

PAY FOR PERFORMANCE IN HEALTH CARE

If a physician make a large incision with an operating knife and cure it, or if he open a tumor (over the eye) with an operating knife, and saves the eye, he shall receive ten shekels in money.

If a physician make a large incision with the operating knife, and kill him, or open a tumor with the operating knife, and cut out the eye, his hands shall be cut off.

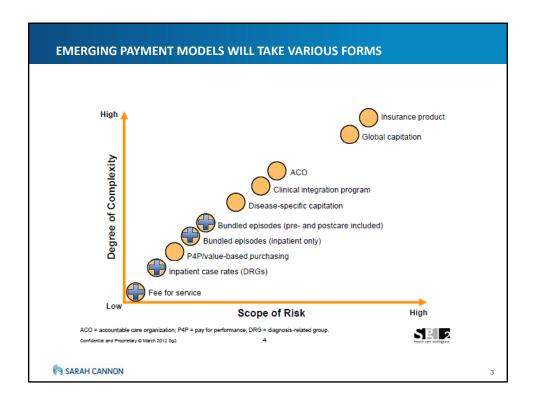




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PAY FOR PERFORMANCE

- Background
 - Arrangement where a portion of payment is based on performance of a defined measure
 - Typically another component of remuneration independent of amount at risk
 - Most current discussions address quality or performance objectives but could also target profitability, volume or patient satisfaction
- Goals
 - Improve quality of care
 - Control rate of growth in health care costs
 - Adoption of health information technology and EMR
 - Promote development of preventive services

Pay for Performance in Health Care CRS Report for Congress Updated December 12,2006

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FOUR TYPES OF PERFORMANCE MEASURES

- Clinical Outcomes (SCTOD)
 - Preferred standard
 - Often difficult to collect
- Process Measures (FACT)
 - Address proper delivery of healthcare services and practice p
 - Easier to collect but may not be consistent with outcomes
 - Often focus on underuse of services and may be cost-increas
- Structural Measures
 - Health information technology
- Patient Satisfaction Measures
 - Can be controversial
 - Easy to collect and may enhance compliance
 - No clear link to satisfaction and technical quality













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PERFORMANCE UNIT FOR MEASUREMENT AND REWARD

- Should reward the agent or decision maker directly for changes in the standard being measured
- · Accountability without responsibility is inappropriate and can be counter-productive
- Does it work?:
 - Attributing cause and effect with regard to improvements in health can be difficult because improvements may result from multiple factors
 - Health care interventions are often collaborative and may not be attributable to a single individual or provider
 - Who should be rewarded?
 - Top-performers
 - · Performers with greatest improvement?
 - All who meet a threshold?





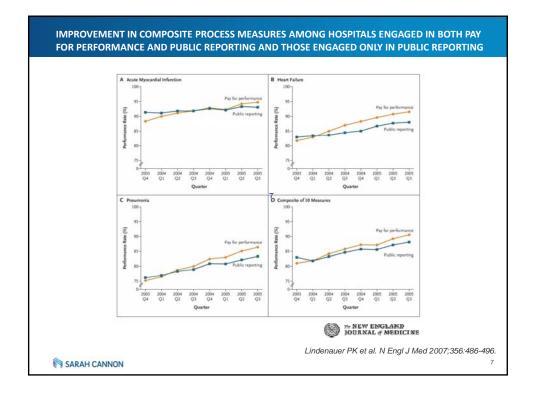


Table 5. Estimates of Incremental Effect of Pay for Performance. **Analytic Approach** Incremental Effect of Pay for Performance Acute Myocardial Infarction Composite of 10 Measures Heart Failure Pneumonia % (95% CI) 4.3 (2.5-6.1) 5.2 (2.8-7.7) Matched for hospital characteristics 2.6 (1.3-3.9) 4.1 (2.6-5.5)

ESTIMATES OF INCREMENTAL EFFECT OF PAY FOR PERFORMANCE.

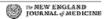
matched and adjusted for baseline perfor-mance, condition-specific volume, and all hospital characteristics; 1.9 (0.8-3.1) 3.8 (2.1-5.5) 3.5 (2.3-4.7)

Unmatched and adjusted for baseline performance, condition-specific volume, and all hospital characteristics; hospitals that declined participation in pay for performance included and grouped with those that agreed to participate¶ 1.8 (0.9-2.8) 2.8 (1.4-4.2) 2.7 (1.7-3.6)

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P-0.001 for all categories, unless otherwise noted, Y Multiple linear regression of matched pairs was adjusted for baseline performance and condition-specific hospital volume. K Multiple linear regression of data for 2490 hospitals that engaged in pay for performance and public reporting was adjusted for hospital size, teaching status, region, location, ownership status, baseline performance, and condition-specific volume. P = 0.002.

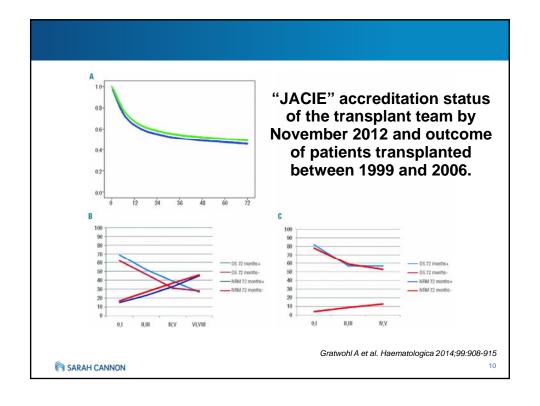
I/=0.0022.
Hultiple linear regression of data for 2490 hospitals that engaged in pay for performance and public reporting was adjusted for hospital size, teaching status, region, location, ownership status, baseline performance, and condition-specific volume; hospitals that declined to participate or withdrew from the Hospital Quality Incentive Demonstration were added to the pay-for-performance group to attempt to account for a volunteer effect.



Lindenauer PK et al. N Engl J Med 2007;356:486-496.

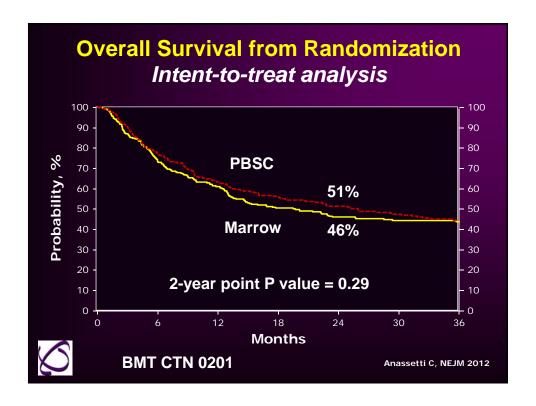


	Meaningful	Measureable	Actionable
1 Yr. OS	Meaningrui	Measureable	Actionable
FACT	+	4	4
100 day OS	+	_	4
Readmission	4	_	_
HAC	-	_	-
cGVHD	4	_	_
Pt. Reported Outcomes	4	-	4
Marrow vs PBSC	4	+	4
G-csf post allo	4	+	+
Survivorship Measures	4	-	-
Data Management	-	4	-

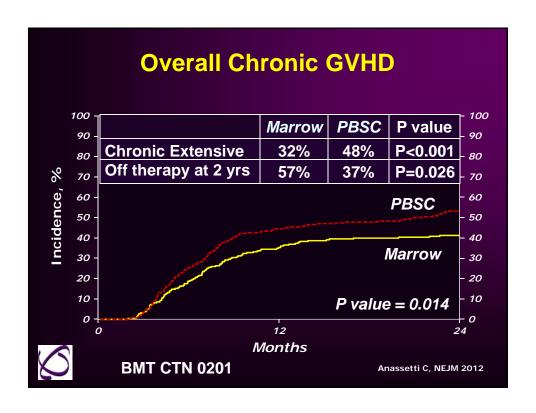




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G-csf post allo	4	+	+
Survivorship Measures	4	-	_
Data Management	4	4	-







Study-author	Transplant population	Stem cell source	N	Comparison	Primary conclusion
CIBMTR-Khoury et al. ⁴³	AML, CML	BM=2110 PBSC=609	2719	Patients who received G- CSF in first 7 days post HSCT versus others	G-CSF shortened time to ANC recovery; no change in D30 or D100 TRM. No changes in GVHD, LFS or OS
EBMT-Ringdenet al. 41	AML	BM=1789 PBSC=434	2223	Patients who received G- CSF in first 14 days post HSCT versus others	G-CSF worsened acute and chronic GVHD, TRM, OS and DFS in BM but not in PBSC transplants
CIBMTR-Eapen et al. ⁴²	Pediatric and adolescent	BM=630 PBSC=143	773	Children who received G or GM-CSF in first 7 days post HSCT versus others	G-CSF worsened TRM, treatment failure and OS
Meta-analysis–Ho et al. ⁴⁵	9 prospective randomized trials, 8 retrospective cohort comparisons, 1 case- controlled study	BM=1056 PBSC=142	1198	Patients who received G or GM-CSF post HSCT versus others	No difference in TRM, GVHD, or 100 day survival
Meta-analysis- Dekker et al. 44	34 randomized controlled trials	BM and PBSC		Patients who received G- or GM-CSF post auto or allo HSCT prior to neutrophil engraftment versus others	Growth factors reduced documented infections but did not impact acute GVHD or TRM



Recommended	6 Months	1 Year	Annually	Recommended	6 Months	1 Vear	Annually
screening/prevention	O INOTITIES	i i cui	Airidully	screening/prevention Skeletal	0 1110111110		, annual,
mmunity				Bone density testing (adult women,			
Encapsulated organism prophylaxis	2	2	2	all allogeneic transplant recipients and	4	1	+
PCP prophylaxis	1	2	2	patients at high risk for bone loss)	•		
CMV testing	2	2	2	Nervous system			
Immunizations	1	1	1	Neurologic clinical evaluation	+	1	1
Ocular				Evaluate for cognitive development		1	1
Ocular clinical symptom evaluation		1	1	Endocrine			
Ocular fundus exam	+	1	+	Thyroid function testing		1	1
Oral complications				Growth velocity in children		1	1
Clinical assessment	1	1	1	Gonadal function assessment			
Dental assessment	+	1	1	(prepubertal men and women)	1	1	1
Respiratory				Gonadal function assessment			
Clinical pulmonary assessment	1	1	1	(postpubertal women)		1	+
Smoking tobacco avoidance	1	1	1	Gonadal function assessment		+	+
Pulmonary function testing	+	+	+	(postpubertal men)		+	+
Chest radiography	+	+	+	Muco-cutaneous			
Cardiac and vascular Cardiovascular risk-factor				Skin self-examination and sun	1	1	1
assessment	+	1	1	exposure counseling Gynecological examination in			
iver				women	+	1	1
Liver function testing	1	1	+	Second cancers			
Serum ferritin testing		1	+	Second cancer vigilance counseling		1	1
ídnev				Screening for second cancers		1	1
Blood pressure screening	1	1	1	Psychosocial			
Urine protein screening	1	1	1	Psychosocial/QOL clinical	1	1	1
BUN/creatinine testing	1	1	1	assessment			
DOT VOI CALITIMIC TOSTING				Sexual function assessment	1	1	1

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cGVHD	4	_	_
Pt. Reported Outcomes	-	-	-
Marrow vs PBSC	4	+	+
G-csf post allo	+	+	+
Survivorship Measures	4	-	-
Data Management	+	4	4



CONCLUSIONS

- Pay for Performance may not make currently make sense for HSCT.
 - FACT/SCTOD
 - Case Rates
 - Narrow networks
- Goals for Pay for Performance programs need to be clearly defined.
 - Quality/Outcomes
 - Financial Alignment
- Careful consideration must be given to metrics:
 - Meaningful, measurable, actionable
 - Metrics that leverage CIBMTR data sets preferable
 - Partnership with payers in determining comparative effectiveness and value going forward
- Incentives need to be aligned with responsible parties.

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