



28 Juni 2014

Kwaliteit versus volume: wat heeft het PROCARE project ons geleerd?

P R O C A R E

PROJECT ON CANCER OF THE RECTUM

Penninckx Freddy
namens PROCARE

Kwaliteit versus volume in de aanpak van rectumkanker

- PROCARE: doel en methode
- Kwaliteit: variabiliteit en volume effect
- Effect size van volume
- ‘Kwaliteitssysteem’ belangrijker dan centralisatie

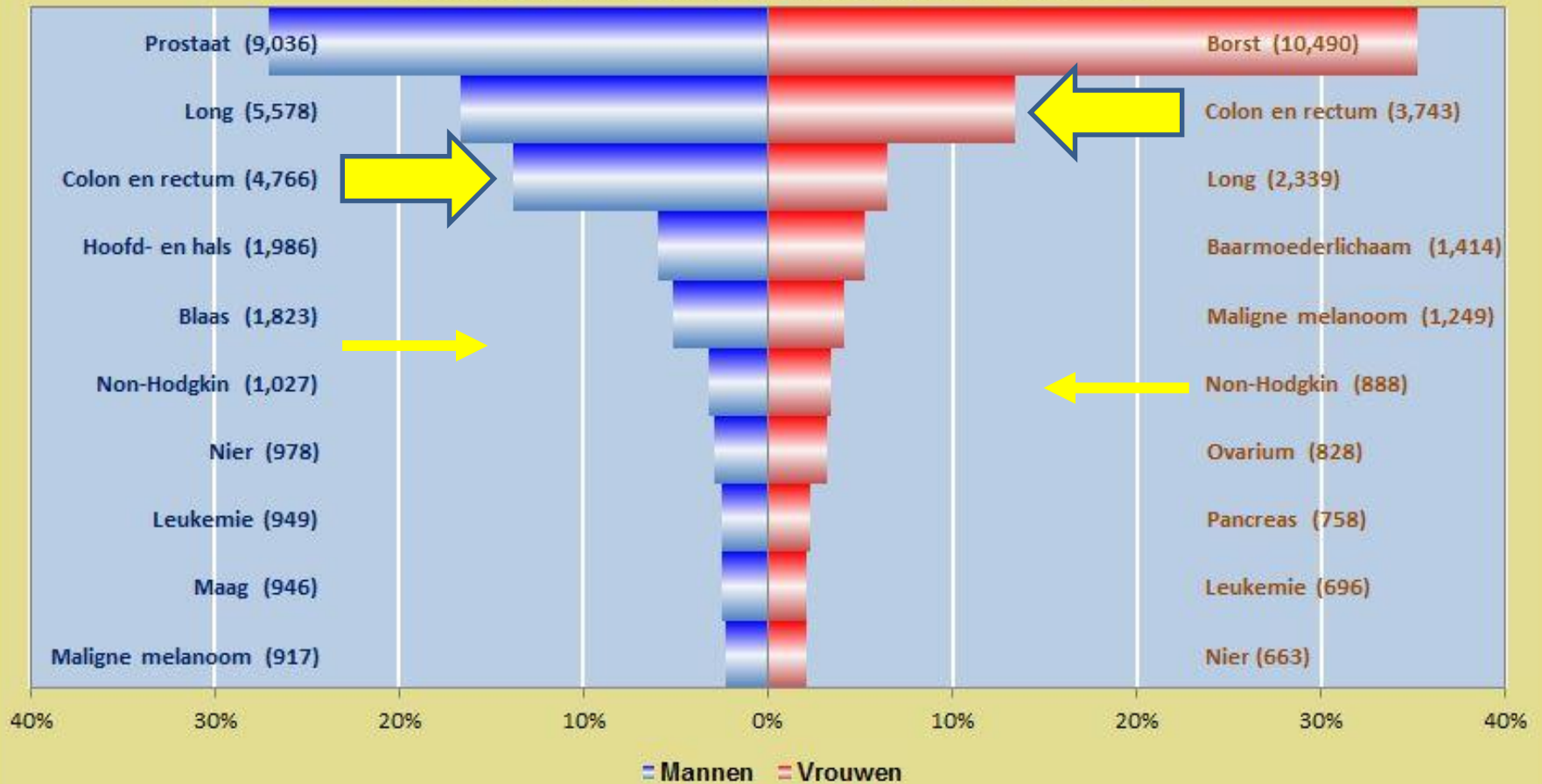
PRO CARE
PROJECT ON CANCER OF THE RECTUM


Belgian Cancer Registry


IMA-AIM

De tien meest frequente tumoren per geslacht, België 2011

Bron: Stichting Kankerregister (www.kankerregister.org/statistieken)

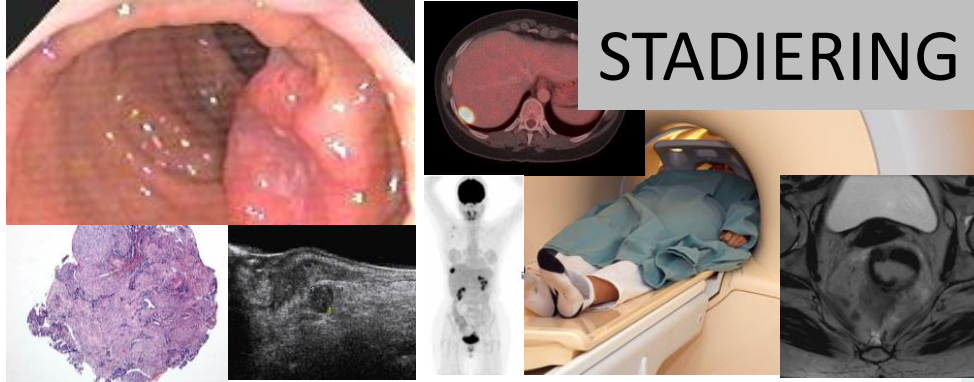


1507 + 886 = 2393 rectumkanker

6116 colonkanker (incl. rectosigmoidale junctie)

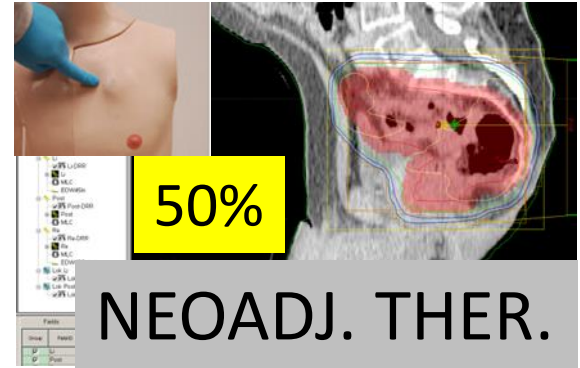
De multimodale aanpak van rectumkanker

STADIERING



50%

NEOAJ. THER.



82%

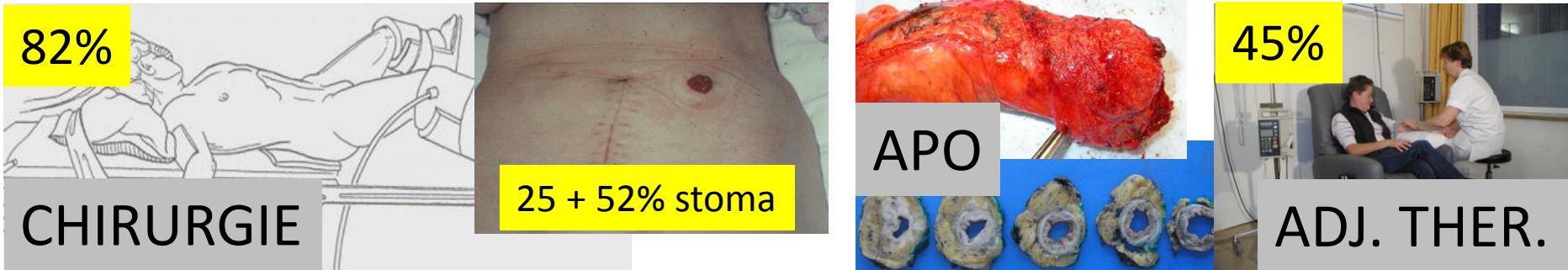
CHIRURGIE

25 + 52% stoma

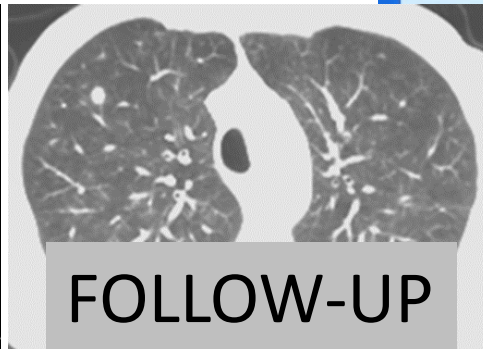
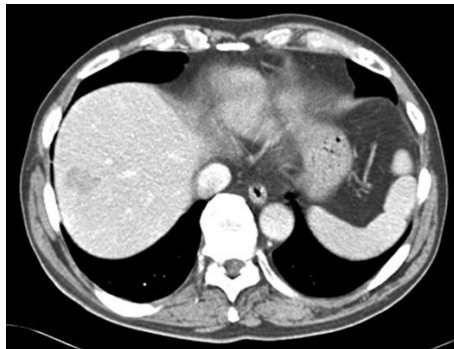
APO

45%

ADJ. THER.



FOLLOW-UP





2004 - 2005

PRO CARE

PROJECT ON CANCER OF THE RECTUM

2006 - 2014



Belgian Cancer Registry

PROCARE AIMS

reduce variability & improve outcome
for all aspects and stages of RC

- Multidisciplinary
- Profession-driven, all centers/teams
 - Voluntary participation
- Educational (confidentiality)

PROCARE METHODS

Guidelines

Implementation of guidelines

Quality of Care Indicators

Registration (151 items) for
Benchmarking and
Feedback

www.kankerregister.org

www.registreducancer.org

Assurance de Qualité pour
le cancer du rectum
– Phase I -
Recommandation de bonne pratique
pour
la prise en charge du cancer rectal

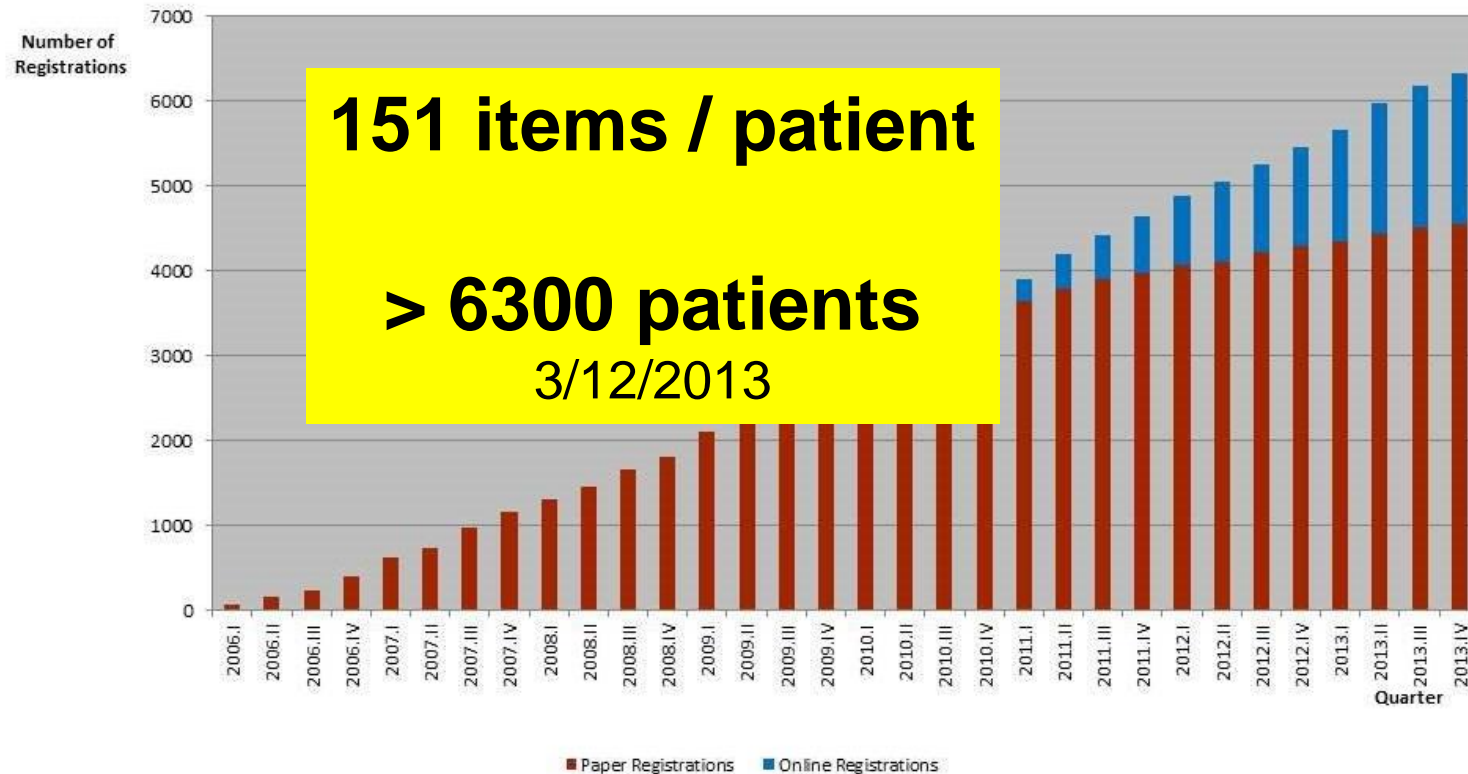
KCE reports 69B

Kwaliteit van rectale
kankerzorg – Fase 2:
ontwikkeling en test van een
set van kwaliteitsindicatoren

KCE reports 81A

Een grote inspanning op vrijwillige basis, maar ...

Cumulative Number of Registrations per Quarter



Registratie in PROCARE t.o.v. BKR/IMA (2006-2008)

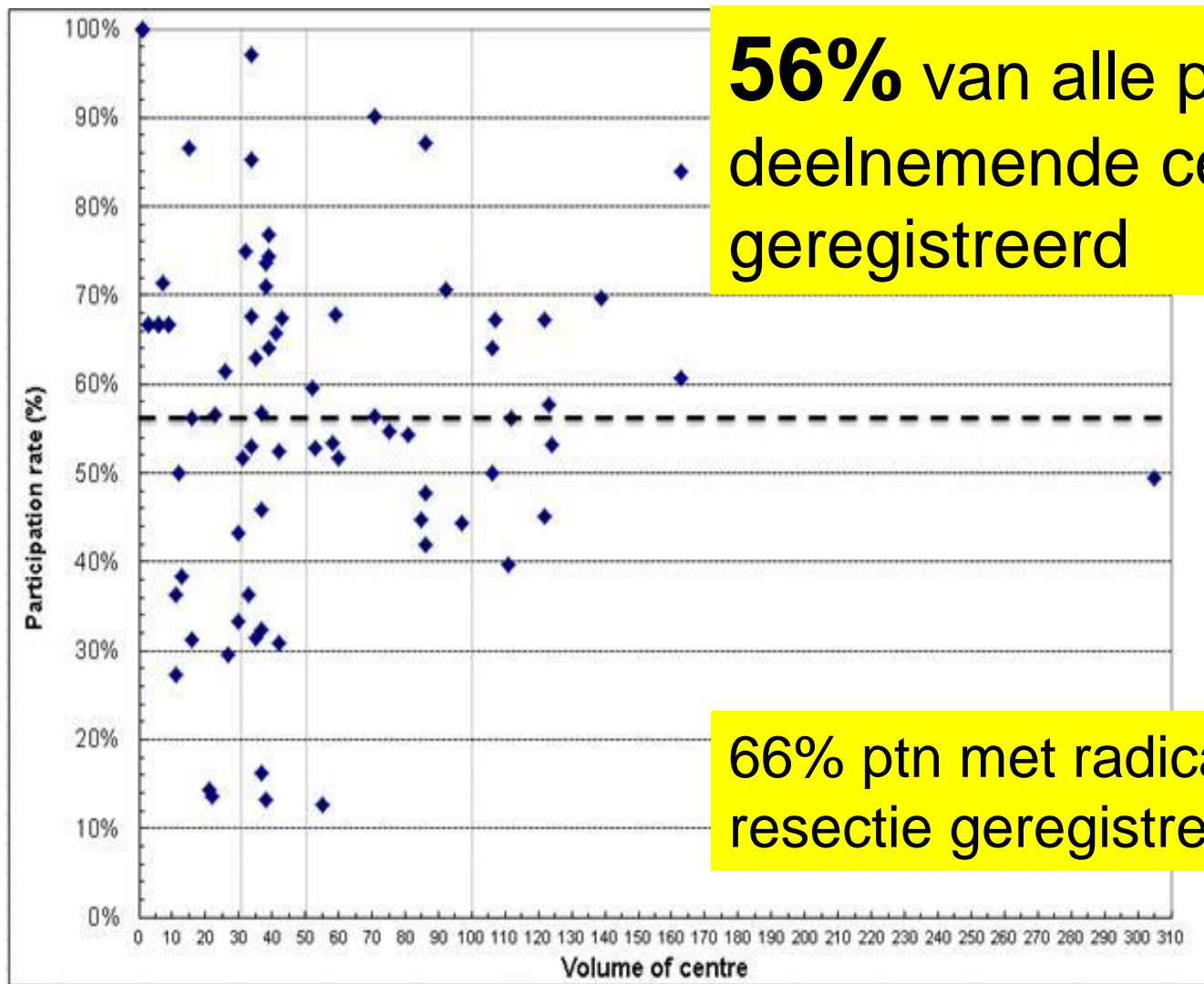
- 37 % van alle rectumkankers (C-20)
- 43 % van ptn met radicale resectie

- Door onvolledige deelname
- Door onvolledige registratie

... onvolledige deelname aan PROCARE (2006-08)

78 (62-67) / 108 centra	Beds	Participation (%)
Type		
Univ. hospitals	15%	100
Gen. hosp. with univ. beds	85%	65
General hospitals		71
Region		
Flanders		75
Wallonia		51
Brussels Capital		93
Volume of centre		
≤ 30 (10/yr)		50
31 – 74		75
≥ 75 (25/yr)		89

... onvolledige registratie door deelnemende centra



Registratie door deelnemende centra is gebiased

<75 yr	61
>75 yr	47
WHO 0	69
WHO 1-2	59
WHO >2	38
cStage I-II	62
cStage III	71
cStage IV	41
Neoadjuvant no	52
Neoadjuvant yes	77

Variabiliteit en volume effect

- Op zorgkwaliteitsindicatoren
PROCARE
 - per domein
 - globale score
- Op bepaalde outcome parameters
BKR/IMA en/vs. PROCARE
 - type radicale resectie (APE ratio)
 - 30 d postop mortaliteit
 - overleving

Effect of hospital volume on 23 QCI (PROCARE)

STAGING

cStage reported

distance to MRF in cStage II-III

staging accuracy

TIME TO FIRST TREATMENT

NEOADJUVANT TREATMENT

RT/RCT for cStage II-III reported

RT/RCT for cStage II-III

idem if < 75 yr

idem if > 75 yr

SURGERY

good TME quality

negative resection margins (R0)

APE rate (adjusted)

major complications after SSO or APE

30-day mortality (adjusted)

PATHOLOGY

TME quality reported

pCRM reported

N nodes if no/short RT

N nodes after RCT

ADJUVANT CHEMOTHERAPY

for pStage III if < 75 yrs: reported

for pStage III if < 75 yrs: administered

FOLLOW-UP

data on LR/DM reported

LOCAL RECURRENCE RATE (adjusted)

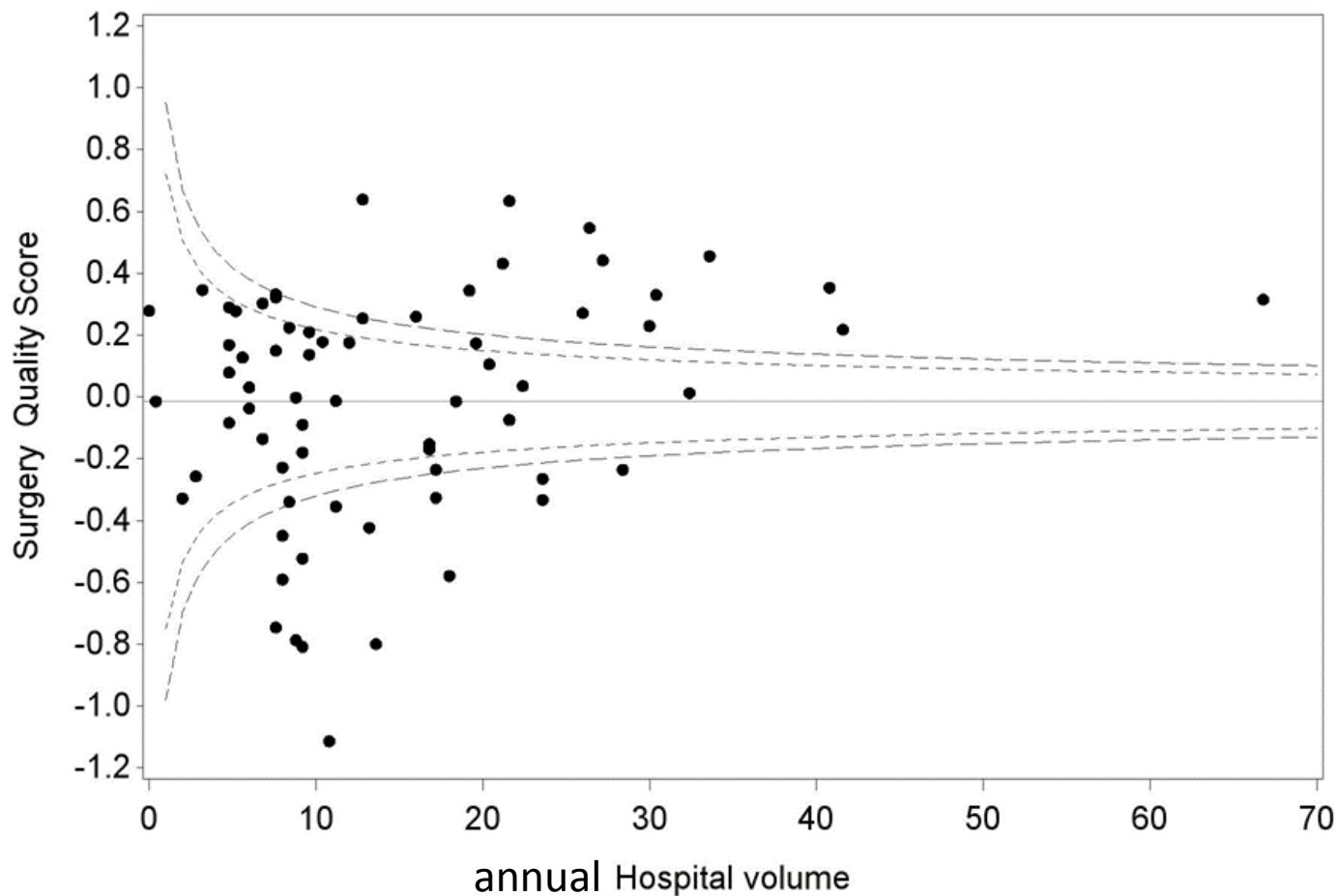
OVERALL RECURRENCE RATE (adjusted)

OVERALL SURVIVAL (adjusted)

Pearson correlations (p-value) between quality scores and hospital volume

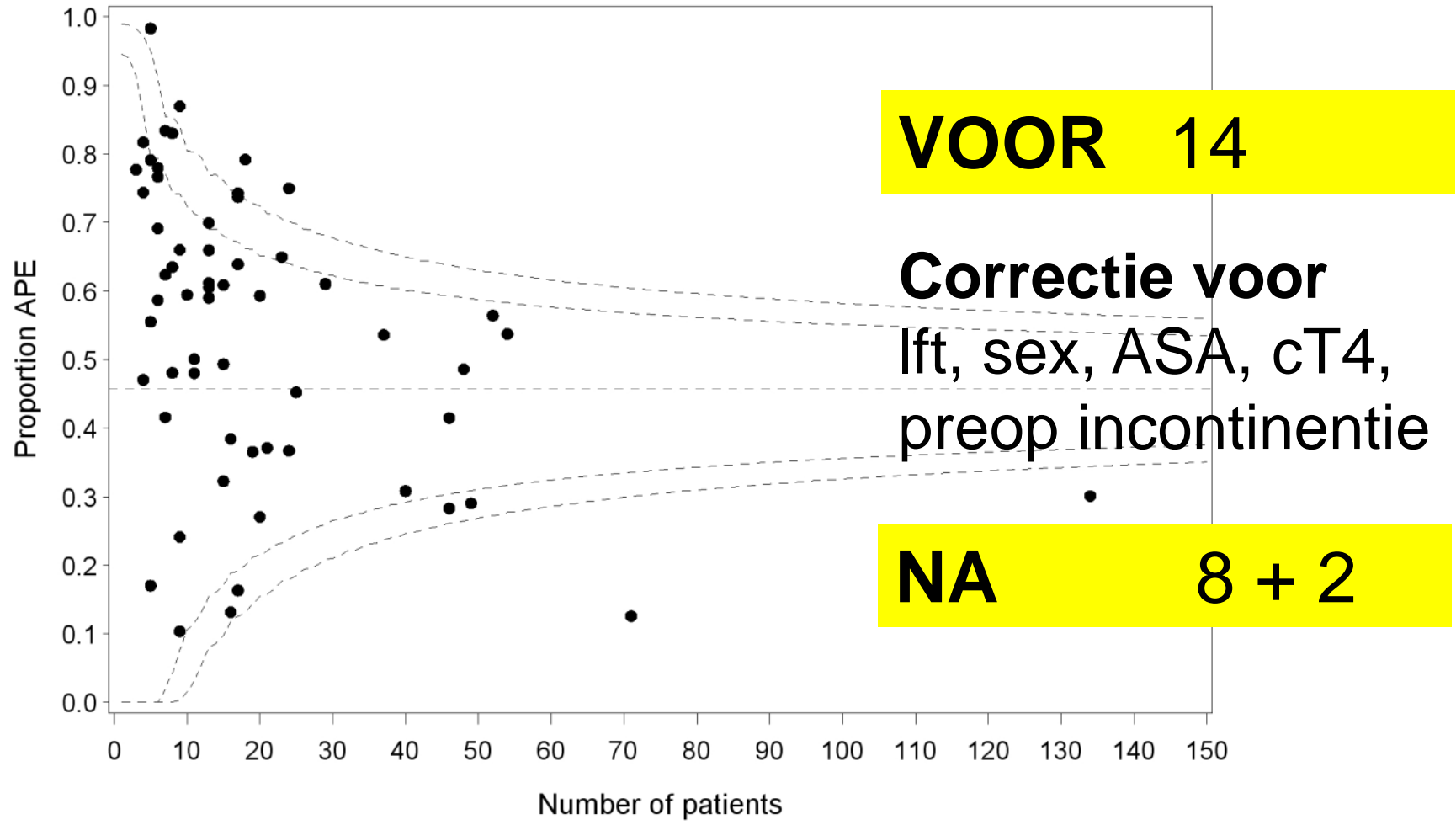
STAGING cCRM, staging accuracy	0.123 (0.319)
NEOADJUVANT R(C)T for cStage II-III	0.119 (0.339)
SURGERY good TME quality, R0 resection, adjusted APE rate	0.265 (0.029)
PATHOLOGY N nodes after RCT	0.139 (0.268)
ADJUVANT CHEMO for pStage III	0.005 (0.975)
GLOBAL SCORE	0.237 (0.052)

Funnel plot of surgery quality score and hospital volume

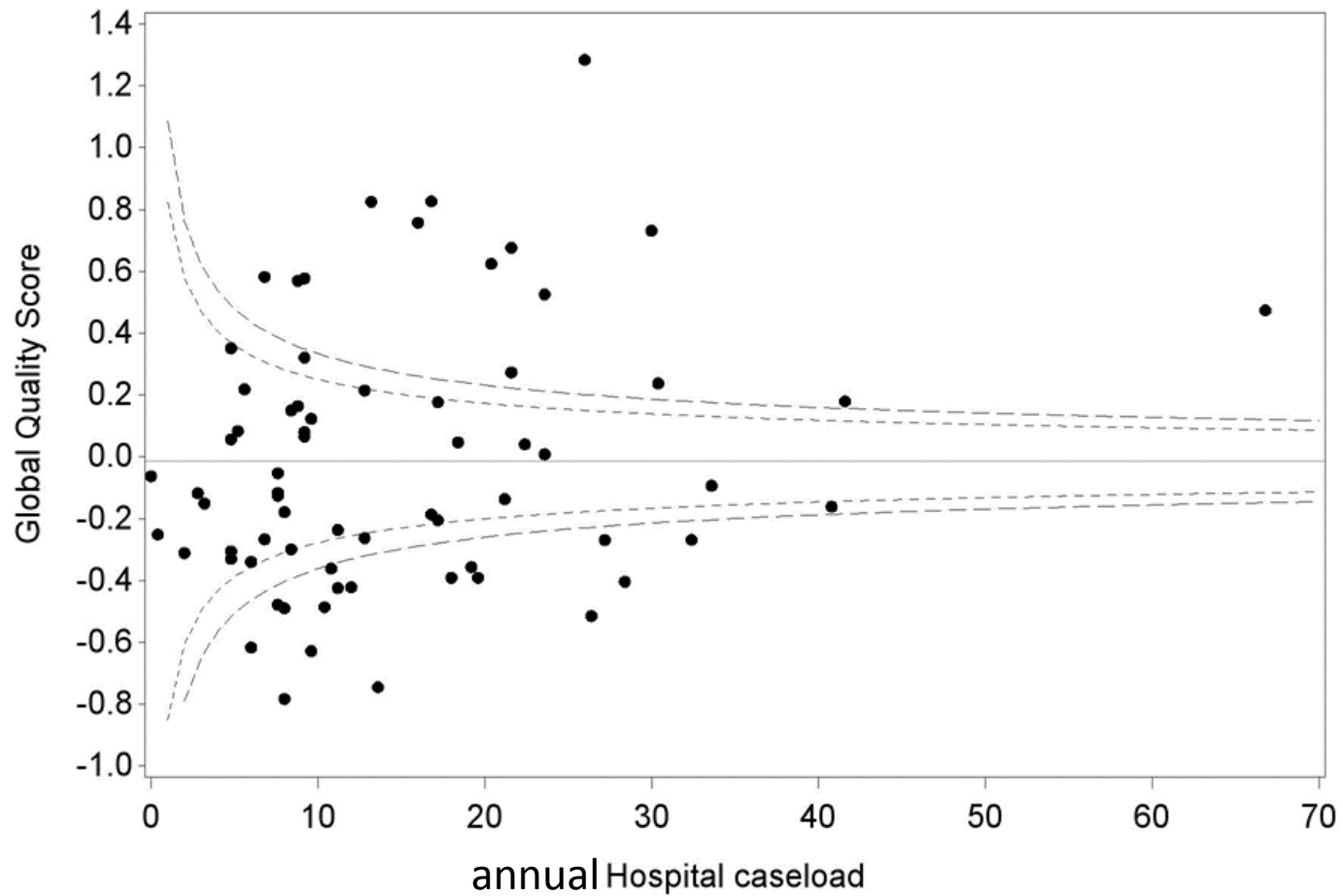


Variabiliteit in PROCARE

Percentage APE bij rectumkanker op 0-5 cm



Funnel plot of global quality score and hospital volume



Volume effect size on SSO rate in PROCARE and in BCR/IMA

	PROCARE N = 1263	BCR/IMA N = 5869
SSO rate (unadjusted)	OR 1.020/case (1.008-1.032)	OR 1.007/case (0.997-1.017)
RR/10 added cases (gender, age, ASA3, cT4, inco. adjusted)	3% increased chance to avoid an APE / 10 cases more per yr	

Volume effect size on 30-day mortality

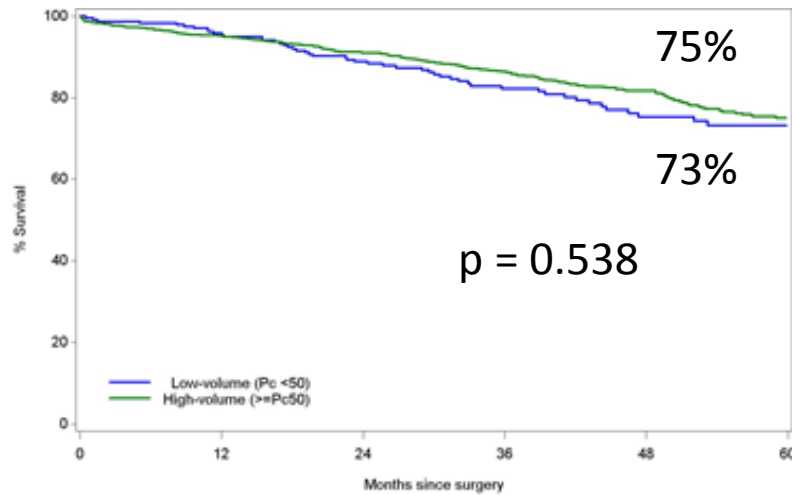
	PROCARE N = 1263	BCR/IMA N = 5869
30 d mortality	1.3 %	2.4 % [p=0.001]
30 d postop mortality (age adjusted)	OR 1.007/case (0.986-1.026)	OR 0.987/case (0.977-0.997)
RR/10 added cases (age adjusted)		14% less risk / 10 cases 14% of 2.4 = 0.3 reduction

Volume effects

	PROCARE N = 1263	BCR/IMA N = 5869
SSO rate (unadjusted)	OR 1.020/case (1.008-1.032)	OR 1.007/case (0.997-1.017)
30 d postop mort. (age adjusted)	OR 1.007/case (0.986-1.026)	OR 0.987/case (0.977-0.997)
OS (age, sex, pStage adjusted)	HR 0.996/case (0.987-1.005)	HR 0.994/case (0.990-0.997)

OS in pStage I-III rectal cancer treated in hospitals with BCR/IMA-based **volume < versus > the median**

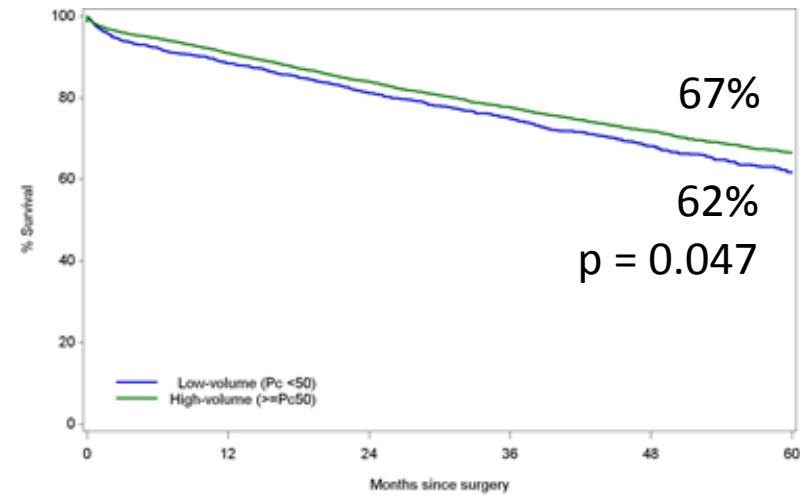
PROCARE



Number at risk	0	12	24	36	48	60
Low-volume	240	230	187	131	83	46
High-volume	1023	972	826	643	434	226

Median volume = 11/yr
Volume <Pc50 = 19% pts

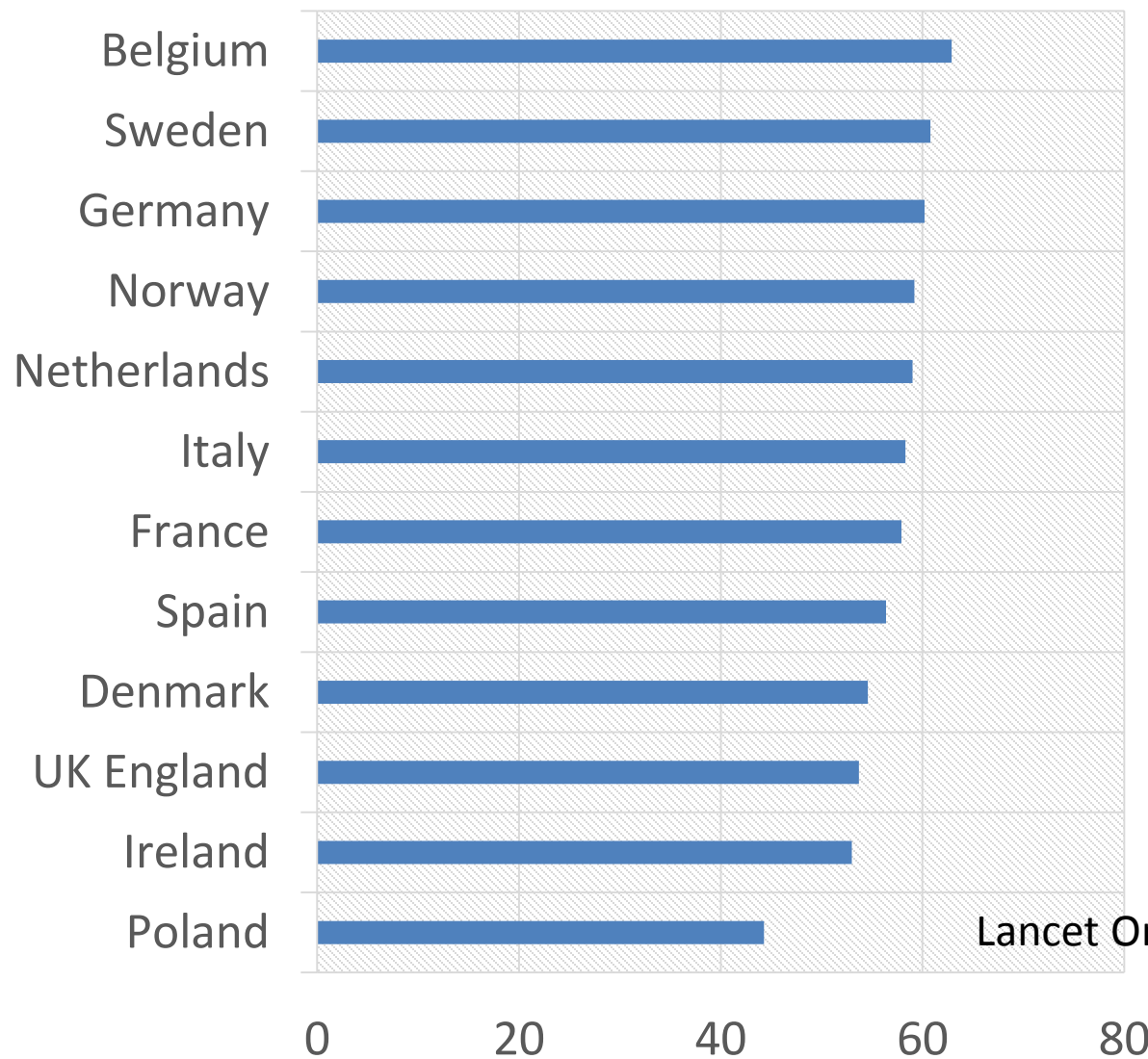
BCR/IMA



Number at risk	0	12	24	36	48	60
Low-volume	1479	1308	1159	844	542	307
High-volume	4390	3990	3546	2621	1781	1042

Median volume = 10/yr
Volume <Pc50 = 25% pts

5-yr age-standardized Relative Survival for rectal cancer 2000-2007



Lancet Oncology 2014, 15: 23-34

Besluiten

- België scoort zeer goed
- Volume effect ja, maar beperkt (welke cut-off?) en
- Centralisatie is geen garantie voor zorgkwaliteit
- Audit van zorgkwaliteit is essentieel
- Kanker-specifieke registratie moet verplicht worden
- Benchmarking en feedback naar teams
- Zorgkwaliteit bewaking door peers en de overheid

A minimum compulsory cancer-specific data entry set

Annexe 93

FORMULAIRE D'ENREGISTREMENT REGISTRE DU CANCER

Patient : Date de naissance :
 N° national/ N° organisme assureur : Sexe :

1. DATE D'INCIDENCE (JJ-MM-AAAA).....
 ordre de priorité décroissante: 1=première confirmation histo/cyto,2= évaluation clinique/hospitalisation, 3= décès
 2. MOYEN DE DIAGNOSTIC (entourer, plusieurs items sont permis):
 1 = autopsie (pas d'application pour la CMO) 5 = examen technique (ex. RX, endoscopie, ...)
 2 = histologie tumeur primitive 6 = examen clinique
 3 = histologie métastase 7 = marqueur tumoral (ex. PSA, HCG, AFP, Ig, ...)
 4 = cytologie / hématologie 9 = inconnu
 3. SCORE OMS DU DIAGNOSTIC (entourer)
 0 = Asymptomatique, activité normale 3 = Symptomatique, alité >50% de la journée
 1 = Symptomatique, mais ambulatoire 4 = Complètement dépendant pour les soins, grabataire
 2 = Symptomatique, alité <50% de la journée
 4. LOCALISATION DE LA TUMEUR PRIMITIVE (remplir) :
 5. LATERALITE uniquement pour organes pairs (entourer) : 1. gauche 2. droite 3. inconnu
 6. DIAGNOSTIC HISTOLOGIQUE (remplir):
 7. DEGRE DE DIFFERENCIATION (entourer): 1 = bien 2 = moyen 3 = peu 4 = indiff/anapl 9 = inconnu
 8. TNM CLINIQUE (UICC 2002): cT..... cN..... cM.....
 9. TNM PATHOLOGIQUE (UICC 2002): pT..... pN..... pM.....
 10. AUTRES TYPES DE STADE CLINIQUE (remplir):
1. Ann Arbor, 3. Figo, 4. Salmon Dury, 5. Clark, 6. Breslow, autre
- Stade (remplir):
11. DATE DEBUT DU PREMIER TRAITEMENT (JJ-MM-AAAA):
 12. TRAITEMENTS DEJA RECUS:
 remplir chronologiquement à partir de la date du premier traitement
- | | | | |
|--|--|--|--|
| | | | |
|--|--|--|--|
- | | | |
|--------------------------------------|---------------------------|---------------------|
| 10: chirurgie | 15: greffe moelle osseuse | 90: pas de thérapie |
| 20: radiothérapie | 30: isotopes | 95: refus thérapie |
| 25: chimioradiothérapie concomitante | 50: hormonothérapie | 99: inconnu |
| 40: chimiothérapie | 60: immunothérapie | |
13. PLAN DE TRAITEMENT ULTERIEUR (intention)
 remplir les codes chronologiquement, voir point 12 - jusqu'à max. 1 an après le début du traitement
- | | | | |
|--|--|--|--|
| | | | |
|--|--|--|--|
14. SYMPTOMATIQUE / PALLIATIF (entourer): OUI NON
 15. NOUVEAU DIAGNOSTIC (1) OU FOLLOW-UP (2) : (entourer) 1 /2
- COORDINATEUR
 Dr :
 Institution :

BCR + IMA
+ 29 additional items

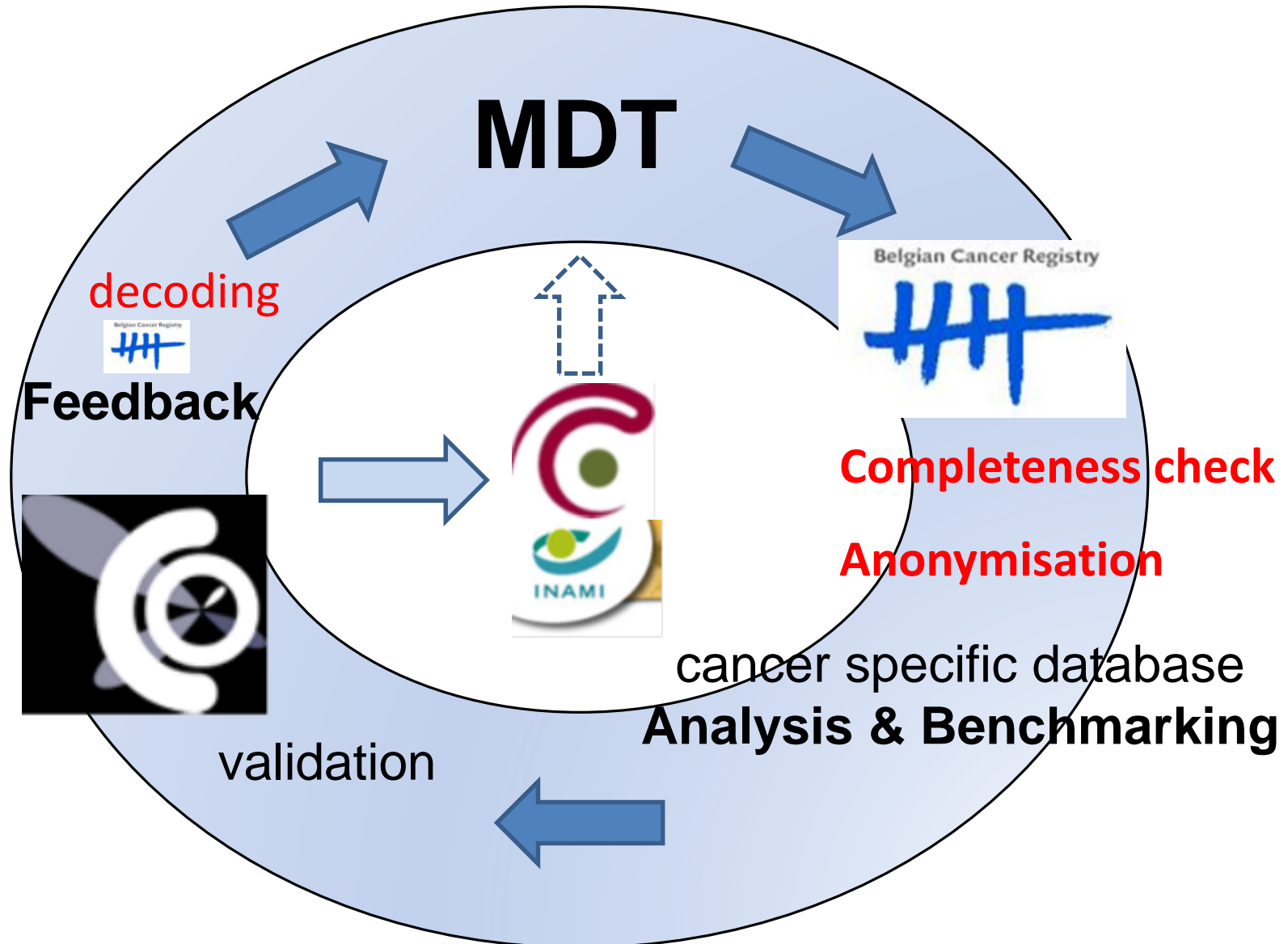
- 8 pre-treatment items
- 8 (post)operative items
- 6 pathology items
- 7 radio(chemo)ther. items

Belgian Cancer Registry



From project to structure

Proposed 'audit loop'



How PROCARE became possible

- Belgian Foundation against Cancer (2006, 2013)
- **RIZIV/INAMI (2007-2012-2014)**
- KCE (2007, 2008, 2010)

- Steering Group (all societies)
- **Participating professionals**
- **Belgian Cancer Registry**

RBSSurgery (BSCRS), BSSO, BGES
BSRadiotherapy – Oncology
BSPathology (Dig Path Club)
BSMOncology, BGDO
RBSRadiology
VVGE
SRBGE
BSGIEndoscopy
BPSA
FBCR