

A Global Perspective on Health System Strengthening

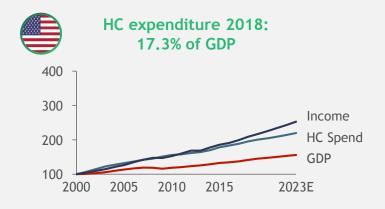
2019 SAAPI Conference - Dr Jonathan Lim

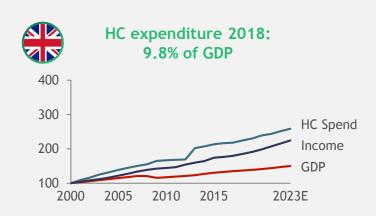


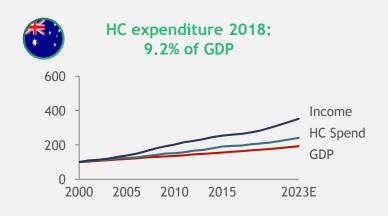


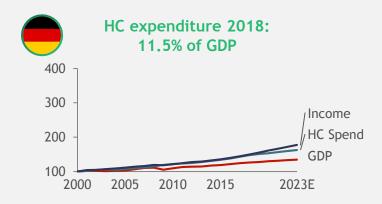


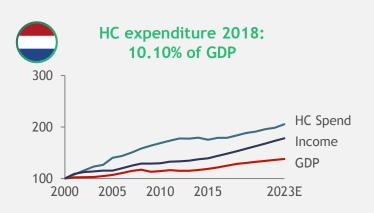
However, this development has come at a cost

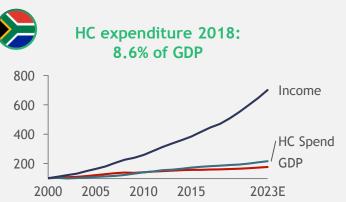












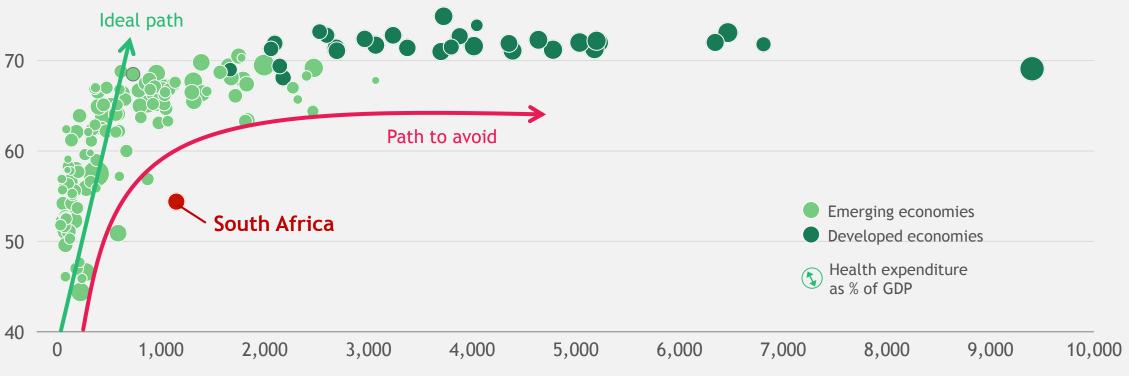
Notes: Index 100 at 2000, based on local currencies; Income = Personal Disposable Income

Source: WHO; EIU (May 2019); BCG analysis

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More importantly, greater spending does not necessarily yield better results

Outcome: Health-adjusted life expectancy¹ (years, 2015)



Input: Health expenditure per capita (PPP US\$, 2014)

^{1.} Health-adjusted life expectancy: Estimates the number of years in full health an individual is expected to live at birth by subtracting the years of ill health (weighted according to severity) from overall life expectancy. Sources: WHO, BCG analysis

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There remains massive variation in health outcomes across OECD

Breast cancer five year survival

Cervical cancer five year survival

Colorectal cancer five year survival

Diabetes lower extremity amputation

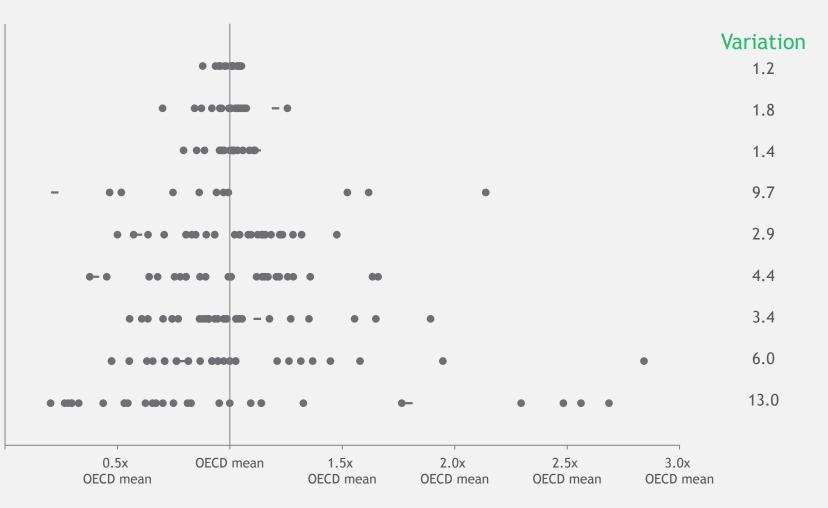
Hemorrhagic stroke 30 day mortality

Ischemic stroke 30 day mortality

AMI¹ 30 day mortality

Infant mortality

Maternal mortality



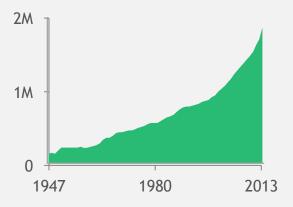
Note: Most recent data from 2011-13 used. Mexico and OECD candidate countries not included Source: OECD Health May 2016, BCG analysis

^{1.} Acute Myocardial Infarction

Part of this is driven by the exponential increase in health care complexity

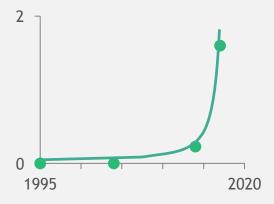
Exponential increase in medical knowledge

of new MEDLINE publications per year



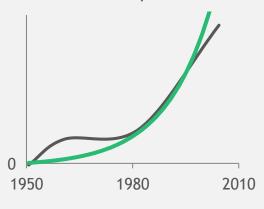
Exponential increase in number of human genomes sequenced

of sequenced human genomes (M)

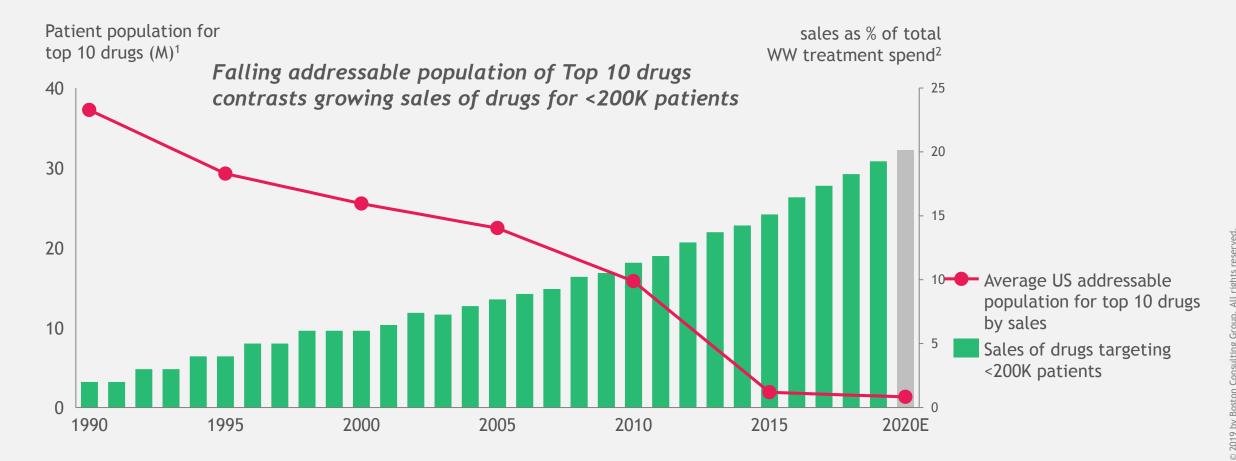


Exponential growth in medical practice complexity

of US MD Subspecialties



For industry and regulators, complexity is evidenced by interventions targeted at ever smaller pop. segments

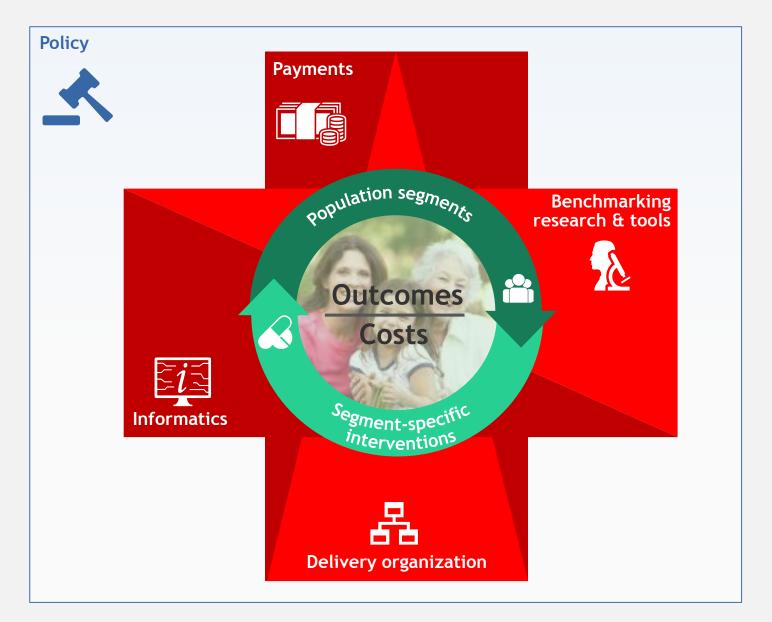


^{1.} For top 10 drugs by sales, addressable population calculated based on prevalence of first/major indication marketed. Where prevalence data not available, incidence rates were used instead 2. % sales of orphan drugs before 2000 was extrapolated from trend; Source: EvaluatePharma®, BCG analysis



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Value-based health care (VBHC):
Designing people-centered health systems



Source: BCG analysis





Bruno Bruins

Health Minister in the Netherlands

"It was agreed that by 2022, outcome information would be available for 50% of the disease burden, and that shared decision-making in the consultation room will be promoted"



Omar Ishrak

Chairman and CEO, Medtronic

"Moving to a value-based healthcare system is the only thing that can keep MedTech from being commoditised"

Leaders around the world are embracing the concept of VBHC



Christophe Weber

CEO, Takeda

"The acquisition of Shire will enable Takeda to significantly accelerate its transformational journey to become a value-based, R&D driven global biopharmaceutical leader..."

The full potential of VBHC to strengthen health systems will require new levels of collaboration



By Pharma and MedTech



By Hospitals and Insurers



By Regulators and Policymakers

Starting point for industry: Relentless focus on improving value for targeted populations

Which segments are underserved?

• Where do we see an unmet need?

- What outcomes to measure?
- How to measure and analyse data?
- What tools are needed to make the process seamless?



- What are the key pain points?
- What interventions can improve value?

Oshi offers clear proposition for all stakeholders



For patients...



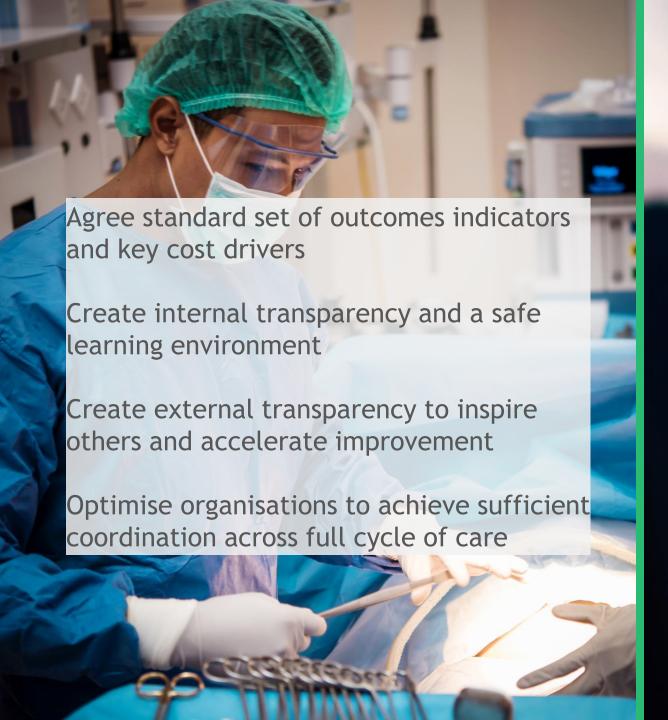
For care providers...



For academic institutions...

For payers...







Take a holistic approach to payment across the full cycle of care

Tailor type of payment to the needs of particular population segments

Leverage outcomes transparency to motivate the right behaviours - do not rely too much on strong financial incentives

ICHOM is a global non-profit defining outcomes standards

Founded in 2012 by Harvard Business School Professor Michael Porter, Boston Consulting Group, and Karolinka Institutet







International focus with involvement of multiple countries around the world



>650 Organisations

>15 National Registries

OECD PaRIS programme



Open source and peer reviewed standard sets



26 Sets covering 50-60% of the global disease burden



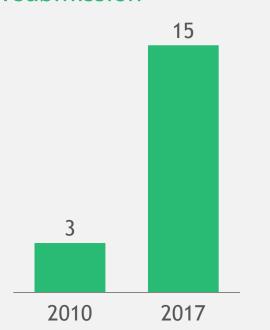
Endorsed by the OECD to map key disease areas



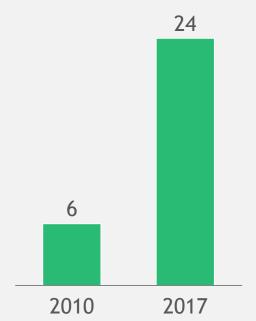
Independent, non-profit

Regulators are responding to increasing HC complexity by encouraging innovative approaches to clinical trials

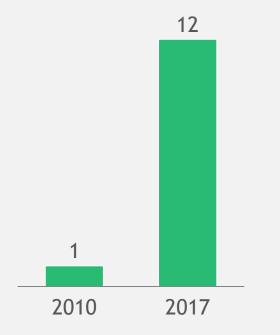
Number of trials where a novel clinical trial design was used in an approved FDA submission



Number of approved submissions which used surrogate endpoints



Number of approved submissions which included studies with PROs and/or Real World Data



Thrombus Aspiration during ST-Segment Elevation Myocardial Infarction

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ABSTRACT

BACKGROUN

The clinical effect of routine intracoronary thrombus aspiration before primary percutaneous coronary intervention (PCI) in patients with ST-segment elevation myocardial infarction (STEMI) is uncertain. We aimed to evaluate whether thrombus aspiration reduces mortality.

METHOD

We conducted a multicenter, prospective, randomized, controlled, open-label clinical trial, with enrollment of patients from the national comprehensive Swedish Coronary Angiography and Angioplasty Registry (SCAAR) and end points evaluated through national registries. A total of 7244 patients with STEMI undergoing PCI were randomly assigned to manual thrombus aspiration followed by PCI or to PCI only. The primary end point was all-cause mortality at 30 days.

RESULTS

No patients were lost to follow-up. Death from any cause occurred in 2.8% of the logarithms of the follopatients in the thrombus-aspiration group (103 of 3621), as compared with 3.0% in the PCI-only group (110 of 3623) (hazard ratio, 0.94; 95% confidence interval [CI], Up, 100 of 2.2; P=0.63). The rates of hospitalization for recurrent myocardial infarction at 30 days were 0.5% and 0.9% in the two groups, respectively (hazard ratio, 0.61; 95% CI, 0.34 to 1.07; P=0.09), and the rates of stent thrombosis were 0.2% and 0.5%, respectively (hazard ratio, 0.47; 95% CI, 0.20 to 1.02; P=0.06). There were no significant differences between the groups with respect to the rate of stroke or neurologic complications at the time of discharge (P=0.87). The results were consistent across all major prespecified subgroups, including subgroups defined according to thrombus burden and coronary flow before PCI.

CONCLUSION

Routine thrombus aspiration before PCI as compared with PCI alone did not reduce 30-day mortality among patients with STEMI. (Funded by the Swedish Research Council and others; Clinical Trials, gov number, NCT01093404.)

From the Department of Cardiology, Öre bro University Hospital, Örebro (O.F., F.C.). Department of Medical Sciences. Cardiology, and Uppsala Clinical Research Center, Uppsala University, Uppsala (B.L., O.O., S.K.J.), Department of Cardiology Lund University Hospital, Lund (G.K.O., D.E., J.H.), Department of Cardiology, Sahlgrenska University Hospital, Gothen burg (E.O., O.A.), Department of Cardiology, Karolinska Institutet, Sodersjukhuset (M.A.), and Cardiology Unit, Department tal (U.J.), Stockholm, Department of Cardiology, Karlstad Hospital, Karlstad (M.D.), Department of Cardiology, Gävle Hospital, Gävle (L.H.), PCI Unit, Sunderby Hospital, Sunderby (A.C.J.), Department of Cardiology Västeräs Hospital, Västeräs (A.K.), Department of Cardiology, Heart Center, Umea University, Umea (LN.). Department of Cardiology, Borås Hospi tal. Borás (L.R.). Department of Radiolo (L.S.), and Department of Cardiology, Faun Hospital, Falun (I.S.) — all in Sweden: Department of Cardiology and Cardiovascular Research Center, Landspitali Univer sity Hospital of Iceland, Revkiavik, Iceland (T.G.); and Department of Cardiology, Aarhus University Hospital, Skejby, Aarhus, to Dr. Fröbert at the Department of Cardiology, Örebro University Hospital, Södra Grev Rosengatan, 701 85 Örebro, Sweden, or at ole.frobert@orebroll.se.

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Unfortunately, too many of the decisions made today about health and healthcare are not supported by high quality evidence. Prospectively designed registries and cohort studies in the context of clinical practice are highly valuable, and randomised trials conducted in the context of clinical practice...may be the most important source of knowledge in the future...

- Former FDA Commissioner Robert M. Califf

Regulatory harmonisation across national borders enables access to medicines that drive better outcomes

West Africa

Established: WAHO, 2009 MRH project, 2015

Ambition:

• Strengthen registration, quality control of medicines, pharma production and pharmacovigilance

Key achievements so far:

- Developed common regulatory submission process
- Review sessions for submissions planned for end 2018

East Africa: EAC

Established: 2012

Ambition:

Develop regional regulatory strategies and common review process

Key achievements so far:

- 27 products evaluated; 4 approvals
- ~ 30-40% faster assessment process than national reviews

ASEAN

Established: 1967 / 1999

Ambition:

 Develop a transparent regulatory process and standardize regulation requirements

Key achievements so far:

Defined ASEAN common technical dossier

PANDRH

Established: 1999

Ambition:

• Strengthen regulatory functions, and develop guidelines for regulation of health technologies

Key achievements so far:

 Developed and adopted 23 technical documents

ZAZIBONA

Established: 2013

Ambition:

 Collaborate in assessment and inspections for medicines

Key achievements so far:

• 179 submissions reviewed: 90 products with final decisions (mean decision time of 9 months)

Gulf: GCC-DR

Established: 1999; reviews from 2002 Ambition:

Review/approve pharmaceutical companies and their drugs via centralised registration and inspections

Key achievements so far:

Approved 413 products (2006-10) with approval timelines between 114-206 days

Low

Now is the time for us all to shape and strengthen the health system of the future





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