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Please cli	ck on the	table heade	E Setting and time	F Methods	Section G Socio-demography	H Phenotypes	I Summar
n work	in work	in work	in work	in work	in work	in work	in work
complete	complete	complete	complete	complete	complete	complete	complete
B Manuscrip	ot classificatio	on according to s	ystematic review aim	ıs			
phenotype/s	of IgE - asso	ciated diseases o	or conditions including	ng asthma/\	fy a clinically expressed wheezing, atopic eczem n childhood to young ad	ıa, rhinoconjuı	
O yes				O no			
2. Do you th	ink this study	y does fulfill the p	protocol inclusion cr	iteria ?			
O yes		0	no		show inclusion criter	ia	
3. Do you th	ink this study	y fulfills one or m	ore protocol exclusi	ion criteria ?			
O yes		0	no		show exclusion criter	ria	
	riteria is not f	ullfilled ?					
3.1 Which ci							
3.1 Which ci							
3.1 Which ci							
3.1 Which ci							

 ☐ The study explores heterogeneity within phenotypes ☐ The study explore overlap between phenotypes ☐ The study appraised or validated methods (test or biological measurements) to better define or classify phenotypes ☐ None of the previous categories 					
C - Study aim/s and hypothesis					
Study hypothesis					
Was the study hypothesis clearly described in the text?					
O yes	O no				
Study aim/s					
2. Please capy the full simila text and pasts into the bay below.					
2. Please copy the full aim/s text and paste into the box below					
D - Disease/s or condition/s included and study design					
1. Which IgE associated diseases or conditions are included in	this study (MR) ?				
Asthma and/or wheeze	☐ Food Allergy				
Rhinitis	☐ Urticaria				
Eczema / Atopic dermatitis	Anaphylaxis				
2. Considering the aim/s of the study, in which of the following					
Description of phenotypes (*)	Treatment response (*)				
☐ Validation of phenotypes (*)	Others				
Prognosis (*)	(*) see tooltips				
3. Which of the following categories of study designs should th	is study be included ?				
O Case series	O Cohort				
O Cross-sectional	O Clinical trial				
○ Case-control					
Cohort study: please answer questions from 4.1 to 4.3					

4.2 Could you indicate the age at the starting and end of individual's follow-up? Age at starting point at end point if in	4.1 Which was the type of fol	low-up ?					
Age at starting point at end point wears if in years years years if in months	O Prospective						
Age at starting point at end point wears if in years years years if in months							
wears was information collected over the period of follow-up including baseline and end point? A.3 How many times was information collected over the period of follow-up including baseline and end point? Neverfunknown	4.2 Could you indicate the ag	ge at the starting and end of i	ndividual's follow-up 1	•			
wears	Age at starting po	oint at end point					
months mo	if in	vears vears					
### A How many times was information collected over the period of follow-up including baseline and end point? Never/unknown	years	, , , , , , , , , , , , , , , , , , , ,					
4.3 How many times was information collected over the period of follow-up including baseline and end point? Never/unknown 1 x 2 x 3 x 4 x 5 x 6 x 7 x 8 x 9 x >= 10 x Clinical trial : please answer questions from 5.1 to 5.10 Randomized		month	IS				
Never/unknown 1x	months						
Never/unknown 1x							
Clinical trial : please answer questions from 5.1 to 5.10 5.1 If the study design was a clinical trial, mark the items that best describe its characteristics (MR)? Randomized		rmation collected over the p	eriod of follow-up incl	uding baseline	and end poi	nt ?	
Clinical trial: please answer questions from 5.1 to 5.10 5.1 If the study design was a clinical trial, mark the items that best describe its characteristics (MR)? Randomized	_	3 × 0 4 × 0 5 ×	06v 07v	∩ 8 v	Ogy	○ >= 10 v	
S.1. If the study design was a clinical trial, mark the items that best describe its characteristics (MR)? Randomized	011 021 03	74 044 034	00% 07%	001	037	0 >= 10 x	
S.1. If the study design was a clinical trial, mark the items that best describe its characteristics (MR)? Randomized							
Randomized Community Intervention Double blind Comparison group Multicenter "Intervention" selected: please answer also questions 5.2 to 5.3 5.2 Was the intervention clearly defined? yes no 5.3 Could you briefly describe the intervention? 5.4 Were the inclusion/exclusion criteria clearly specified? yes no 5.5 How many groups were compared? groups 5.6 Overall, how long was the period of follow-up? years: OR months: 5.7 How many times was information collected over the period of follow-up including baseline and end point? Never/unknown	Clinical trial : please answer	questions from 5.1 to 5.10					
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Double blind Comparison group Multicenter "Intervention" selected: please answer also questions 5.2 to 5.3 5.2 Was the intervention clearly defined? yes		clinical trial, mark the items			(MR) ?		
Population-based	□			ntion			
5.2 Was the intervention clearly defined? yes no 5.3 Could you briefly describe the intervention? 5.4 Were the inclusion/exclusion criteria clearly specified? yes no 5.5 How many groups were compared? groups 6.6 Overall, how long was the period of follow-up? years: OR months: 5.7 How many times was information collected over the period of follow-up including baseline and end point? Never/unknown							
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○ Never/unknown							
○ Never/unknown	5.7 How many times was info	ormation collected over the p	eriod of follow-up incl	uding baseline	and end poi	nt ?	
1x 02x 03x 04x 05x 06x 07x 08x 09x 010x	O Never/unknown	<u> </u>	•	-	•		
	•	3x	○ 6 x ○	7 x 0 8	х 🔾 9	x 0 10 x	

	at analysis described in t	the methods section ?		
○ yes ○ no				
5.9 Was intention to tre	eat analysis really carried	out ?		
O yes		O no		
5.10 Did the authors of	the manuscript mention	that the study design com	plied with the CONSORT statement ?	
O yes	O no	see CONS	SORT statement	
E - Study setting and tin	me			
	ribe the study population			
O Population based		Clinica	l population	
1.1 Could you indicate	which was the setting fro	om which the individuals	were recruited (MR) ?	
☐ Hospital	Outpatients clinic	Disease registry	Primary care Others	
2 Was the study conduct One country	cted only in one country o		ational level	
0.4 la salais la sassata	4144			
2.1 in which country wa	as the study conducted ?			
3. Was the study period	l available ?			
3. Was the study period yes	l available ?	O no		
O yes				
yes 3.1 Period of recruitme		o no	rear:	
O yes			rear:	
yes 3.1 Period of recruitme Starting year :	ent ?	◯ no Finishing y	rear:	
yes 3.1 Period of recruitme Starting year :		◯ no Finishing y	rear:	
yes 3.1 Period of recruitme Starting year: 4. Which was the total r	ent ?	◯ no Finishing y	rear:	
3.1 Period of recruitme Starting year : 4. Which was the total r Number of individuals :	number of individuals incl	◯ no Finishing y	rear:	
3.1 Period of recruitme Starting year : 4. Which was the total r Number of individuals : 5. Which was the partic	number of individuals incl	◯ no Finishing y	rear:	
yes3.1 Period of recruitmeStarting year :4. Which was the total rNumber of individuals :	number of individuals incl	◯ no Finishing y	rear:	
yes 3.1 Period of recruitme Starting year: 4. Which was the total r Number of individuals: 5. Which was the partic Rate (in %):	number of individuals incl	ono Finishing y		
yes 3.1 Period of recruitme Starting year: 4. Which was the total r Number of individuals: 5. Which was the partic Rate (in %):	number of individuals incl	ono Finishing y	rear:	

F - Methods used for individuals assessment					
Questionnaires					
Were questionnaires used for the individuals assessment?	0				
O yes	O no				
1.1 Was a reference given for the questionnaire ?					
O yes	○ no				
4.0 Which was the automore annual and					
1.2 Which was the reference number ? Reference number :					
Reference number .					
1.3 How was the questionnaire administered (MR) ?					
Self-administered Telephone	Personal interview Clinical examination				
Biological measurements					
2.1 Was total IgE measured?					
O yes	O no				
2.2 Were specific IgEs measured ?					
O yes	O no				
2.3 Which type of allergens were tested ?					
Food 2.3.1 How many allergens were tested?					
☐ Indoor 2.3.2 How many allergens were tested?					
Outdoor 2.3.3 How many allergens were tested?					
, ,					
2.4 Was the skin prick test used ?	O 20				
O yes	O no				
2.5 Which type of allergens were tested ?					
Food 2.5.1 How many allergens were tested?					
☐ Indoor 2.5.2 How many allergens were tested?					
Outdoor 2.5.3 How many allergens were tested?					
2.6 Was the patch test used ?					
	O no				
	,				
2.6.1 How many allergens were tested ?					
Number of allergens :					

2.7 Was spirometry test used to measure lung function ?		
O yes	O no	
2.7.1 Weeks most broughed ileter test performed 2		
2.7.1 Was a post-bronchodilator test performed ? O yes	O no	
<u> </u>	0 110	
2.8. Was peak flow test used to measure lung function?	0.50	
O yes	O no	
2.9. Was a BHR test performed ?	0	
O yes	O no	
2.9.1 Which was the method used ?		
☐ Methacoline ☐ Hypertonic Saline ☐ Exercise	Histamine	EVH (Eucapnic voluntary hyperpnoea)
2.10 Were other biomarkers measured ?		
O yes	O no	
2.10.1 Which biomarkers were measured ?		
1) 2)		
3)		
4)		
5)		
2.10.2 In which of the following media were they measured?		
Serum Plasma Broncheoalveolar lavage	Sputum	Breath condensate Urine
2.11 Which other tests / measurements / examinations were per	formed ?	
(please enter 'none' if not other were performed !)		
1)		
2) 3)		
4)		
5)		
7,]		
Severity assessment		
3.1 Was phenotype/s severity assessed ?		
O yes	O no	
3.1.1 Could you briefly describe which guidelines criteria or m	etnods were used t	o assess severity ?

G Socio-demographic characteristics and family history		
, ,		
Socio-demographic characteristics		
1.1 Which is the individuals age range for the overall populatio	n ?	
in Months : to OR in Years :	to	
In the case of cohort studies or clinical trials with a period of fol D5.6)	low-up the age range at baseline should be inc	dicated (see D4.4 /
53.0)		
1.2 Which was the sex distribution ?		
Males (in %):		
Females (in %):		
1.3 Was ethnicity/ race collected ?		
O yes	○ no	
<u></u>	<u> </u>	
1.4 Was parent education collected ?		
O yes	O no	
1.5 Were any of the socio-economic indicators collected ?		
O yes	O no	
Family bisks m.		
Family history		
Mother		
2.1 Was mother's allergic disease/s history collected ?		
O yes	O no	
2.1.1 Which condition/c 2		
2.1.1. Which condition/s ?		
1)		
2)		
3)		
4)		

2.1.2. How was/were this/these cond	lition/s assessed (MR) ?			
Questionnaire	gE / Blood	Skin Prick Test	Not clear	
Father				
2.2 Was fathers's allergic disease/s h	nistory collected ?			
O yes		O no		
2.2.1. Which condition/s ?				
1)				
2)				
3)				
4)				
2.2.2. How was have this it has a sound	litianle accessed (MD) 2			
2.2.2. How was/were this/these cond Questionnaire	gE / Blood	Skin Prick Test	Not clear	
Questioninane	gc / 5100u	SKIIT HOR TOST	1 Not cical	
H Phenotypes name				
Phenotype			Was the phen definition pro-	
1.				
			O yes	O no
2.			o yes	O no
3.			_	_
			O yes	O no
4.			O yes	O no
5.			0,	0
J.			o yes	O no
6.			o yes	O no
			O yes	O III
7.			o yes	O no
8.			_	_
			O yes	O no
9.			o yes	O no
10.				_
			o yes	O no
,				
I - Summary of study conclusions				

4. Please include any additional comments which might be relevant for the review