Supplementary file 1. Diagnostic labels of chILD

Classification of chILD (based on Deutsch et al, 2007)

DISORDERS MORE PREVALENT IN	DISORDERS LESS PREVALENT IN
INFANCY	INFANCY
Diffuse developmental disorders of the	Disorders related to systemic disease
lung:	processes:
Acinar dysplasia	Îmmune mediated/collagen- vascular
Congenital alveolar dysplasia	disorders
Alveolar capillary dysplasia with	Storage disease
misalignment of pulmonary veins	Sarcoidosis
	Langerhans cell histiocytosis
	Malignant infiltrates
Lung growth abnormalities reflecting	Disorders of the normal host:
deficient alveolarisation:	Related to infections
Pulmonary hypoplasia	Related to environmental agents
Chronic neonatal lung disease	- hypersensitivity pneumonitis
Related to chromosomal disorders	- toxic inhalation
Related to congenital heart disease	Aspiration syndromes
	Eosinophilic pneumonia
Specific conditions of undefined cause:	Disorders masquerading as ILD:
Neuroendocrine cell hyperplasia of infancy	Arterial hypertensive vasculopathy
Pulmonary interstitial glycogenosis	Congestive changes related to cardiac
	dysfunction
	Veno-occlusive disease
	Lymphatic disorders
Inherited surfactant disorders:	Disorders of the immunocompromised
Surfactant protein B mutations	host:
Surfactant protein C mutations	Opportunistic infection
ABCA3 mutations	Disorders related to therapeutic intervention
Histology consistent with surfactant	Disorders related to transplantation and
dysfunction without a yet recognised	rejection syndromes
genetic	Diffuse alveolar damage of unknown
aetiology	aetiology
- pulmonary alveolar proteinosis	
- chronic pneumonitis of infancy	
- desquamative interstitial pneumonitis	
- non-specific interstitial pneumonia	

Deutsch GH, Young LR, Deterding RR, Fan LL, Dell SD, Bean JA, Brody AS, Nogee LM, Trapnell BC, Langson CL. Diffuse lung disease in young children. Application of a novel classification system. Am J Resp Crit Care Med 2007:176;1120-1128