

# Nursing Perspective on irAEs: Patient Education, Monitoring and Management

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## Disclosures

- No relevant financial relationships to disclose
- I will be discussing non-FDA approved indications during my presentation.









## Objectives

- Improve the early recognition, education and management of immune-related side effects in cancer immunotherapy patients
- Identify strategies for the management of toxicities
- Determine key points for patient education on the management of side effects

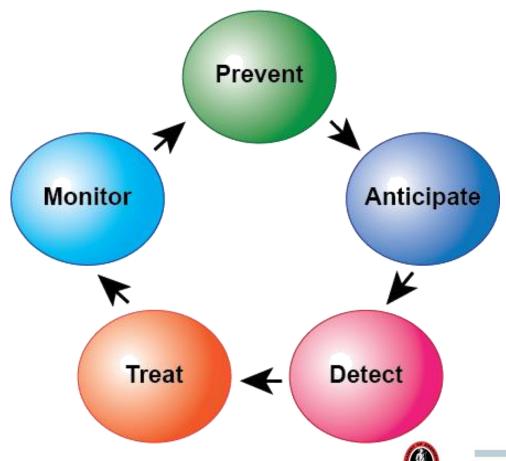








## The Five Pillars of Toxicity Management

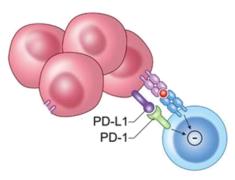








## Case Study



T cell

- Mr. M.C. is a 65-year-old male with a recent diagnosis of stage IV melanoma to the lungs. Patient has consented to start pembrolizumab (checkpoint inhibitor) at 2mg/kg every 3 wks.
- Mr. M.C and family would like to know what are the most common adverse events with this immunotherapy?





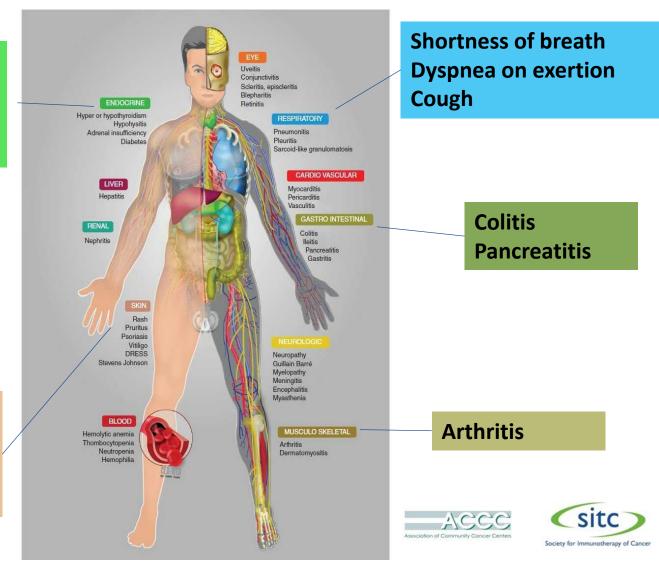




### Toxicity Spectrum: Immune Related **Adverse Events**

**Hypothyroid Hypophysitis Adrenal insufficiency Diabetes** 

Maculopapular rash Vitiligo (positive factor)



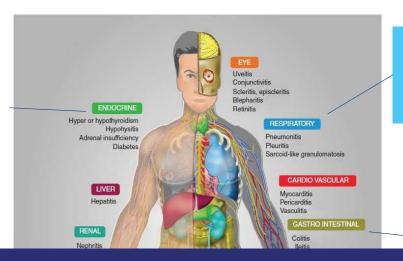
**Pruritus** 

**DRESS** 



## Toxicity Spectrum: Immune Related Adverse Events

Hypothyroid
Hypophysitis
Adrenal insufficiency
Diabetes

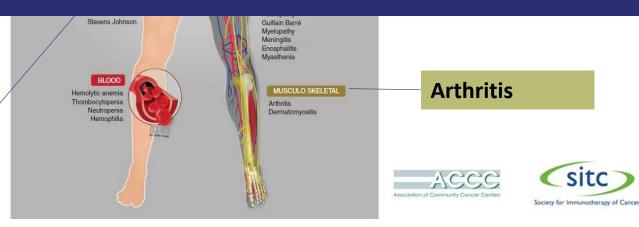


Shortness of breath Dyspnea on exertion Cough

**Colitis** 

These are some of the most common; HOWEVER; immune-related side effects do not discriminate.

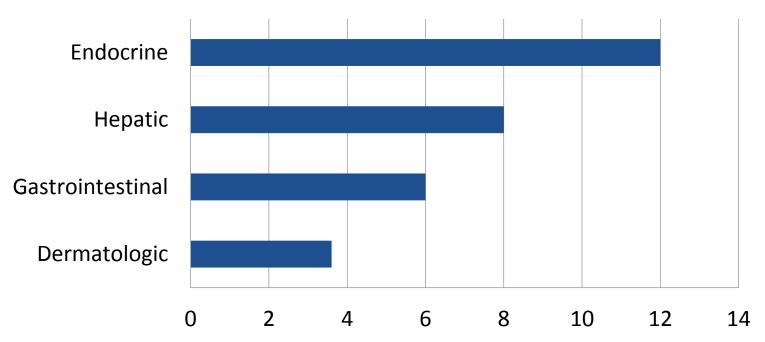
Maculopapular rash
Pruritus
DRESS
Vitiligo (positive factor)





### Immune checkpoint inhibitors-irAEs

### Median time to development (weeks)











## Symptoms to look for with immune check-point inhibitors

### Ipilumamab:

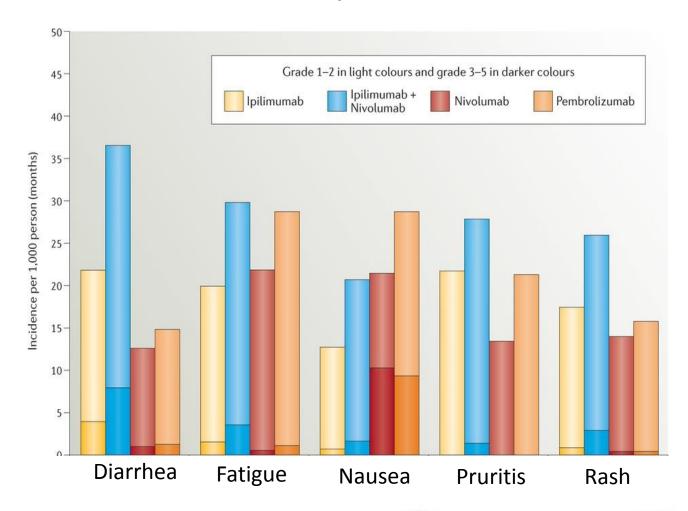


#### Nivolumab:



### Pembrolizumab:





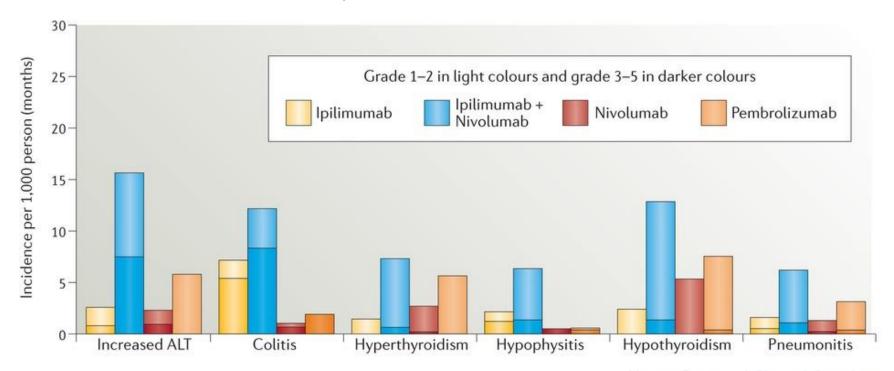








# Clinical features for adverse events with immune check-point inhibitors



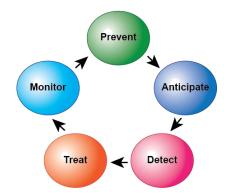
Nature Reviews | Clinical Oncology











### Nurse's Role: Prior to Immunotherapy

- Review & assess
  - Co-morbidities (dermatologic, endocrinopathies, gastrointestinal)
  - Medications
- Patient & family education
  - Most common side effects, including variability in the timing of onset
  - Importance of early & ongoing communication regarding side effects
  - Appropriate skin care during immunotherapy treatment, initiate now





## Case Study #2 – Dermatologic irAEs

 69yo M with Hodgkin Lymphoma, s/p 5 cycles of Pembrolizumab.
 Presents to clinic with grade 2 rash to BUE.











## Managing irAEs

Table 4. Ty	pical management of i	rAEs		
Severity— CTCAE grade	Ambulatory versus inpatient care	Corticosteroids	Other immunosuppressive drugs	Immunotherapy
1	Ambulatory	Not recommended	Not recommended	Continue
2	Ambulatory	Topical steroids or Systemic steroids oral 0.5–1 mg/kg/day	Not recommended	Suspend temporarily <sup>a</sup>
3	Hospitalization	Systemic steroids Oral or i.v. 1–2 mg/kg/day for 3 days then reduce to 1 mg/kg/day	To be considered for patients with unresolved symptoms after 3–5 days of steroid course Organ Specialist referral advised	Suspend and discuss resumption based on risk/benefit ratio with patient
4	Hospitalization consider intensive care unit	Systemic steroids i.v. methylprednisolone 1–2 mg/kg/day for 3 days then reduce to 1 mg/kg/day	To be considered for patients with unresolved symptoms after 3–5 days of steroid course Organ specialist referral advised	Discontinue permanently

CTCAE = Common Terminology Criteria for Adverse Events

Champiat S, et al, Ann Oncol, 2016









## Managing irAEs

Table	<b>4.</b> Typical management of	irAEs			
Severity-	•	Corticosteroids	Other immunosuppressive dr	rugs Immunotherapy	
CTCAE	grade inpatient care				
1	Dringinle	o of Monoo	ing in A Co.		
2	_	s of Manag			
	<ul> <li>Hold im</li> </ul>	munothera	py for grade > 2	2	
	• Initiate	corticoster	oids (e.g.,1–2 m	a/ka of	
3			Jiao (Jigi, i – iii		ion based
	predniso	•			atient
	<ul> <li>Conside</li> </ul>	er other the	rapies (example	e: infliximab if	
4	gastroint	estinal toxi	city or mycoph	enolate if	
			improvement w		
	<del>-</del>		improvement w	VILII	
	corticost	eroids)			
(	CTCAE = Commo	n Terminólogy Ci	iteria for Champi	at S, et al, Ann Oncol,	<b>20</b> 16

Society for Immunotherapy of Cancer

**Adverse Events** 



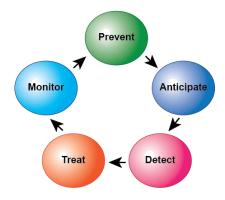
### Nurse's role: rash

### **Anticipate/Prevent**

- Skin toxicities can be seen in up to 58% of cases
- Autoimmune conditions can worsen
- Occupational/recreational activities (exposure to outdoors/high temps can worsen skin AEs)
- Possibility of developing hypopigmentation (vitiligo correlated to positive outcome)

#### **Monitor**

- New onset of rash
- New lesions
- Itching
- Sunburn
- Photosensitivity



### Manage

- Educate patient about potential side effects
- Grade 1: topical OTC hydrocortisone / oral diphenhydramine
- Grade 1/2: triamcinolone or clobetasol cream, diphenhydramine or hydroxyzine (if and when)
- Grade 2: hold treatment, oral corticosteroids
- Grade 3/4: discontinue agent









### Nurse's role: rash

### **Anticipate/Prevent**

- Skin toxicities can be seen in up to 58% of cases
- Autoimmune conditions can worsen
- Occupational/recreational

#### **Monitor**

- New onset of rash
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### Manage

Treat

Prevent

**Anticipate** 

Detect

 Educate patient about potential side effects

Monitor

 Grade 1: topical OTC hydrocortisone / oral diphenhydramine

Crada 1/2, riamainalana

## **MOST IMPORTANT:**

## CONTACT HEALTH CARE PROVIDERS IMMEDIATELY!! COME IN NOW!!!

outcome)

Grade 3/4: discontinue agent









## Case Study #3 - Colitis

- 26 yo Female with Hodgkin Lymphoma. h/o salvage treatment with combo therapy brentuximab vedotin and nivolumab; ASCT and is now s/p 2 cycles pembrolizumab consolidation treatment.
- Presents to triage c/o grade 3 diarrhea x3 days; abdominal cramping, nausea, fatigue. Pt reports no relief with at home loperamide and other supportive measures.

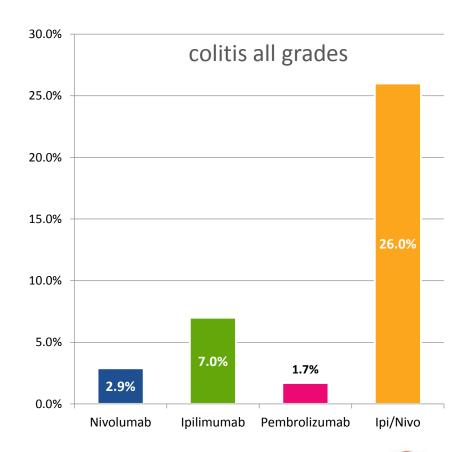








### Immune-Mediated Colitis











# Educate patients: constant communication of symptoms is essential sooner rather than later

 Updated safety information with 9 additional months of follow-up were consistent with the initial report

	NIVO+IPI (N=313)		NIVO (N=313)		IPI (N=311)	
Patients reporting event, %	Any Grade	Grade 3-4	Any Grade	Grade 3-4	Any Grade	Grade 3-4
Treatment-related adverse event (AE)	95.8	56.5	84.0	19.8	85.9	27.0
Treatment-related AE leading to discontinuation	38.7	30.7	10.5	7.3	15.4	13.5
Treatment-related death*	d death* 0		0.3		0.3	

 68.8% of patients who discontinued NIVO+IPI due to treatment-related AEs achieved a response

\*One reported in the NIVO group (neutropenia) and one in the IPI group (colon perforation)

Database lock Nov 2015

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Treatment-related adverse	05.8	56.5	840	10.8	85 Q	27.0

## Grade 3/4 is life-threatening

Treatment-related death* 0	0.3	0.3
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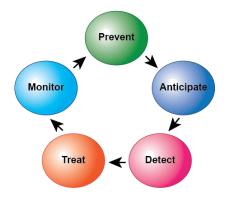












### Nurse's role: GI toxicities

### **Anticipate/Prevent**

- Diarrhea can be seen in up to 48% of cases
- Autoimmune conditions can worsen
- Avoid foods that cause loose stools
- Rule out infections (c-diff)
- Remain well-hydrated

#### **Monitor**

- Worsening loose stools
- Dehydration
- Abdominal pain/cramping
- Bloody stools

#### Manage

- Educate patient about potential side effects
- Grade 1: hydration, loperamide, bland diet
- diphenoxylate/atropine
  QID, budesonide, stool
  studies, possible
  sigmoidoscopy/colonoscop
  y & steroid taper
- Grade 3/4: discontinue agent, IV steroids and fluids (if not effective, infliximab)



## Case Study #1 - Pneumonitis

- 66 yo Female with Hodgkin Lymphoma, s/p 2 cycles of nivolumab treatment. Routine CT to assess response to therapy shows improvement in disease sites, but new bilateral pulmonary infiltrates.
- Patient c/o intermittent grade 1 cough and some mild dyspnea. Per patient report, h/o seasonal asthma with similar symptoms.



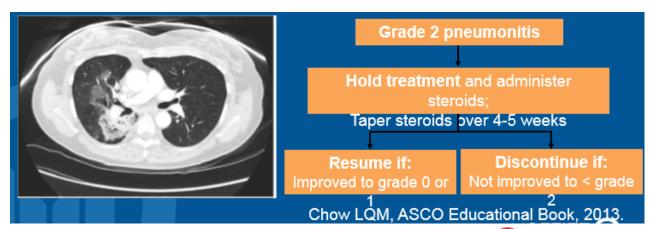






# Pneumonitis is more common with anti-PD1/CTLA-4 combination therapy

- Important to address respiratory symptoms and check oxygen saturations at each visit
- On any patients where pneumonitis is suspected based on H&P or clinical exam, provider will hold treatment and order a CT scan of the chest.
- Specific management is necessary for grade 2 or greater pneumonitis.



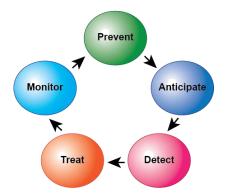








## Nurse's role: pneumonitis



### **Anticipate/Prevent**

- Pneumonitis on single vs combination immunotherapy
- Exposure to heavy smoke areas / smoking cessation
- Vaccinations (flu + pneumonia
- Pneumonia vs PE vs CHF

#### **Monitor**

- SOB, DOE, CP, persistent cough, fevers, worsening fatigue
- Pulse-ox at rest and with ambulation

### Manage

- Educate patient about potential side effects
- Grade 1: asymptomatic
- Grade 2: chest x-ray or CT, anticipate steroid taper
- Grade 3/4: discontinue agent, IV steroids and fluids (if not effective, infliximab), oxygen therapy









## Case study

- J.C. is a 75-year-old male with metastatic melanoma currently on nivolumab/ipilimumab combination therapy. He reports that for the past five days he has had:
  - Moderate headaches, severe fatigue, weakness and nausea.
  - Endocrine labs revealing low cortisol, low ACTH and low testosterone levels. Free T4 and TSH were normal.
- As the nurse you see the patient first in clinic and alert the doctor of his symptoms and current labs.



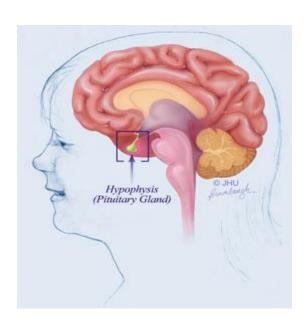


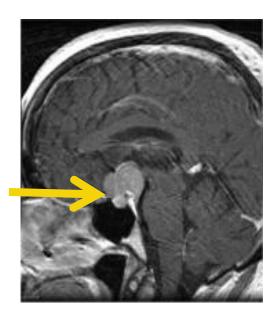




## Case study

Oncologist orders an MRI of the brain which shows inflammation of the pituitary gland













## Immune-mediated endocrinopathies

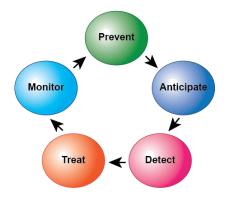
- More common with anti-PD-1 than anti CTLA-4
- Hypophysitis with nivo/ipi median time to onset was about 2.7 months. All grades 9%
- Hypothyroidism
- Hyperthyroidism
- Adrenal insufficiency
  - Rule out brain metastasis
  - Hold for symptoms and/or any Grade 3/4
  - Give steroids (IV followed by PO 1-2mg/kg) tapered over four weeks and replace appropriate hormones
    - Hormone replacement may be required for life in ~50% of patients











## Nurse's role: endocrinopathies

### **Anticipate/Prevent**

- Hypothyroidism
- Hyperthyroidism
- Hypophysitis
- Adrenal insufficiency
- Especially in combination ipi/nivo

#### **Monitor**

- Labs: Free T4, TSH, ACTH, cortisol and testosterone (in males)
- Worsening fatigue
- Constipation
- Headaches
- Dizzy episode(s)
- Muscle weakness

### Manage

 Hormonal replacement therapy or steroid taper accordingly









## Immune checkpoint inhibitors irAEs

- Rare toxicities
  - Type I and II diabetes mellitus
  - Pancreatitis-usually asymptomatic amylase/lipase elevations (hold for grade 3/4)
  - Myositis
  - Renal toxicity (acute interstitial nephritis)
  - Autoimmune myocarditis
  - Bullous pemphigoid









### Immune checkpoint inhibitors irAEs

## Rare toxicities Bullous pemphigoid



- Myasthenia-like syndrome-motor paralysis, intravenous immune globulins
- Optic neuritis-photophobia, pain, blurred vision, may correlate with colitis
- Sarcoidosis-lymphadenopathy, increased angiotensin converting-enzyme level, biopsy is granulomata, PET positive
- Hematologic
- Cardiotoxicities: Myocarditis









### Immune-mediated toxicities

- General principles of toxicity management
  - Reversible toxicities when recognized quickly and treated appropriately
  - Treatment may include dose delay, omission, or discontinuation, corticosteroids, tumor necrosis alfa (TNF- $\alpha$ ) antagonists, and mycophenolate mofetil
  - Corticosteroids may require a long tapering duration to prevent recurrence of symptoms
    - Rechallenge with checkpoint inhibitor may only be done, if clinically appropriate, once a patient is receiving 10 mg of oral prednisone or equivalent or less.
    - Prolonged use of steroids predisposes patients to systemic infection so prophylaxis may be indicated.

Villadolid J and Amin A. Transl Lung Cancer Res 2015; 4 (5): 560-575









### Conclusions

- Nurses have an ESSENTIAL role in monitoring and managing patients undergoing treatment with immunotherapy.
- Potential irAEs grade 2 and above require frequent visits, drug hold/discontinuation and corticosteroids.
- Combination anti-PD-1/CTLA-4 immunotherapy significantly increases the grade 3-4 AE rate.
- Close monitoring for irAEs is mandatory for prevention of serious adverse events, decreased ER visits and improve patient outcomes.
- As immunotherapies indications broaden, our understanding of toxicity identification and management is essential to make the risk-benefit ratio favorable.





