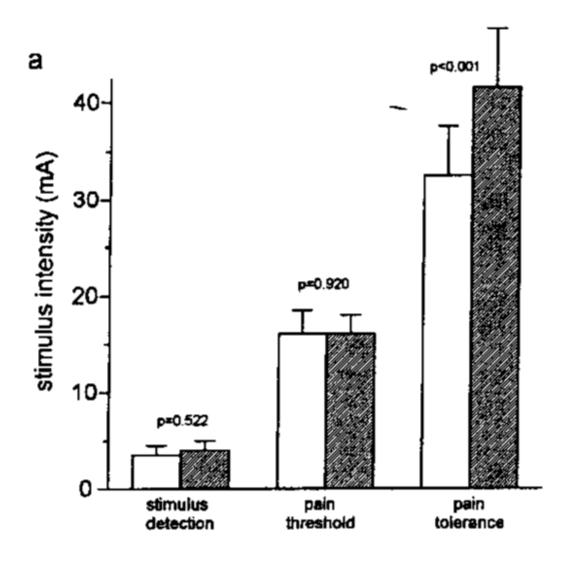
MILD to MODERATE STAGE

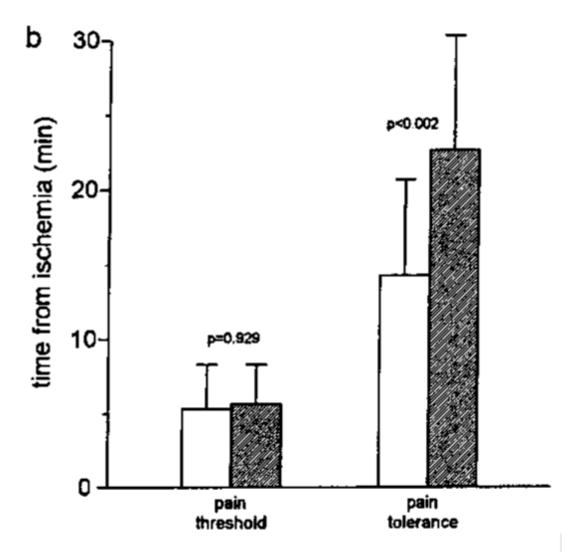




#### ALZHEIMER DISEASE AND PAIN

Pain in normal subjects (white) and in AD patients (shadow)

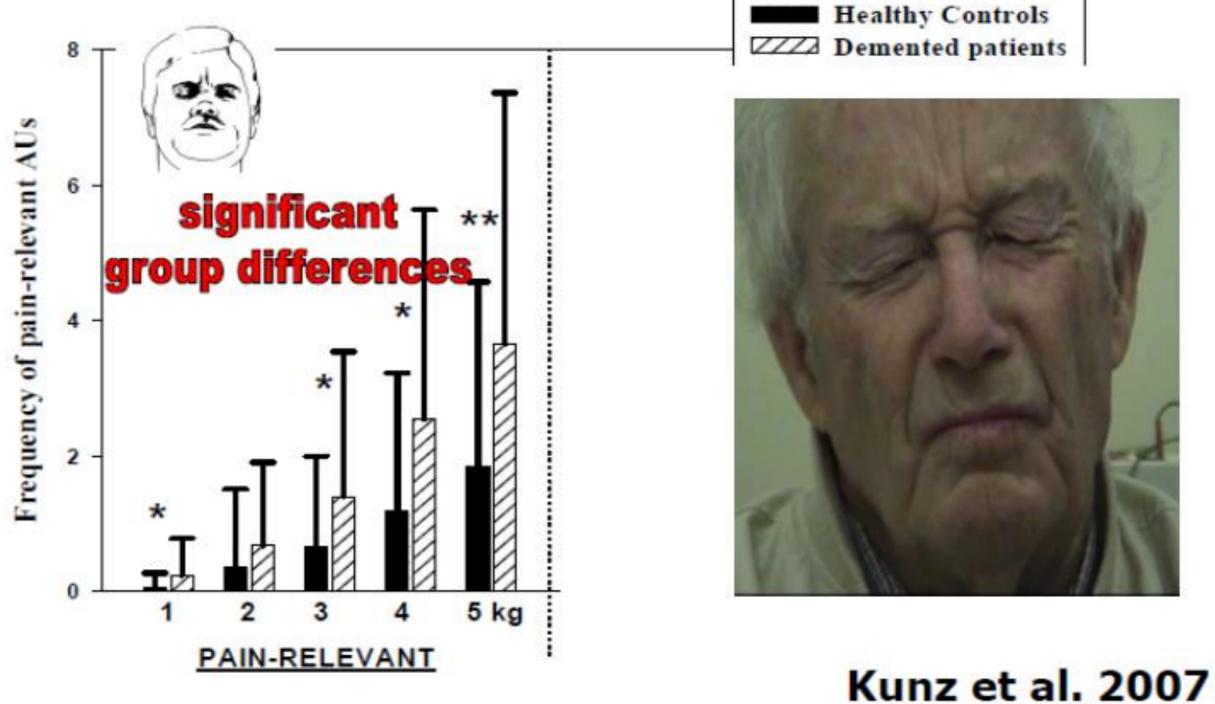








#### FACIAL EXPRESSIONS

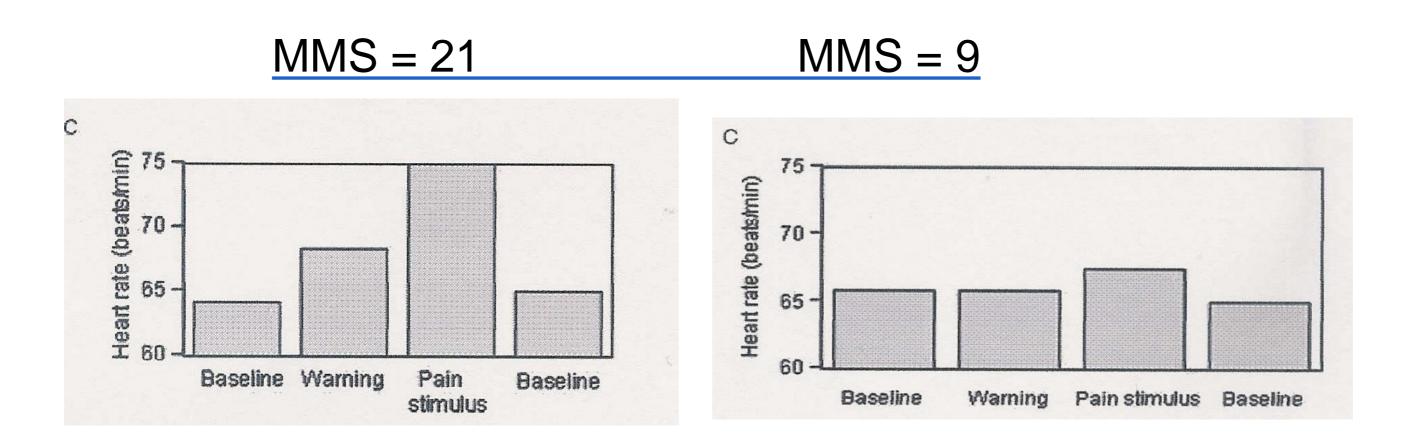




Pain 133 (2007) 221-228

ahead 4 6

## <u>AUTONOMOUS RESPONSE</u>

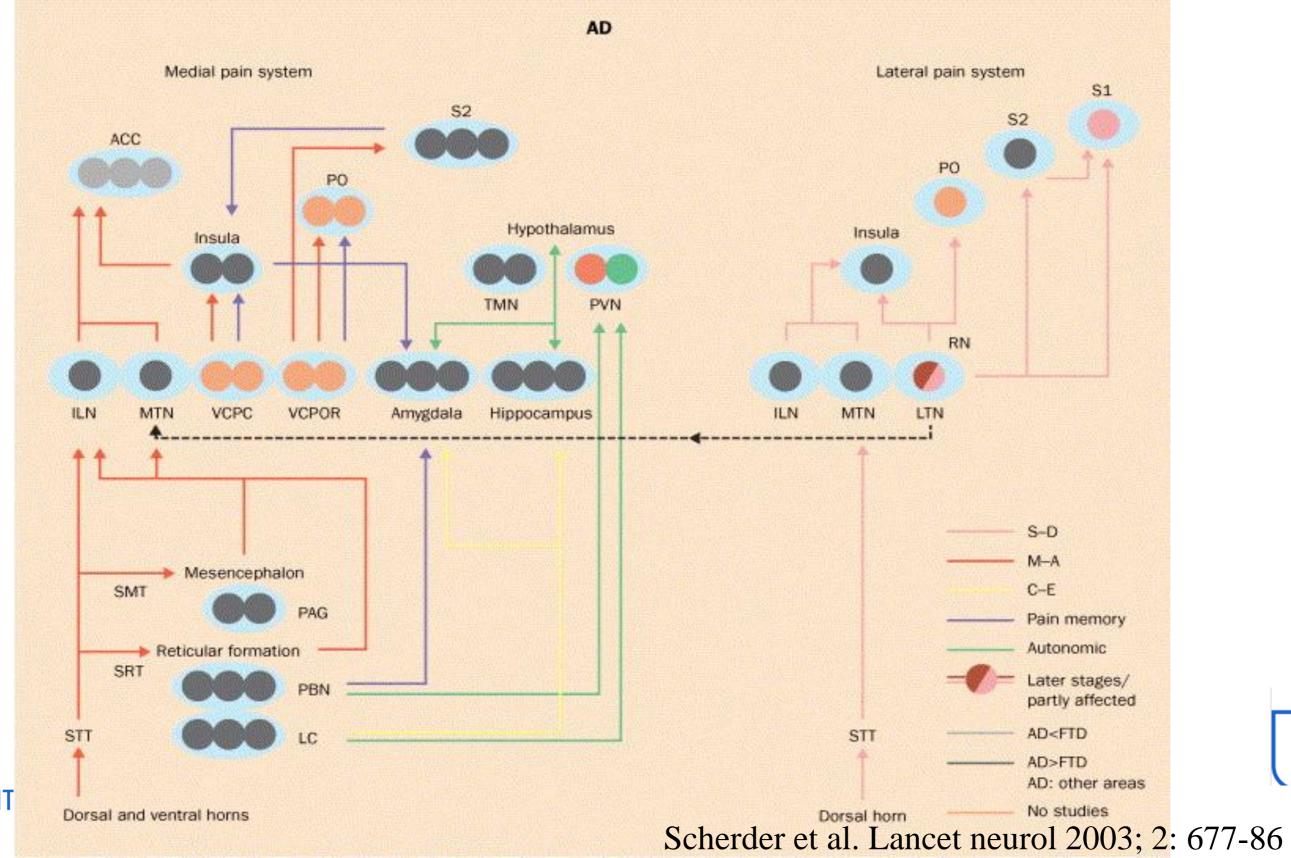


Decrease in autonomic response





#### ALZHEIMER DISEASE AND PAIN







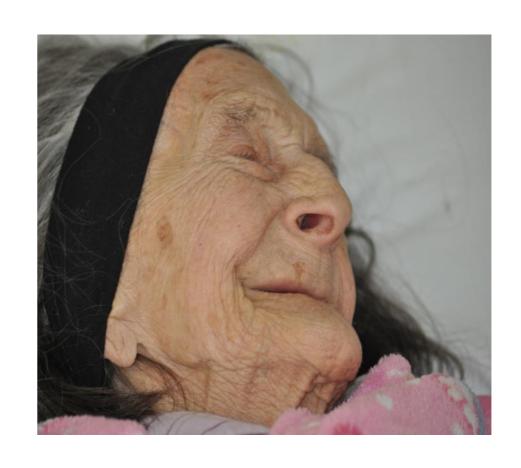
### **ROSA**

88 years old

Moderate to severe dementia

#### Since three days:

- Restlessness, crying and shouting especially when moved out of the room
- Hitting and biting nurses during toileting







## **ROSA**









**Table 3**Association of pain severity with behavioral and psychiatric symptoms.<sup>a</sup>

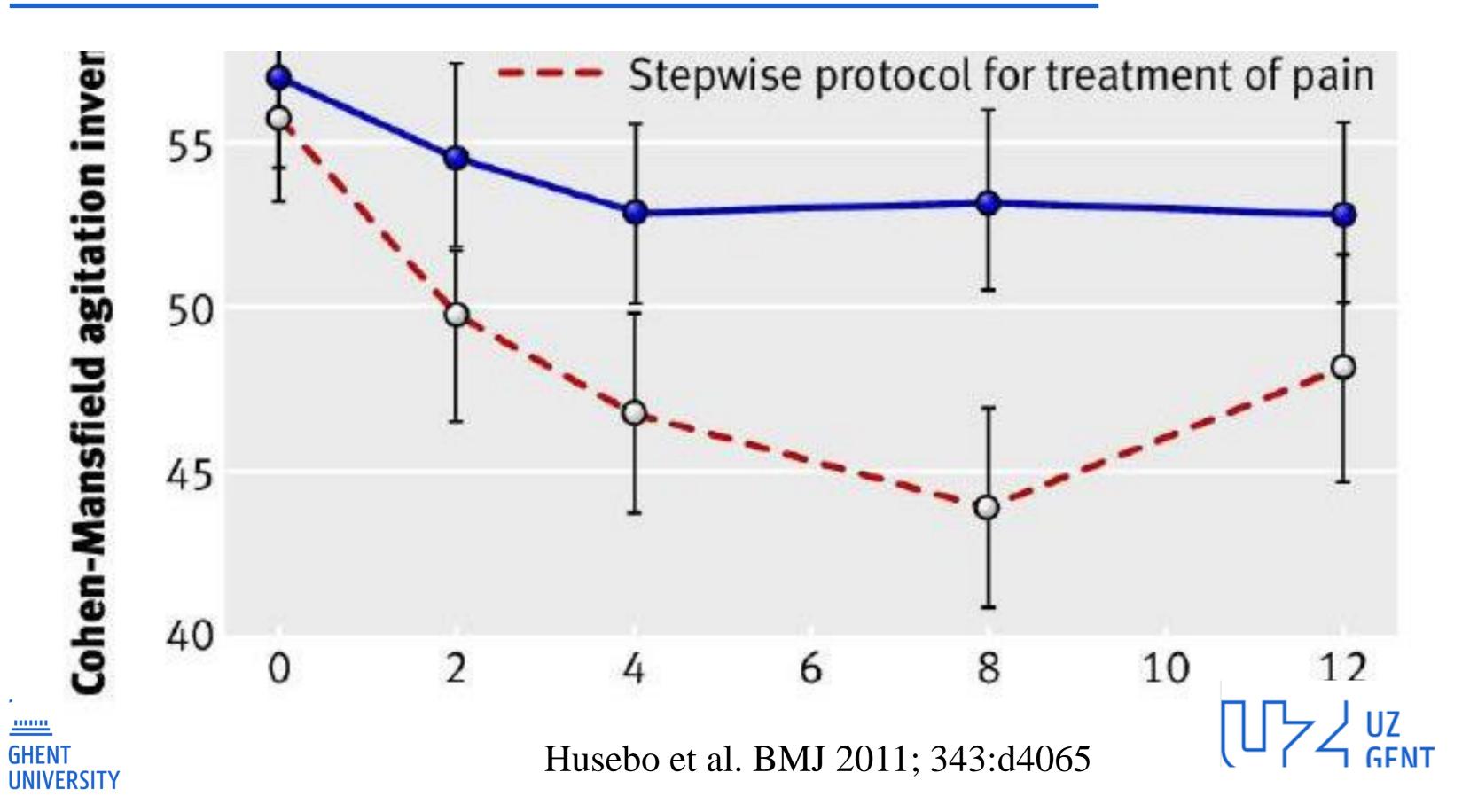
Symptom	No pain $(n = 2284)$	Mild (n = 159), OR (95% CI)	Moderate/severe/excruciating (n = 374) OR (95% CI)
Behavioral symptoms Wandering Verbal abuse Physical abuse Socially inappropriate behave Resists care	Socially inappro	opriate beha	aviour (.89) (.41) (.62) (.70) (.95)
1 or more behavioral symp  Psychiatric symptoms  Abnormal thought process	Delu Abnormal th	usions ought proce	.24) .97)
Delusions Hallucinations 1 or more psychiatric symptoms	Ref.	1.36 (0.68-2.72) 1.35 (0.90-2.00)	1.40 (0.87-2.26) 1.49 (1.14-1.95)

Ref., reference; OR, odds ratio; CI, confidence interval.

<sup>&</sup>lt;sup>a</sup> Data are adjusted for age, gender, country, cognitive impairment, number of diseases, ischemic heart disease, stroke, falls, communication problems, and a flare-up of a chronic or recurrent condition. Data on pain severity were not collected in 5 participants.



#### INFLUENCE OF TREATMENT ON BD



#### ALZHEIMER DISEASE AND PAIN: CONCLUSION

- Severity of dementia is probably related to the tolerance of pain
- Change in anticipation and reaction on pain
  - Decrease in recognition and understanding of pain
  - Behavioural changes can be the result of pain
- Decrease in autonomic response in AD
- Increase in facial expression in AD





#### PAIN AND DEMENTIA

	Experimental and clinical results			
Condition	Motivational-affective aspects of pain	Presence or intensity of pain		
Alzheimer's disease	<b>1</b>	Relatively unaffected		
Vascular dementia	<b>†</b>	Not examined		
Frontotemporal	<b>↓</b>	Not examined		
Parkinson (no cogn)	<b>†</b>	Not examined		





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#### PREVALENCE OF PAIN IN OLDER PEOPLE

Table 2. Prevalence of Pain in Participants with and without Dementia						
All Respondents, n = 802		Self-Report, n = 395 <sup>b</sup> Proxy Report, n = 407		Matched Cohort All Respondents, n = 80		
Pain	% (95% CI)					
Bothersome Activity limiting	63.5 (60.5–66.4) 43.3 (40.2–46.5)	62.7 (58.7–66.6) 40.1 (35.7–44.6)	64.4 (59.8–68.7) 46.6 (42.5–50.7)	54.5 (51.4–57.7) <sup>c</sup> 27.2 (25.2–29.2) <sup>c</sup>		





## PREVALENCE OF OLDER HOSPITALISED PEOPLE WITH DEMENTIA

**680** E.L. Sampson et al. • 156 (2015) 675–683

PAIN®

#### Table 2

Prevalence of pain in 230 older people with dementia and unplanned acute medical admission.

Pain	Time during admission, number (%)					
	At baseline, $n = 230$	At least once during admission, n = 230	All assessments n = 965*	Persistent, n = 138†		
Self-reported	54/200 (27.0)	84/218 (38.5)	196/821 (23.9)	8/117 (6.8)		
95% CI	(20.8, 33.2)	(32.0, 45.0)	(18.6, 27.5)	(2.2, 11.5)		
PAINAD scale ≥2						
Pain during rest	22/229 (9.6)	43/230 (18.7)	68/950 (7.2)	0/135 (0.0)		
95% CI	(5.8, 13.5)	(13.6, 23.8)	(5.3, 9.8)	-		
Pain during movement	97/229 (42.4)	131/230 (57.0)	331/946 (35.0)	21/135 (15.6)		
95% CI	(35.9, 48.8)	(50.5, 63.4)	(29.4, 39.0)	(9.4, 21.7)		

<sup>\*</sup> Prevalence for all assessments combined, estimated by generalised estimating equations.

<sup>†</sup> Defined in the population with 3 or more assessments, as in pain in at least 75% of the occasions.

Cl, confidence interval; PAINAD, Pain Assessment in Advanced Dementia scale.

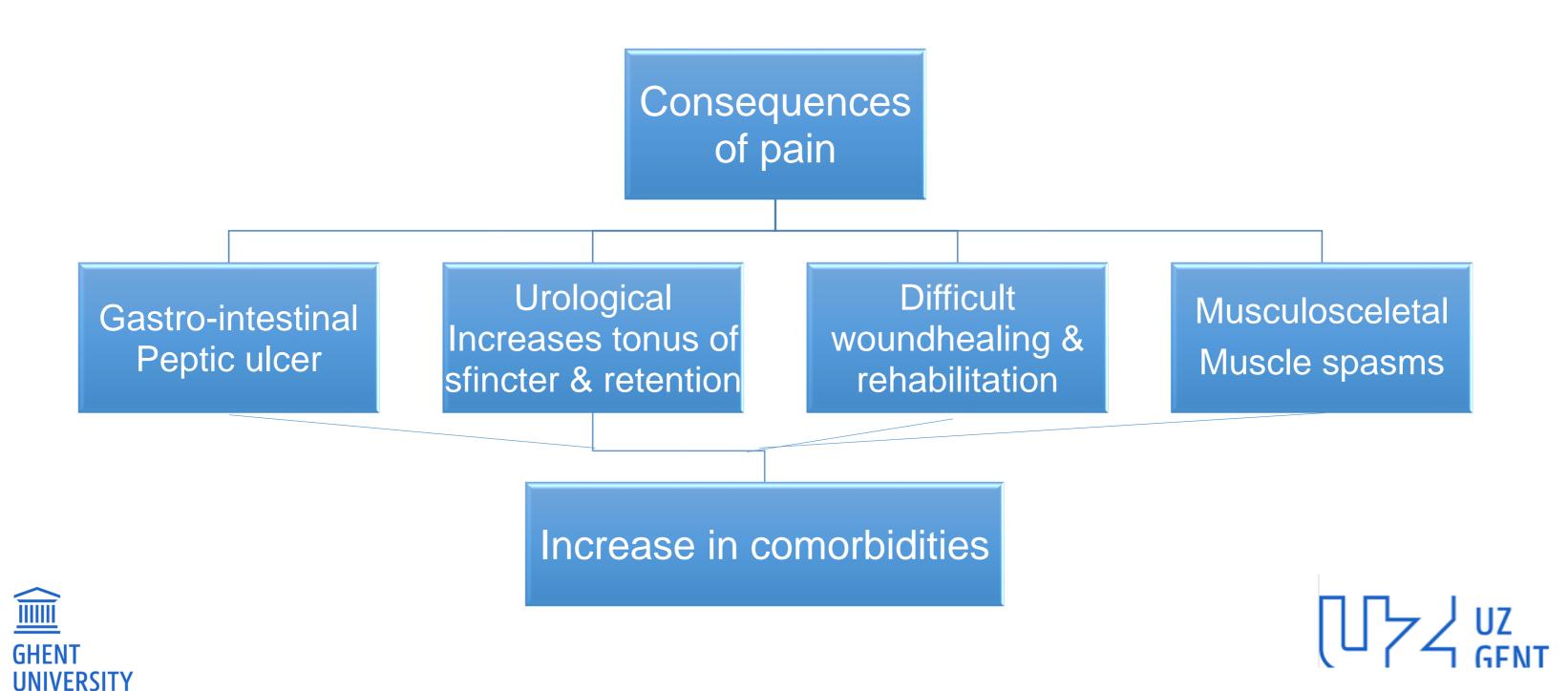
# ETIOLOGY OF PAIN IN PEOPLE WITH DEMENTIA

Number of patients reporting pain, n (%)	57 (44)
Etiology of pain, n (%)	
Osteoarthritis of joints	39 (68)
Back pain (osteoporosis or osteoarthritis)	3 (5)
Skin lesion	7 (12)
Other causes	8 (14)

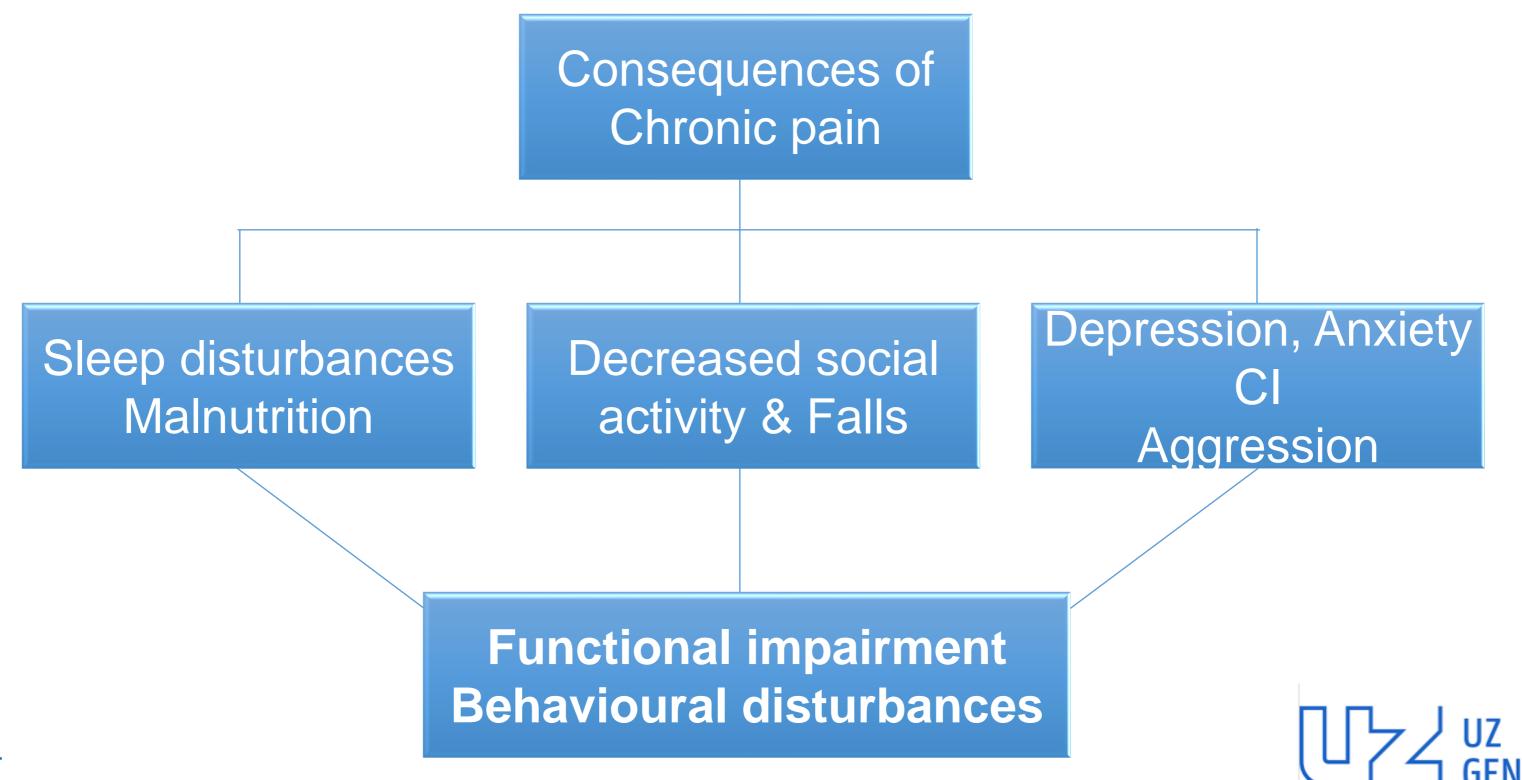




## Pain: consequences



#### CHRONIC PAIN: CONSEQUENCES





### **SUMMARY**

Influence of ageing and cognitive impairment on perception of

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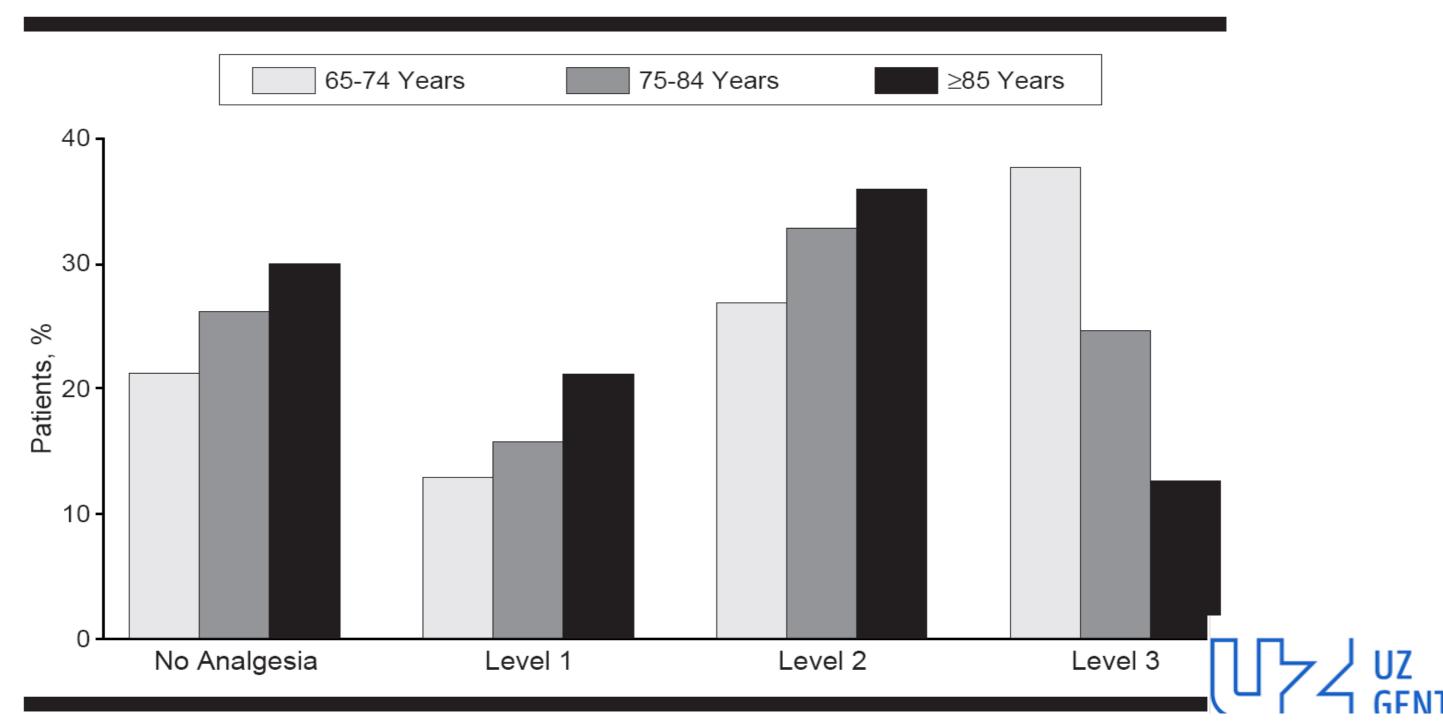
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## USE OF ANALGESICS IN THE OLDER PERSON WITH CHRONIC MALIGN PAIN





#### UNDER-TREATMENT OF PAIN

Table 2. Use of Analgesia and Report of Pain by Persons with Dementia (N = 115)

	Report of Pain <sup>†</sup> n (%)		
Analgesic Use*	Yes (n = 62)	No (n = 53)	
None	(42 (68))	45 (85)	
Any	20 (32)	8 (15)	
Acetaminophen	12 (60)	4 (50)	
Nonsteroidal antiinflammatory drug	5 (25)	4 (50)	
Weak opioid	3 (15)	О	
Strong opioid	0	0	





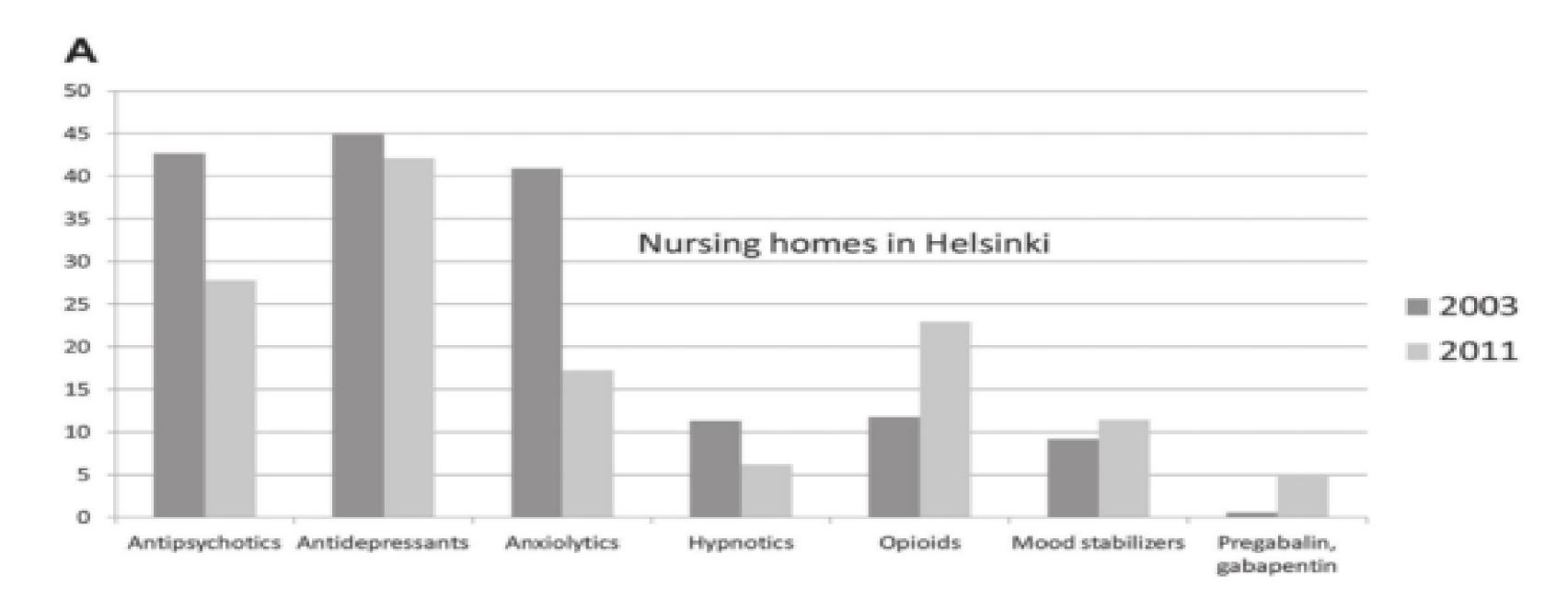
#### UNDER-TREATMENT OF PAIN

Table 4. Logistic Regression of Potentially Insufficient Analgesia on Pain Management Index According to Patient Demographics and Variables that Significantly Added to the Model

Characteristic	Odds Ratio	95% Confidence Interval	<i>P</i> -value
Age	1.07	1.01-1.14	.03
Male	1.06	0.40 - 2.82	.91
Lower education	1.04	0.71-1.52	.84
Advanced dementia*	3.08	1.05-9.10	.04
Impaired function†	2.50	1.01-6.25	04
Depression <sup>‡</sup>	2.13	0.82 - 5.52	.12





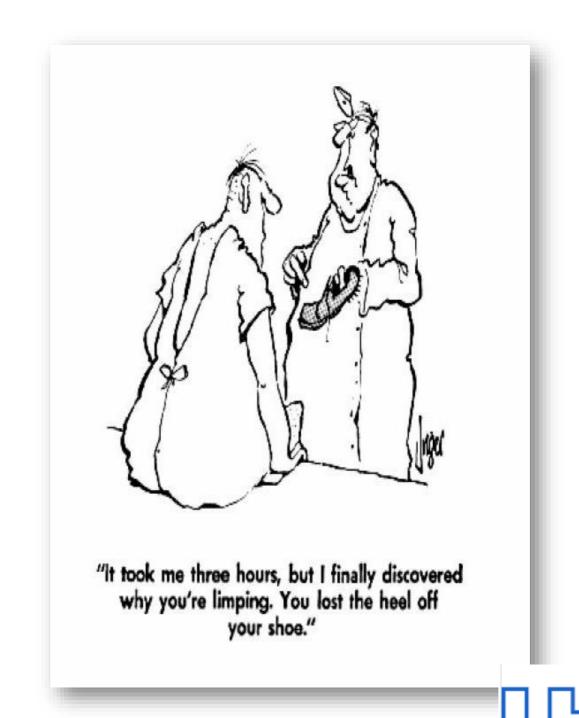






#### **UNDER-TREATMENT OF PAIN: AETIOLOGY**

- Pain is perceived as unavoidable with ageing
- High incidence of adverse drug reactions (ADR)
- Pain is not recognised
  - Communication of pain (especially in mild to advanced dementia) is diminished
  - No regular assessment for pain by the caregiver





## SYSTEMATIC EVALUATION

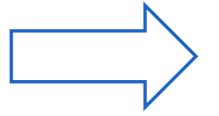
Geriatric problems	Before MGST	After MGST	Gain	р
ADL-IADL	26%(21-31)	89%(86-93)	63%(59-69)	< 0.0001
Incontinence	4%(3-9)	60%(55-65)	56%(48-59)	< 0.0001
Falls	35%(30-40)	46%(41-52)	11%(7-26)	0.1497
Cognition	34%(29-39)	68%(67-77)	34%(27-48)	< 0.0001
Depression	13%(9-17)	49%(43-54)	46%(33-43)	< 0.0001
Social	7%(5-11)	45%(44-55)	38%(35-50)	< 0.0001
Nutritional	17%(13-21)	65%(60-71)	48%(45-57)	< 0.0001
Pain	8%(5-11)	43%(38-49)	35%(32-42)	< 0.0001
Total of suspected problems / patient (mean±SD)	1.5±1.2	4.7±1.7	3.2±1.8	<0.0001



#### WHEN TO SCREEN FOR PAIN?

Assessing pain is as measuring temperature, tension

To be done as older person go worse



Pain = Fifth vital sign





#### HOW TO ASSESS PAIN?

1/ Evaluation of pain and intensity

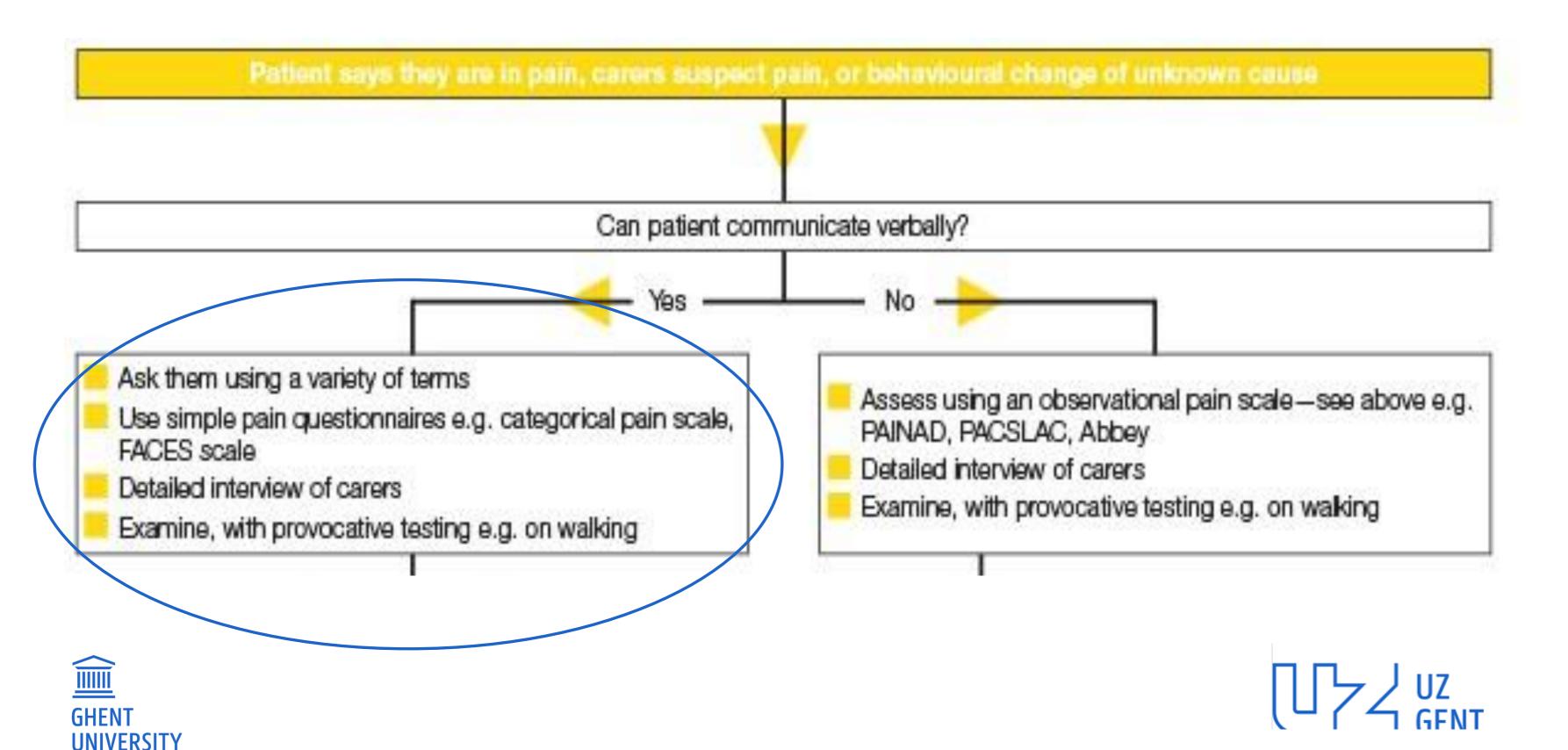


- 2/ Evaluation of consequences of pain
  - ADL/IADL/mobility
  - Eating/ rest at night
  - Memory/concentration/mood





#### ASSESSMENT OF PAIN



# ASSESSMENT OF PAIN IN PEOPLE WITH DEMENTIA – SCALES

Table 3. Number and Percentage of Patients Understanding Each Scale According to the Level of Dementia (CDR)

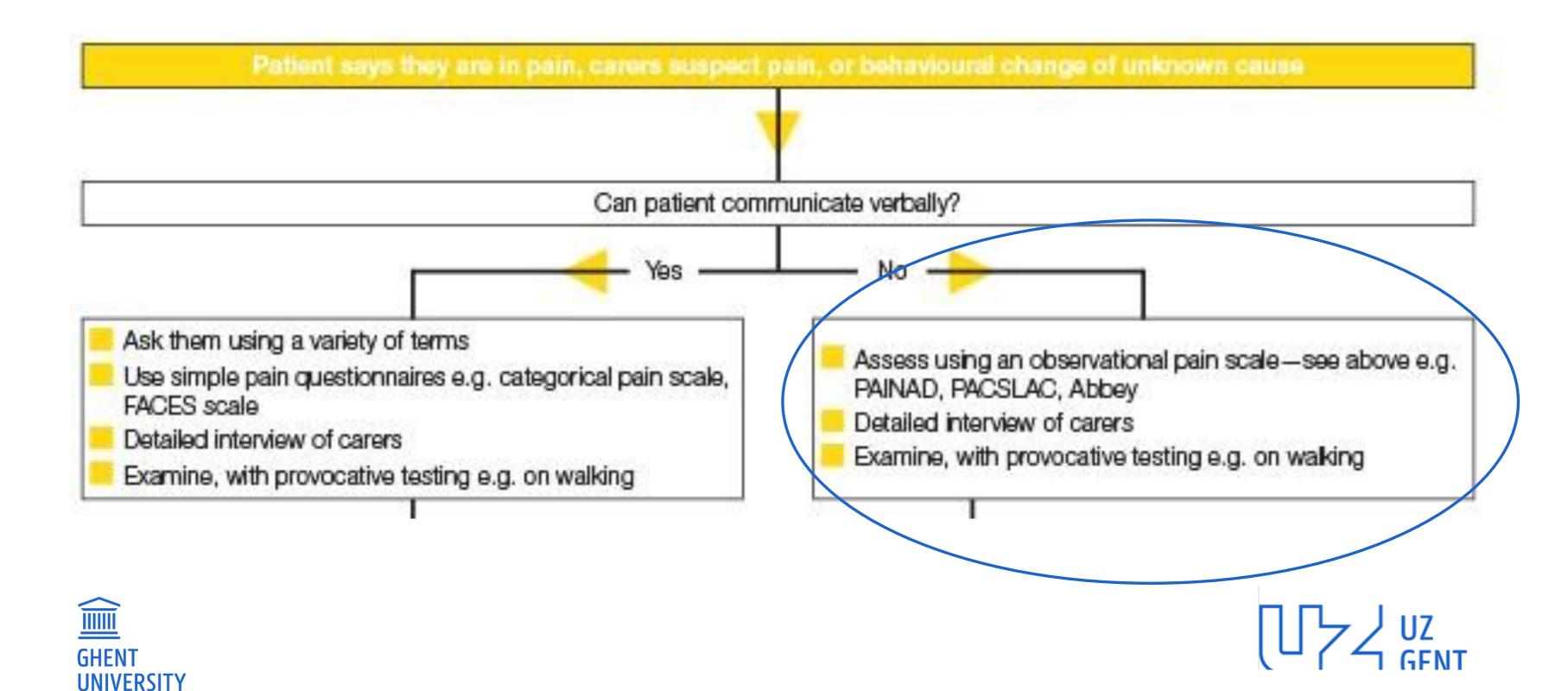
	CDR = 1		CDR = 3	Total
Scale	(64 Cases) N (%)	(81 Cases) N (%)	(15 Cases) N (%)	(160 Cases) N (%)
Verbal Rating Scale (VRS)	58 (91)	59 (73)	5 (33)	122 (76)
Horizontal Visual Analong Scale (HVAS)	62 (97)	64 (79)	4 (27)	130 (81)
Vertical Visual Analog Scale (VVAS) Faces Pain Scale (FPS)	59 (92) 57 (89)	60 (74) 53 (65)	4 (27) 4 (27)	123 (77) 114 (72)

*Notes*: For each scale, comprehension is significantly associated with the CDR (Clinical Dementia Rating) scale.

p < .001 (Fisher's exact test).



#### ASSESSMENT OF PAIN



#### NON VERBAL PAIN INDICATORS

#### Common Pain Behaviours in Cognitively Impaired Elderly Persons

#### **Facial expressions**

Slight frown, sad frightened face
Grimacing, wrinkled forehead, closed or tightened eyes
Any distorted expression
Rapid blinking

#### **Changes in interpersonal interactions**

Aggressive, combative, resisting care Decreased social interactions
Withdrawn

#### Verbalisations, vocalisations

Sighing, moaning, groaning Grunting, chanting, calling out Noisy breathing Asking for help Verbally abusive

#### Changes in activity patterns or routines

Refusing food, appetite change Increase in rest periods Sleep, rest pattern changes Increased wandering

#### **Body movements**

Rigid, tense body posture, guarding Fidgeting Increased pacing, rocking Restricted movement Gait or mobility changes

#### Mental status changes

Crying or tears
Increased confusion
Irritability or distress





### Panel 1: Instruments suitable for the assessment of pain in the elderly adult with dementia

- Abbey Pain Scale<sup>7283-84</sup>
- Checklist of Non-Verbal Pain Indicators (CNPI)788485
- Certified Nursing Assistant Pain Assessment Tool (CPAT)<sup>7586</sup>
- DOLOPLUS-2<sup>87,88-90</sup>
- Discomfort Scale in Dementia of the Alzheimer's Type (DS-DAT/DS-DAT modified)<sup>93-95</sup>
- EPCA-2<sup>96</sup>
- Mahoney Pain Scale<sup>97</sup>
- Mobilization-Observation-Behaviour-Intensity-Dementia (MOBID and MOBID-2) Pain Scale<sup>74:98:99</sup>
- Non-Communicative Patient's Pain Assessment Instrument (NOPPAIN)5373,85100
- Pain Assessment in the Communicatively Impaired (PACI)<sup>103-103</sup>
- Pain Assessment Checklist for Seniors with Limited Ability to Communicate (PACLSAC and PACSLAC-II)<sup>273,85,104-107</sup>
- Pain Assessment for the Dementing Elderly (PADE)<sup>85 108</sup>
- Pain Assessment in Advanced Dementia (PAINAD)<sup>53,71,83,85,109</sup>
- Pain Assessment in Noncommunicative Elderly Persons (PAINE)<sup>76</sup>
- The Rotterdam Elderly Pain Observation Scale (REPOS)<sup>110</sup>





### PAIN ASSESSMENT IN ADVANCED DEMENTIA SCALE (PAINAD)

Items	0	1	2	Score
Breathing independent of vocalization	Normal	Occasional laboured breathing. Short period of hyperventilation	Noisy laboured breathing. Long period of hyperventilation. Cheyne-Stokes respirations.	
Negative vocalization	None	Occasional moan or groan. Low- level speech with a negative or disapproving quality	Repeated trouble calling out. Loud moaning or groaning. Crying.	
Facial expression	Smiling or inexpressive	Sad. Frightened. Frown.	Facial grimacing.	
Body language	Relaxed	Tense. Distressed pacing. Fidgeting.	Rigid. Fists clenched. Knees pulled up. Pulling or pushing away. Striking out.	
Consolability	No need to console	Distracted or reassured by voice or touch.	Unable to console, distract or reassure.	
			Total**	1 [



Warden et al, 2001.

	Abbey Pain Scale For measurement of pain in people with dementia who cannot verbalise.					
How	How to use scale: While observing the resident, score questions 1 to 6.					
Name	of resident :					
Name	and designation of person completing the s	cale :				
Date	Time :					
Lates	t pain relief given was		at.	hrs.		
Q1.	Vocalisation eg whimpering, groaning, crying Absent 0 Mild 1 Moderate 2 Sev	ere 3	C	21		
Q2.	Facial expression eg looking tense, frowning, grimacing, loo Absent 0 Mild 1 Moderate 2 Sev	king frightened ere 3	C	Q2		
<b>Q</b> 3.	Change in body language eg fidgeting, rocking, guarding part of bod Absent 0 Mild 1 Moderate 2 Sev		¢	Q3		
Q4.	Behavioural Change eg increased confusion, refusing to eat, al Absent 0 Mild 1 Moderate 2 Sev		patterns	24		
Q5.	Physiological change eg temperature, pulse or blood pressure o perspiring, flushing or pallor Absent 0 Mild 1 Moderate 2 Sev		nits,	Q5		
Q6.	Physical changes eg skin tears, pressure areas, arthritis, cor previous injuries Absent 0 Mild 1 Moderate 2 Sev	ntractures, ere 3	¢	Q6		
	Add scores for 1 - 6 and record here					
Tota	Pain Score 0 - 2 No pain	3 - 7 Mild	8 - 13 Moderate	14 + Severe		
	ly, tick the box which matches ype of pain	Chron	ic Acute	Acute on Chronic		
	Abbey, J; De Bellis, A; Piller, N; Estern Funded by the JH & JD Gunn Medi					





### <u>ALGOPLUS</u>

HOIII - I TOHOIII.

Date de l'évaluation de la douleur											
Heure											
	oui	non									
1 Visage											
Froncement des sourcils, grimaces, crispation, mâchoires serrées, visage figé.											
2 Regard											
Regard inattentif, fixe, lointain ou suppliant, pleurs, yeux fermés.											
3 Plaintes											
« Aie », « Ouille », « J'ai mal », gémissements, cris.											
4 Corps											
Retrait ou protection d'une zone, refus de mobilisation, attitudes figées.											



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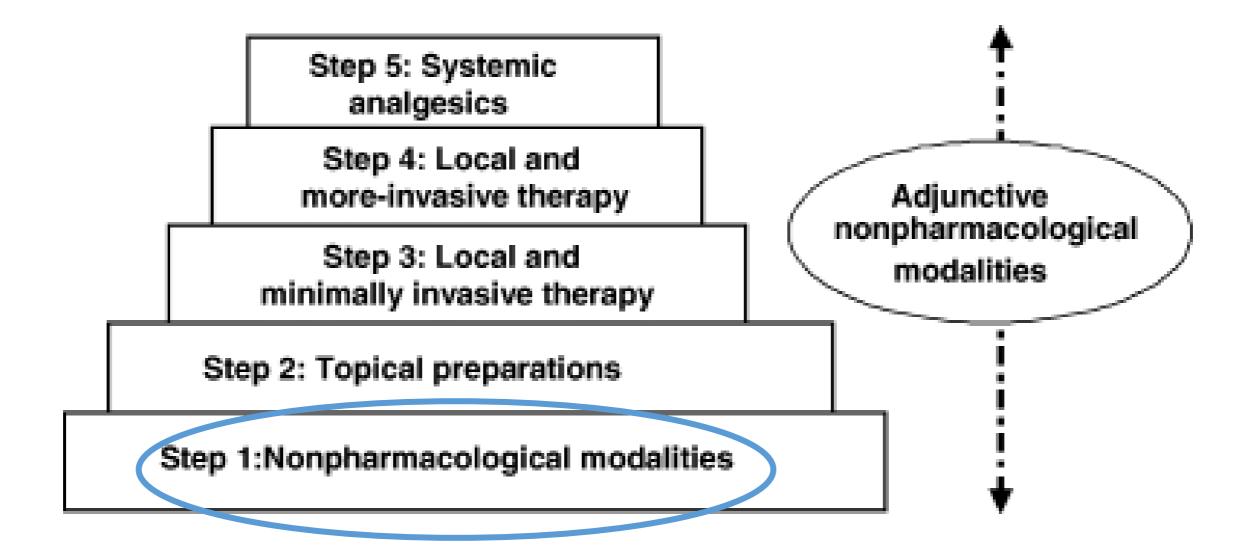
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# MANAGEMENT OF PAIN: STEPWISE APPROACH





### NON-PHARMACOLOGICAL APPROACHES

Approach	Considerations	
Physical therapy		
Exercise	Recommended pain management strategy	
	Inconsistent evidence whether one type of exercise is better than another	
	Patient preference is the primary consideration	
	Focus on strengthening, flexibility, endurance, and balance	
	Individual capacity limits options	
Foot orthotics, patellar taping	Foot orthotics may change gait pattern/muscle activation and reduce joint loading	
Manual therapy	Requires significant levels of skill and care	
TENS	Consider for persistent pain when patient can provide accurate feedback	
Physical modalities (eg heat)	Beneficial for acute pain as effects are transient	
	Monitor for safety if used for patients with dementia	





### NON-PHARMACOLOGICAL APPROACHES

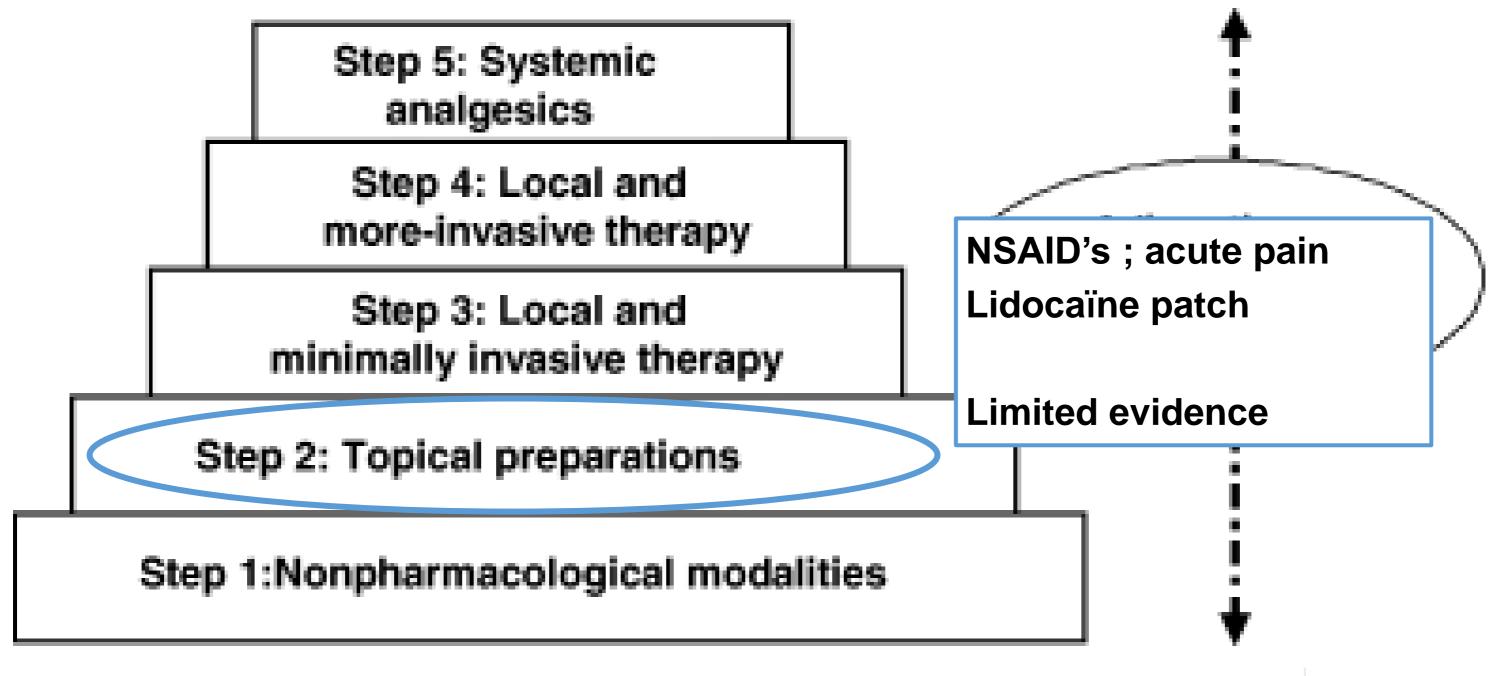
Occupational therapies			
Assistive devices (eg walking frames)	Some evidence of reducing functional decline and pain interpoleS		
Psychological approaches	Can increase pain it used incorrectly  of these there		
Cognitive behaviour therapy	Some evidence of reducing functional decline and pain interapies  Can increase pain if used incorrectly  Demonstrated benefit for some care Recommended in professional  Enteraprical can interfere with professional  Enteraprical care  Can interfere with some care  Recommended in professional  Can interfere with some care  Recommended in professional  Can interfere with some care  Recommended in professional  Can increase pain if used incorrectly  Enteraprical care  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Demonstrated benefit for professional  Can increase pain if used incorrectly  Can increase pain if used incorrectly  Can increase pain if used incorrectly  Can increase pain if used incorrect		
Complementary and alternative medicine	can interfer		
Acupuncture	ration or for older people as adjunctive therapy		
1 of	May improve function and pain relief		
aitive dec	Duration of long-term effects are uncertain		
Massage, Tai Chi, yoga CO9	Consider for older people as adjunctive therapy		
	Massage may have some benefit for non-specific lower back pain		
Nutritional supplements	Some evidence that chondroitin and glucosamine improve pain and function in osteoarthritis		

TENS, transcutaneous electrical nerve stimulation





#### TREATMENT OF PAIN

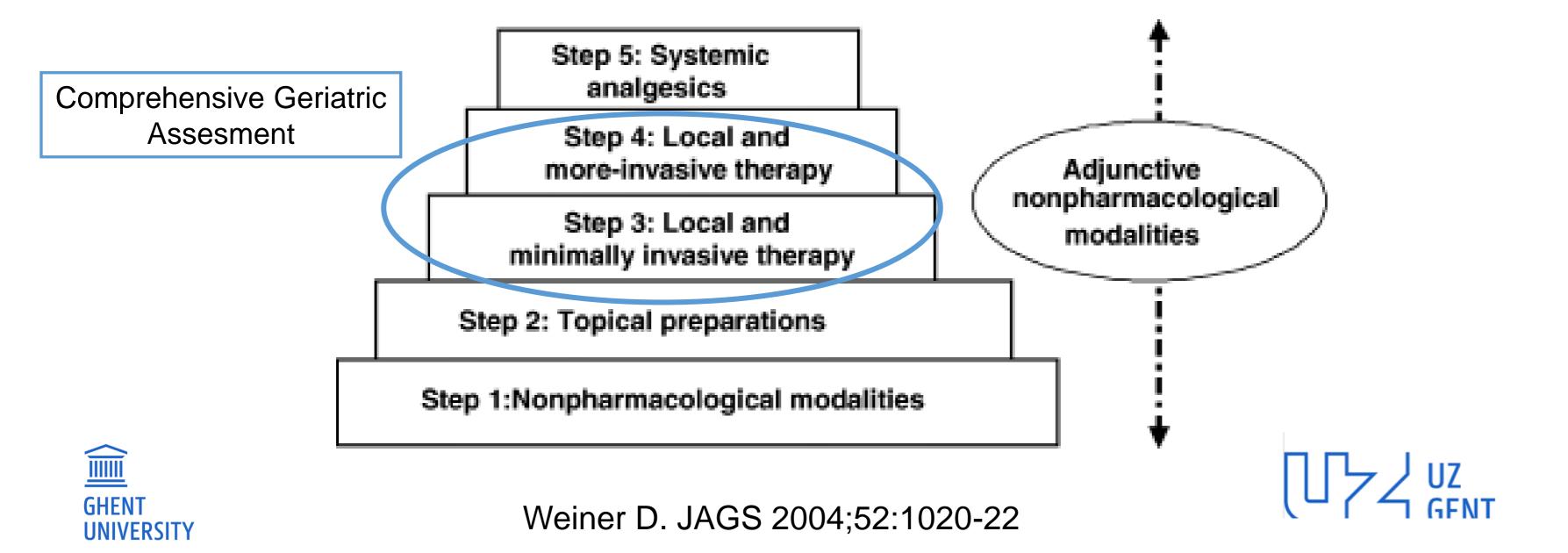




Massey T,et al. Topical NSAIDs for acute pain in adults. *Cochrane Database of Systematic Reviews* 2010, Issue 6. Art. No.: CD007402. Wolff et al. Acta Neurol Scand 2011;123:295-309.



# MANAGEMENT OF PAIN: STEPWISE APPROACH



# MANAGEMENT OF PAIN: STEPWISE APPROACH

No trials in persons with dementia Step 5: Systemic analgesics Step 4: Local and more-invasive therapy Adjunctive nonpharmacological Step 3: Local and modalities minimally invasive therapy Step 2: Topical preparations Step 1:Nonpharmacological modalities Weiner D. JAGS 2004;52:1020-22



# UNDER-TREATMENT OF PAIN: ADVERSE DRUG REACTIONS

#### Adverse drug reactions

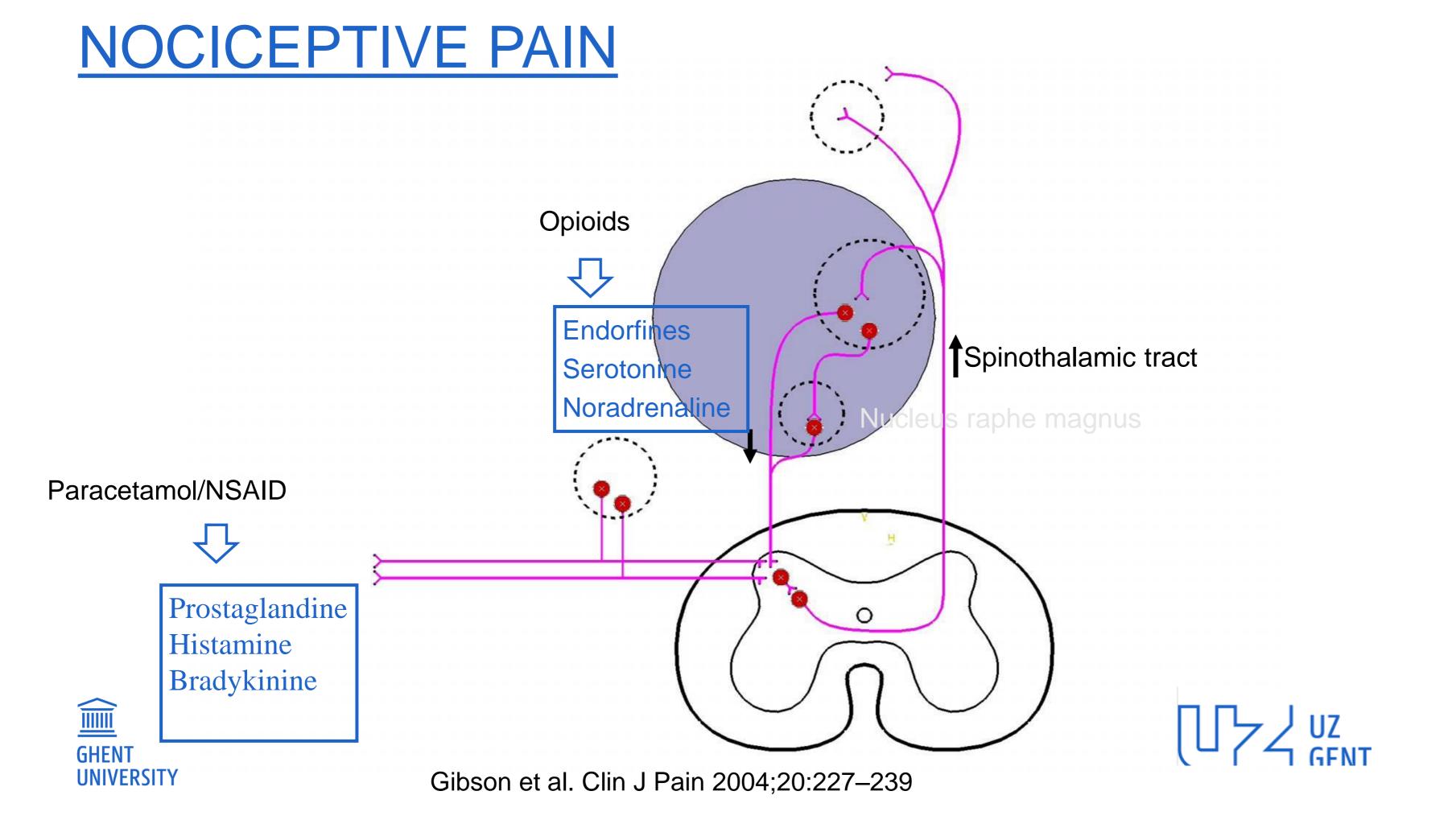
Pharmacokinetic and –dynamic changes with ageing
 -higher risk for drug drug interactions

Avoidance of prescription

Prescription and higher surveillance







### **WHO-LADDER**

Persistant Pain

Minimal invasive technic **4** 

Morfine

Oxycodone

Fentanyl

Buprenorphine

(Hydromorphone)

Tapentadol

Persistant Pain Strong-opioid **±** Non-opioid

± adjuvant

Persistant

Pain

Weak-opioid

**±** Non-opioid

± adjuvant

Hydrocodone

Tramadol

Non-opioid ± adjuvant (anti-epileptic, anti-depressive)

1

Paracetamol NSAID



UZ GENT

System	Change with ageing	Clinical consequence	
Absorption and function of the GI tract	Reduced:  • Motility of the large intestine  • Vitamin absorption by active transport mechanisms  • Splanchnic blood flow  • Bowel surface area	Passive diffusion-little change in absorption with age	
	Delayed gastric emptying and reduced peristalsis	Increased risk of GI-related side effects	
	Decreased body water	Reduced Morphine, tramadol, oxycodone	
Distribution	Increased body fat and accumulation of lipid-soluble drugs	Lipid-soluble drugs have longer effective ha phentanyl.	
	Decreased serum albumin and altered protein binding	Increased potential for drug-drug interactions	
II	Decreased hepatic blood flow	First-pass metabolism can be less effective morphine	
Hepatic-biliary	Reduced liver mass	Phase I metabolism of some drugs might be slightly impaired	
	Reduced renal blood flow	Morphine, tramadol, oxycodone,	
Renal excretion	Reduced glomerular filtration	Reduced ex phentanyl, gabapentine,	
	Reduced tubular secretion	Ridiicy	
Pharmacodynamic	Decreased receptor density	Increased sensitivity to the therapeutic and side effects	
changes	Increased receptor affinity	opioids,	

### **RECOMMENDATIONS**

- Start low, go slow
- Around the clock (24hour)
- Compliance is important; inform caregiver
- For neuropathic pain, combine different classes of drugs
- Know pharmacokinetics and dynamics
- Take into account interactions and polypharmacy
- Inform caregiver regarding the side effects and advise regular monitoring



	Pitfalls	Recommendations
Paracetamol	Liver failure in malnutrition	Low dose; max 3g/24h
NSAID	Gastro-intestinal bleeding	Associate PP
	Fluid retention	Monitor arterial tension and weight – stop treatment if there is a substantial increase
	Acute kidney failure	Monitor serum creatinine after three days





### EVIDENCE OF WEAK/STRONG OPIOIDS?

No well performed studies in the elderly

 Extrapolation from studies in younger patients and specific patient groups

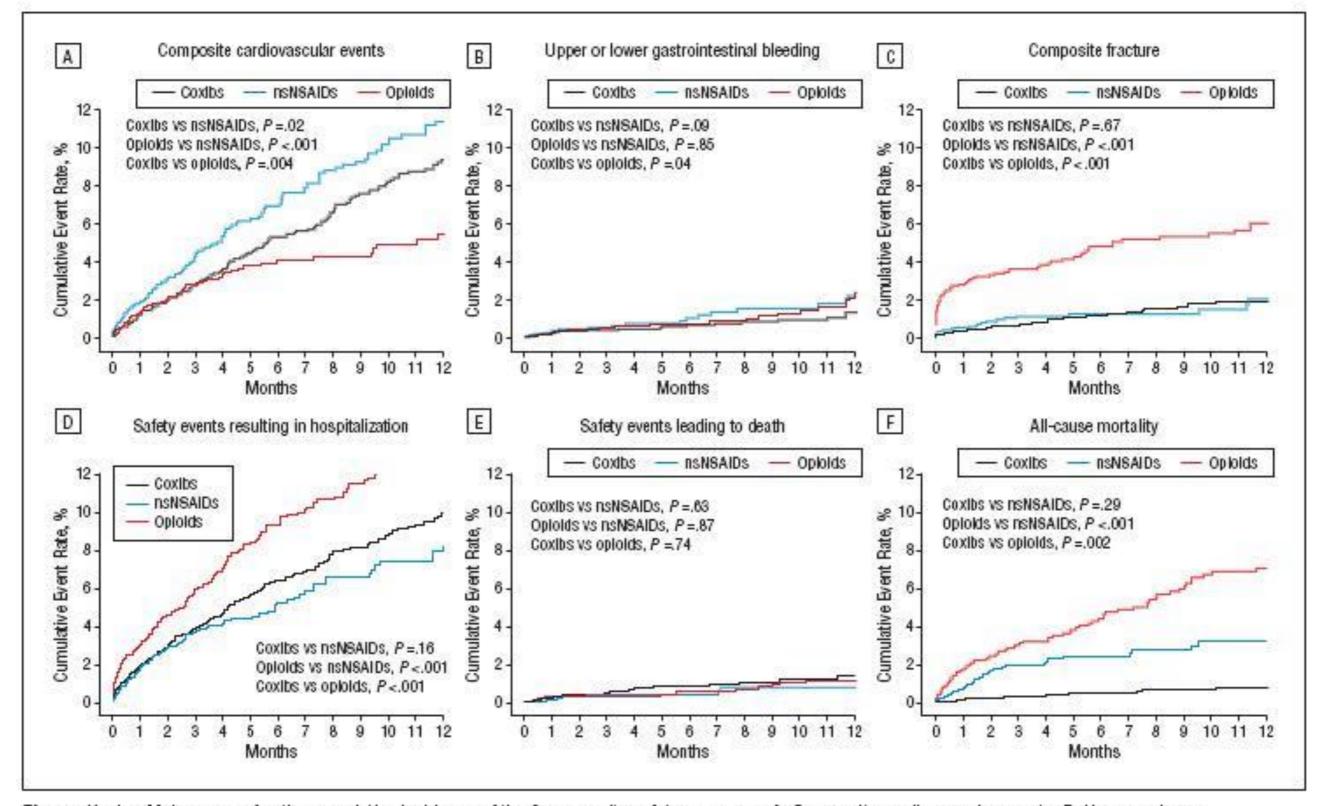
 Some open label studies available in older patient population showing acceptable safety





#### SAFETY OF OPIOIDS?

cyclooxygenase-2 inhibitors; nsNSAIDs, nonselective nonsteroidal anti-infla









	Pitfalls	Recommendations
Tramadol	Renal clearance Diminish seizure threshold Bind to serotonin receptors	Reduce dose in CKD** Avoid use in seizure patients Monitor serotonergic syndrome
Morphine	Higher plasma concentrations Renal clearance	Start low dose (2.5mg OR; 1mg IV) Reduce dose in CKD stage 3 Avoid in CKD stage 4 and 5
Fentanyl Buprenorphine	Take 2à3 patches to achieve stable plasma concentrations	Wait one week before increasing dose – foresee IR morphine for breakthrough pain (1/10 of equivalent dose)



Symptoms	Consider				
Temporally side effects – disappear normally after a few days					
Sedation / sleepiness	Dose reduction of concomitant medication as anxiolytics				
Hallucinations/ delirium  Dose reduction of concomitant medication as  Gabapentin, Pregabaline, Amitriptyline  Association of low dose haloperidol					
Nausea/vomiting	Association of gastro-kineticum  Domperidon, Metoclopramide, Alizapride  If persistent: association of low dose haloperidol				
Urinary retention	Intermittent catheterization				
Persistent side effects					
Constipation	Non pharmacological therapy (fluid, exercise, fibers)  To start laxatives together with the initiation of opioid therapy  Macrogol, Osmotic laxatives  If persistent: associate stimulating laxatives or enemas  Bisacodyl, Picosulfaat  If persistent: Methylnaltrexone or naloxone in combination with opioid				
Risk of falling/fracture	Fall assessment and prevention  Adding walking aids				
UNIVERSITY					

#### TREATMENT OF NEUROPATHIC PAIN

- Inhibition of the action potential in neuron: Anti-epileptics
  - (carbamazepine, natriumvalproaat)
  - Gabapentine, Pregabaline
- Support of the non-opioïd modulating neurons (serotonine en noradrenaline)
  - Tricyclic antidepressants (amitryptiline)
  - SNRI (duloxetine en venlafaxine)
- Weak and strong opioids
  - Tramadol; buprenorfine; fentanyl, oxycodon





	Pitfalls	Recommendations
Tricyclic antidepressant	Anticholinergic side effects	Monitor urinary retention; glaucoma worsening cognition Avoid in patients with cardiac
	Cardiac arrhythmias	arrhytmias
SNRI	Syndrome of Inappropriate ADH	Monitor natremia Dizziness, sedation, Arterial hypertension, tachycardia
Anticonvulsant therapy	Renal clearance  Side effects Takes 2-3 weeks before clinical effect	Dose reduction CKD3: 50%; CKD4: 25% of dose Avoid in CKD stage 5 Monitor: Dizziness, sedation Inform patient/family

### <u>THOM'S</u>

- Pain is important in the older person
- Think about and Assess pain
- Discuss treatment plan put achievable goals and evaluate
- Take into account changing pharmacokinetics with ageing
- Know most important side effects in the older person and teach them
- If therapy is not working, consider other influencing factors (psychosocial, financial etc ...)



