

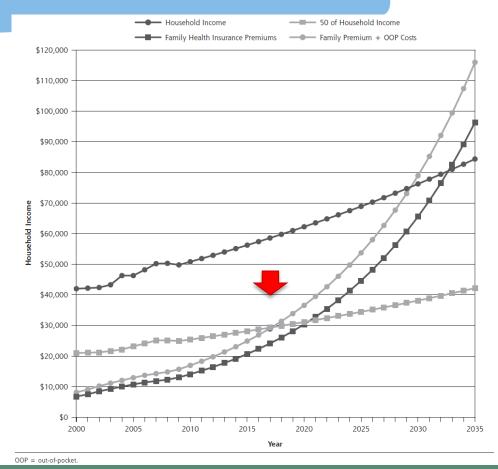
### **BCBS Georgia Cancer Care Quality Program**

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GASCO September 6, 2014



### **Rising Healthcare Costs Are Unsustainable**



In 3 years, premiums and out of pocket health care costs for a family are projected to equal half the median household income



## The cost of cancer treatment is a financial burden

Huge Costs	Like new drugs	Impact productivity	and create hardships
\$267 billion	\$100,000	36%	\$26,860
total cost of cancer in the U.S Includes medical costs and costs from lost productivity. <sup>1</sup>	the average yearly cost for newer oncology products <sup>2</sup>	of employees do not return to work after cancer treatment. <sup>3</sup>	the mean amount of debt for cancer patients. <sup>4</sup>

1-Source: National Business Group on Health: A Purchaser's Guide to Clinical Preventive Services: Moving Science into Coverage http://www.businessgrouphealth.org/pub/f2f59214-2354-d714-5198-3a8968092869 (accessed February 2013).

<sup>2-</sup> Source: Chase B. What's a Cancer Drug Worth? August 23, 2011. http://www.minyanville.com/businessmarkets/articles/seattle-genetics-adcetris-dendreon-provengebristol/8/23/2011/id/36512#ixzz2Jg5uaa2F

<sup>3-</sup>Source: American Cancer Society: Workplace Solutions. www.acsworkplacesolutions.com/ceocancerimpact.asp (accessed February 2013).

<sup>4-</sup>Source: Shankaran V, Jolly S, Blough D, Ramsey SD. Risk Factors for Financial Hardship in Patients Receiving Adjuvant Chemotherapy for Colon Cancer: A Population-Based Exploratory Analysis. Journal of Clinical Oncology 30:1608-1614.



### Why is change necessary?



\*\* Source: Ramsey S, Blough D, Kirchhoff A. Cancer Patients Found to be at Greater Risk for Bankruptcy than People Without a Cancer Diagnosis. Health Affairs, 32: 1143-1152. 2013.

### Incidence of bankruptcy one year after a cancer diagnosis

Cancer Type**	Incidence*
Thyroid	9.3
Lung	9.1
Uterine	6.8
Leukemia/lymphoma	6.2
Colorectal	5.9
Melanoma	5.7
Breast	5.7
Prostate	3.7

\*Per 1000 Person-Years

### Quality of cancer care is inconsistent

- Up to 1 in 3 people treated with chemotherapy do not receive a treatment regimen that is consistent with current medical evidence and best practices<sup>1</sup>
- People are often hospitalized during treatment because of side-effects which could be avoided by using less toxic treatment regimens and appropriate supportive care<sup>2</sup>
- People frequently receive tests and treatment that they do not need, putting them at risk of side-effects, as well as imposing an additional care burden and cost<sup>2</sup>

1 J Clin Oncol 2011, 30:142-50; J Clin Oncol 2012, 30:3800-09; J Clin Oncol 2006, 24:626-34; Oncologist 2011;16:378-87; 2 J Clin Oncol 2002 20:4636-42. JACR 2012, 9:33-41; JAMA 2013, 309:2587-95; J Clin Oncol 2013; 31:epub. 3 Barr et al. J Oncol Pract. 2011;7: 2s-15s.



## Widespread Variation in use of CSF with Cancer Rx

#### with CSF Prescribed by Practice 2009-2011 250 Number of Practices 200 150 100 50 0 <5% 6-14% 15-24% 25-34% 55-64% >64% 35-44% 45-54%

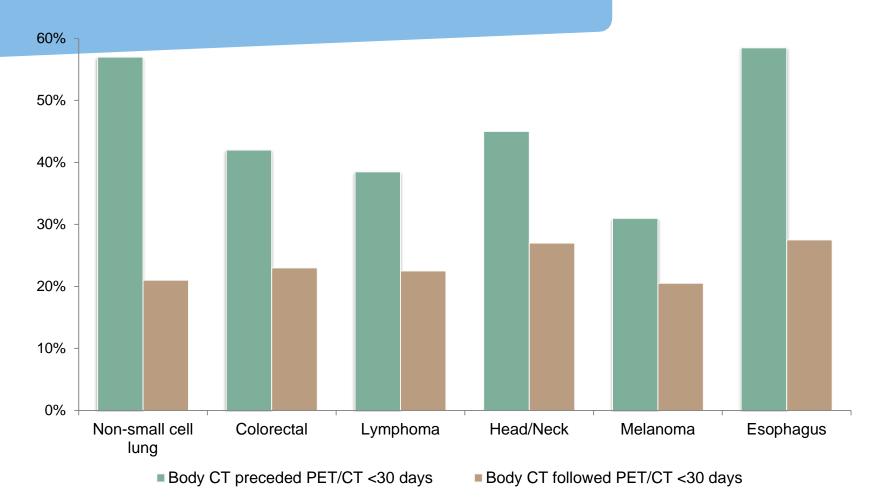
Percent of Chemotherapy Episodes with CSF Prescribed

Variation in Percent of Chemotherapy Episodes

Source: WellPoint affiliated health plans internal data 2012



### Imaging often repeated





## ASCO Choosing Wisely highlights unnecessary care

Choosing Wisely®

An initiative of the ABIM Foundation

Don't give **anti-nausea drugs** (anti-emetics) to patients starting on chemotherapy regimens that have low or moderate risk of causing nausea and vomiting.

Don't use **combination chemotherapy** (multiple drugs) instead of single-drug chemotherapy when treating an individual for metastatic breast cancer unless the patient needs urgent symptom relief.

Avoid using **advanced imaging technologies** — positron emission tomography (PET), CT and radionuclide bone scans to monitor for a cancer recurrence in patients who have finished initial treatment and have no signs or symptoms of cancer.

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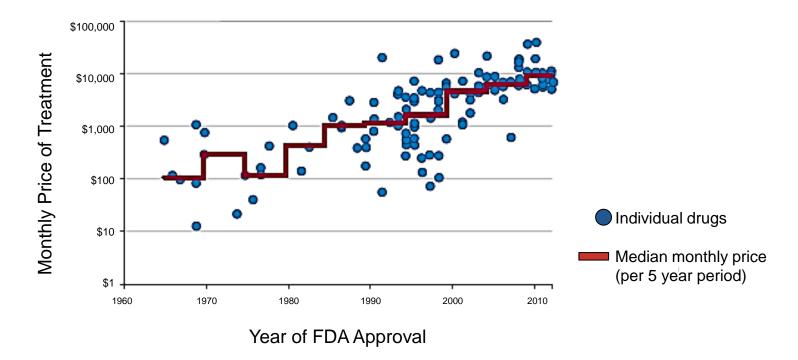
ASCOS<sup>®</sup> American Society of Clinical Oncology Don't perform **PSA testing** for prostate cancer screening in men with no symptoms of the disease when they are expected to live less than 10 years.

Don't use a **targeted therapy** intended for use against a specific genetic abnormality unless a patient's tumor cells have a specific biomarker that predicts a favorable response to the targeted therapy.



## New cancer drugs are becoming more expensive . . .

Monthly and median cost of cancer drugs at the time of FDA approval (1965 – 2013)





### and often not producing value

#### 13 new cancer treatments approved by FDA in 2012



Source: Emanuel et al, New York Times, A Plan To Fix Cancer Care, March 23, 2013, retrieved from http://opinionator.blogs.nytimes.com/2013/03/23/a-plan-to-fix-cancer-care/?\_r=0



## Charting a New Course for Cancer Care

# 2013

Institute of Medicine report recommends measures to improve quality and affordability of cancer care

#### DELIVERING HIGH-QUALITY CANCER CARE

Charting a New Course for a System in Crisis



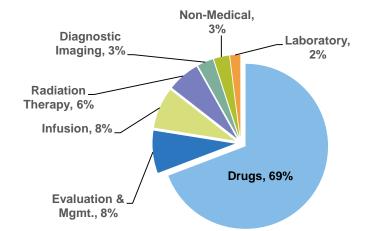
INSTITUTE OF MEDICINE

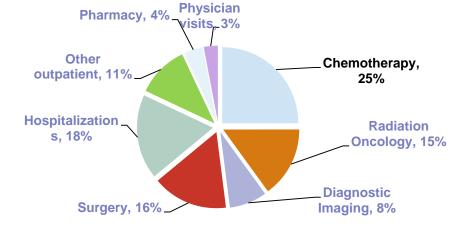


### Together, we can transform cancer care

#### Oncology Practice Revenue Sources Barr et al. J Oncol Pract. 2011;7: 2s-15s.

#### Chemotherapy Accounts for 25% of Health Plan Cost of All Cancer Care





WellPoint affiliated health plans internal data 2012

Reimbursement model must change so that focus shifts to providing cancer care that is value-based and patient-centered.



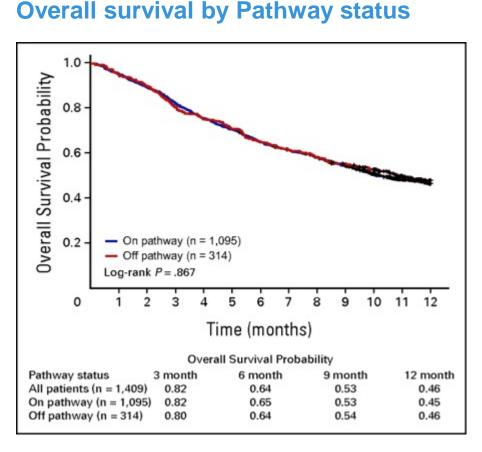


### **Our Model: a Quality Initiative**

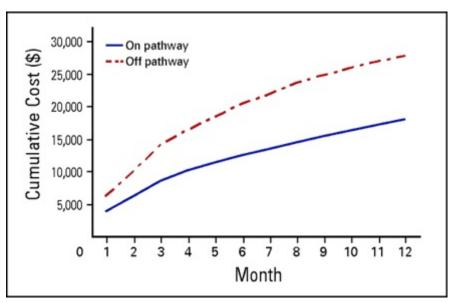
- BCBS Cancer Care Quality Program provides a framework for rewarding high quality cancer care
- Oncologists participating in the Cancer Care Quality Program will receive additional payment for treatment planning and care coordination when they select a treatment regimen that is on Pathway
- Practices participating in the Program can gain efficiency through synchronization with Health Plan Medical Policy and Clinical Guidelines



## US Oncology found pathways associated with same overall survival and 30% lower cost



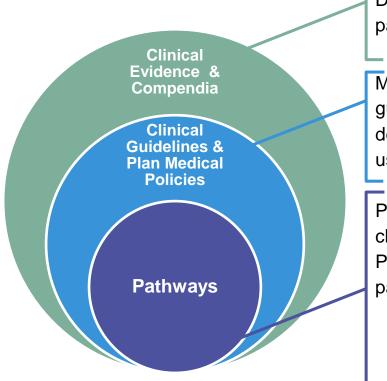
### 12-month cumulative cost by Pathway status



Neubauer M A et al. JOP 2010;6:12-18



### WellPoint Approach to Pathway Development



WellPoint's external advisors include ~10 oncologists from geographically diverse academic and community oncology practices who have specific interest in quality of care; 4 are affiliated with NCIdesignated cancer centers, 6 with Blue Centers of Distinction, and 6 have served on national committees for organizations such as NQF, ASCO, and IOM to improve the quality of cancer care. Data from trials, publications, and compendia for many different patient populations are extracted, reviewed, and analyzed.

Medical evidence is synthesized by national experts into clinical guidelines. Evidence is also used by health plan committees to develop medical policies and utilization management guidelines used in making benefit coverage determinations.

Pathways are a subset of regimens supported by evidence and clinical guidelines and aligned with health plan medical policies. Pathways are intended to be applicable for 80%-90% of patients and are selected based on:

- 1. Clinical benefit (efficacy)
- 2. Side effects/toxicities (especially those leading to hospitalizations & impact quality of life)
- 3. Strength of national guideline recommendations
- 4. Cost of regimens

WellPoint Pathways are developed through a rigorous evidence based medicine process and reviewed by external advisors.



## Variation in outcomes for 1<sup>st</sup> line platinum regimens for lung cancer\*

	Estimated Survival (months)	Grade 3-4 Adverse Events	Any serious AE (Hospitalization)	Deaths on Rx (Deaths due to Rx)
Rx A	13.0 (NR) mos.	N/V risk: Moderate* FN + infection:1% Neuropathy: 11% Debilitating fatigue: 6% <sup>,</sup>	53% (**)	<1% (<1%)
Rx B	10.4 (9.6-11.2) mos.	N/V risk: High FN + infection:4% Neuropathy: ND Debilitating fatigue: 5%	35% (**)	7% (1%)
Rx C	11.8 (10.4-13.2) mos.	N/V risk: High FN + infection:1% Neuropathy: ND Debilitating fatigue: 7%	37% (**)	7% (1%)
Rx D	13.1 (NR) mos.	N/V risk: Moderate FN + infection:1% Neuropathy: 3% Debilitating fatigue: 4%	** (**)	<1% (<1%)
Rx E	13.4 (11.9-14.9) mos.	N/V risk: Moderate FN + infection:4% Neuropathy: 4% Debilitating fatigue: 5% Bleeding 4%	75% (19%)	5% (4%)
Rx F	12.6 (11.3- 14.0) mos.	N/V risk: Moderate FN + infection:2% Neuropathy:0% Debilitating fatigue:11%	** (20%)	** (2%)

\* Non-squamous histology; first line platinum based chemotherapy indicated when no EGFR or ALK mutation present \*\* Not reported Socinski JCO 2012; Sandler NEJM 2006:355; Scagliotti JCO 2008:26; Reck Annals of Oncology 2010; Patel 2012



## Variation in outcomes across 1<sup>st</sup> line platinum regimens for lung cancer\*

		Estimated	Grade 3-4 Adverse	Anv serious AE	Deaths on Rx
	•	What is more in	nportant to t	he persor	n with
Rx		cancer – overal	l survival, su	urviving b	eyond
Rx		initial few mont when not receiv			
Rx		hospital?			
Rx		Do people have treatment plan i			n their
Rx		-			

Which regimen would you choose?

Rx F 12.6 (11.3- 14.0) mos.

FN + infection:2% Neuropathy:0% Debilitating fatigue:11%

\*\* (20%)



\* Non-squamous histology; first line platinum based chemotherapy indicated when no EGFR or ALK mutation present \*\* Not reported Socinski JCO 2012; Sandler NEJM 2006:355; Scagliotti JCO 2008:26; Reck Annals of Oncology 2010; Patel 2012



## Little variation in patient outcomes marked variation in treatment cost

	Estimated Survival (months)	Deaths on Rx (Deaths due to Rx)	Cost (4 cycles)
Carbo/Paclitaxel	13.0 (NR) mos.	<1% (<1%)	\$452
Gem/Cis	10.4 (9.6-11.2) mos.	7% (1%)	\$886
Cis/Pemetrexed	11.8 (10.4-13.2) mos.	7% (1%)	\$25,619
Carbo/nab-Paclitaxel	13.1 (NR) mos.	<1% (<1%)	\$24,740
Carbo/Paclitaxel/Bev	13.4 (11.9-14.9) mos.	5% (4%)	\$39,770
Carbo/Pemetrexed/Bev	12.6 (11.3- 14.0) mos.	** (2%)	\$64,988



### Four regimens included in Pathway

		Estimated Survival (months)	Deaths on Rx (Deaths due to Rx)	Cost (4 cycles)
$\checkmark$	Carbo/Paclitaxel	13.0 (NR) mos.	<1% (<1%)	\$452
$\checkmark$	Gem/Cis	10.4 (9.6-11.2) mos.	7% (1%)	\$886
$\checkmark$	Cis/Pemetrexed	11.8 (10.4-13.2) mos.	7% (1%)	\$25,619
	Carbo/nab-Paclitaxel	13.1 (NR) mos.	<1% (<1%)	\$24,740
$\checkmark$	Carbo/Paclitaxel/Bev	13.4 (11.9-14.9) mos.	5% (4%)	\$39,770
	Carbo/Pemetrexed/Bev	12.6 (11.3- 14.0) mos.	** (2%)	\$64,988

- Pathway is specific for patients who do not have mutations such as EGFR, ALK Pathways are personalized to tumor biology and genomics
- If the oncologist or patient determines that a different regimen is better for their unique circumstances, they are still treated according to their preference – Pathway adherence does not impact coverage determination

Socinski JCO 2012; Sandler NEJM 2006:355; Scagliotti JCO 2008:26; Reck Annals of Oncology 2010; Patel 2012



# Pathways include breast, lung and colorectal, ovarian, pancreas cancer, NHL and myeloma

		Cancer Treatment Pathways Worksheet
	Cancer Treatment Pathways Worksheet	Breast Cancer
	Lung Cancer	Patient name: Date of birth:
Cancer Treatment Pathways Work		Member number:
Colorectal Cai	Patient name: Date of bir	Breast Cancer, adjuvant, HER 2 negative AC weekly P: Doxorubicin and Cyclophosphamide (Every 3 Weeks) followed by Weekly Pacilitaxel
	Member number:	TC: Docetaxel (Taxotere) and Cyclophosphamide (Cyctoxan)
atient name:	1st Line Therapy for Metastatic Non-Small Cell Lung Car	ddAC weekly T: Dose Dense Doxorubicin and Cyclophosphamide Followed by Weekly Paclitaxel
ember number:	Crizotinib	AC: Doxorubicin and Cyclophosphamide
olorectal Cancer, Adjuvant	1st Line Therapy for Metastatic Non-Small Cell Lung Car	Breast Cancer, adjuvant, HER 2 positive
FULV: Fluorouracil (5FU) (Bolus) and Leucovor	Erlotinib	AC TH : Doxorubicin and Cyclophosphamide Followed by Paclitaxel and Trastuzumab (Herceptin)
FOLFOX-6 - Fluorouracii (5-FU), Leucovorin ar	Afatinib	TCH: Docetaxel (Taxotere), Carboplatin and Trastuzumab (Herceptin)
FLOX: Fluorouracil (5-FU), Leucovorin and Oxa Capecitabine	1st Line Therapy for Metastatic Non-Small Cell Lung Car with ECOG performance status = 0, 1, 2	Breast Cancer, HER 2 negative, metastatic disease, first and subsequent lines of therapy (1st line+)
t Line or 2nd Line Therapy for Metastatic following regimens are options for patients regardle	Carboplatin+Paclitaxel (Allow substitution of Cisplatin for Carb Cisplatin+Pemetrexed (Allow substitution of Carboplatin for Cit	Adriamycin (Doxorubicin)
FOLFOX (Fluorouracil, Leucovorin and Oxalipla	Cisplatin+Gemcitibine (Allow substitution of Carboplatin for Cis	Epirubicin
FOLFOX (Fluorouracil, Leucovorin and Oxalipla	Paclitaxel + Carboplatin+	Gemzar (Gemcitabine)
FOLFIRI (Fluorouracil, Leucovorin and Irinoteca	Bevacizumab	Navelbine (Vinorelbine)
FOLFIRI (Fluorouracil, Leucovorin and Irinoteca	1st Line Therapy for Metastatic Non-Small Cell Lung Car	Taxol (Paclitaxel)
FULV (Fluorouracil and Leucovorin)	ECOG performance status = 0, 1, 2	Xeloda (Capecitabine)
FULV (Fluorouracil and Leucovorin) with Bevac	Carboplatin+Paclitaxel (Allow substitution of Cisplatin for Cart	Supportive Care
Line or 2nd Line Therapy for Metastatic Idition to the above, the following regimens are also	Cisplatin+Gemcitibine (Allow substitution of Carboplatin for Cis	Breast Cancer, HER 2 positive, metastatic disease, First and subsequent lines of therapy (1st line+)
FOLFIRI (Fluorouracil, Leucovorin and Irinoteca	Maintenance Therapy in patients with Metastatic Non-Sn Non-squamous histology and ECOG performance status	Pertuzumab, Trastuzumab and Docetaxel
Irinotecan (Camptosar) and Vectibix (Panitumu	Continuation Bevacizumab	Pertuzumab, Trastuzumab and Paclitaxel
Line+ Therapy for Metastatic Colorects	Continuation	Capecitibine (Xeloda) and Trastuzumab
following regimens are options for patients regardle	Pemetrexed	Trastuzumab and Gemcitabine
Regorafenib (Stivarga)	Supportive Care	Vinorelbine and Trastuzumab
(KRAS mutant ONLY)	Switch	
Supportive care addition to the above, the following regi- t are KRAS-WT:		
Vectibix (Panitumumab) Monotherapy		
Irinotecan (Camptosar) and Vectibix (Panitumuma	d)	

#### Available at www.cancercarequalityprogram.com



### Treatment planning payments support cost-effective care



Enhanced reimbursement for treatment planning and care coordination will be provided when patient is registered with the Cancer Care Quality Program and treatment regimen in on pathway



S0353 reimbursed \$350 once at the onset of treatment S0354 reimbursed \$350 no more than monthly while managing care for an established patient\*

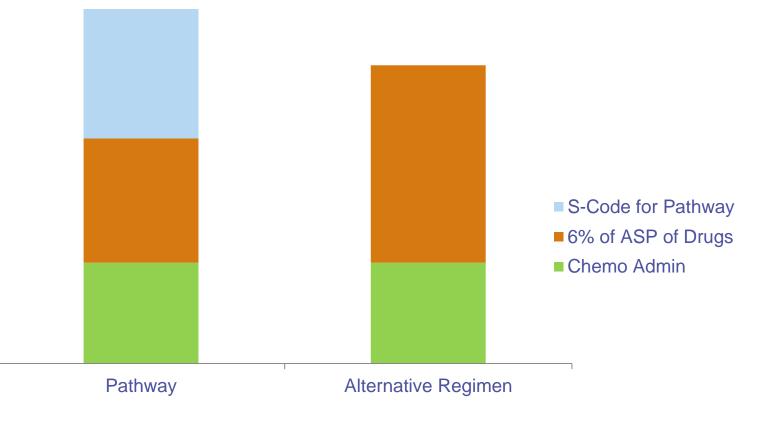


S-code billing authorization is triggered through AIM *ProviderPortal* when practice selects a regimen that aligns with WellPoint Cancer Treatment Pathways



### Impact of enhanced reimbursement and support for Pathways

Example of Pathway regimen where enhanced reimbursement (s-code) offsets higher revenue associated with more expensive regimen





### Cancer Care Quality Program administered by AIM Specialty



#### **CLINICAL REQUEST**

Request is made by a Provider via AIM's web self-service tools

#### TREATMENT REVIEW

Treatment request reviewed against an evidence-based regimen library for alignment with health plan medical policy for members in that health plan

Wellpoint's Pathways are based on efficacy - toxicity and cost are also highlighted

#### **DECISION RENDERED**

Immediate approval is granted if consistent with plan medical policy

Clinical experts available as necessary for peer-to-peer discussion

Notified if Pathway option available

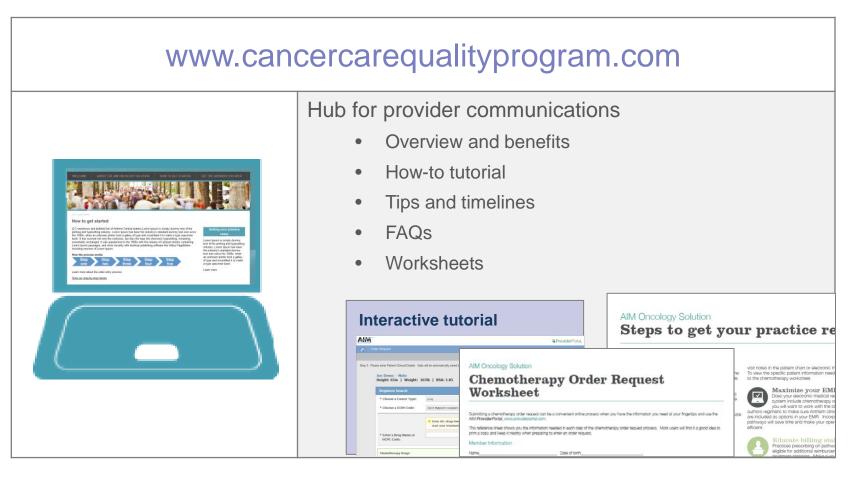
#### PATHWAY ADHERENCE

Practice authorized to bill S0353 and S0354 for *Treatment Planning and Care Coordination* when regimen is on pathway

Quarterly Analytics and Reporting are available



#### **Provider Website**



### Discussion



