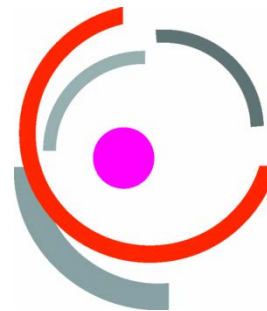


# Monitoring of equity in access to CRC screening and CRC care: two sides of the same coin

Ondřej MÁJEK

Institute of Biostatistics and Analyses, Masaryk University



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Institute of Biostatistics and Analyses, Masaryk University, Brno

- **Cancer screening programmes are effective in reducing cancer burden**
- **Continuous monitoring should be included to ensure high-quality cancer screening (benefits > harms)**
- **How should we set the health services?**
  - **understand the cancer burden**
  - **promote equity (population-based approach)**
  - **optimise the human, physical, technical and financial resources (already available & build capacity)**
  - **treatment services must be available**

# Sources of data for colorectal cancer screening information support

## MONITORING OF CANCER BURDEN

- Epidemiology of cancer in target population
- Long-term impact indicators

Source of data: CZECH NATIONAL CANCER REGISTRY

## MONITORING OF SCREENING PROCESS USING CLINICAL DATA

- Early performance indicators at screening centres
- Detection of cancer and precancerous lesions in screening

Source of data: SCREENING CENTRES

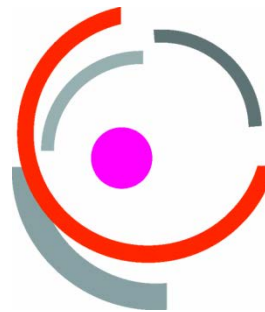
## MONITORING OF SCREENING PROCESS USING ADMINISTRATIVE DATA

- Population-based early performance indicators
- Monitoring of programmes accessibility by target population

Source of data: HEALTH INSURANCE COMPANIES – NATIONAL REFERENCE CENTRE

Information Support Provider  
MASARYK UNIVERSITY, INSTITUTE OF BIOSTATISTICS AND ANALYSES

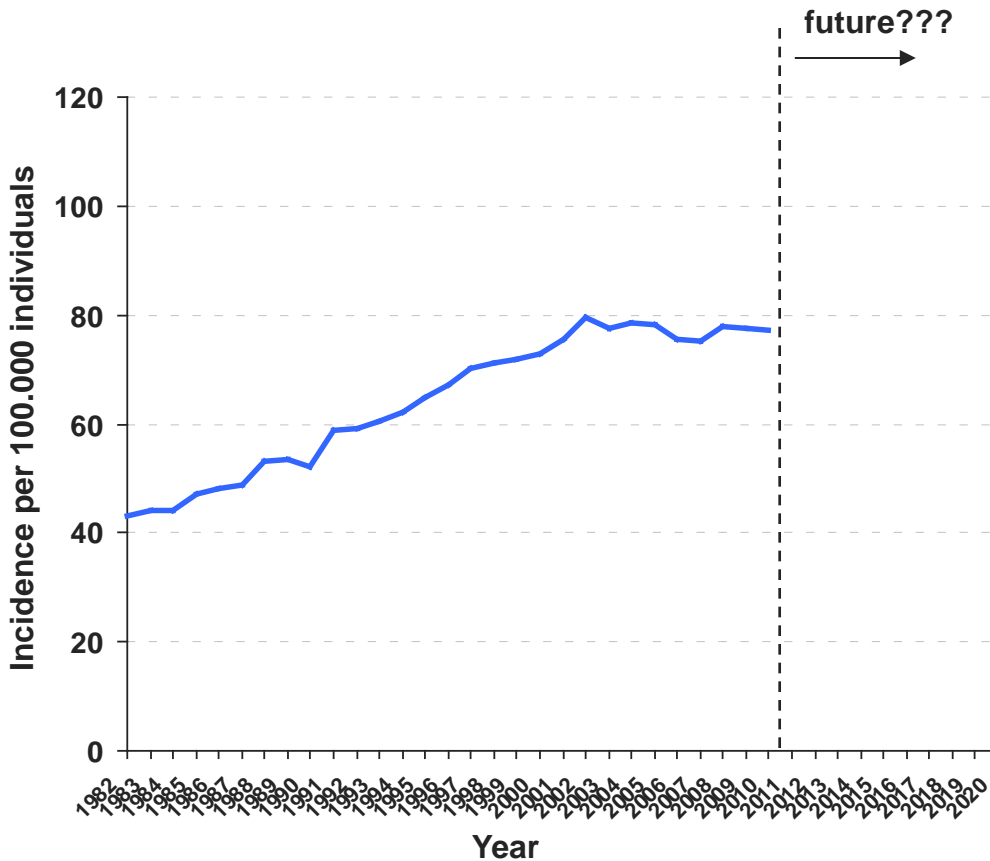
# Understanding the cancer burden (to population and healthcare system)



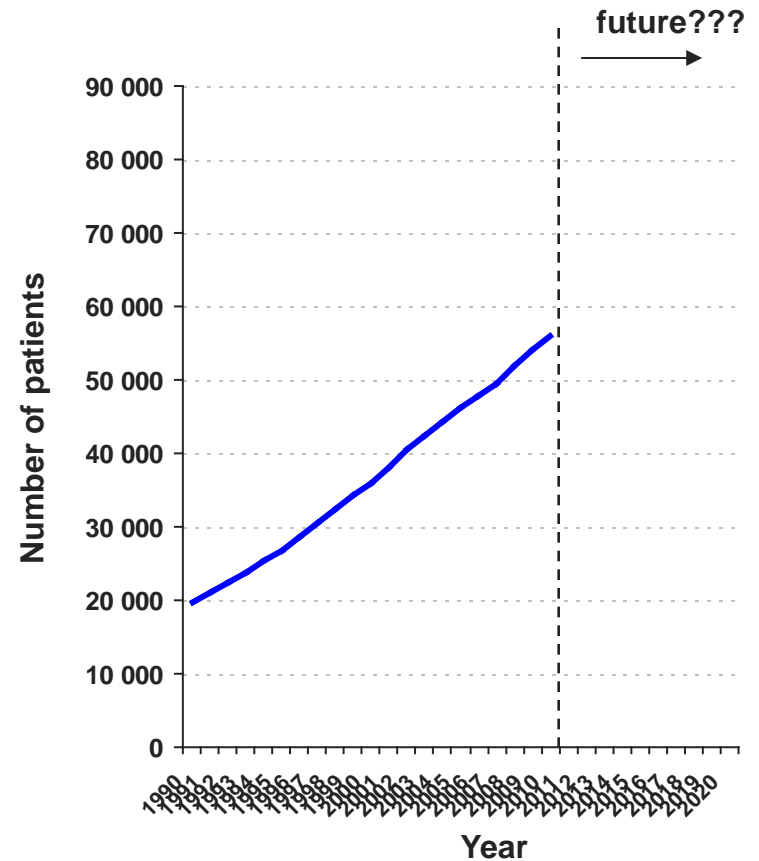
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# Future prediction of CRC incidence and mortality

## Time trends in CRC incidence

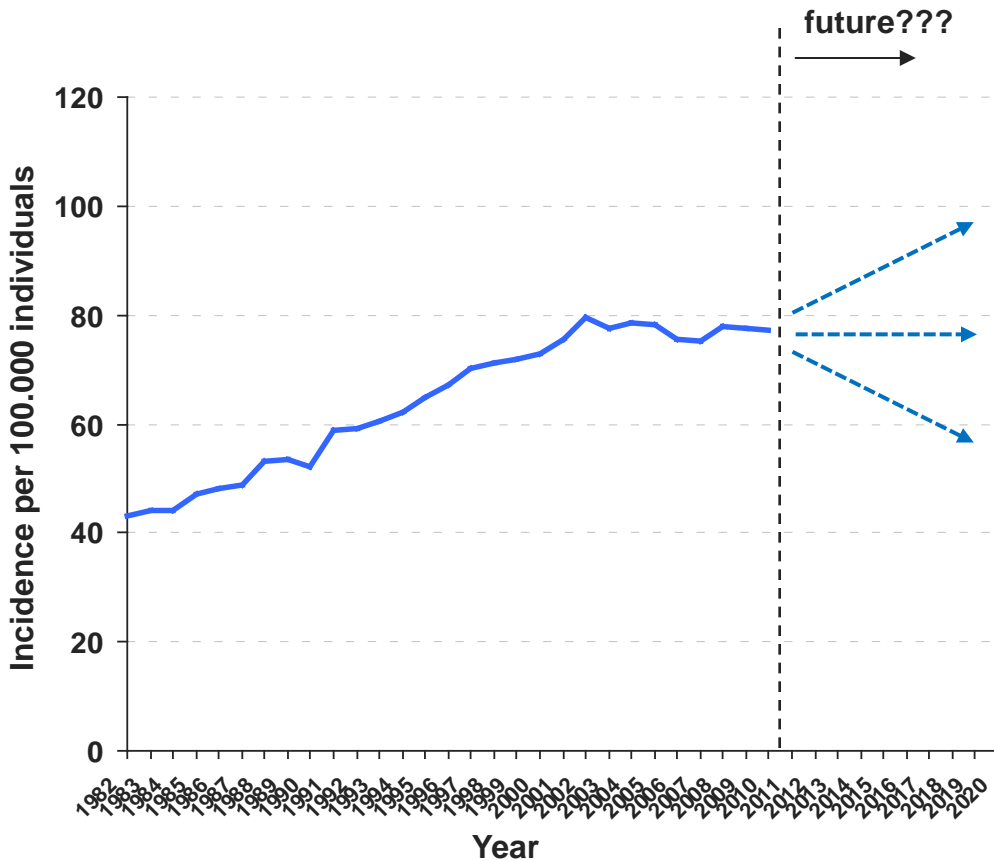


## Time trends in CRC prevalence

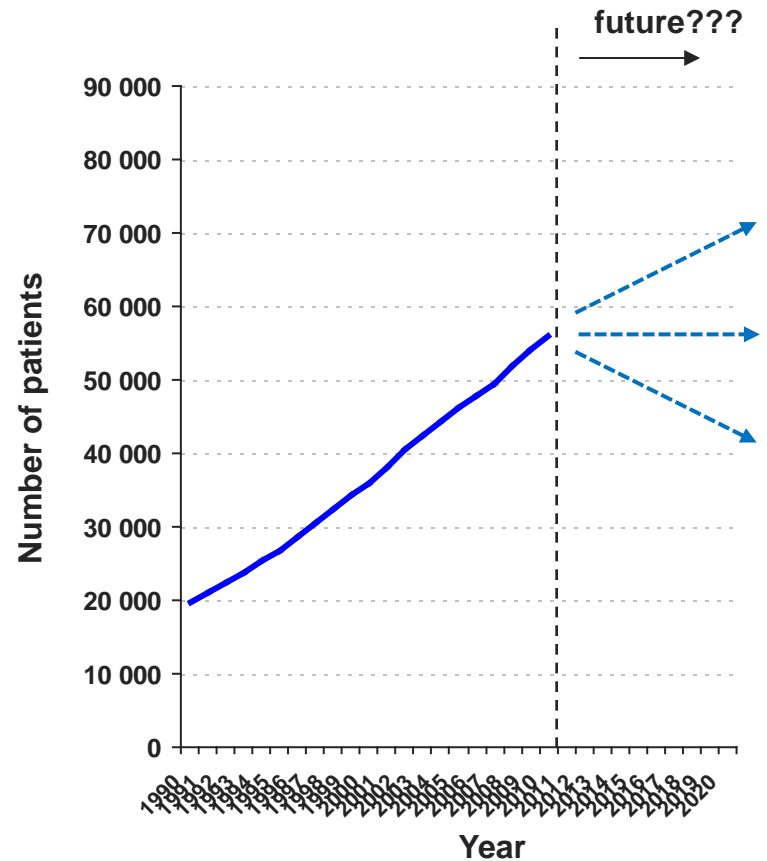


# Future prediction of CRC incidence and mortality

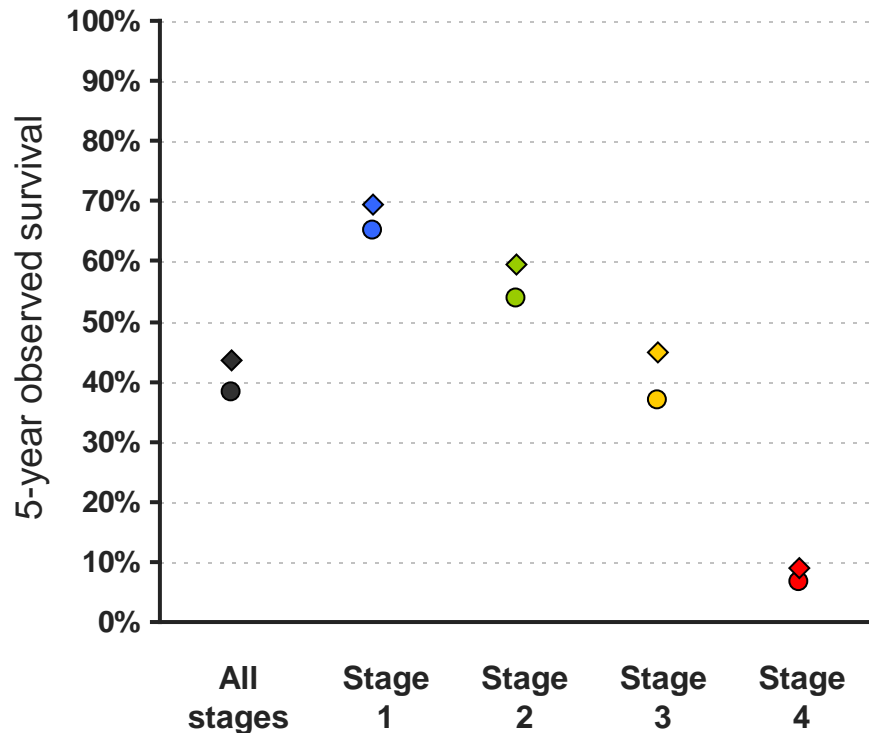
## Time trends in CRC incidence



## Time trends in CRC prevalence



# Time trend in 5-year observed survival of CRC patients



- ◇ Period analysis 2006-2010
- Period analysis 2001-2005

Colorectal cancer	5-year observed survival (95% confidence interval) period analysis	
	2001-2005	2006-2010
<b>All patients</b>	38.1 (37.7-38.5)	43.4 (43.0-43.8)
<b>Stage 1</b>	65.2 (64.1-66.2)	69.5 (68.6-70.4)
<b>Stage 2</b>	53.9 (53.0-54.7)	59.4 (58.5-60.2)
<b>Stage 3</b>	36.8 (35.8-37.8)	44.7 (43.8-45.6)
<b>Stage 4</b>	6.7 (6.3-7.2)	9.1 (8.6-9.6)

# Different scenarios in predictive modelling

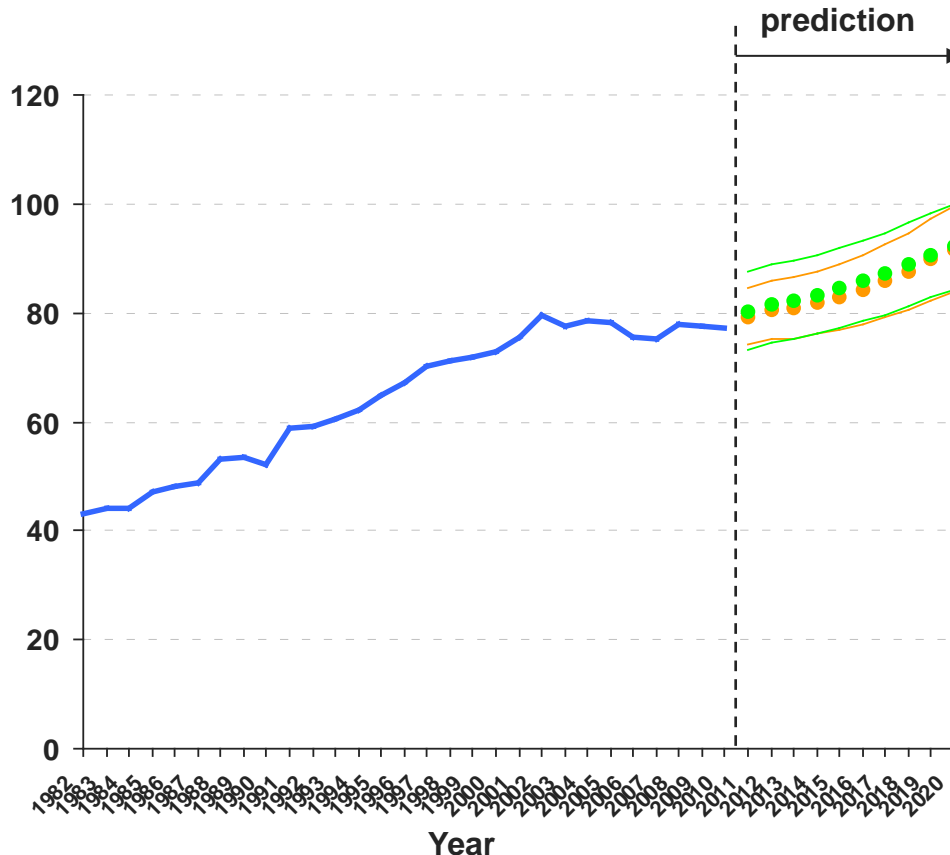
- Predictive modelling allows to consider different scenarios of future development of population characteristics → this leads to diverse predictions of incidence, prevalence and number of potentially treated patients

<b>Cancer incidence (period 2011-2020)</b>	<b>Survival of cancer patients (period 2011-2020)</b>	
	<b>Survival in 2011-2020 unchanged from values observed in 2010</b>	<b>Assumed improvement in survival</b>
<b>Incidence rate extrapolated based on data from 2001-2010</b>	Scenario 1	Scenario 2
<b>Assumed stable incidence (based on years 2006-2010)</b>	Scenario 3	Scenario 4



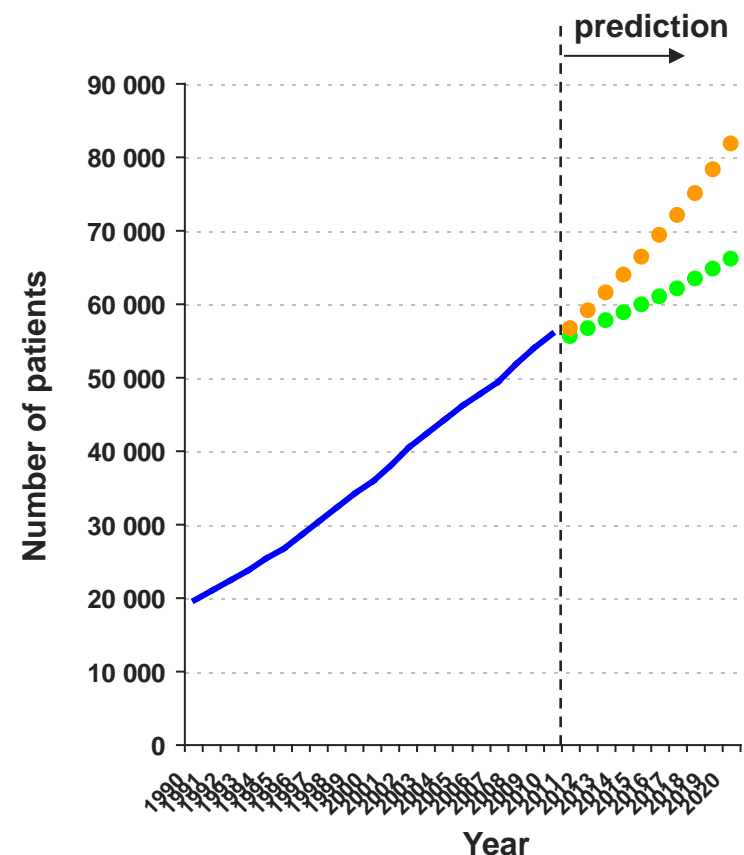
# Future prediction of CRC incidence and mortality

## Time trends in CRC incidence



- modelled incidence
- assumed stable incidence

## Time trends in CRC prevalence



- modelled survival
- assumed stable survival

# CRC incidence and prevalence predictions

C18-C20 Stage	Incidence – year 2015		Incidence – year 2020	
	n	90% CI	n	90% CI
1	2166	2017 - 2316	2609	2398 - 2819
2	1930	1823 - 2037	1971	1829 - 2114
3	2341	2209 - 2473	2802	2611 - 2993
4	2151	2002 - 2301	2362	2145 - 2579
Unknown	292	185 - 400	177	91 - 263
<b>Total</b>	<b>8880</b>	<b>8236 - 9527</b>	<b>9921</b>	<b>9074 - 10 768</b>

C18-C20 Stage	Prevalence – year 2015		Prevalence – year 2020	
	n	90% CI	n	90% CI
1	21 387	21 146 - 21 628	27 242	26 971 - 27 513
2	18 936	18 710 - 19 162	21 336	21 096 - 21 576
3	15 476	15 271 - 15 681	21 295	21 055 - 21 535
4	8329	8179 - 8479	9840	9677 - 10 003
Unknown	2484	2402 - 2566	2157	2081 - 2233
<b>Total</b>	<b>66 612</b>	<b>66 187 - 67 037</b>	<b>81 870</b>	<b>81 399 - 82 341</b>

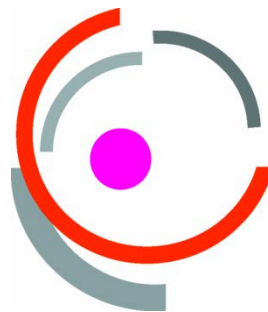
What about number of patients to be treated?

# Number of patients treated with CRC - predictions

<b>C18-C20 Year 2015</b>	<b>Newly diagnosed cancer patients</b>	<b>Recurrences from previous years</b>	<b>Disseminated recurrences from previous years</b>
Stage1	1915 (1783 - 2047)	798 (752 - 844)	
Stage 2	1803 (1703 - 1903)	1015 (963 - 1067)	
Stage 3	2196 (2072 - 2320)	1339 (1279 - 1399)	
Stage 4	1444 (1344 - 1544)	661 (619 - 703)	1779 (1710 - 1848)
Stage unknown	68 (43 - 94)	22 (14 - 30)	
<b>Total</b>	<b>7426 (6945 - 7908)</b>	<b>3835 (3733 - 3937)</b>	<b>1779 (1710 - 1848)</b>
		<b>13 040 (12 852 - 13 228)</b>	

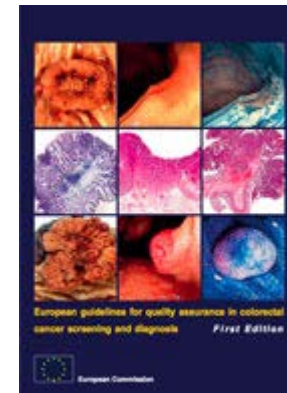
<b>C18-C20 Year 2020</b>	<b>Newly diagnosed cancer patients</b>	<b>Recurrences from previous years</b>	<b>Disseminated recurrences from previous years</b>
Stage1	2306 (2120 - 2492)	997 (945 - 1049)	
Stage 2	1841 (1709 - 1975)	1058 (1004 - 1112)	
Stage 3	2628 (2449 - 2808)	1729 (1661 - 1797)	
Stage 4	1585 (1440 - 1731)	815 (768 - 862)	1797 (1727 - 1867)
Stage unknown	41 (21 - 61)	16 (9 - 23)	
<b>Total</b>	<b>8401 (7739 - 9067)</b>	<b>4615 (4503 - 4727)</b>	<b>1797 (1727 - 1867)</b>
		<b>14 813 (14 613 - 15 013)</b>	

# Monitoring of cancer screening: equitable coverage and quality



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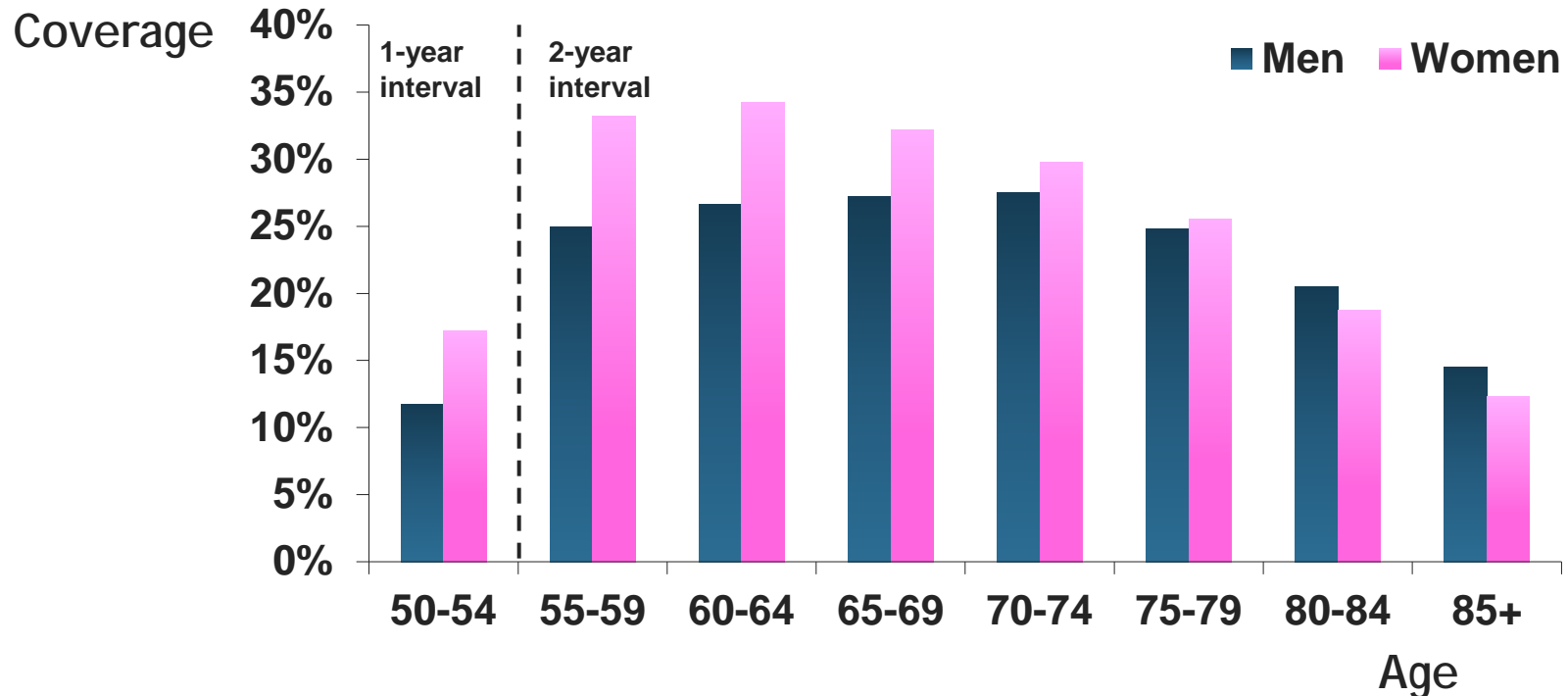
- **Programme coverage and uptake**
  - Coverage by invitation  
Recommendation: 95%
  - Coverage by examination
  - Uptake (participation) rate  
Recommendation : 45% / 65%



Source: European Guidelines  
Chapter 3

# Coverage by FOBT screening by age

2011-2012, N = 986 111 examinations, source of data: NRC



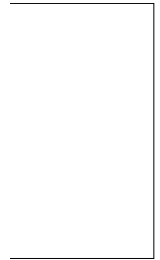
Total coverage of Czech population (2011-2012): **25.5%**

# Coverage by FOBT screening by district

2011-2012, N = 986 111 examinations, source of data: NRC

Ind

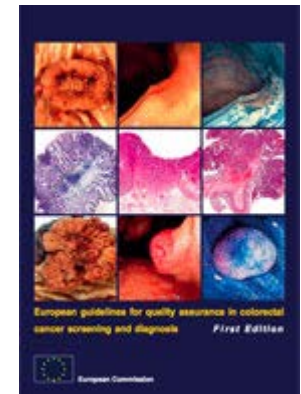
range in  
districts



Total coverage (2011-2012): **25.5 %** (range in districts: 16.0-38.9 %)

- **Outcomes with colonoscopy (CS) as primary screening test**
  - Inadequate CS rates
  - Complete CS rate
  - Positive CS rate
  - Detection rates of CS screening programmes
  - Referral to follow-up colonoscopy after CS
  - Follow-up colonoscopy compliance rate after screening CS
  - Completion of follow-up colonoscopy after CS
  - Endoscopic complications of CS screening programmes

**Recommendation : monitor the rate carefully**

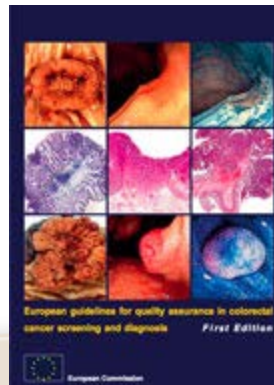


Source: European Guidelines



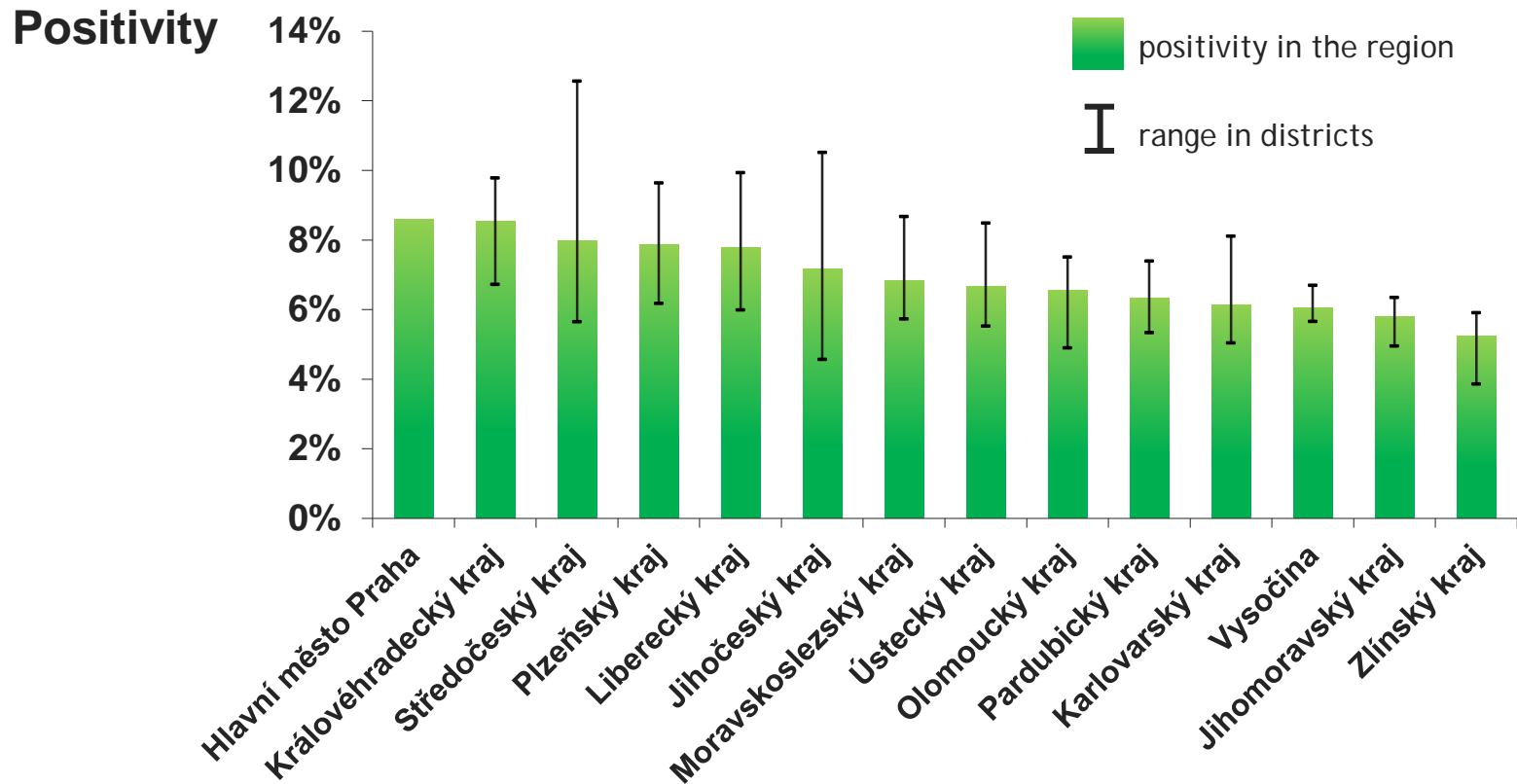
- **Outcomes with FOBT for primary screening**
  - Inadequate FOBT rate
  - **Positive FOBT rate**
  - Referral to follow-up colonoscopy after FOBT
  - Follow-up colonoscopy compliance rate
  - Completion of follow-up colonoscopy after FOBT  
Recommendation : 90% / 95%
  - Detection rates of FOBT screening programme
  - Stage of screen-detected cancers  
Recommendation : favourable compared to clinically diagnosed
  - Positive predictive values for FOBT screening programmes
  - Endoscopic complications in FOBT screening programme  
Recommendation : monitor the rate carefully

Source: European Guidelines



2012, N = 551 883 examinations, source of data: NRC

Individuals aged over 50

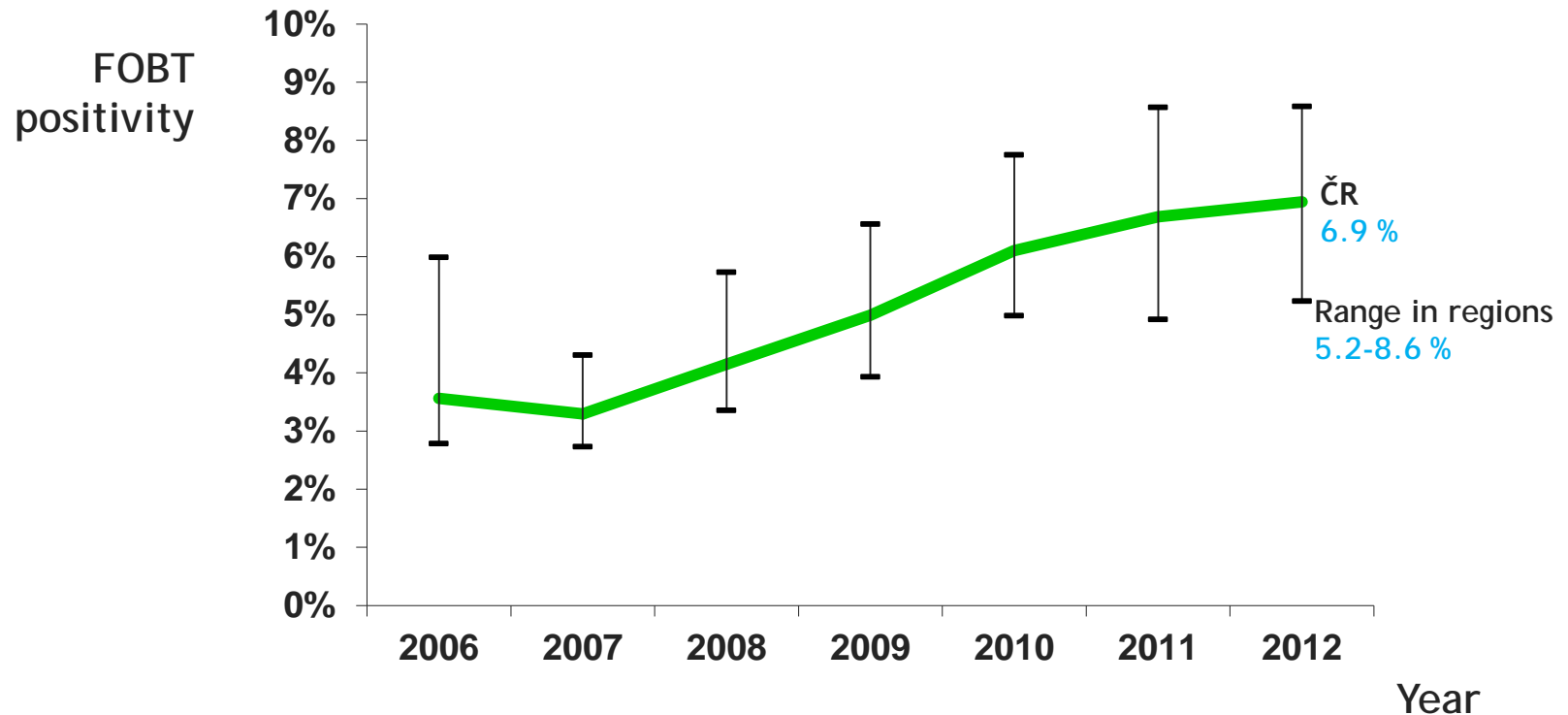


Total positivity (2012): 6.9 % (range in regions: 5.2-8.6 %)

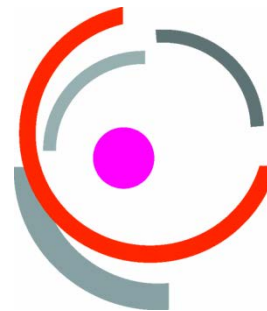
# Time trends in FOBT positivity

Source of data: NRC

## Individuals aged over 50

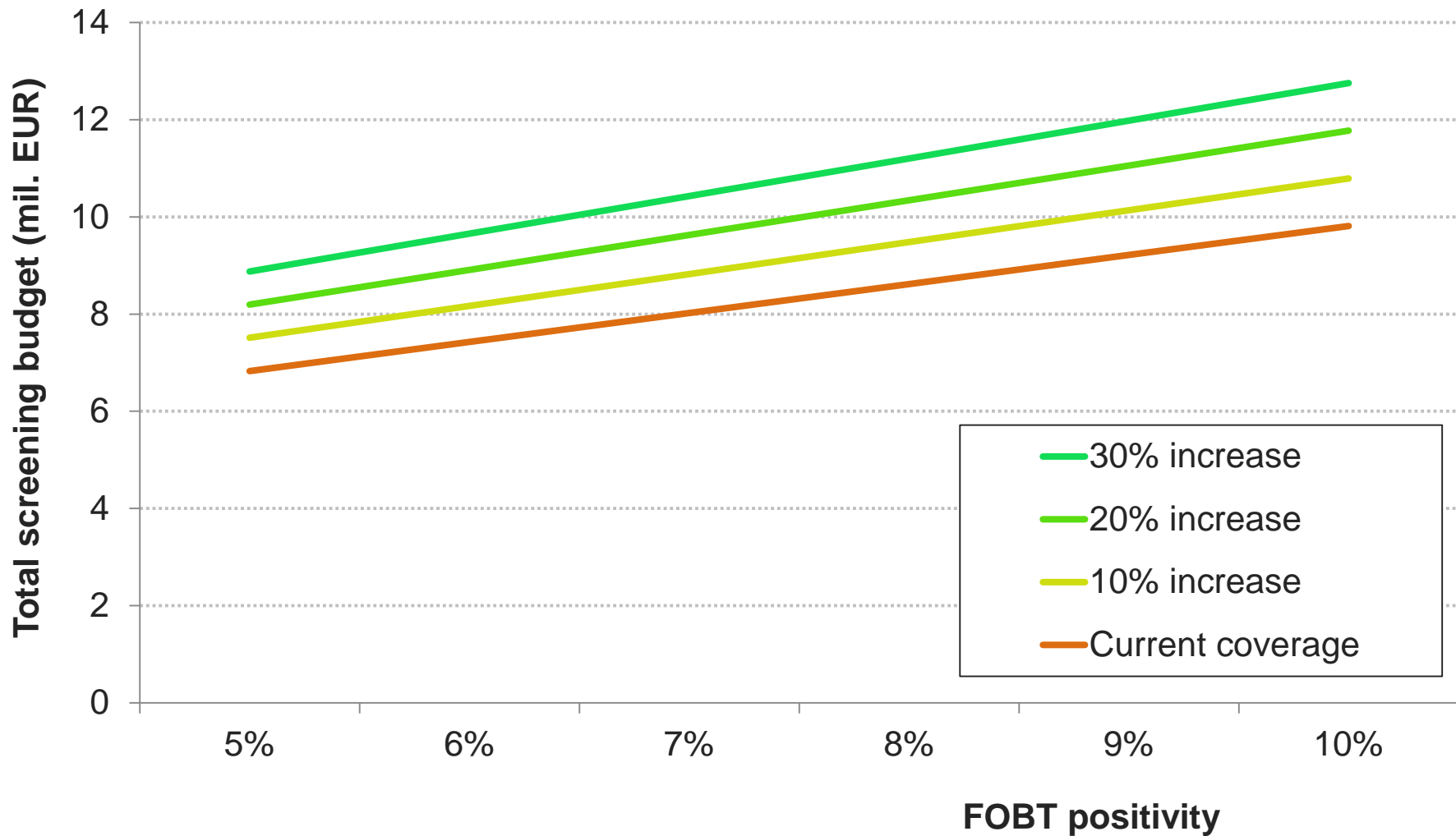


# Optimisation of resources: capacity of the network

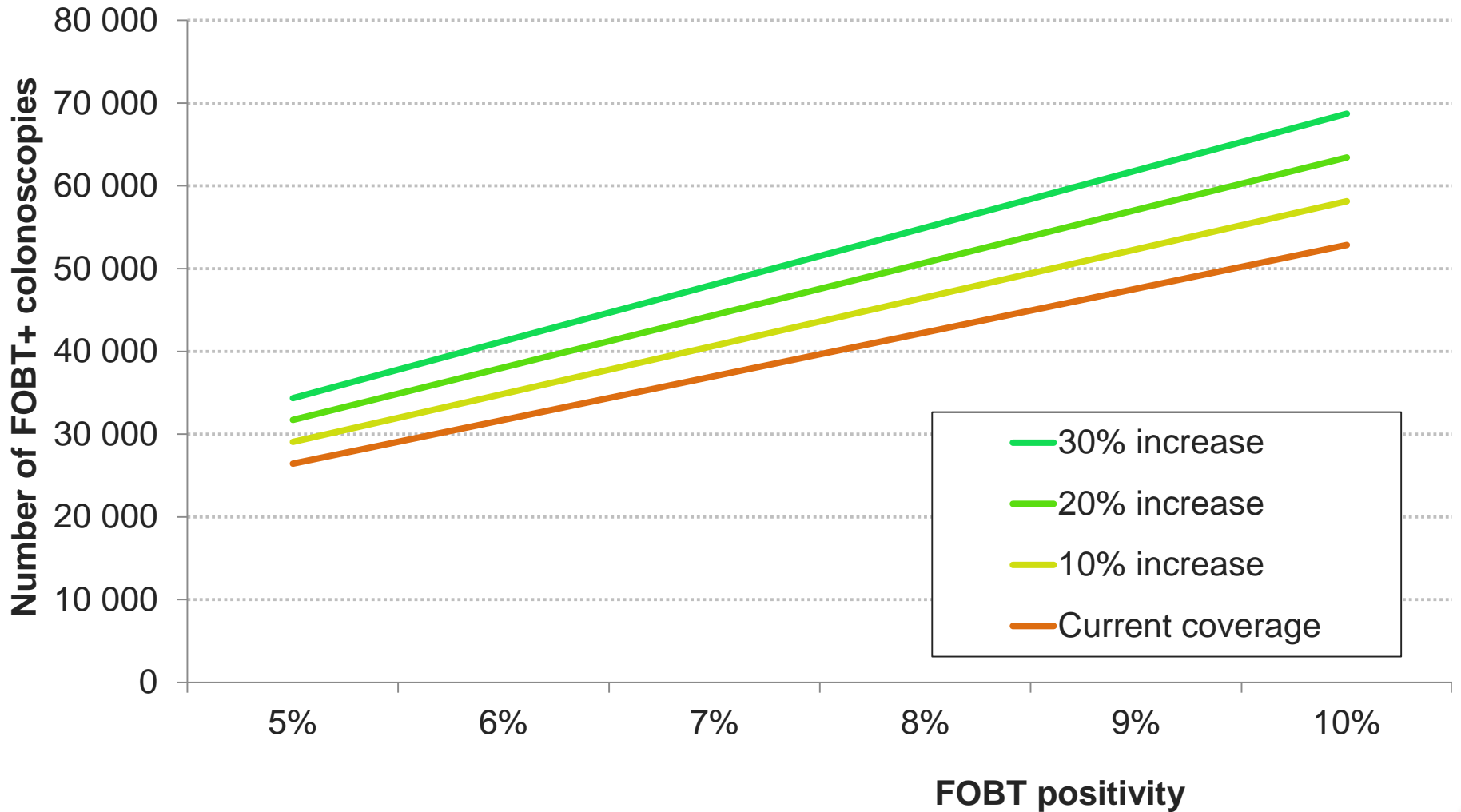


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# Changing system: impact on budget

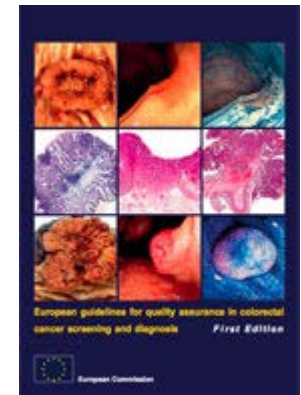


# Changing system: impact on colonoscopy volume



- **Screening organisation**

- Time interval between completion of test and receipt of results
- **Time interval between positive test and follow-up colonoscopy**  
Recommendation : 90% / 95% within 31 days
- Time interval between positive endoscopy and start of definitive management
- Time interval between consecutive primary screening tests



Source: European Guidelines

# Waiting time for FOBT+ colonoscopy

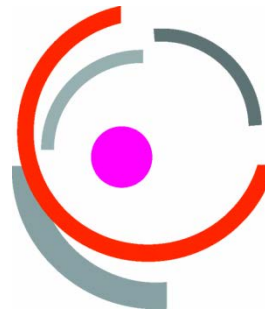
<b>Year</b>	<b>FOBT+ colonoscopies</b>	<b>% recorded FOBT date</b>	<b>Average waiting time (months)</b>
2006	5 334	94%	0,82
2007	5 679	92%	0,89
2008	7 458	96%	0,90
2009	11 711	96%	0,94
2010	18 327	95%	1,12
2011	20 132	94%	1,17
2012	21 137	95%	1,17
2013	21 837	95%	1,27



# Waiting time for FOBT+ colonoscopy in regions, 2012

Region	FOBT+ colonoscopies	% recorded FOBT date	Average waiting time (months)
Pardubický	1542	100%	0,92
Vysočina	882	83%	0,93
Olomoucký	1698	98%	1,00
Ostravský	2732	100%	1,11
Zlínský	1430	99%	1,12
Plzeňský	1281	98%	1,12
Praha - východ	1720	92%	1,25
Královéhradecký	1347	82%	1,26
Praha - jih	1232	94%	1,35
Praha - západ	1259	93%	1,38
Karlovarský	711	99%	1,38
Jihomoravský	2240	100%	1,51
Ústecký	1150	99%	1,52
Jihočeský	1539	99%	1,66
Liberecký	1074	78%	1,67
<b>Total</b>	<b>21 837</b>	<b>95%</b>	<b>1,27</b>

# Availability of the treatment

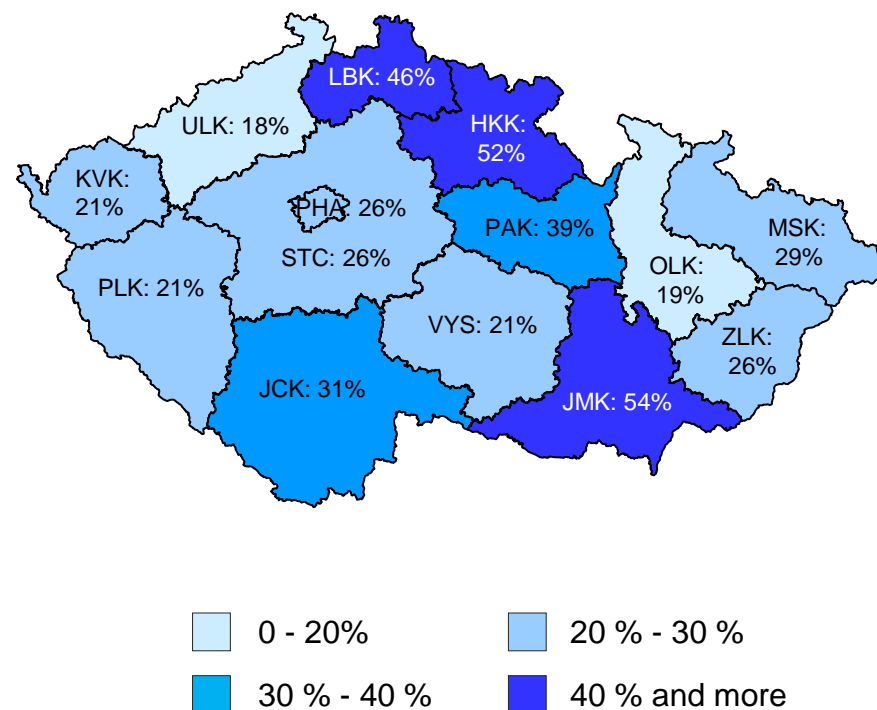


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# Number of patients initiating treatment: comparison of reality in CORECT registry and predictions - year 2012

Regions	No. of patients initiating treatment: CORECT registry <sup>A</sup>	Prediction <sup>B</sup>
PHA + STC	179 (26%)	700 (656; 744)
JCK	59 (31%)	193 (170; 216)
PLK + KVK	65 (21%)	315 (286; 344)
ULK	43 (18%)	235 (210; 260)
LBK	57 (46%)	124 (106; 142)
HKK	83 (52%)	159 (138; 180)
PAK	54 (39%)	140 (121; 159)
VYS	31 (21%)	151 (131; 171)
JMK	178 (54%)	329 (299; 359)
OLK	37 (19%)	191 (168; 214)
ZLK	44 (26%)	171 (149; 193)
MSK	115 (29%)	392 (359; 425)
<b>Czech Republic</b>	<b>945 (30%)</b>	<b>3100 (3008; 3192)</b>

Percentage of predicted number of patients, who initiated treatment



<sup>A</sup> Number of patients in registry and percentage of predicted number of patients.

<sup>B</sup> Estimate is accompanied with 90% confidence interval.

# Conclusion



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- **System is changing**
  - ageing population
  - trends in risk factors (period & cohort effects)
  - improved survival
  - organisation of the programme
  - performance parameters of screening process
- **Good data are needed for planning and monitoring**
  - to ensure accessible and high quality health services and thus promote equity
- **Cancer prevention is not only screening testing**
  - primary, secondary and tertiary prevention

- **Screening colonoscopy centres, for participation at data collection**



- **Providers of administrative and cancer registry data**



**KSRZIS**

