Testicular Cancer

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Clinical Risk Factors

Medical Conditions

Cryptorchidism
Spermatic/Testicular dysgenesis
HIV (seminoma)

Family History

Genetic Risk Factors

Isochromosome 12p

Klinefelters

Presentation

Primary Testicular Mass (90-95%)
Primary Mediastinal Mass

Work Up

Ultrasound

Orchiectomy

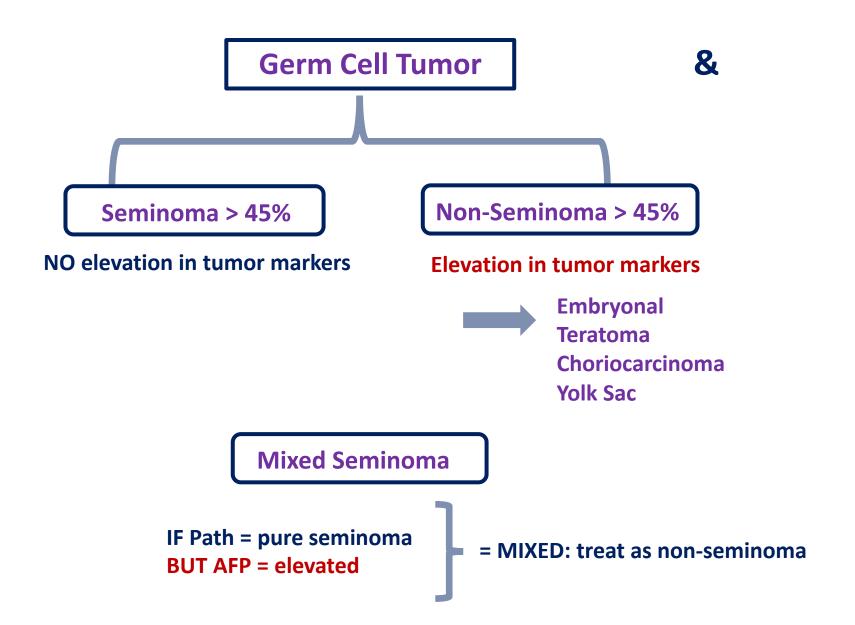
Tumor Markers

LDH

HCG

AFP

Testicular Cancer



Non-Germ Cell Tumor

SEX CORD STROMAL TUMORS:
Leydig Cell
Sertoli Cell
Granulosa Cell

Testicular Cancer: Staging

Seminoma

Non-Seminoma or Mixed

NO elevation AFP

MAYBE elevation AFP

Tumor Markers (half lives)

AFP (7-10D)

HCG (2-3D)

LDH (2-3D)

check markers pre and post-op

1 month post-op markers used for staging

STAGING

T Stage

T1 = intra-testicular

T2 = tunica vaginalis, epididymis, rete testes, LVI

T3 = spermatic cord

T4 = scrotum

Serum Marker Staging

S0 = normal tumor markers

S1 = AFP < 1K HCG < 5K, LDH < 1.5 ULN

S2 = AFP 1-10K, HCG 5-50K, LDH 1.5-10x ULN

S3 = AFP > 10K, HCG > 50K, LDH > 10x ULN

Stage

Stage IS = S1-3, any T, N0, M0 (tx as Stage II-III)

Stage I = NO, SO

Stage 2 = N+, S0-1

Stage 3 = S2-3

* No stage 4

Testicular Cancer: Risk Stratification

NON-SEMINOMA
MIXED SEMINOMA

YES elevation in tumor markers YES Poor Risk Group

SEMINOMA

NO elevation in tumor markers NO Poor Risk Group

GOOD RISK

Can have pulmonary or mediastinal mets

INTERMEDIATE RISK

Non-pulmonary visceral mets

GOOD RISK

S1 level tumor markers

AFP < 1K

HCG <5K

LDH <1.5x ULN

Stage IS (S1-S3, TxN0M0)

INTERMEDIATE RISK

S2 level tumor markers

AFP 1-10K

HCG 5-50K

LDH 1.5-10x ULN

POOR RISK

S3 level tumor markers

AFP >10K

HCG >50K

LDH >10x ULN

Non-pulmonary visceral mets

Mediastinal mass

PROGNOSIS

Good Risk 5Y OS = 90%

Intermediate Risk 5Y OS = 80%

Poor Risk 5Y OS = 50%

Testicular Cancer: Treatment (Stage I-IIA)

IIA = N1 (LN < 2cm)

SEMINOMA

NO elevation in tumor markers NO Poor Risk Group

Orchiectomy +	
Stage I NO	Observation Consider RT OR Consider chemo
Stage IIA N+ IIA = N1 (LN < 2cm)	Adjuvant RT OR Adjuvant Chemo (Carbo x 1-2C)
NO RPLND for seminomas! no surgery for seminomas	

NON-SEMINOMA
MIXED SEMINOMA

YES elevation in tumor markers
YES Poor Risk Group

Orchiectomy +		
Stage I NO	Observation Consider RLPND OR Consider chemo (BEP x 1-2C)	
Stage IIA N+ IIA = N1 (LN < 2cm)	Adjuvant RPLND OR Adjuvant Chemo (BEP x1-2C)	
NO RT for non-seminomas!		

Testicular Cancer: Treatment (Stage IIB-IV)

IIB = N2+ (LN > 2cm) or S1+ any TNM

SEMINOMA

NO elevation in tumor markers
NO Poor Risk Group

Good Risk can include pulmonary mets or mediastinal mass	BEP x3 EP x4
Intermediate Risk non-pulmonary visceral mets	BEP x4 VIP x4

NON-SEMINOMA
MIXED SEMINOMA

YES elevation in tumor markers YES Poor Risk Group

Good Risk S1 (AFP <1K, HCG <5K, LDH <1.5 ULN), IS (S1-S3, TxN0M0)	BEP x3 EP x4
Intermediate Risk	BEP x4
Poor Risk	VIP x4
S2, S3, non-pulmonary visceral mets, or mediastinal mass	

Chemotherapy Regimens

BEP: (B) Bleomycin

(E) Etoposide

(P) Cisplatin

EP:

(E) Etoposide

(P) Cisplatin

VIP:

(V) Etoposide

(I) Ifosfamide + Mesna

(P) Cisplatin

^{*} If residual disease after chemo → resection

Testicular Cancer: Chemotherapy

Chemotherapy Regimens

BEP: (B) Bleomycin

(E) Etoposide

(P) Cisplatin

EP:

(E) Etoposide

(P) Cisplatin

VIP:

(V) Etoposide

(I) Ifosfamide + Mesna

(P) Cisplatin

TIP:

(T) Paclitaxel

(I) Ifosfamide + Mesna

(P) Cisplatin

* TIP used for RD or 2nd line therapy

Notable Side Effects

Bleomycin → Pneumonitis (PFTs prior to tx), Rash (flagellate erythema)

Cisplatin → Neuropathy, nephropathy, ototoxicity

Ifosfamide → Hemorrhagic cystitis, CNS toxicity (tx with methylene blue)

Paclitaxel → Neuropathy

Testicular Cancer Reference Handout

Testicular Cancer Dx Germ Cell Tumor Non-Germ Cell Tumor

Seminoma

Non-Seminoma/Mixed

NO elevation AFP

Maybe elevation AFP

IF Path = pure seminoma
BUT AFP = elevated = MIXED
treat as non-seminoma

Tumor Markers (half lives)

AFP (7-10D) HCG (2-3D) LDH (2-3D) **STAGING**

Stage IS = S1-3, any T, N0, M0 Stage I = N0, S0 Stage 2 = N+, S0-1 Stage 3 = S2-3

Testicular Cancer Risk

SEMINOMA

GOOD RISK

Can have pulmonary mets
Can have mediastinal mets

INTERMEDIATE RISK

Non-pulm visceral mets

Prognosis

Good Risk 5Y OS = 90%
Intermediate Risk 5Y OS = 80%
Poor Risk 5Y OS = 50%

NON-SEMINOMA

GOOD RISK

S1 level tumor markers

AFP <1K HCG <5K LDH <1.5x ULN Stage IS

INTERMEDIATE RISK

S2 level tumor markers

AFP 1-10K HCG 5-50K LDH 1.5-10x ULN

POOR RISK

S3 level tumor markers

AFP >10K HCG >50K LDH >10x ULN Non-pulmonary visceral mets

Mediastinal mass

Testicular Cancer Tx Stage I-IIA

SEMINOMA NO elevation in tumor markers NO Poor Risk Group

Orchiectomy +	
Stage I NO	Observation Consider RT OR Consider chemo
Stage II N+	Adjuvant RT OR Adjuvant Chemo (Carbo x 1-2C)

NO RPLND for seminomas!

NON-SEMINOMA YES elevation in tumor markers
MIXED YES Poor Risk Group

Orchiectomy +		
Stage I NO	Observation Consider RLPND OR Consider chemo (BEP x 1-2C)	
Stage II N+	Adjuvant RPLND OR Adjuvant Chemo (BEP x1-2C)	
NO RT for non-seminomas!		

Testicular Cancer Tx Stage IIB-IV

SEMINOMA

NO elevation in tumor markers NO Poor Risk Group

Good Risk	BEP x3 EP x4
Intermediate Risk	BEP x4 VIP x4

NON-SEMINOMA/MIXED SEMINOMA

YES elevation in tumor markers YES Poor Risk Group

Good Risk	BEP x3 EP x4
Intermediate Risk	BEP x4
Poor Risk	VIP x4

Chemo Regimens

BEP: (B) Bleomycin

(E) Etoposide

(P) Cisplatin

Chemo Side Effects

EP: (E) Etoposide

(P) Cisplatin

Bleomycin → Pneumonitis, Rash
Cisplatin → Neuronathy, nephror

Cisplatin → Neuropathy, nephropathy, ototoxicity **Ifosfamide** → Hemorrhagic cystitis, CNS toxicity

Paclitaxel → Neuropathy

VIP: (V) Etoposide

(I) Ifosfamide

(P) Cisplatin