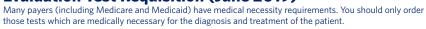
## Athena Diagnostics NewbornDx™ Advanced Sequencing Evaluation Test Requisition (July 2019)





Fields in red indicate required information		Patient Identification	1
Complete this requisition for direct billing to hospitals, I clinics.	aboratories or	NOTE: Two forms of pat Patient Name First	tient ID must be listed on EACH specimen.
Name			ble)
Phone		DOB	Sex:
Fax		1	☐ Female☐ Unknown
Email		Mailing Address	
Test Options	Test code		State Zip
NewbornDx <sup>™</sup> Advanced Sequencing Evaluation			Day 🗌 Eve 🗌 Cell
☐ Proband	2052	Phone #2	
Proband Specimen Type*: ☐ DBS ☐ Whole	e blood	Authorization to Use De-ide	<b>ntified Specimen for Research</b> . Test results are confidential and prized by the patient or the patient's authorized representative or
☐ Family Member 1	2053	consistent with applicable stat better health insights, Quest D	orized by the patient or the patient's authorized representative or team federal law. To promote medical understanding and develop Diagnostics requests the patient's permission to use the specimen it identifying the test subject) for research, educational studies, publication. Your name or other personal identifying information of the results of any studies or publications. You are not required set, and the decision to consent to the use of the specimen for such ext processing or testing of the specimen, the test results or the services in connection with this testing. Please indicate your choice regarding ecimen by checking the line next to the appropriate option below.
Family Member Name		commercial purposes and/or will not be used in or linked t	publication. Your name or other personal identifying information of the results of any studies or publications. You are not required
Family Member DOB		to consent to any of these use purposes will not in any way affe	es, and the decision to consent to the use of the specimen for such each processing or testing of the specimen, the test results or the services in consent in with the testing of the specimen.
Specimen Type*: ☐ DBS ☐ Whole blood ☐	☐Saliva		
☐ Family Member 2	2053	I consent to the use of my de-	identified specimen for research: Yes No
Family Member Name		Signature of Patient, Parent or Le	gally Authorized Representative Date
Family Member DOB		Printed Name of Patient, Parent of	or Legally Authorized Representative
Specimen Type*: $\square$ DBS $\square$ Whole blood $\square$	☐Saliva	Relationship to Patient if Signator	ry is Someone Other than Patient
► ICD Code (Required):		Authorized Result Re	,
Hospital/Laboratory Billing Information		Required Physician II	
(Hospital billing is required for all Medicare patients – both inpatients and outpatients.)			Athena Acct #
Athena Account # (if assigned)		NameFirst	t Last
CLIA #		Address	
Purchase Order # (if available)		City	State Zip
Billing Contact		Phone	Fax
Email		Email	
Phone		Laboratory Informati	ion
Fax		Lab Name	
Hospital/Lab Name		Address	
Address			Fax
City		Email	
State	_ Zip		
*2.0 mL (1.0 mL minimum) whole blood EDTA (lavender top) tub *Do not eat, drink, smoke or chew gum 30 minutes prior to saliva *One blood spot card required with 5 blood spots preferred (2 sp NOTE: Specimen(s) must be labeled with two of the following for These same two forms of ID must be indicated on the test requise	a collection. Saliva mus oots minimum). Please rms of identification: n	allow the spots to dry fully be ame, date of birth, last four di	fore placing in any packaging, approximately three hours gits of SS#, patient ID no.
Physician Attestation of Informed Consent In accordance with Massachusetts General Law Chapter 111, Section 70G, testing laboratories located in Massachusetts require a signed acknowledgr have not previously signed a blanket Physician Attestation of Informed Conse	nent from the ordering me	dical practitioner. The signed acknowledge	wledgment is required to complete the genetic testing ordered if yo
I warrant that I have obtained both oral and written consent using the <b>Patien</b> is the subject of the test (or if that person lacks capacity to consent, signed by			nena Diagnostics. This written consent was signed by the person wh
Medical Practitioner Signature	Da	rte Printed Name of Medica	l Practitioner NPI
Patient Informed Consent Form for Genetic Testing is available at Ather	naDiagnostics.com/cons	ent.	

## Athena Diagnostics NewbornDx™ Advanced Sequencing Evaluation Test Requisition (June 2019)





Family member information	Previous genetic testing		
If ordering trio or duo testing, please provide the names and relationship to proband of additional family members:	Please attach any relevant results to this form for review.  Chromosomes/Microarray analysis   No results/not performed  Yes/results:		
Family Member Name 1:	Chromosomes/ Microardy analysis1101esuns/hot performed1es/resules.		
Relationship to Proband:			
Family Member Name 2:			
Relationship to Proband:			
Proband Information			
Reason(s) for referral for NewbornDx™ Advanced Sequencing			
Evaluation, ie, what was the initial presenting symptom?	Biochemical studies ☐ No results/not performed ☐ Yes/results:		
	Other molecular studies, including prenatal testing:		
Age of onset of initial presenting symptom:	Please fill out the below information based on the history of the patient being tested. Clinic notes should also be submitted. This information is essential to correlate results found using the NewbornDx <sup>TM</sup> Advanced Sequencing Evaluation. The laboratory may contact the ordering cliniciar if more information is necessary.		
☐ Central or South American ☐ French Canadian ☐ Hispanic or Latino	Clinical details		
☐ Middle Eastern ☐ Pacific Islander or Native Hawaiian	Date of last clinical exam:		
☐ Sephardic Jewish ☐ Western/Northern European	Biological sex: ☐ Male ☐ Female		
☐ Other (please specify):	☐ Other (please specify):		
	Head circumference:%tile		
Is there a family history of a similar or related disorder?	Weight:%tile		
Consanguinity (related by blood, eg, parents related by blood)?	Height: %tile		
Yes ☐ No ☐ Unsure ☐ If yes, please specify:	Common diagnoses (please provide more information about these common diagnoses using the check boxes on the next page):		
History of a bone marrow transplant:  Yes* No *If yes, please contact 1.800.394.4493 ext. 2031 to speak to a Genomic Science Specialist before sending any samples.	<ul> <li>☐ Ambiguous genitalia</li> <li>☐ Congenital heart defect</li> <li>☐ Developmental delay</li> <li>☐ Failure to thrive</li> <li>☐ Hypotonia</li> <li>☐ Multiple congenital anomalies</li> <li>☐ Structural brain abnormalities</li> <li>☐ Developmental delay</li> <li>☐ Failure to thrive</li> <li>☐ Metabolic acidosis</li> <li>☐ Seizures</li> </ul>		

## Athena Diagnostics NewbornDx™ Advanced Sequencing **Evaluation Test Requisition (June 2019)**Many payers (including Medicare and Medicaid) have medical necessity requirements. You should only order those tests which are medically necessary for the diagnosis and treatment of the patient.



Cancer	Endocrine	Hair & skin (continued)	Pre/perinatal history
☐ Type of cancer:	☐ Adrenal gland abnormality	☐ Cafe-au-lait spots	☐ Conceived via artificial
	☐ Adrenal insufficiency	Hair	reproductive technology
☐ Age of diagnosis:	☐ Cushing syndrome	Alopecia	Congenital diaphragmatic hernia
	☐ Diabetes insipidus	☐ Brittle	☐ Cystic hygroma
Family history of cancer and	☐ Diabetes mellitus	☐ Coarse ☐ Hypopigmentation	☐ Encephalocele ☐ Increased nuchal translucency
affected relatives:	☐ Growth hormone deficiency	☐ Hemangioma	☐ Intrauterine growth restriction
	☐ Hirsutism	☐ Hyperextensible skin	☐ Oligohydramnios
Cardiovascular	☐ Immunologic abnormality	☐ Hyperextensible skill ☐ Hyperpigmented macule	☐ Omphalocele
Anemia	Specify:	☐ Hypopigmented macule	☐ Polyhydramnios
☐ Aortic root dilation	☐ Obesity	☐ Hypertrichosis	☐ Prematurity
	☐ Pancreatic insufficiency	☐ Ichthyosis	☐ Terratogen exposure
Arrhythmia	☐ Parathyroid dysfunction	☐ Infections	Specify:
Atrial septal defect	☐ Thyroid dysfunction	Lipoma	Other:
☐ Bicuspid aortic valve	Other:	☐ Nail abnormality	Respiratory
☐ Cardiomyopathy	Eye defects & vision	☐ Neurofibroma	Apnea
☐ Coarctation of aorta	☐ Amblyopia	Rash	□ Asthma
☐ EKG abnormality	☐ Aniridia	Other:	☐ Bronchiectasis
☐ Mitral valve prolapse	☐ Anophthalmia	Metabolic	Hyperventilation
☐ Patent ductus arteriosus	☐ Blue sclerae	Acidosis	Hypoventilation
☐ Patent foramen ovale	<del></del>	□ Lactic	☐ Pneumothorax
☐ Teratology of Fallot	☐ Cataracts	☐ Metabolic	Recurrent infections
☐ Thrombocytopenia	☐ Congenital	☐ Creatinine Phosphokinase (abnormal)	Respiratory failure
· ·	☐ Postnatal	☐ CSF lactate level (abnormal)	☐ Respiratory insufficiency
Thrombosis	☐ Cherry red spot	☐ Dicarboxylic aciduria	☐ Other:
Tortuosity	Coloboma	☐ Hyperammonemia	Structural brain abnormalities
☐ Truncus arteriosus	☐ Corneal abnormality	Hyperglycemia	☐ Aplasia/hypoplasia of the
☐ Ventricular abnormality	☐ Ectopia lentis	☐ Hypoglycemia	cerebellar vermis
☐ Ventricular septal defect	☐ Microphthalmia	Hypoammonemia	☐ Aplasia/hypoplasia of the
Other:	☐ Nystagmus	☐ Ketosis	cerebellum
Craniofacial	☐ Optic Atrophy	Organic aciduria	☐ Basal ganglia abnormality
☐ Bifid uvula	☐ Ptosis	☐ Plasma carnitine (abnormal)	☐ Brain atrophy
☐ Cleft lip	☐ Retinitis pigmentosa	☐ Serum creatine (abnormal)	☐ Brainstem abnormality
·	☐ Strabismus	☐ Serum pyruvate (abnormal)	☐ Cerebral dysmyelination
Cleft palate	☐ Visual impairment	Other:	□ Cerebral hypomyelination
Craniosynostosis	□ Blind	Musculoskeletal	☐ Cerebral white matter abnormality
☐ Epicanthal folds	☐ Cortical	☐ Contractures	Corpus callosum abnormality
☐ Hypertelorism	☐ Myopia	☐ Fractures	Cortical dysplasia
☐ Hypotelorism	☐ Other:	☐ Hemihypertrophy	☐ Encephalocele
☐ Macrocephaly	Gastrointestinal	☐ Hyperlordosis	Holoprosencephaly
Microcephaly	☐ Anal malformation	☐ Hypermobility	Hydrocephalus
☐ Micrognathia	☐ Constipation (chronic)	☐ Hypertonia	Leukoencephalopathy
☐ Nose abnormality	☐ Crohn's disease	☐ Hypotonia	Leukodystrophy
☐ Palpebral fissures	☐ Diarrhea (chronic)	☐ Kyphosis	Lissencephaly
☐ Philtrum abnormality	☐ Esophageal atresia	☐ Muscle atrophy	☐ Neuronal migration abnormality
	☐ Gastroesophageal reflux	☐ Muscular dystrophy	☐ Pachgyria
☐ Teeth abnormality	☐ Gastroesophiagearrenax	☐ Myopathy	☐ Polymicrogyria
☐ Tongue abnormality	☐ Hepatic failure	☐ Myotonia	<ul><li>☐ Ventriculomegaly</li><li>☐ Other:</li></ul>
☐ Other:	☐ Hepatic failure ☐ Hepatomegaly	Oligodactyly	Other
Cognitive development	☐ Hirschsprung disease	Overgrowth	☐ Allergies (severe)
☐ ADD/ADHD		Polydactyly	Fever (episodic)
☐ Autism spectrum disorder	☐ Inflammatory bowel disease	☐ Rib defects	☐ Failure to thrive
☐ Developmental delay	☐ Intestinal pseudo-obstruction	Scoliosis	☐ Heterotaxy
☐ Developmental regression	☐ Pancreatitis	☐ Short stature	Lethargy
☐ Intellectual disability	☐ Pyloric stenosis	Skeletal dysplasia	☐ Organomegaly
☐ Mild	☐ Splenomegaly	☐ Spina bifida	☐ Pain (chronic)
☐ Moderate	☐ Vomiting (episodic/cyclic)	Syndactyly	Other:
<del></del>	☐ Tracheoesophageal fistula	☐ Talipes equinovarus	
Severe	☐ Other:	☐ Tall stature	Attach any imaging or laboratory
☐ Profound	Genitourinary	☐ Vertebral anomalies	results, including but not limited to:
☐ Motor milestones delayed	☐ Ambiguous genitalia	Other:	prenatal testing, biopsies,
☐ Speech delay	☐ Cryptorchidism	Neurologic  ☐ Areflexia	EMG, EEG, imaging, metabolic,
☐ Other:	☐ Hypogonadism	☐ Ataxia	, , , , , , , , , , , , , , , , , , , ,
Ear & hearing	☐ Hypospadias	☐ Chorea	molecular, etc.
☐ Deafness	☐ Kidney abnormality	☐ Dystonia	Overtions 2 To an act with a
☐ Acquired	☐ Agenesis	☐ Epileptic encephalopathy	Questions? To speak with a
☐ Congenital	☐ Horseshoe	☐ Neuropathy	Genomic Science Specialist, please
☐ Bilateral	☐ Partially duplicated	☐ Neuropatny ☐ Seizures	call: 1.800.394.4493 ext. 2031
<del></del>	Polycystic	☐ Seizures ☐ Absence	
☐ Unilateral	☐ Ovarian streak	☐ Absence	
☐ Conductive	Polycystic ovarian syndrome	☐ Febrile	
Sensorineural	☐ Testicular abnormality	☐ Generalized clonic	
☐ Low-set ears	☐ Ureter abnormality	☐ Generalized clothic ☐ Generalized myoclonic	
☐ Pinna abnormality	☐ Urethra abnormality	☐ Generalized triyocionic	
☐ Preauricular pit	Other:	Generalized tonic	
☐ Preauricular skin tag	Hair & skin	☐ Infantile spasms	
Other:	☐ Albinism	☐ Spasticity	
	☐ Rlistering	Other:	

Any additional comments?					
Form Submission Instructions					
When the form is complete please include a printed copy with the samples.					
Ordering Checklist					
☐ Completed requisition form					
☐ Clinical history completed					
$\square$ Any clinical notes should be sent with the specimen(s)					
☐ Patient sample either blood spot card or whole blood in a lavender top tube					
☐ Family member(s) samples either blood spot cards, whole blood in a lavender top tube, or saliva					
☐ Family member(s) name, date of birth, and relationship					

NOTE: Specimen(s) must be labeled with two of the following forms of identification: name, date of birth, last four digits of SS#, patient ID #. These same two forms of ID must be indicated on the test requisition.

Athena Diagnostics Client Service Representatives are available from 8:30AM to 9:00PM Eastern Time (US). Customers in the US and Canada please call toll-free

1.800.394.4493

(Non-US customers please call 508.756.2886 or fax 610.271.6085.)



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