

Project Name: ITOTS Expansion



**Infant & Toddler
Connection of Virginia**

Lookup List Data

Project Name: ITOTS Expansion
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Version Control

Version	Date	Author	Change Description
1.0	12/03/09	Wendy Cary	Document created
1.1	12/04/09	Wendy Cary	<ul style="list-style-type: none">• Added values for Developmental Screen Tools• Modified Risk Factors lists to combine developmental, hearing, and vision lists.

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ID	Category	Description	Value 1	Value 2
LK-001	Relationships	Father Foster Father Foster Mother Grandfather Grandmother Guardian ad litem Legal Guardian Mother Other Social Worker Step-Parent Surrogate Parent		
LK-002	Living Situations	CPS Custody and Living with Family CPS Foster Care Group/Institutional Setting Homeless Living with Family Other Other Foster Care Unknown		
LK-003	How Hear	Advertising, Billboard Advertising, Other Advertising, Print Advertising, Radio Advertising, TV Central Directory Community Services Board Program Day Care Provider Discharge Planners in Hospital Doctor's Office DSS Friend Head Start		

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ID	Category	Description	Value 1	Value 2
		Health Department Healthy Families Other Early Intervention Program Parent Public Schools Relative		
LK-004	Referral Source Types	Central Directory Community Services Board Program Day Care Provider Discharge Planners in Hospital Doctor's Office DSS DSS CAPTA Friend Head Start Health Department Healthy Families Other Early Intervention Program Parent Public Schools Relative		
LK-005	Referral Dispositions	Additional Referral Deceased Lost to Follow-up Parent Declined to Proceed Proceed to Intake		
LK-006	Referral Disclosures	Assessment Reports Current IFSP Contact Notes Discharge Summaries Other Results of Eligibility Determination		

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ID	Category	Description	Value 1	Value 2
		Services Being Provided to Child/Family Status of Initial Family Contact		
LK-007	Provider Types	Community Health Aide Practitioner Family/General Practitioner No Medical Home Nurse Practitioner Other Pediatrician Physician's Assistant Public Health Nurse		
LK-008	Genders	Female Male Unknown (inactive)		
LK-009	FIPS Codes	(see XREF_FIPS)		
LK-010	Races	American Indian or Alaskan Native Asian Asian or Pacific Islander (inactive) Black or African American Hispanic/Latino (inactive) Other (inactive) Pacific Islander or Hawaiian Native Two or More Races Unknown White		
LK-011	School Districts	(get from DOE)		
LK-012	Languages			
LK-013	Contact Methods	Email Fax Mail Phone		
LK-014	Risk Factors	An illness or condition requiring admission of 48	H	Present during the

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		hours or greater to a NICU		neonatal period
		Apgar Score of 0-3 at 5 minutes	D	
		Birth weight – Low (1500 g to <2500 g or 3.25 lbs to 5.5 lbs)	D	
		Birth weight – Very Low (<1500 g or <3.25 lbs)	D	
		Cerebral palsy	V	
		CHARGE syndrome	V	
		Craniofacial anomalies, including those with morphological abnormalities of the pinna and ear canal	H	Present during the neonatal period
		Craniofacial or external ear anomalies	H	Delay may develop
		Documented systemic infection, congenital or acquired	D	
		Down Syndrome	V	
		Encephalitis	V	
		Environmental or social risk factor – Domestic violence	D	
		Environmental or social risk factor – Lack of adequate shelter	D	
		Environmental or social risk factor – Lack of familial support	D	
		Family history of childhood Blindness	D	
		Family history of childhood Deafness	D	
		Family history of hereditary vision loss (such as Retinoblastoma, Albinism)	V	
		Family history of permanent childhood hearing loss	H	Present during the neonatal period
		Family history of permanent childhood sensorineural hearing loss	H	Delay may develop
		Fetal Alcohol Syndrome	V	
		Head trauma	H, V	Delay may develop
		Hearing loss	V	

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		Hydrocephalus	V	
		Hyperbilirubinemia requiring exchange transfusion	D	
		Hypoxia, anoxia, birth trauma	V	
		In-utero infection such as cytomegalovirus, herpes, toxoplasmosis, or rubella	H	Present during the neonatal period Delay may develop
		In-utero infections including cytomegalovirus, herpes, toxoplasmosis, or rubella	H	
		In utero infections, such as cytomegalovirus (CMV), rubella, herpes, toxoplasmosis, or syphilis	V	
		Intrauterine drug exposure	V	
		Intraventricular hemorrhage (IVH grade I – III), stroke	V	
		Lack of well-child care	D	
		Lead poisoning	D	
		Low Apgar Score	V	
		Low birth weight	V	
		Major congenital anomalies (see instructions)	D	
		Maternal age 15 or less	D	
		Maternal conditions during pregnancy such as accidents, phenylketonuria (PKU), maternal diabetes or sickle cell	D	
		Meningitis	D,V	
		Neurodegenerative disorders including Hunder Syndrome, Friedreich’s ataxia and Charcot-Marie-Tooth Syndrome	H	Delay may develop
		Neurofibromatosis Type II	H	Delay may develop
		Other (please list)	D	
		Other syndromes such as Goldenhar, Hurler, Marfan, Norrie, Refsum, Trisomy 13, Tay-Sachs, neurofibromatosis, Lowe’s, Stickler	V	

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		Periventricular leucomalacia	D	
		Persistent pulmonary hypertension	D	
		Persistent pulmonary hypertension associated with mechanical ventilation, hyperbilirubinemia requiring exchange transfusion, or conditions requiring extracorporeal membrane oxygenation (ECMO)	H	Delay may develop
		Postnatal infections associated with sensorineural hearing loss including bacterial meningitis	H	Delay may develop
		Prematurity	V	
		Prematurity – Gestational Age: 28-31 weeks	D	
		Prematurity – Gestational Age: 32-37 weeks	D	
		Prematurity – Gestational Age: Less than 28 weeks	D	
		Recurrent or persistent otitis media with effusion for at least 3 months	H	Delay may develop
		Seizure disorder – excluding recurrent febrile seizures	D	
		Seizures	V	
		Severe parenting risk factor – Intellectual disability	D	
		Severe parenting risk factor – Mental illness	D	
		Severe parenting risk factor – Physical disability	D	
		Severe parenting risk factor – Substance abuse	D	
		Shaken Baby Syndrome	V	
		Small for gestational age (10 th percentile or less)	D	
		Stigmata of syndromes known to have sensorineural or conductive hearing loss	H	Delay may develop
		Stigmata or other findings associated with a syndrome known to include a sensorineural and/or conductive hearing loss	H	Present during the neonatal period

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		Syndromes associated with progressing hearing loss including neurofibromatosis, osteopetrosis, Usher’s Syndrome, Goldenhar Syndrome, Brancio-Oto-Renal Syndrome, CHARGE Association, Pendred Syndrome, Pierre Robin Syndrome, Trisomy 21 (Down) Syndrome, Waardenburg Syndrome, choanal atresia, Stickler Syndrome and Rubinstein-Taybi Syndrome	H	Delay may develop
LK-015	Third Party Coverage Types			
LK-016	Developmental Screen Tools	Ages and Stages Questionnaire (ASQ) Bayley Infant Neurodevelopmental Screen (BINS) Parents’ Evaluation of Developmental Status (PEDS)		
LK-017	Developmental Screen Areas	Adaptive Cognitive Communication – Expressive Communication – Receptive Motor – Fine Motor – Gross Social/Emotional		
LK-018	Intake Dispositions	Deceased Lost to Follow-up Parent Declined to Proceed Proceed to Eligibility		
LK-019	Eligibility Methods	Comprehensive developmental screening Formal/informal observation Informed clinical opinion Parent Report Part C Vision Screening		

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		Part C Hearing Screening Review of pertinent medical records less than six (6) months old from the primary care physician and other sources related to the child's current health status, physical development (including vision and hearing), and medical history Review of other records such as birth records, newborn screening results and early medical history, with parent consent, even if those records are more than six (6) months old.		
LK-020	Eligibility Dispositions	Parent Declined to Proceed Proceed to Service Planning		
LK-021	Developmental Delays	Adaptive Cognitive Communication Physical: including fine & gross motor Social or emotional		
LK-022	Atypical Developments	Abnormal or questionable sensory-motor responses Behavioral disorders that interfere with acquisition of developmental skills Identified affective disorders Impairment in social interaction and communication skills along with restricted and repetitive behaviors		
LK-023	Diagnosed Conditions	Autism Spectrum Disorder Brain or spinal cord trauma, with abnormal neurologic exam at discharge Chromosomal abnormalities Cleft lip or palate Congenital or acquired hearing loss		

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		Effects of toxic exposure including fetal alcohol syndrome, drug withdrawal, exposure to chronic maternal use of anticonvulsants, antineoplastics, and anticoagulants Endocrine Disorders (with a high probability of resulting in developmental delay) Failure to thrive Hemoglobinopathies (Sickle Cell) (with a high probability of resulting in developmental delay) Inborn errors of metabolism Meningomyelocele (spina bifida) Microcephaly None Other physical or mental conditions (please list) Seizures with significant encephalopathy Severe attachment disorder Severe Grade 3 intraventricular hemorrhage with hydrocephalus or Grade 4 intraventricular hemorrhage Significant central nervous system anomaly (e.g. cerebral palsy) Symptomatic congenital infection Visual disabilities		
LK-024	Adaptive Skill Areas			
LK-025	Service Types	Assistive Technology Audiology Family Training & Counseling Health Services Medical Services (diag/eval) Nursing Services Nutrition Services		

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ID	Category	Description	Value 1	Value 2
		Occupational Therapy Other Physical Therapy Psychological Services Respite Care Service Coordination Sign Language and Cued Language Services Social Work Services Special Instruction Speech/Language Pathology Transportation Vision		
LK-026	Service Occurs	As Needed Every Only		
LK-027	Service Methods	a Coaching, including hands-on as appropriate b Consultation c Assessment d Provision of assistive technology device		
LK-028	Service Settings	1 Program Designed for Children with Development Delays or Disabilities 2 Program Designed for Typically Developing Children 3 Home 4 Hospital (inpatient) 5 Residential Facility 6 Service Provider Location (center/clinic/hospital) 7 Other		
LK-029	Surrogate Needed Reasons	Child is a ward of the state residing in a residential facility/group home/etc. Parent/Guardian cannot be identified		

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		Parent/Guardian whereabouts are unknown after reasonable efforts have been made by the early intervention system to locate the parent/guardian.		
LK-030	Surrogate No Longer Needed Reasons	Child is a ward of the state residing with foster parent(s) Child resides with “person acting in the place of a parent” (grandparent, relative, stepparent, etc.) Child resides with parent/guardian		
LK-031	Surrogate Termination Reasons	Other The child’s circumstances have changed such that a surrogate parent is no longer required. You have chosen to relinquish this responsibility. You have indicated you are no longer able to advocate effectively for the child. You no longer meet the criteria established for being a surrogate parent.		
LK-032	Hearing Screening Results	Born outside Virginia, no newborn hearing screening or unknown results Missed Pass Pass but at-risk Refer		
LK-033				
LK-034	Hearing Behavioral Observations	Coos Makes some sounds Quiets when talked to or with soothing sounds Startles or cries at loud, sudden noises Turns eyes or head toward source of sound Attends to music or singing Makes strings of sounds; babbles (ba-ba-ba, ga-	6 6 6 6 6 9 9	

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		ga-ga) Stops or pays attention when told “no” or name called Turns head when called from behind Babbles using variety of sounds and intonation patterns Begins to repeat some of the sounds other make Responds to own name Uses 3 – 20 or more words Follows simple commands (e.g. “Come here.”) Indicates wants/needs with words/vocalizations & gestures Begins using 2-word “sentences” Enjoys listening to stories Points to some body parts Understands 300 or more words Uses 50-100 or more words Follows 2-stage commands Notices different sounds (doorbell, phone, etc.) Speaks so understood 50 – 75% of time Understands most things that are said to him/her Uses 3 to 4-word phrases Uses 50 – 250 or more words	9 9 12 12 12 12 18 18 24 24 24 24 24 36 36 36 36 36 36	
LK-035	Hearing Screening Findings	There are no components of the Virginia Part C Hearing Screening that would indicate the need for referral for full audiological evaluation. One or more of the components of the Virginia Part C Hearing Screening indicate the need for monitoring of the child’s hearing status (please describe recommended frequency and type of monitoring).		

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	Eyes	<p>Drooping eyelid which obscures the pupil</p> <p>High sensitivity to bright light, indicated by squinting, closing eyes, or turning head away</p> <p>Obvious abnormalities in the shape or structure of the eyes</p> <p>Persistent redness of the conjunctiva (normally white)</p> <p>Absence of eyes moving together or sustained eye turn after four to six months of age</p> <p>Involuntary rhythmic or jerky eye movements (nystagmus)</p> <p>Averts gaze or seems to be looking beside, under, or above the object of focus</p> <p>Holds object close to eyes</p> <p>Tilts or turns head in certain positions when looking at an object</p> <p>Accurate reaching for objects by six months</p> <p>Eye contact by age three months</p> <p>Visual fixation or following by three months</p>	<p>appearance of the eyes</p> <p>Atypical appearance of the eyes</p> <p>Atypical appearance of the eyes</p> <p>Atypical appearance of the eyes</p> <p>Atypical appearance of the eyes</p> <p>Unusual eye movements</p> <p>Unusual eye movements</p> <p>Unusual gaze or head positions</p> <p>Unusual gaze or head positions</p> <p>Unusual gaze or head positions</p> <p>Absence of the following behaviors</p> <p>Absence of the following behaviors</p> <p>Absence of the following</p>	

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			behaviors	
LK-040	Vision Screening Findings	<p>There are no components of the Virginia Part C Vision Screening that would indicate the need for referral for full vision evaluation.</p> <p>One or more components of the Virginia Part C Vision Screening indicate the need for monitoring of the child’s vision status (please describe recommended frequency and type of monitoring)</p> <p>One or more components of the Virginia Part C Vision Screening indicate the need for referral to a physician or eye care specialist for full vision evaluation.</p>		