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Current Guidelines in Cardio-Oncology
 2016 ESC Position Paper on cancer treatments and cardiovascular toxicity developed under the auspices of the ESC Committee of Practice Guidelines
 The Task Force for cancer treatments and cardiovascular toxicity of the European Society of Cardiology and the European Association of Cardiovascular Imaging
 Expert Consensus for Multimodality Imaging Evaluation of Adult Patients during and after Cancer Therapy: A Joint Statement by the American Society of Echocardiography and the European Association of Cardiovascular Imaging

What is the optimal strategy?

ACC/AHA Heart Failure guidelines (2013)

Stage A: Recommendations (Class I, LOE C)

Conditions that may lead to or contribute to HF, such as ... cardiotoxic agents, should be controlled or avoided.

- “...it may be reasonable to evaluate those who are receiving (or who have received) cardiotoxic chemotherapy agents for LV dysfunction.”
- “The use of advanced echocardiographic techniques or biomarkers to identify increased HF risk in those receiving may be useful ...”

Yancy, et al. Circulation 2013

ESMO guidelines (2012)

- Patients receiving anthracyclines and/or trastuzumab in the adjuvant setting should perform serial monitoring of cardiac function at baseline, 3, 6, and 9 months during treatment, and then at 12 and 18 months after the initiation of treatment.
- Monitoring should be repeated during or following treatment as clinically indicated.

Curigliano, et al. Ann Oncol 2012

Cohort	QTc Stratification by Group (msec)			
	< 450	≥ 450 to 499	≥ 500	
Overall				
Baseline	n=71	73% (n=52)	23% (n=16)	4% (n=3) p=0.75
On-Treatment	n=49	67% (n=33)	27% (n=13)	6% (n=3)
Pre- and On-Treatment Group				
Baseline	n=35	80% (n=28)	20% (n=7)	0% (n=0) p=0.34
On-Treatment	n=35	71% (n=25)	20% (n=7)	9% (n=3)
On-Treatment Group Only				
First ECG	n=30	80% (n=24)	20% (n=6)	0% (n=0) p=0.09
Longest QTc	n=30	56% (n=17)	33% (n=10)	10% (n=3)

WHO/ISH risk chart

WHO/ISH risk chart			Total
<10%	10%–20%	>20%	

UKPDS engine

WHO/ISH risk chart				
<10%		10%–20%	>20%	Total
Count	1,221	39	14	1,274
Expected count	1,128.1	96.3	49.6	1,274.0
10%–20%				
Count	565	74	38	677
Expected count	599.5	51.2	26.4	677.0
>20%				
Count	124	50	32	206
Expected count	182.4	15.6	8.0	206.0
Total				
Count	1,910	163	84	2,157
Expected count	1,910.0	163.0	84.0	2,157.0
Value	SE	Approx T ^b	Approx sig	
Measure of agreement	0.144	0.014	11.236	0.000

Primary Cardioprotection

Assessment & Optimization of risk +/- extent of CV Disease

- Age
- Co-morbid conditions (CAD, HTN, HLP)
- Modifiable lifestyle risk factors (deconditioning, obesity)

Primordial Prevention

Treat all high cardiac risk patients

Advantages

Minimize screening

Disadvantages

Treatment despite:

-Normal function

-Low (~5%) overall incidence of overt HF

Primary Prevention

Screen for subclinical dysfunction (Imaging, blood biomarkers)

Advantages

Minimize unnecessary treatment

Disadvantages

Intensive / frequent screening

Optimal screening method uncertain

Potential delayed diagnosis / treatment

Advantages

Early intervention → Limit development of cardiotoxicity

Disadvantages

Duration of treatment uncertain

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