



*Information required for testing

Patient Information

LAST NAME*		FIRST NAME*		MI	DOB* MM/DD/YYYY	SEX	GENDER
ADDRESS		CITY	STATE	ZIPCODE	PHONE NUMBER	EMAIL ADDRESS	

Billing Information (Please include a copy of insurance card(s) for billing purposes.)

*☐ CLIENT BILL ☐ INSURANCE ☐ SELF PAY ☐ MEDICARE/MEDICAID (☐ PRIMARY ☐ SECONDARY) RELATIONSHIP: ☐ SELF ☐ SPOUSE ☐ DEPENDENT

INSURANCE NAME	MEMBER/POLICY ID	GROUP #
POLICY HOLDER NAME	POLICY HOLDER DOB MM/DD/YYYY	TEST INDICATION/ICD-10 CODE(S)*

Account Information

FACILITY/PRACTICE NAME*	PHONE NUMBER	FAX NUMBER	ORDERING PHYSICIAN NAME*
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Specimen Information PREFERRED SPECIMEN IS BUCCAL SWAB

☐ PERIPHERAL BLOOD ☐ BUCCAL SWAB COLLECTION DATE: MM/DD/YYYY COLLECTION TIME: 00:00 AM/PM

Test(s) Requested*

☐ MICROARRAY ☐ FRAGILE X SYNDROME (FXS) ☐ CYSTIC FIBROSIS (CF) ☐ SPINAL MUSCULAR ATROPHY (SMA) ☐ DUCHENNE MUSCULAR DYSTROPHY (DMD)

FISH Studies FISH STUDIES WILL NOT BE RUN UNLESS BOX IS CHECKED AND OPTIONS BELOW HAVE BEEN CHOSEN

FISH FOR ANEUPLOIDY	FISH FOR SEX CHROMOSOME ABNORMALITIES:	FISH FOR MICRODELETION SYNDROMES	
<input type="checkbox"/> AneuVysion) X/Y/18/13/21 <input type="checkbox"/> X/Y/18 ONLY <input type="checkbox"/> 13/21 ONLY <input type="checkbox"/> Trisomy 21 - Down Syndrome <input type="checkbox"/> Trisomy 18 - Edwards Syndrome <input type="checkbox"/> Trisomy 13 - Patau Syndrome	<input type="checkbox"/> Sex Determination (X/SRY) <input type="checkbox"/> Turner Syndrome (CEPX/CEPY) <input type="checkbox"/> Klinefelter Syndrome (CEPX/CEPY)	<input type="checkbox"/> Angelman syndrome (15q11-13) <input type="checkbox"/> Cri du Chat-syndrome (5p15.2) <input type="checkbox"/> DiGeorge/VCFS/CATCH22 (22q11.2) <input type="checkbox"/> Kallmann syndrome (Xp22.3) <input type="checkbox"/> Miller-Dieker syndrome (17p13.3) <input type="checkbox"/> Prader-Willi syndrome (15q11-13) <input type="checkbox"/> Smith-Magenis syndrome (17p 11.2)	<input type="checkbox"/> Soto syndrome (5q35) <input type="checkbox"/> SRY (Yp11.3) <input type="checkbox"/> Steroid Sulfatase (STS) (Xp22.3) <input type="checkbox"/> Wolf-Hirschhorn syndrome (4p16.3) <input type="checkbox"/> Williams Beuren syndrome (7q11.23) <input type="checkbox"/> XIST (Xq13.2) <input type="checkbox"/> Other _____

ICD-10 Code(s)*

<input type="checkbox"/> F70	Mild intellectual disabilities	<input type="checkbox"/> R47.89	Other speech disturbances
<input type="checkbox"/> F71	Moderate intellectual disabilities	<input type="checkbox"/> R62.0	Delayed milestone in childhood
<input type="checkbox"/> F72	Severe intellectual disabilities	<input type="checkbox"/> R62.5	Other unspecified lack of expected normal physiological development in childhood
<input type="checkbox"/> F73	Profound intellectual disabilities	<input type="checkbox"/> R62.59	Other lack of expected normal physiological development in childhood
<input type="checkbox"/> F81.81	Disorder of written expression	<input type="checkbox"/> Q18.9	Congenital malformation of face and neck, unspecified
<input type="checkbox"/> F81.89	Other developmental disorders of scholastic skills	<input type="checkbox"/> Q79.9	Congenital malformation of musculoskeletal system, unspecified
<input type="checkbox"/> F81.9	Developmental disorder of scholastic skills, unspecified	<input type="checkbox"/> Q89.7	Multiple anomalies
<input type="checkbox"/> F82	Specific developmental disorder of motor function	<input type="checkbox"/> Q89.8	Other specified congenital malformations
<input type="checkbox"/> F79	Unspecified intellectual disabilities	<input type="checkbox"/> Q89.9	Congenital malformation, unspecified
<input type="checkbox"/> F84.0	Autism spectrum disorder	<input type="checkbox"/> Z84.3	Consanguinity
<input type="checkbox"/> F88	Developmental delay	<input type="checkbox"/> Z84.89	Family history of disease or disorders
<input type="checkbox"/> F89	Unspecified Disorder of Development	<input type="checkbox"/> Other	_____
<input type="checkbox"/> K55.30	Necrotizing enterocolitis, unspecified		

Patient Authorization and Consent

I understand that I am responsible for providing accurate information about my insurance to Genesys Diagnostics Inc. I understand that Genesys Diagnostics Inc. will be providing testing service and billing my insurance. However, I understand that charges that are not covered by my insurance, including any applicable co-payments and deductibles are my responsibility and I agree to pay such charges promptly.

Patient/Guardian Signature: _____ Date: _____

☐ I do not consent to having my deidentified DNA sample used for internal research purposes.

Healthcare Provider Authorization

I certify that (i) this test is medically necessary, (ii) the patient (or authorized representative on the patient's behalf) has given informed consent (which includes written informed consent or written authorization when required by law) to have this testing performed, and (iii) the informed consent obtained from the patient meets the requirements of applicable law. I agree to provide Genesys, or its designee, any and all additional information reasonably required for this testing to be performed.

Healthcare Provider Signature: _____ Date: _____

Medical Necessity Statement: Tests ordered on Medicare patients must follow CMS rules regarding medical necessity and FDA approval guidelines and must include diagnosis, symptoms and reason for testing as indicated in the medical record. If testing does not come under Medicare guidelines for payment a 'signed' Advanced Beneficiary Notice must be included.



Developmental Delay Microarray



Requisition Form