

Verbier 24 January 2010 Marc-Claude Marti Lecture

P R O C A R E

PROJECT ON CANCER OF THE RECTUM

**The Belgian national
rectal cancer project**





Rien de grand ne se fait sans passion ... assortie de quelques qualités purement humaines

- serenity
- wisdom, knowledge, contacts
- visionary (amb. procto, ECCP 1984, ...)
- kind
- patient
- humour

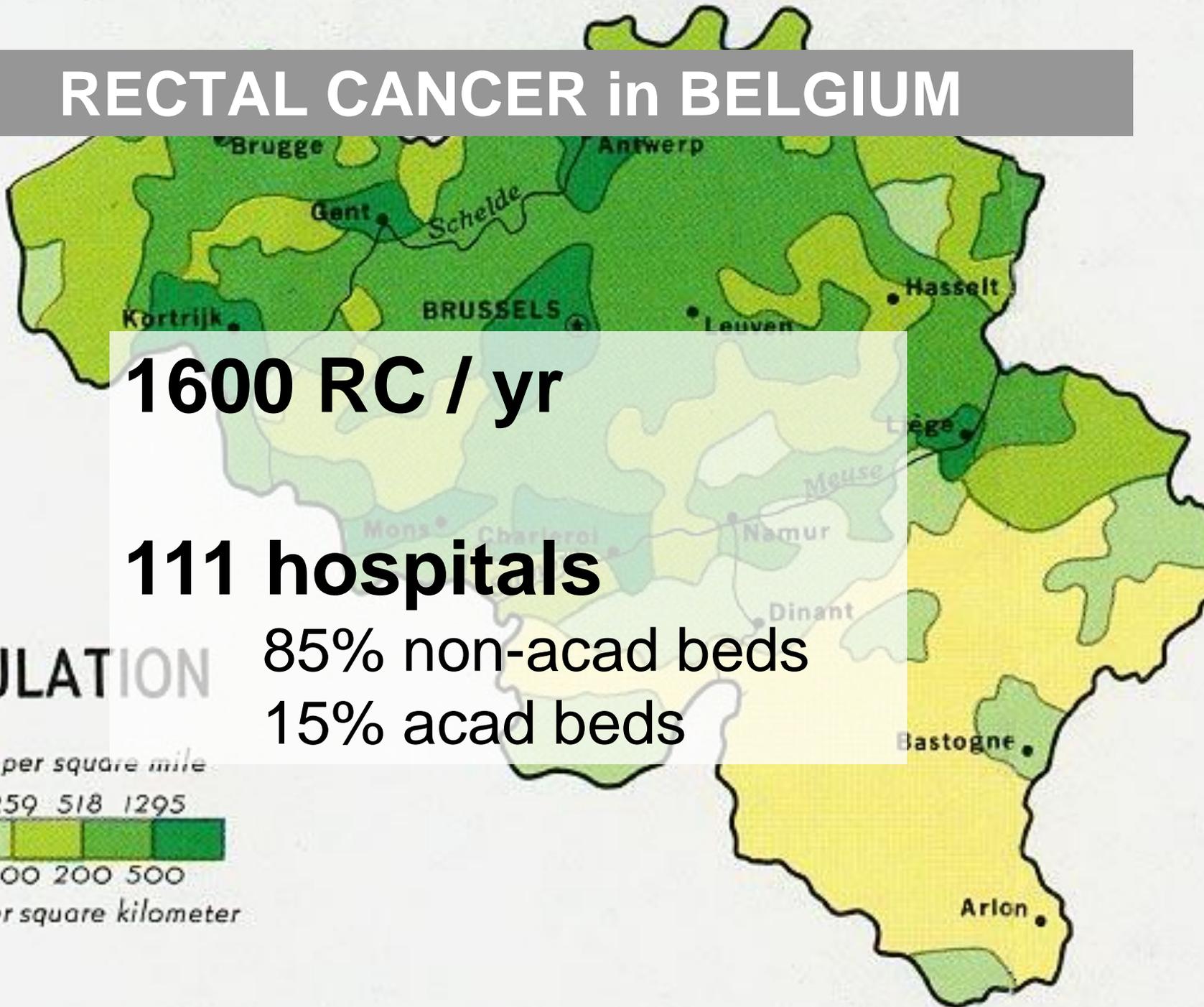


“KOE-KOEK”



zaza

RECTAL CANCER in BELGIUM



1600 RC / yr

111 hospitals

85% non-acad beds

15% acad beds

POPULATION

Persons per square mile

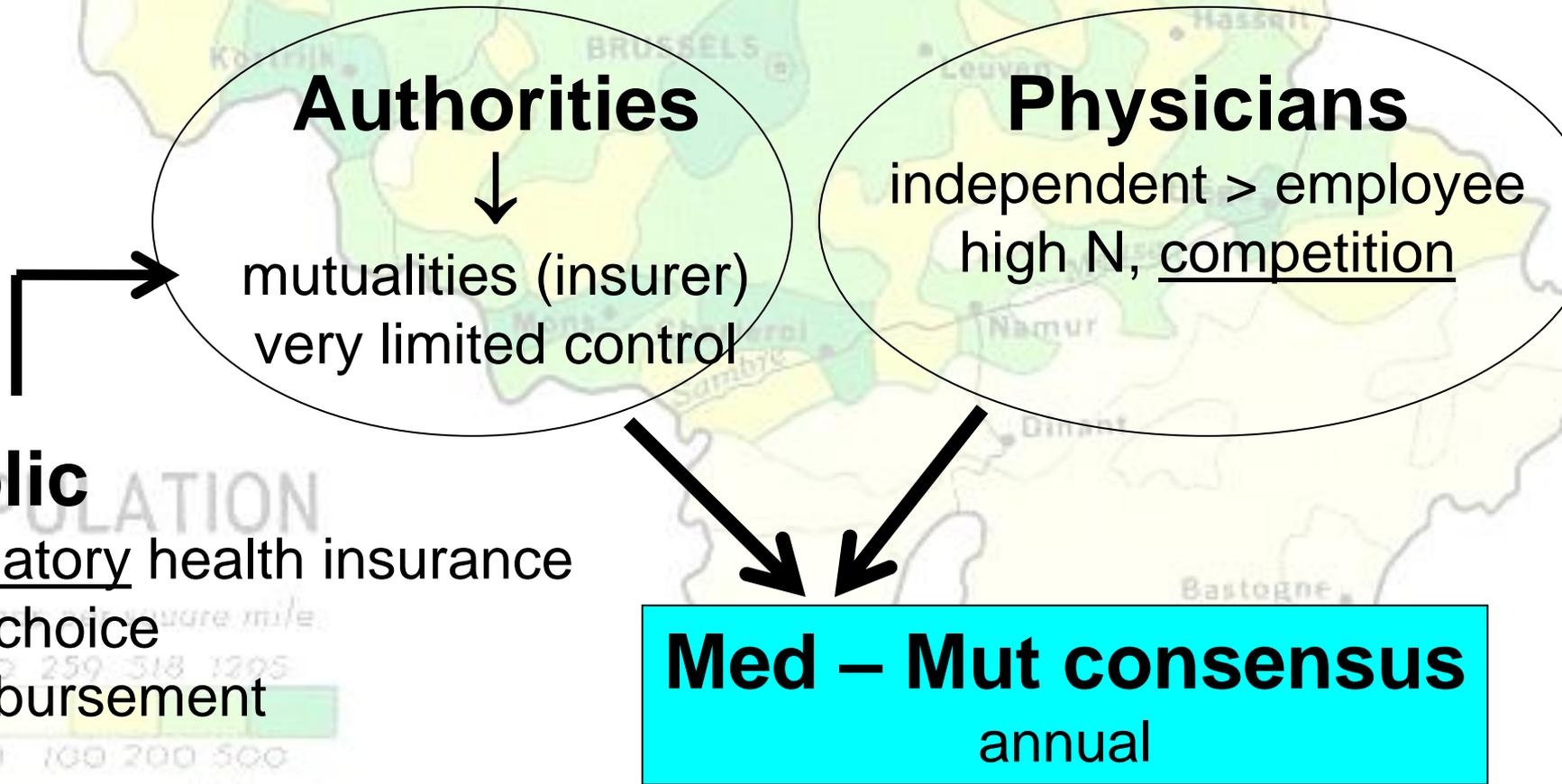
0 130 259 518 1295



0 50 100 200 500

Persons per square kilometer

Public Health in Belgium



Public

Mandatory health insurance

Free choice

Reimbursement

0 50 100 200 500

Persons per square kilometer

PRO CARE

PROJECT ON CANCER OF THE RECTUM

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

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

PROCARE METHODS

- multidisc. EB **Guidelines and QCI** (2005, 07, 08)
- quality assurance (**implementation** of GL)
 - training (radiology, RT, TME, pathology)
 - registration of 151 items (>1/2006)
 - feedback / benchmarking (2008, 2009)

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

Federaal Kenniscentrum voor de Gezondheidszorg
Centre fédéral d'expertise des soins de santé
2007



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

KCE reports 81A



BELGIAN
CANCER
REGISTRY

[NL](#) - [FR](#) - [D](#) - [ENG](#)

- Home
- Het Kankerregister
- Statistieken
- Registratie
- Bijscholing
- Publicaties
- **PROCARE**
 - Contact
 - Presentation
 - Working
 - Statistics
 - Publications
 - Archives
- Links
- Online applicaties
- Vacatures
- Contact

www.kankerregister.org
www.registreducancer.org

PROCARE

Welcome to the PROCARE

PROCARE, a multidisciplinary website presents details ever since. You can all

If you are interested under the heading "Statistics". The working of the presentation entry forms and the T

Latest news

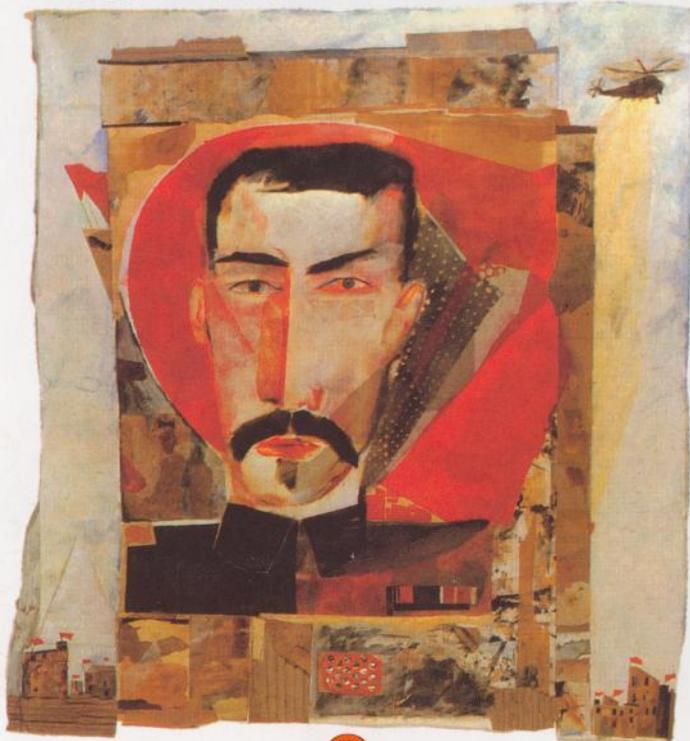
Quality of Care Indicators : 40

PROCARE vs. ADMINISTRATIVE DATABASES

	PROCARE	ADMIN
General (level 1)	3	2
Diagnosis and staging	7	2
Neoadjuvant treatment	7	1
Surgery	6	3
Pathology	6	0
Adjuvant treatment	5	0
Follow-up	3	0
Palliative treatment	2	1
	39	9

GEORGE ORWELL

NINETEEN EIGHTY-FOUR



Big Brother is watching you

The Daily

Friday, November 27, 2009

No 48,051

BRITAIN'S BEST-SELLING

Failing hospital condemns hundreds to death

- Lack of basic hygiene in A&E
- Nurses neglect to feed patients
- Wrong medication handed out

Where were all the P



FUNDING
for
training and
central data registration

Belgian Federation against Cancer (2006)

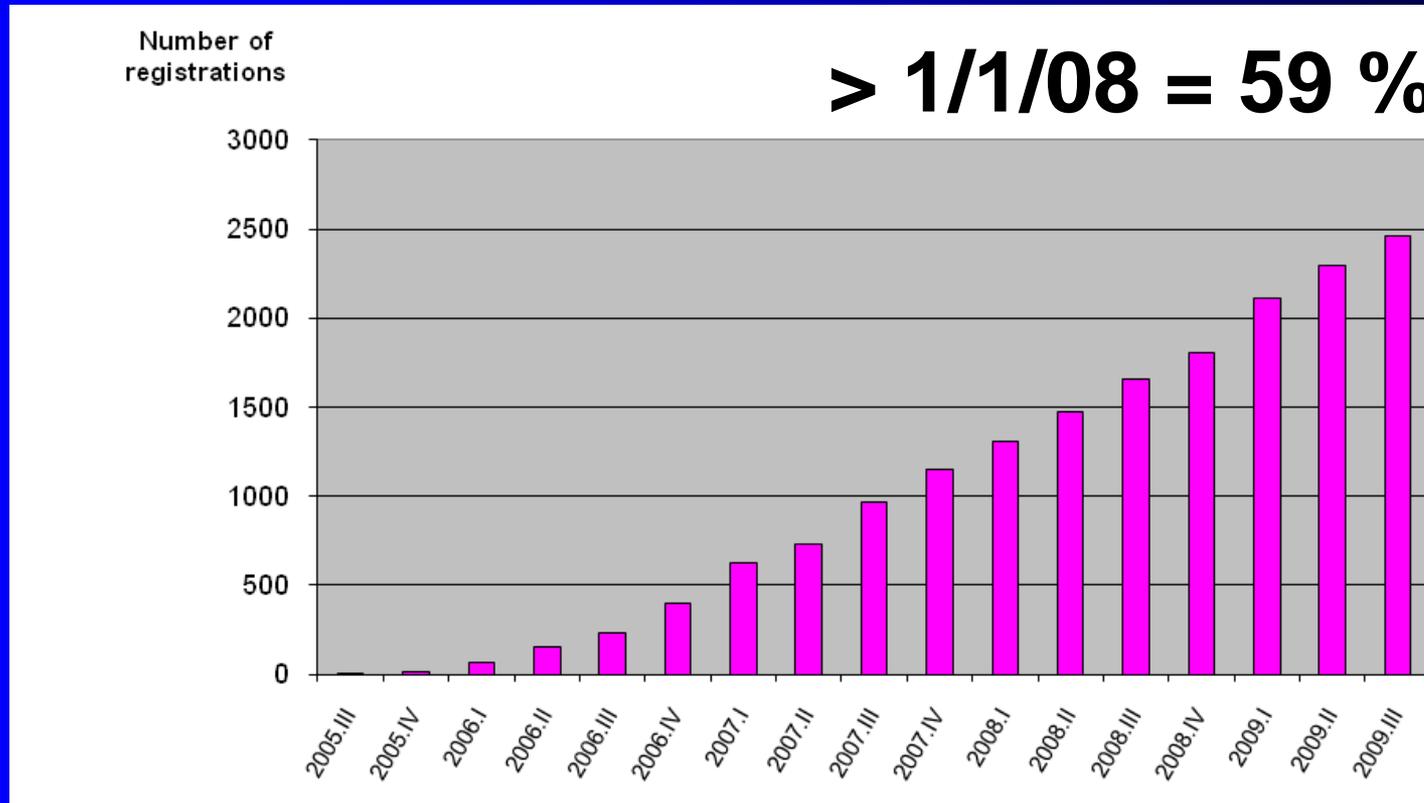
KCE

RIZIV / INAMI (2007 – 2012)

TRAINING

- **PRETREATMENT STAGING (radiologists)**
 - central review CT / MRI images 2010
- *RADIOTHERAPY*
- **TME : 177 / 225 surgeons interested (2005)**
 - 43 candidate-trainers → 25 trainers (18 NL / 7 FR)
 - 6 trained (since 8/2008)
- **PATHOLOGY**
 - TME reviews from candidate trainers
 - > 11/2009 TME review ad random (44% correct material)

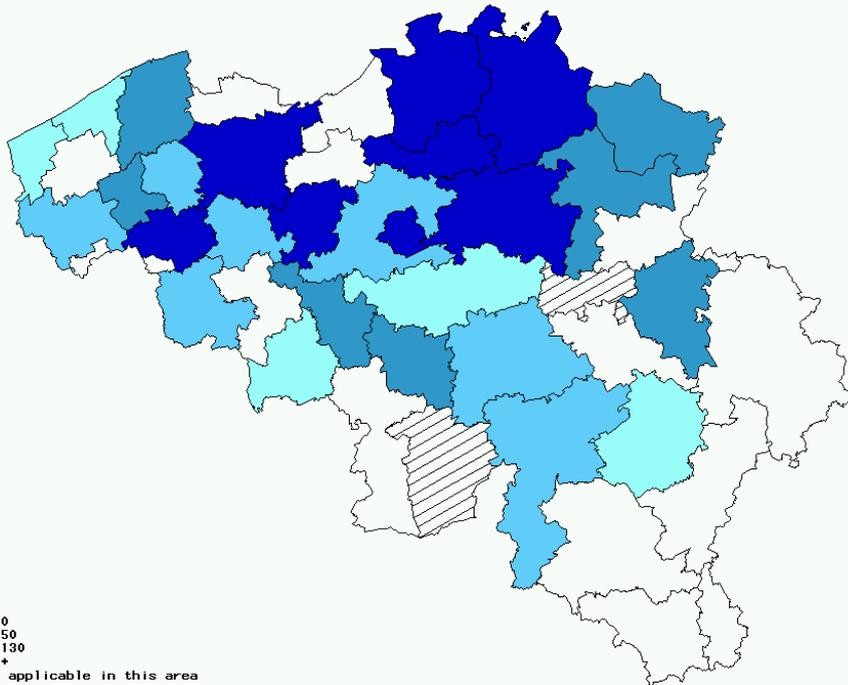
2947 patients registered (Dec 4 2009)



Who submitted patients ?

70 / 111 = 63 % hospitals

Procare registrations in Belgium by residence hospital, by district, status on 28/10/2009 (N=2699)

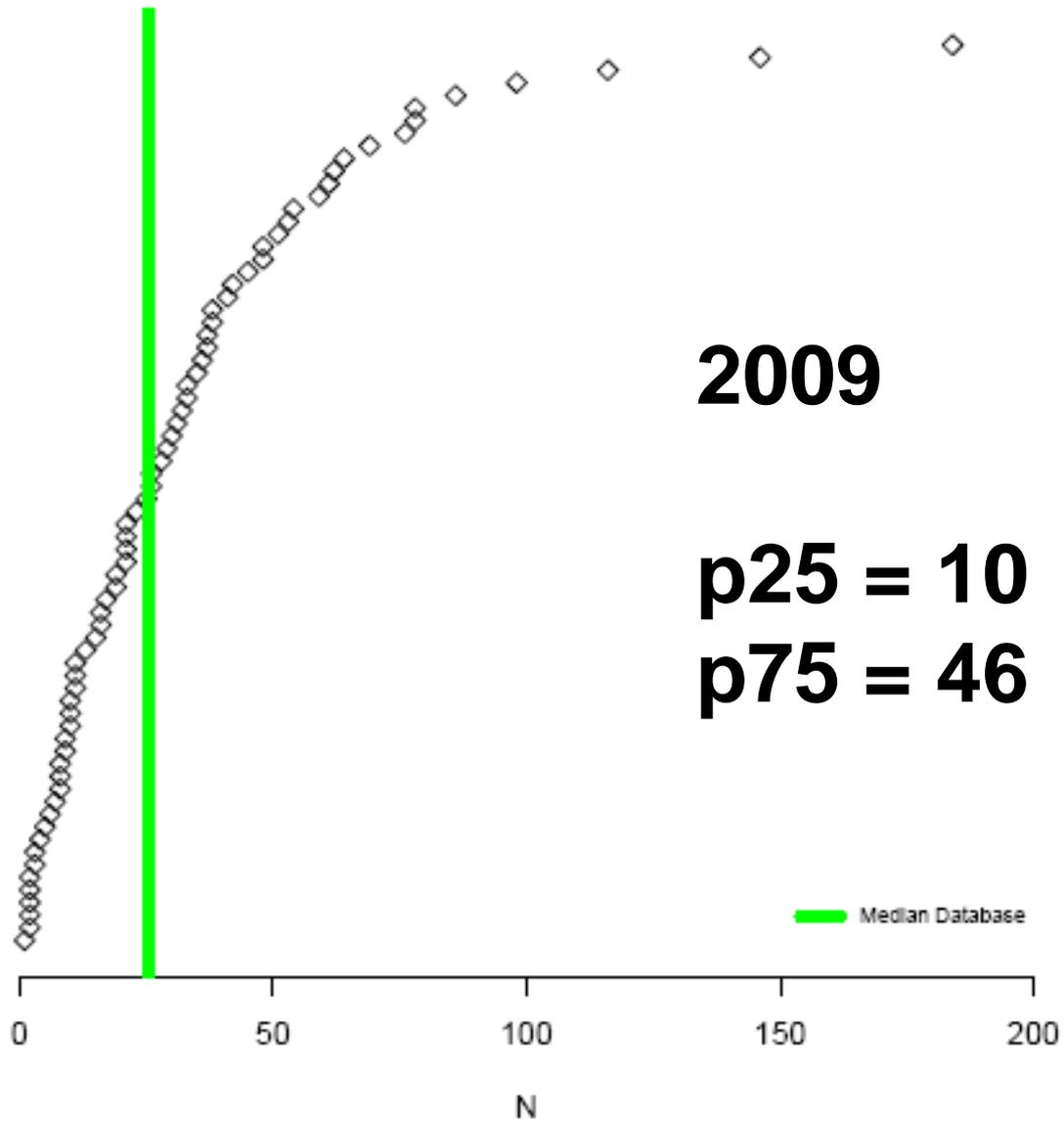


1-20
21-50
51-130
130+
Not applicable in this area

© Procare

West Vlaanderen	12/14
Oost Vlaanderen	7/14
Antwerpen	19/19
Limburg	6/ 8
Vlaams Brabant	4/ 6
Brussel/Bruxelles	9/14
Brabant Wallon	1/ 2
Hainaut	7/16
Namur	2/ 6
Liège	2/11
Luxembourg	1/ 3

Number of patients registered



How to stimulate participation?

NON-PARTICIPATING HOSPITALS

letter to directors, surgeons (Oct 16 2009)

15/41 will participate (Nov 3 2009)

2 submitted patients (Nov 3 2009)

PARTICIPATING HOSPITALS

reminder incl. training (Nov 2009)

list of participating hospitals on website

newsletters, feedback

Analysis for second feedback

N patients	2439
Male/Female (%)	61/39
Age (mean yrs)	68
Lower level of tumour	
High (>10 cm)	17.7%
Mid (>5 - ≤ 10 cm)	38.4%
Low (≤ 5 cm)	43.9%

PME 15.8 %

TME 83.4 %

Pretreatment diagnosis and staging (1)

Complete large bowel examination if elective	98.2%
Use of CT (any stage)	57.4%
Use of TRUS (any stage)	42.4%
Use of TRUS if cT1-2	45.3%
Use of MRI (any stage)	37.8%
Use of MRI if cT3-4	43.5%
TRUS + CT and/or MRI	31%
cCRM if stage II-III	27.3%
CEA before treatment	82.6%

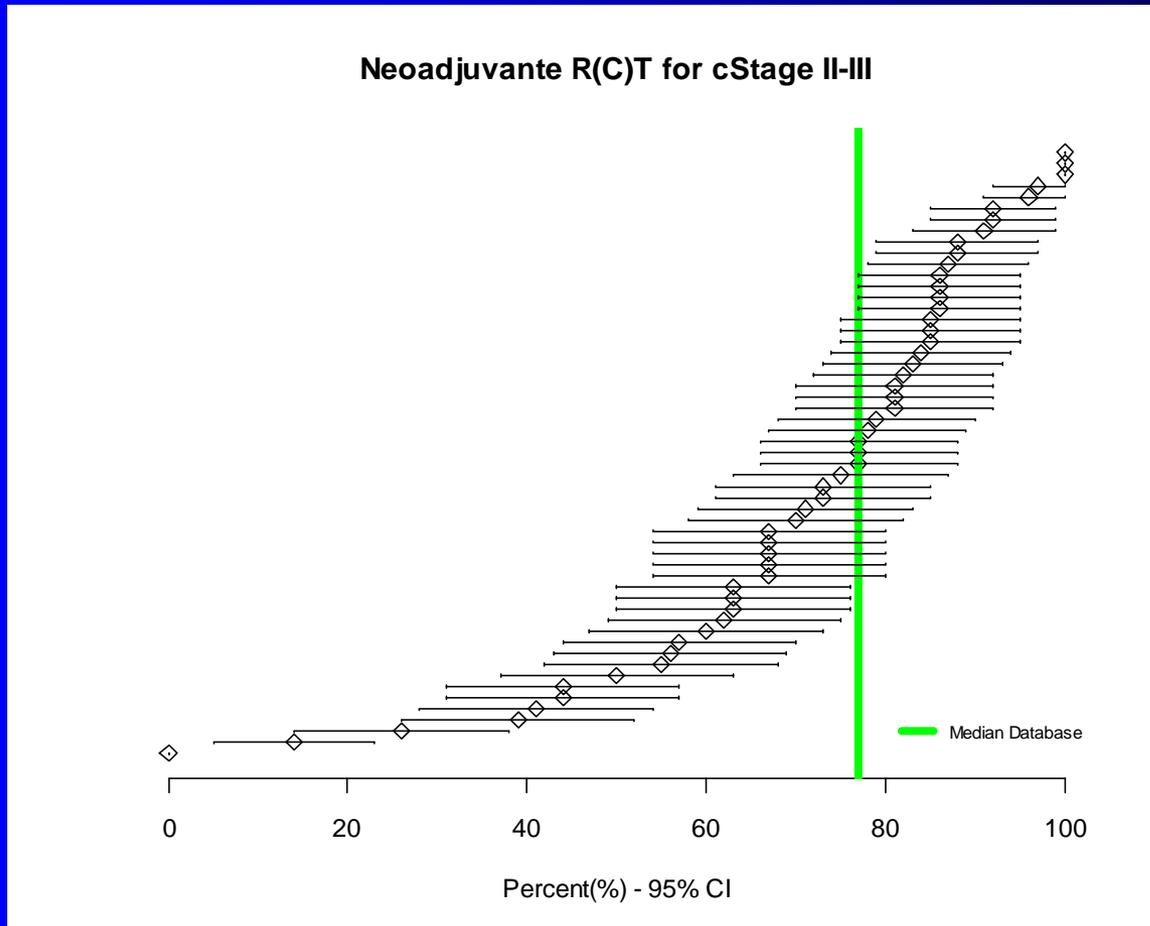
Pretreatment diagnosis and staging (2)

cStages	
cStage 0	0.5%
cStage I	13.6%
cStage II	18.3%
cStage III	50.4%
cStage IV	14.3%
cStage X	2.9%
cStage missing	306 = 12.5%

Neoadjuvant treatment

Short course for cStage II-III	9.9 %
Long course for cStage II-III	63.4 %
RT for high cStage II-III	39.8 %
RT for mid cStage II-III	78.1 %
RT for low cStage II-III	86.2 %
Long course if cCRM \leq 2 mm	69.8 %
Surgery 6-8 wks after long course	61.3 %
Surgery 4-12 wks after long course	97.4 %

Neoadjuvant (chemo)radiotherapy for cStage II or III (if > 10 pts)



Neoadjuvant treatment

	N your hospital	%your hospital	N Procure	%procure	p25	median	p75
QCI: Short course RT for cStage II-III	3	2.5	118	9.9	0	0	7.1
QCI: Long course (C)RT for cStage II-III	104	87.4	758	63.4	50	64.3	83.3
QCI: Long course (C)RT without Interruption	103	99	744	98.2	100	100	100
NEOADJUVANT TREATMENT FOR cSTAGE II-III							
-> For high RC	11	64.7	80	39.8	0	33.3	66.7
-> For mid RC	40	95.2	421	78.1	60	80.5	100
-> For low RC	68	97.1	551	86.2	71.4	92.6	100
Long course (C)RT if cCRM <= 2mm	59	79.7	164	69.8	0	87.5	100

Surgery (1)

Elective/scheduled	98.1 %
Open radical resection	71.5 %
Lap radical resection	24.9 %
Lap-converted rad resection	3.6 %
R0 after radical resection	75.7 %
R1 after radical resection	10.4 %
R2 after radical resection	13.9 %
Rectal perforation	7.7 %

Surgery (2)

Type of resection and reconstruction

Local excision/TEM	1.3 %	28
APER/Hartmann	22.2 %	470
AR + CRA	21.5 %	454
TME + CAA	54.3 %	1148
Other types of resection	0.5 %	11
	100 %	2111
Missing data	6.4 %	145



Unacceptable variation in abdominoperineal excision rates for rectal cancer: time to intervene?

E Morris, P Quirke, J D Thomas, et al.

Gut 2008 57: 1690-1697 originally published online June 5, 2008

Rectal cancer surgery: is restoration of intestinal continuity the primary aim?

C R Selvasekar, G David, D J Corless, et al.

Gut 2009 58: 311

Statistics, damned statistics and time to intervene

N A Scott, P Sagar and and the 30 co-signatories listed below

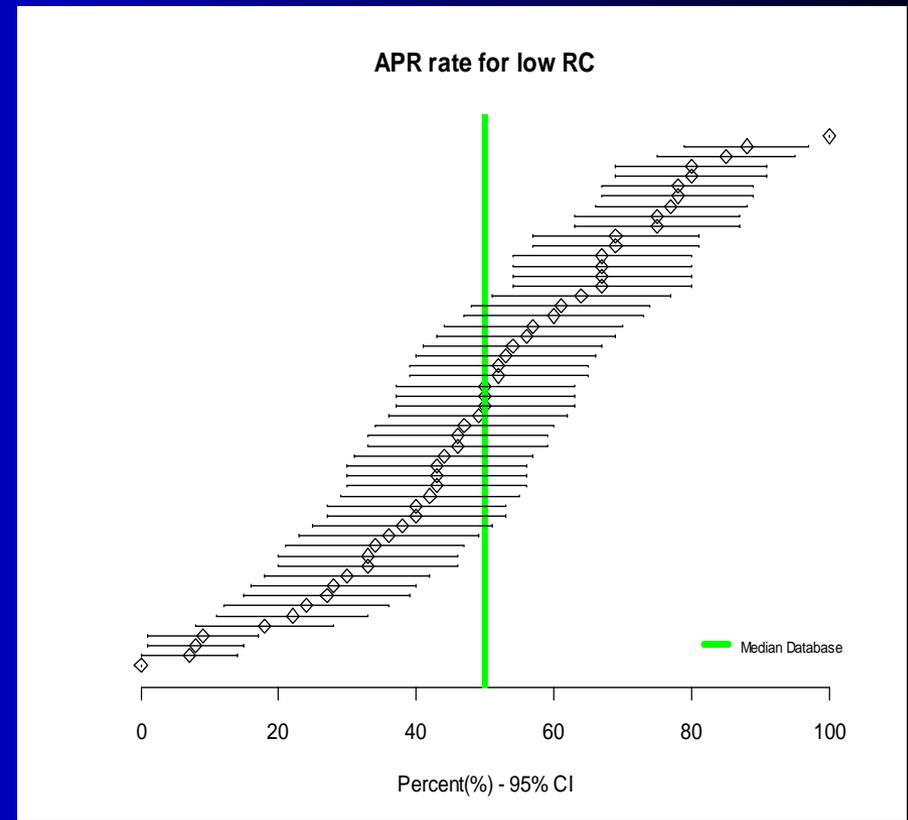
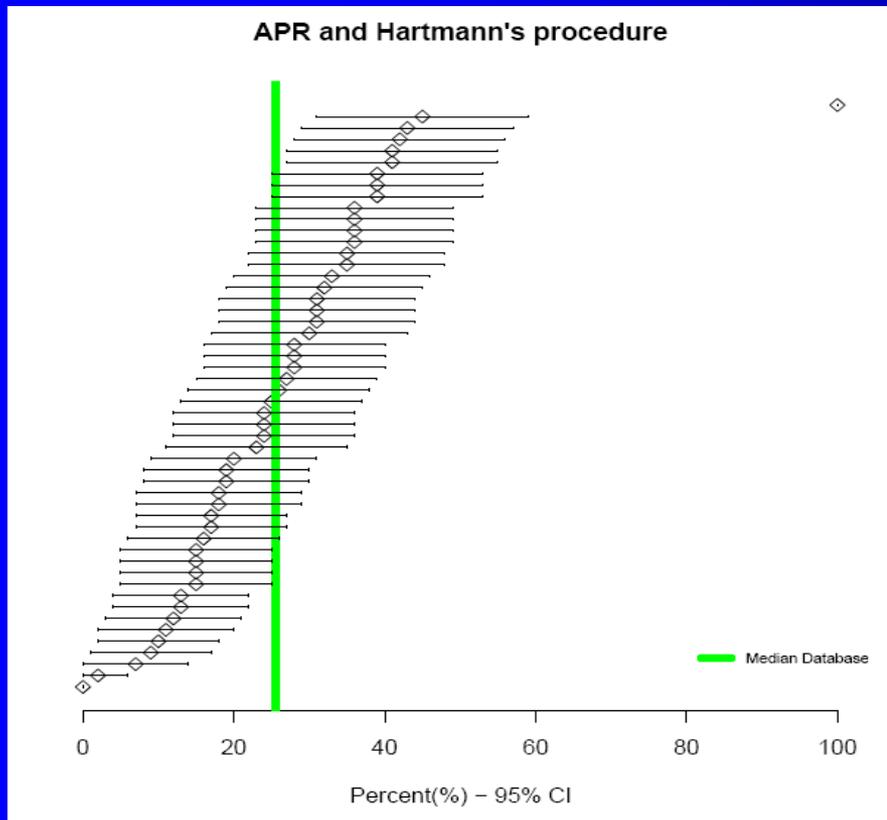
We question the underlying agenda of this type of publication. It is our collective view that incomplete data, naive reasoning and flawed conclusions neither represent good science nor promote and protect the health of patients.

quality. In addition, inferring surgical excellence from low APE rates without adjusting for factors such as tumour height and stage may lead to inappropriate conclusions. Despite considerable efforts by Morris *et al*, this work was unable to adjust these data fully for such confounding factors, demonstrating that the necessary infrastructure to achieve this is not currently available in the UK at the national level. Therefore, APE rates in isolation are unlikely to be a useful benchmark to audit surgical performance at present.

APR and Hartmann (2009)

0 – 15 cm

0 – 5 cm

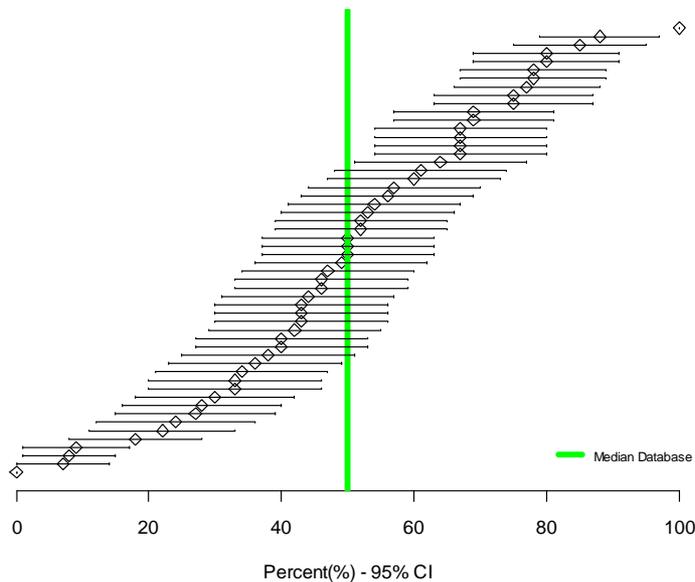


APR and Hartmann (2009) for rectal cancer at 0 – 5 cm

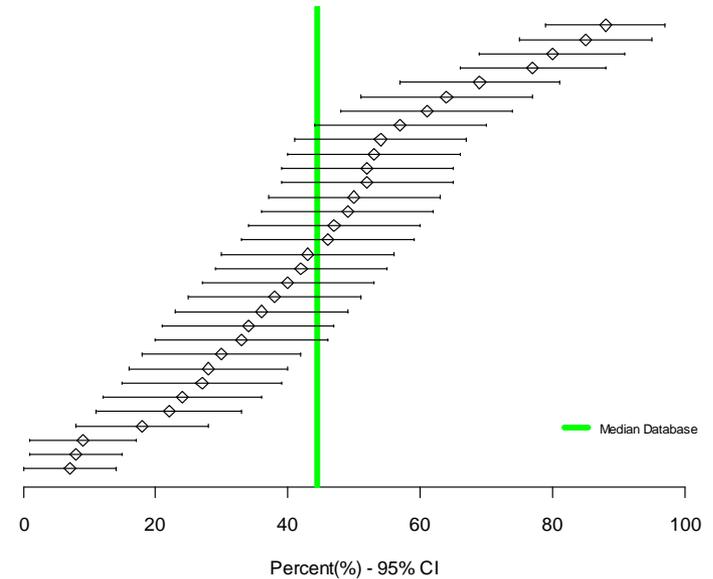
Teams > 10

Teams > 30

APR rate for low RC



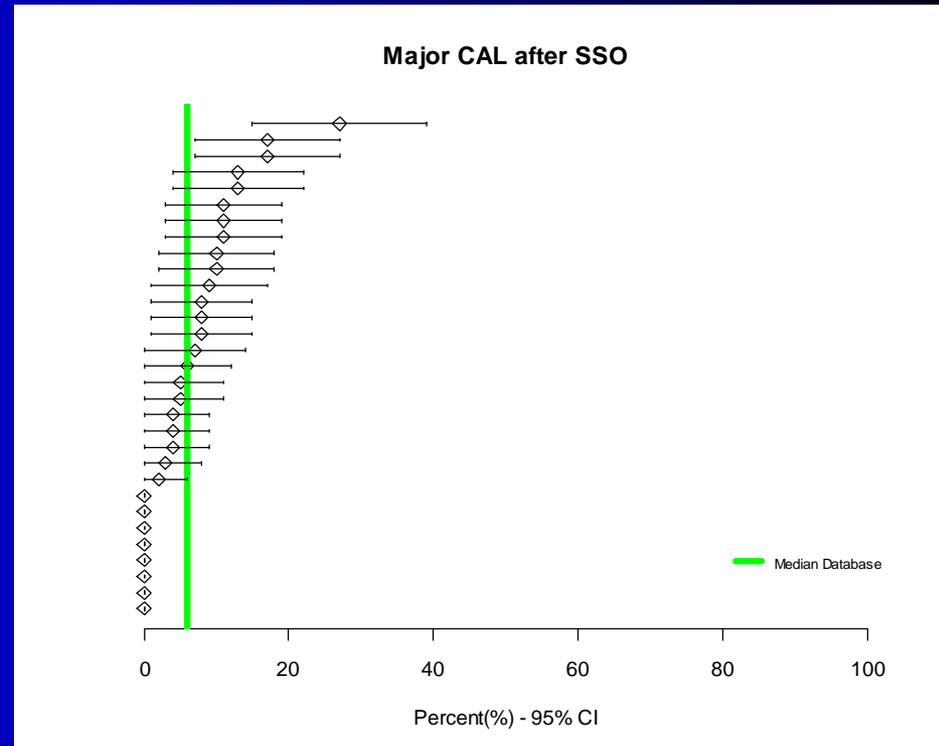
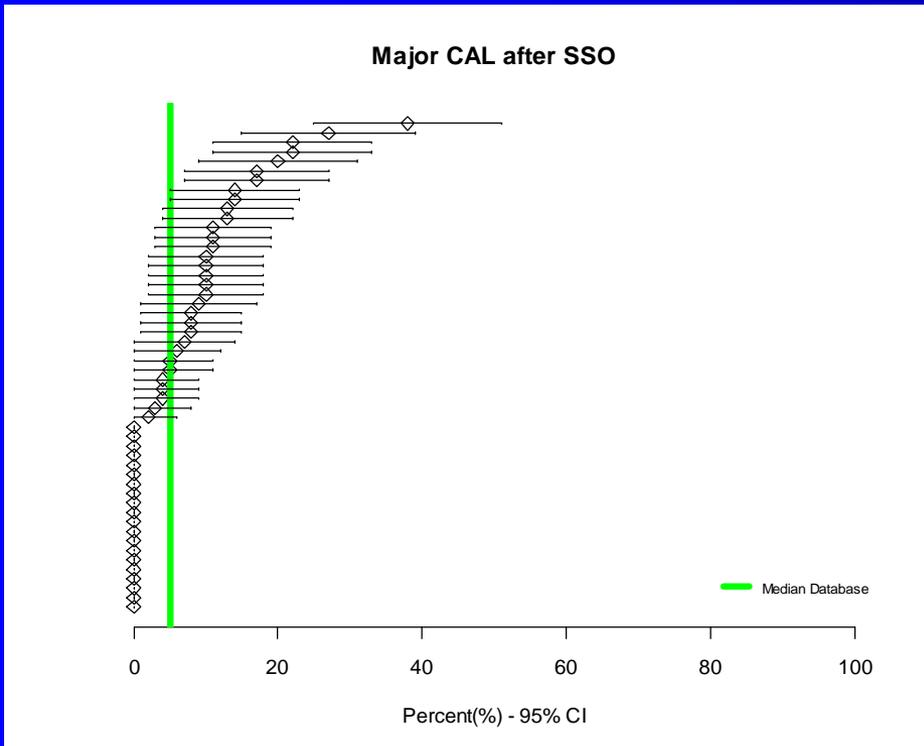
APR rate for low RC



Major leak after SSO with/without DS

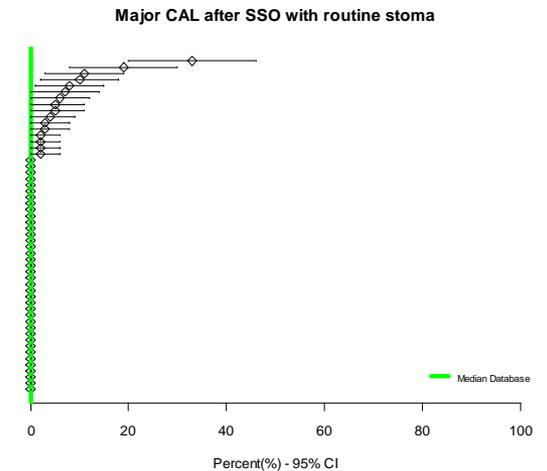
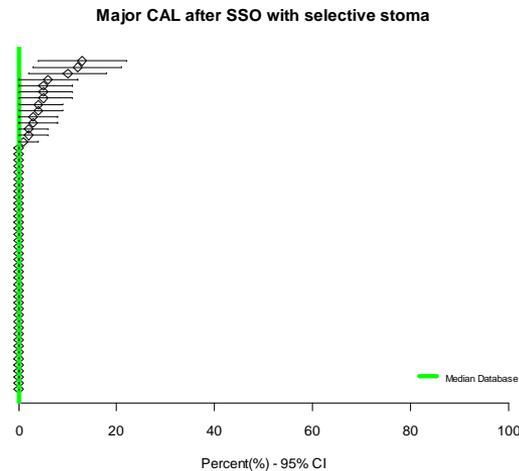
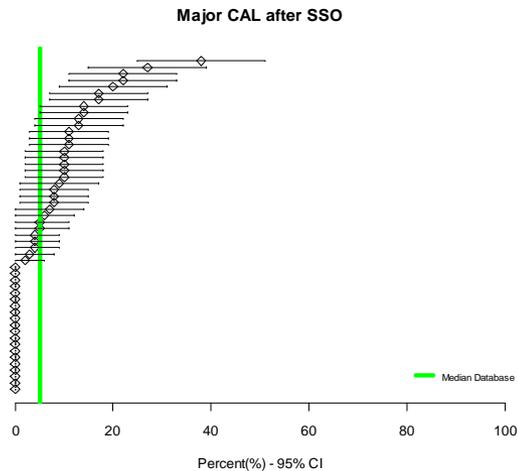
> 10

> 30



Major leak after SSO (if > 10 pts)

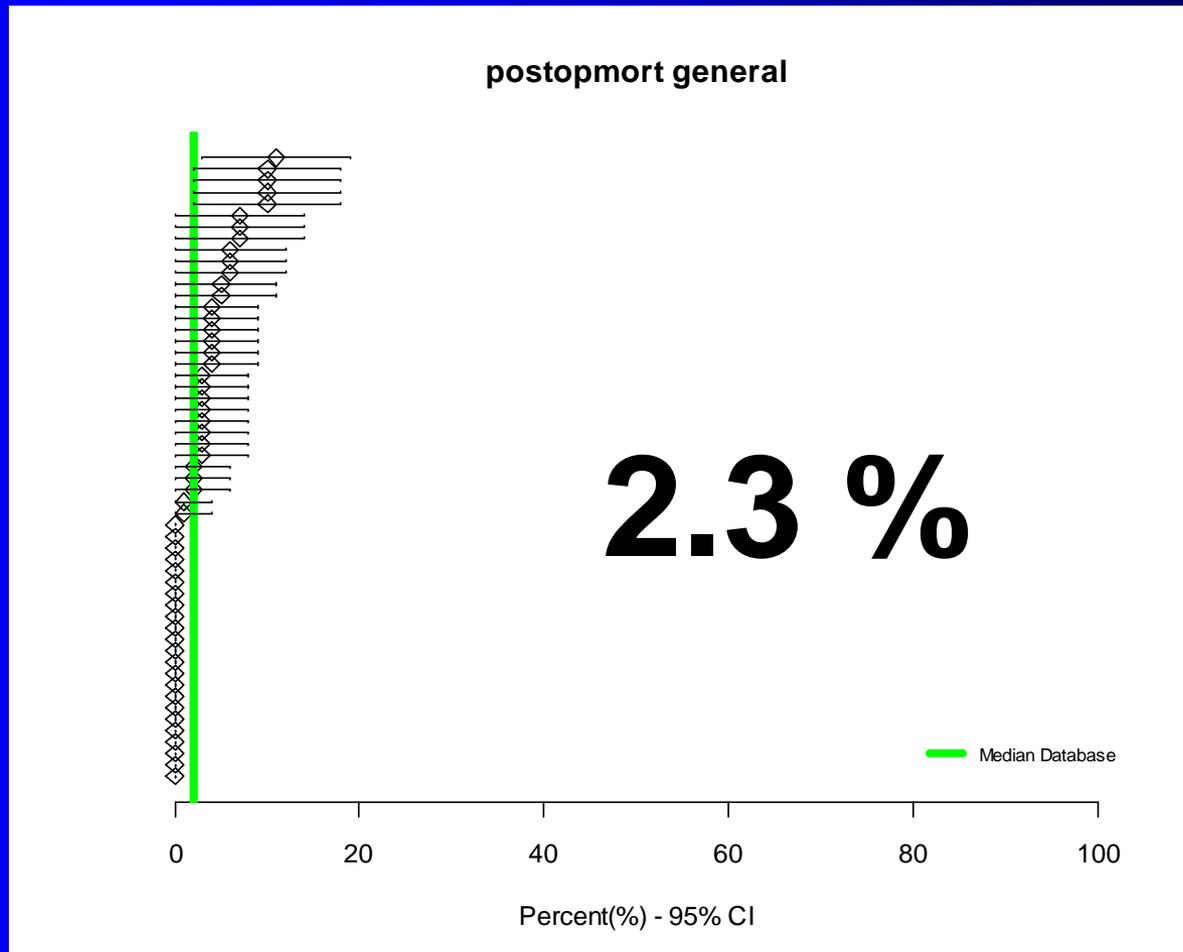
64 %
no DS selective DS 36 %
routine DS



9.5 % leak

5.5 % leak

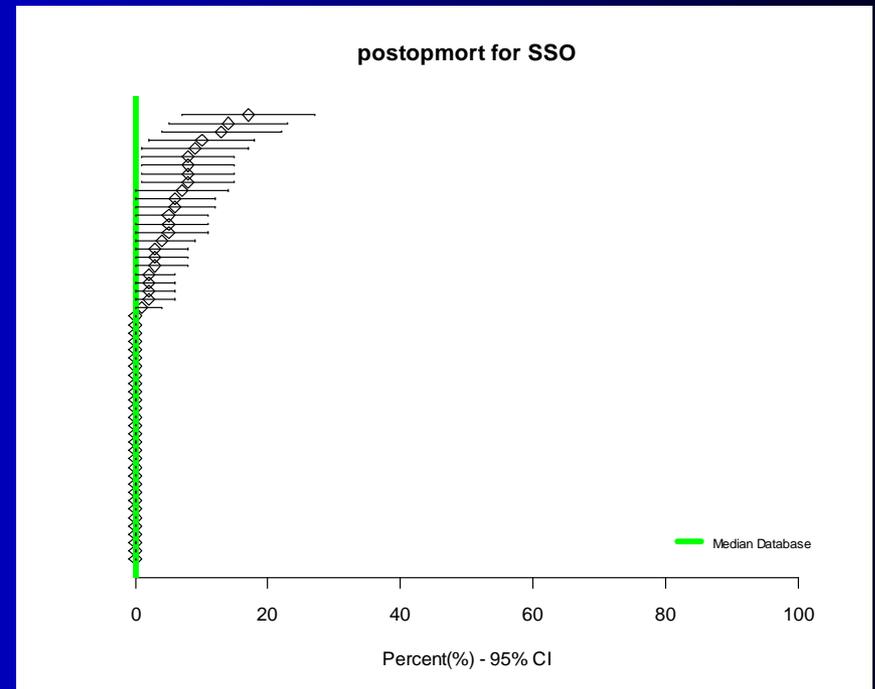
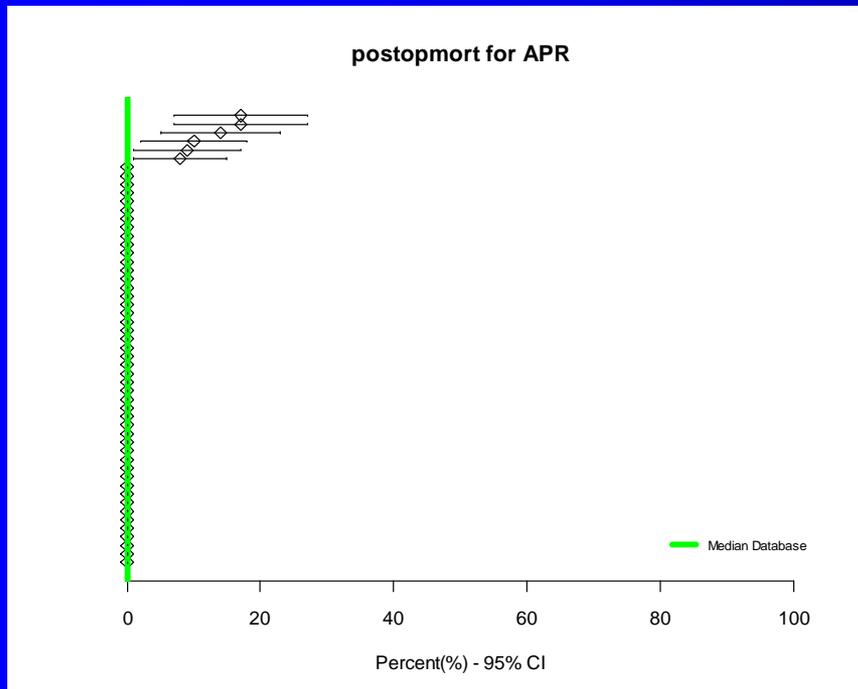
In hospital mortality after elective radical resection (if > 10 pts)



In hospital mortality after elective radical resection (if > 10 pts)

after APR

after SSO



No
 Yes (why?) _____
 reaction of other organ
 No
 Yes
 Ovariotomy
 Metastasectomy (specify) _____
 perforation of the rectum?
 Yes *hole in mesorectum*
 No
 Complete transection of the sigmoid?
 Yes
 No

Yes
 Irrigation of the rectum stump before reanastomosis
 No
 Yes (specify fluid) _____
 Type of reconstruction
 endoscopic polypectomy
 Local excision (disc excision)
 TEMS (transanal microsurgical resection)
 APR
 Hartmann (specify distal transection level):

 High anterior resection = CRA (anastomosis above peritoneal)

	1	2	3
ASA 1	19	55	5
ASA 2	67	38	47
ASA 3	14	7	48
In hosp mortality	0.6	1.8	0

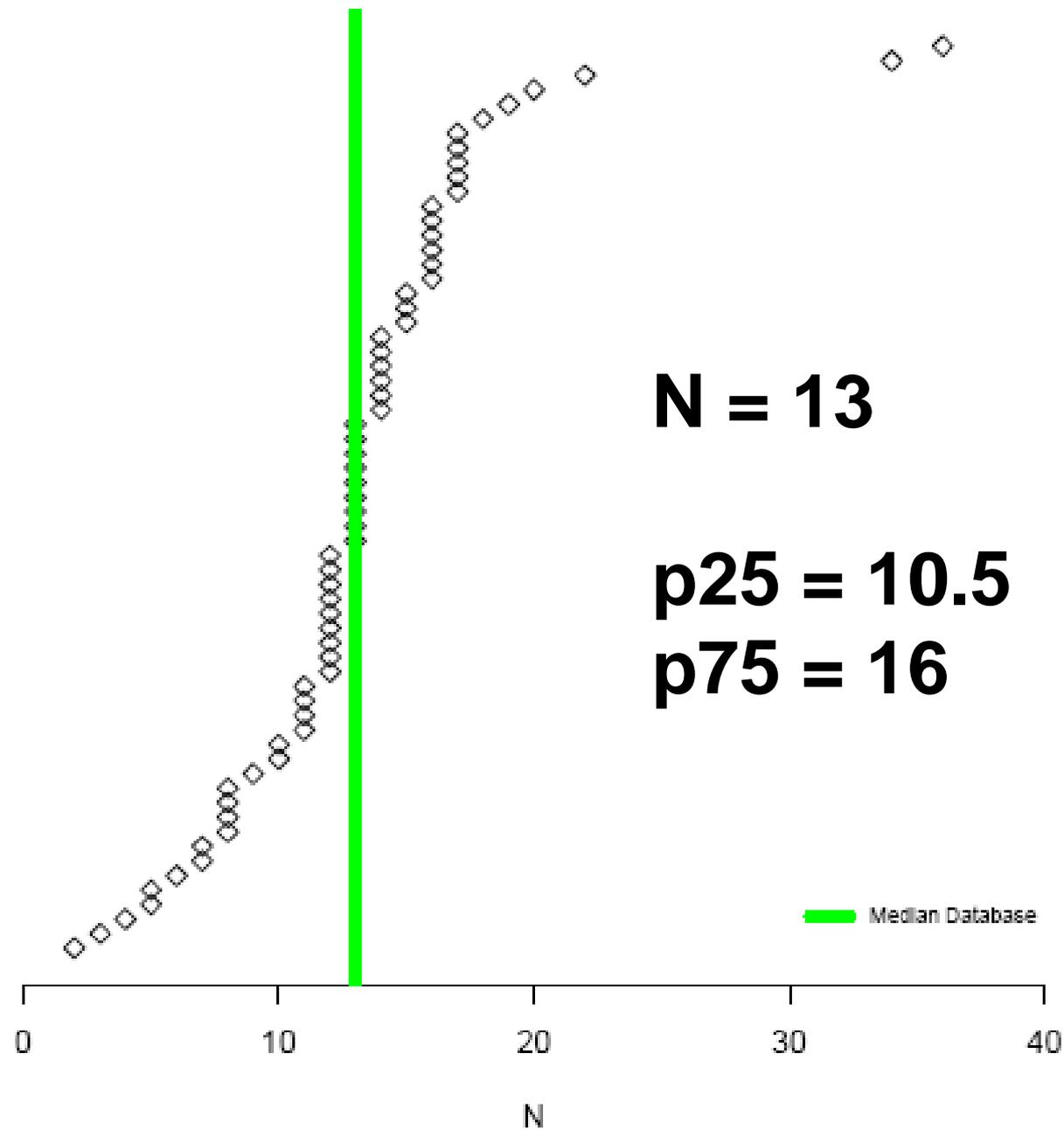
10 Surgical exploration
 Approach:
 Laparotomy
 Laparoscopy
 Converted laparoscopy

Ileum
 Other _____
 Type:
 loop
 terminal
 Reason(s)
 Routine

Pathology (1)

Report sheet used	90.9 %
Quality of TME reported	59.2 %
(y)pCRM reported	71.4 %
Distal margin status reported if low	95.3 %
Dworak regression grade reported after long course RT	70.8 %

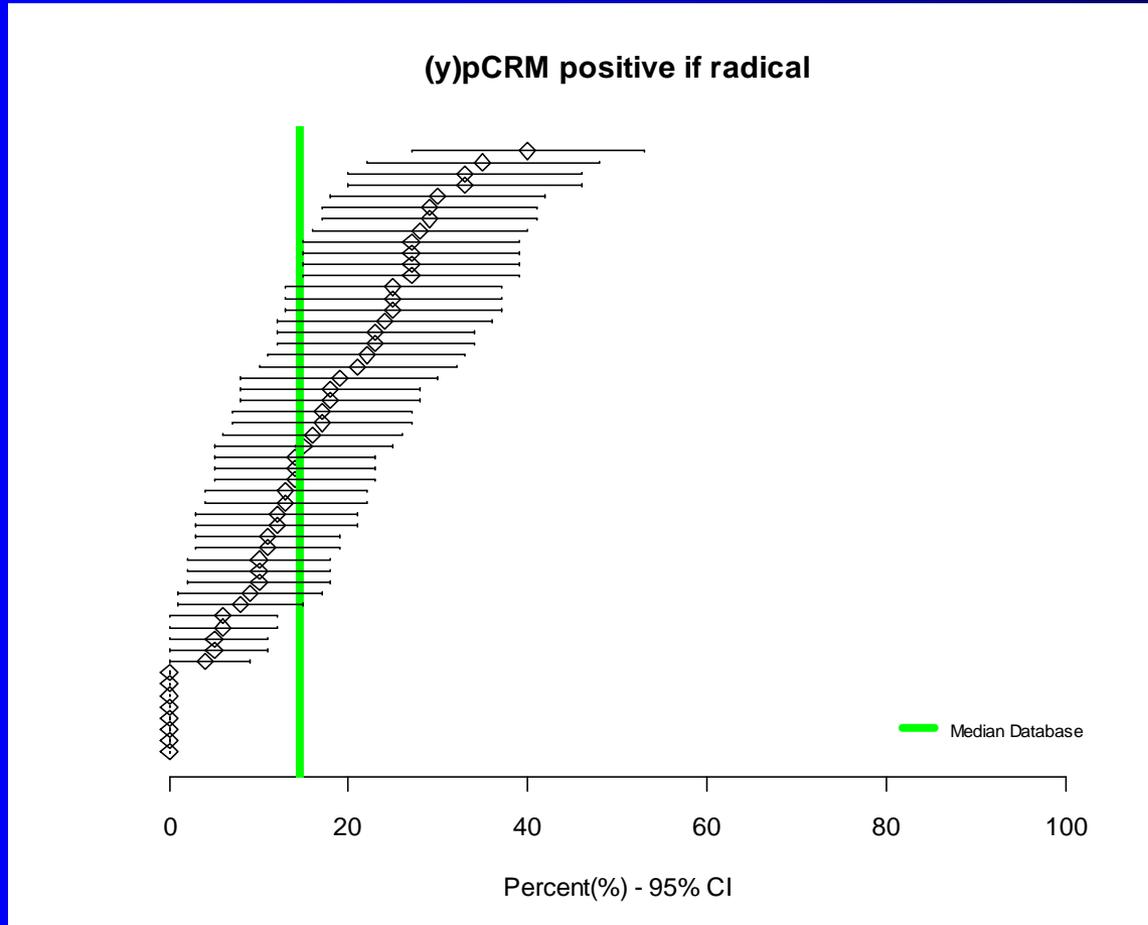
Median number of lymph nodes exam in no or short course neoadj RT



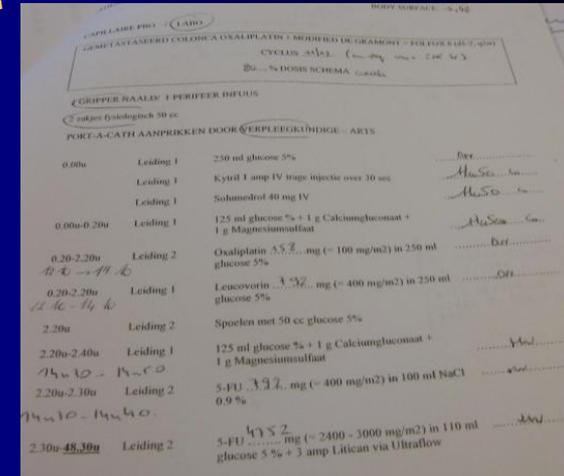
Pathology (2)

Quality of TME poor	10.9 %
(y)pCRM \leq 1 mm if radical resection	16.6 %
Distal margin invaded if SSO for low	2.4 %

Positive (y)pCRM after elective radical resection (if > 10 pts)



Adjuvant treatment



Data on adj chemo if (y)pStage III, R0

47

Data on adj R(C)T if pStage II-III, R0

7

Data on adj chemo if (y)pStage II-III, R0
started within 3 months

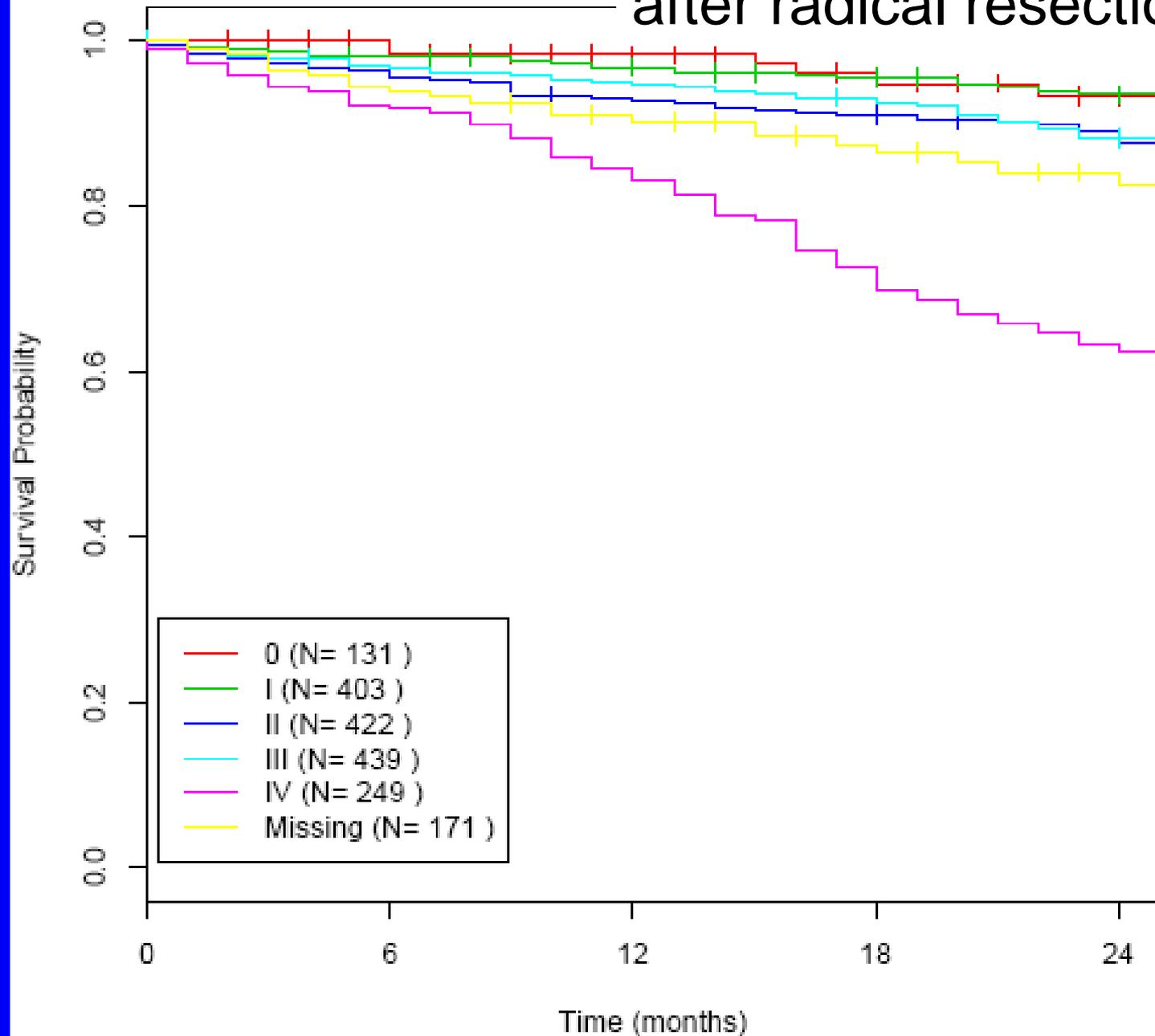
71

Data type of adj chemo for (y)pStage II-III, R0

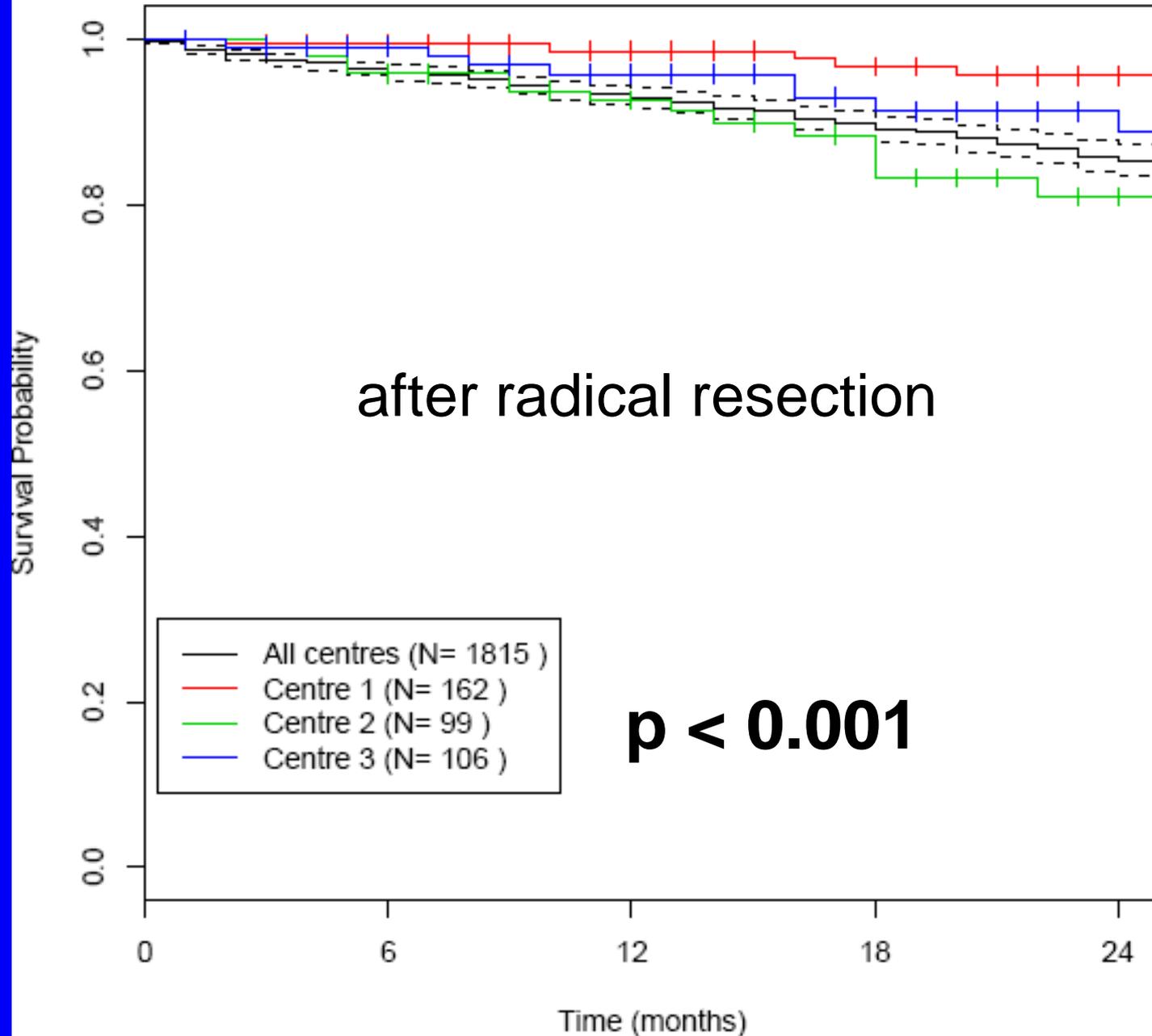
71

Observed survival by (y)pStage (N= 1815)

after radical resection



Observed Survival - all (y)pstages by centre



The project - CONCLUSIONS

- Profession-driven = voluntary participation
- Educational (re-action) not repressive (sanction)
- Multidisciplinary = teams, not individuals
- Open for all teams at any time
- Funding (government)
- Risk adjusted benchmark (peers, statisticians)
- Evolution of 'performance'
- Definition of targets / outliers (clinical > statist.)

What 'target value' for improvement ?

Median with CI 95%: mediocre progress

The 'top 10' teams ? with CI 95% or CI 90% ?
For every QCI or for a set of QCIs ?

How to improve in the 'top 10' ?

Statistical vs clinically relevant targets/differences

The participating teams - CONCLUSIONS

- Burden of registration
- Motivation of all team-players (intention vs practice)
- Quality of data (application of definitions, ...)
- Completeness of 'data' (patients, data)
- Fear for audit ('slow' but progressive particip.)
- Educational risk-adjusted benchmarking with re-action
- Variability always present
- Improvement always possible (low & high vol.)

