SUPPLEMENTARY DATA PRISMA CHECKLIST

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Selection /		T		
b topic		Checklist item	Reported	Assessment
TITLE				
Title	1.	Identify the report as a systematic	No	Not applicable
	1.	review, meta-analysis, or both	110	This is an original
3		Teview, meta unarysis, or oom		research article, not a
ŀ				systematic review
_D ABSTRACT				systematic review
Structured	2		Yes	Not applicable
summary			103	This is not a
1 Summary				systematic review
S				abstract
M aintroductio	N			dobitaet
Rationale	3	Describe the rationale for the review in	Yes	Partialy – provides
C		the context of what is already known	103	backroung on
C		the context of what is already known		childhood renal
n				tumors and
e				SIOP/NWTSG
c -				protocols
K Objectives	4	Provide an explicit statement of	Yes	Yes – states aim to
	-	questions being addressed with reference	103	analyze neoadjuvant
ľ		to participants, interventions,		chemotherapy
S 4		comparisons, outcomes, and study		induced histological
τ		design (PICOS)		changes
METHODS		design (11005)		changes
Protocol and	5	Indicate if a review protocol exist, if and	Not	Not applicable – this
registration		where it can be accessed, and, if	mentioned	is not a systematic
registration		available, provide registration	mentioned	review
		information		TOVIOW
Eligibility	6	Specify study characteristics and report	Yes	Not applicable – this
criteria		characteristics used as criteria for		is a single center
		eligibility		study
Information	7	Describe all information sources and last	Not	Not applicable – this
sources	,	date of search	applicable	is not a systematic
		32 33 33 33		review
Search	8	Present full elecctronic search strategy	Not	Not applicable – this
		for at least one database	applicable	is not a systematic
		222 33 233 333 333 333		review
Study selection	9	State the process for selecting studies	Yes	Not applicable -
		state the process for selecting states		describes patient
	1		<u> </u>	accertoes patient

				selection criteria
				instead
Data collection	10	Describe methods of data extraction	Yes	Partially – describes
process		from reports and any processes for		data collection
		obtaining and confirming data from		methodology
		investigators		
Data items	11	List and define all variables for which	Yes	Yes – describes
		data were sought		histological
				evaluation criteria
				and staging
Risk of bias in	12	Describe methods used for assessing risk	Not	Not mentioned
individual		of bias of individual studies	mentioned	
studies				
Summary	13	State the prinicpal summary measures	Yes	Yes – describes
measures				categorization into
				risk groups and
				staging
Synthesis of	14	Describe the methods of handling data	Not	Not applicable –
results		and combining results of studies	applicable	single study
Risk of bias	15	Specify any assessment of risk of bias	Not	Not applicable –
across studies		that may affect the cumulative evidence	applicable	single study
Additional	16	Describe methods of additional analyses	Not	Not mentioned
analyses		if done	mentioned	
RESULTS				
Study selection	17	Give numbers of studies screened,	Not	Not applicable –
		assessed for eligibility, and included in	applicable	single study
		the review, with reasons for exclusions at		
		each stage		
Study	18	For each study, present characteristics	Yes	Yes – presents patient
characteristics		for which data were extracted		characteristics and
				outcomes
Risk of bias	19	Present data on risk of bias of each study	Not	Not mentioned
within studies			mentioned	
Results of	20	For all outcomes considered, present, for	Yes	Yes – presentrs result
individual		each study: simple summary data for		for both SIOP and
studies		each intervention group		NWTSG groups
Synthesis of	21	Present results of each meta-analysis	Yes	Partially-provides
results		done, including confidence intervals and		statistical analysis of
		measures of consistency		tumor size reduction
Risk of bias	22	Present results of any assessment of risk	Not	Not applicable –
across studies		of bias across studies	aapplicable	single study

Additional	23	Give results of additional analyses, if	Yes	Yes – provides
analysis		done		comparative analysis
				between protocols
DISCUSSION				
Summary of	24	Ssummarize the main findings including	Yes	Yes – summarizes
evidence		the strength of evidence for each main		key findings and
		outcome		comparative
				advantages
Limitation	25	Discuss limitations at study and outcome	Yes	Partially – discusses
		level, and at review level		limitations of each
				protocol
Conclusions	26	Provide a general interpretation of the	Yes	Yes – provides
		results in the context of other evidence		conclusion favoring
				SIOP protocol
FUNDING				•
Funding	27	Describe sources of funding for the	Not	Not mentioned
		systematic review and other support	mentioned	

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b Selection / topic		Checklist item	Reported	Assessment
TITLE				
Title 4 P	1.	Identify the report as a systematic review, meta-analysis, or both	Not reported	Title does not identify study as systematic review. This is a retrospective single-center study
RABSTRACT				
I Structured Ssummary M	2		Not applicable	Abstract present but not structured as systematic review format
AINTRODUCTIO	N			
Rationale C h	3	Describe the rationale for the review in the context of what is already known	Yes	Introductiion provides rationale for comparing SIOP vs NWTS protocols
c Objectives k l i s	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS)	Partially	Objective stated but not in formal PICOS format
t METHODS				

Protocol and	5	Indicate if a review protocol exist, if	Not	Not mention of
registration		and where it can be accessed, and, if available, provide registration	mentioned	protocol or registration
		information		
Eligibility	6	Specify study characteristics and report	Not	Single center study, not
criteria		characteristics used as criteria for eligibility	applicable	systematic review
Information	7	Describe all information sources and	Not	
sources		last date of search	applicable	
Search	8	Present full electronic search strategy	Not	
		for at least one database	applicable	
Study selection	9	State the process for selecting studies	Not	
			applicable	
Data collection	10	Describe methods of data extraction	Yes	
process		from reports and any processes for		
		obtaining and confirming data from		
		investigators		
Data items	11	List and define all variables for which	Not	Limited description of
		data were sought	detailed	variables collected
Risk of bias in	12	Describe methods used for assessing	Not	No assessment of study
individual		risk of bias of individual studies	mentioned	quality/bias
studies				
Summary	13	State the prinicpal summary measures	Yes	Survival rates,
measures				recurrence rates,
				mortality
Synthesis of	14	Describe the methods of handling data	Not	Descriptive statistics
results		and combining results of studies	applicable	only
Risk of bias	15	Specify any assessment of risk of bias	Not	Single study, not
across studies		that may affect the cumulative evidence	applicable	multiple studies
Additional	16	Describe methods of additional	Not	No additional analyses
analyses		analyses if done	mentioned	described
RESULTS				
Study selection	17	Give numbers of studies screened,	Not	Patient selection flow
		assessed for eligibility, and included in	applicable	chart provided (Figure
		the review, with reasons for exclusions		1)
		at each stage		
Study	18	For each study, present characteristics	Not	Single study design
characteristics		for which data were extracted	applicable	
	l			
Risk of bias	19	Present data on risk of bias of each	Not	No risk of bias

Results of	20	For all outcomes considered, present,	Yes	Results presented for
individual		for each study: simple summary data		SIOP vs NWTS
studies		for each intervention group		
Synthesis of	21	Present results of each meta-analysis	Not	No meta-analysis
results		done, including confidence intervals	applicable	conducted
		and measures of consistency		
Risk of bias	22	Present results of any assessment of	Not	Not applicable – single
across studies		risk of bias across studies	aapplicable	study
Additional	23	Give results of additional analyses, if	Not	No additional analyses
analysis		done	reported	
DISCUSSION				
Summary of	24	Ssummarize the main findings	Yes	Discussion summarizes
evidence		including the strength of evidence for		findings and compares
		each main outcome		with literature
Limitation	25	Discuss limitations at study and	Yes	Limitations mentioned:
		outcome level, and at review level		retrospective design,
				lack of
				multidisciplinary
				approach
Conclusions	26	Provide a general interpretation of the	Yes	Clear conclusions
		results in the context of other evidence		provided
FUNDING				
Funding	27	Describe sources of funding for the	Not	Not mentioned
		systematic review and other support	mentioned	

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Selection /		Checklist item	Reported	Assessment
topic		Checking item	Reported	rissessment
TITLE				
5 Title	1.	Identify the report as a	Not	Title does not identify study as
		systematic review, meta-	reported	systematic review. This is a
Ī		analysis, or both		retrospective single-center
P				study
_R ABSTRACT	•			
Structured	2		Partial	This study provide structured
summary				abstracts with objective,
M				methods, result, and
A				conclusions. However, they
				lack elements specific to
C				systematic reviews (data
h				sources, study selection, data

				extraction, synthesis methods) as they are primary research studies
INTRODUCTIO			T	
Rationale	3	Describe the rationale for the review in the context of what is already known	Yes	This studies adequately describe rationale, highlighting disparities in Wilms tumor outcomes between developed and developing countries, and the need for evidence-based approaches in resource limited settings
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS)	Yes	This study focuses on the role of pediatric oncologist in LMIC settings
METHODS				
Protocol and registration	5	Indicate if a review protocol exist, if and where it can be accessed, and, if available, provide registration information	Not applicable	Not applicable as this is primary research study, not systematic review requiring protocol registration
Eligibility criteria	6	Specify study characteristics and report characteristics used as criteria for eligibility	Yes	This study clearly define inclusion criteria patient ≤18 years with Wilms tumor diagnosis during specified time periods. Exclusion criteria are also mentioned (treatment abandonment, incomplete data)
Information sources	7	Describe all information sources and last date of search	Partial	This study describe their single center hospital databases as information sources with specific time periods. However, don't describe comprehensive literature research as expected in systematic review
Search	8	Present full electronic search strategy for at least one database	Not	No search strategies provided as these are retrospective cohort studies using hospital records, not systematic literature reviews
Study selection	9	State the process for selecting studies	Not applicable	This is a single center study analyzing own patient cohort

				rather than selecting studies from literature
Data collection process	10	Describe methods of data extraction from reports and any processes for obtaining and confirming data from investigators	Yes	This study describe data collection methods from hospital records, including clinical, surgical, pathological, and follow-up data extraction processes
Data items	11	List and define all variables for which data were sought	Not detailed	This study comprehensively ist variables collected including demographics, staging, histology, treatment modalities, complications, and outcomes
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies	Not mentioned	No assessment of study quality/bias
Summary measures	13	State the prinicpal summary measures	Yes	This study clearly state primary outcome measues
Synthesis of results	14	Describe the methods of handling data and combining results of studies	Partial	This studies describe statistical analysis methods, but don't involve meta-analysis or systematic synthesis as analyze single cohort
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence	Not applicable	Single study, not multiple studies
Additional analyses	16	Describe methods of additional analyses if done	Yes	This study describe subgroup analyses by stage, treatment protocol, and risk factor. Statistical significance testing and survival analysis stratification are well described
RESULTS			•	
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage	Partial	This study report patient numbers included and excluded with reasons (treatment abandonment, incomplete data) but this applies to patients rather than studies
Study characteristics	18	For each study, present characteristics for which data were extracted	Yes	This study comprehensively present patient characteristic demographics, staging distribution, treaatment

				modalities, and follow up data in detailed tables
Risk of bias within studies	19	Present data on risk of bias of each study	Not applicable	No risk of bias assessment
Results of individual studies	20	For all outcomes considered, present, for each study: simple summary data for each intervention group	Yes	This study present comprehensive outcome data including survival rates, confidence intervals, statistical comparisons, and subgroup analyses with appropriate tables and figures
Synthesis of results	21	Present results of each meta- analysis done, including confidence intervals and measures of consistency	Not applicable	No meta-analysis conducted
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies	Not aapplicable	Not applicable – single study
Additional analysis	23	Give results of additional analyses, if done	Yes	This study provide extensive subgroup analyses including stage specific outcomes, treatment protocol comparisons, and prognostic factor analyses with statistical significance testing
DISCUSSION				
Summary of evidence	24	Ssummarize the main findings including the strength of evidence for each main outcome	Yes	This study effectively summarize the main findings, compare results with international data, and discuss the clinical significance of their outcomes in the context of resource limited settings
Limitation	25	Discuss limitations at study and outcome level, and at review level	Yes	This study acknowledge significant limitations including small sample sizes, single center design, retrospective nature, missing data, and challenges in pathological diagnosis and staging
Conclusions	26	Provide a general interpretation of the results in the context of other evidence	Yes	This study provide clear conclusions with clinical implications for th=reating Wilms tumor in developing

				countries, emphasizing the
				importance of multidisciplinary
				care and adapted protocols.
FUNDING				
Funding	27	Describe sources of funding for	Not	Not mentioned
		the systematic review and other	mentioned	
		support		

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Selection / topic		Checklist item	Reported	Assessment		
TITLE						
Title 6	1.	Identify the report as a systematic review, meta-analysis, or both	Not reported	Title does not identify study as systematic review. This is a retrospective single-center study		
P ABSTRACT						
RStructured I summary S M A C h	2		Partial	This study provide structured abstracts with objective, methods, result, and conclusions. However, they lack elements specific to systematic reviews (data sources, study selection, data extraction, synthesis methods) as they are primary research studies		
kINTRODUCTIO	ON					
Rationale i s	3	Describe the rationale for the review in the context of what is already known	Yes	This studies adequately describe rationale, highlighting disparities in Wilms tumor outcomes between developed and developing countries, and the need for evidence-based approaches in resource limited settings		
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS)	Yes	This study aims to compare NWTS5 and SIOP2001 protocols to determine the most appropriate approach for resource-limited settings		

METHODS				
Protocol and registration	5	Indicate if a review protocol exist, if and where it can be accessed, and, if available, provide registration information	Not applicable	Not applicable as this is primary research study, not systematic review requiring protocol registration
Eligibility criteria	6	Specify study characteristics and report characteristics used as criteria for eligibility	Yes	This study clearly define inclusion criteria patient ≤18 years with Wilms tumor diagnosis during specified time periods. Exclusion criteria are also mentioned (treatment abandonment, incomplete data)
Information sources	7	Describe all information sources and last date of search	Partial	This study describe their single center hospital databases as information sources with specific time periods. However, don't describe comprehensive literature research as expected in systematic review
Search	8	Present full electronic search strategy for at least one database	Not	No search strategies provided as these are retrospective cohort studies using hospital records, not systematic literature reviews
Study selection	9	State the process for selecting studies	Not applicable	This is a single center study analyzing own patient cohort rather than selecting studies from literature
Data collection process	10	Describe methods of data extraction from reports and any processes for obtaining and confirming data from investigators	Yes	This study describe data collection methods from hospital records, including clinical, surgical, pathological, and follow-up data extraction processes
Data items	11	List and define all variables for which data were sought	Not detailed	This study comprehensively ist variables collected including demographics, staging, histology, treatment modalities, complications, and outcomes
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies	Not mentioned	No assessment of study quality/bias
Summary measures	13	State the prinicpal summary measures	Yes	This study clearly state primary outcome measues

Synthesis of results	14	Describe the methods of handling data and combining results of studies	Partial	This studies describe statistical analysis methods, but don't involve meta-analysis or systematic synthesis as analyze single cohort
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence	Not applicable	Single study, not multiple studies
Additional analyses	16	Describe methods of additional analyses if done	Yes	This study describe subgroup analyses by stage, treatment protocol, and risk factor. Statistical significance testing and survival analysis stratification are well described
RESULTS	•			
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage	Partial	This study report patient numbers included and excluded with reasons (treatment abandonment, incomplete data) but this applies to patients rather than studies
Study characteristics	18	For each study, present characteristics for which data were extracted	Yes	This study comprehensively present patient characteristic demographics, staging distribution, treaatment modalities, and follow up data in detailed tables
Risk of bias within studies	19	Present data on risk of bias of each study	Not applicable	No risk of bias assessment
Results of individual studies	20	For all outcomes considered, present, for each study: simple summary data for each intervention group	Yes	This study present comprehensive outcome data including survival rates, confidence intervals, statistical comparisons, and subgroup analyses with appropriate tables and figures
Synthesis of results	21	Present results of each meta- analysis done, including confidence intervals and measures of consistency	Not applicable	No meta-analysis conducted
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies	Not aapplicable	Not applicable – single study

Additional	23	Give results of additional	Yes	This study provide extensive
analysis		analyses, if done		subgroup analyses including
				stage specific outcomes,
				treatment protocol
				comparisons, and prognostic
				factor analyses with statistical
				significance testing
DISCUSSION				significance testing
Summary of	24	Ssummarize the main findings	Yes	This study effectively
evidence	2 1	including the strength of	103	summarize the main findings,
Cvidence		evidence for each main		compare results with
		outcome		international data, and discuss
		outcome		the clinical significance of their
				outcomes in the context of
				resource limited settings
Limitation	25	Discuss limitations at study and	Yes	
Limitation	23	Discuss limitations at study and	res	This study acknowledge
		outcome level, and at review level		significant limitations including
		level		small sample sizes, single
				center design, retrospective
				nature, missing data, and
				challenges in pathological
				diagnosis and staging
Conclusions	26	Provide a general interpretation	Yes	This study provide clear
		of the results in the context of		conclusions with clinical
		other evidence		implications for th=reating
				Wilms tumor in developing
				countries, emphasizing the
				importance of multidisciplinary
				care and adapted protocols.
FUNDING				
Funding	27	Describe sources of funding for	Not	Not mentioned
		the systematic review and other	mentioned	
		support		

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