

Learning Disability Read Codes

Version 4

Produced by the Strategic Health Facilitation Team

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The learning disability read codes list is a document consisting of read codes associated with specific syndromes linked with diagnoses of a learning disability.

This was established to support a development of a learning disability Miquet Query which was to be used in General Practice to identify those patients with a learning disability that will need to be offered an annual health check.

(The Learning Disability Miquet Query is an excel workbook within which a range of diagnostic codes, used by the Clinical Terminology Browser, is used to identify patients to whom these codes are assigned.)

This list could be used without the Miquet Query to conduct a manual search on patients to whom these codes are assigned to, if required.

This list has not been designed to be a diagnostic tool.

Please be mindful of circumstances where a syndrome code has been allocated to a parent and not their child which is an error.

Once patients with a learning disability have been identified, they would ideally be allocated an **administrative code** (used by EMIS or SystemOne) and a **diagnostic code** when possible to support their diagnosis. (This could be a syndrome code):

EMIS WEB – QOF Learning Disability Register Read Code: 918e

SYSTEMOne – QOF Learning Disability Register Read Code: XaKYb

| | Version 2 (5 byte) | Version 3 |
|---|--------------------|----------------------------|
| On learning disability register | 918e. | XaKYb |
| Mental retardation | E3 % | E3 % |
| [x]Mental retardation | Eu7 % | E3 % |
| [x]Developmental disorder of scholastic skills unspecified Learning Disability NOS | Eu81z | Eu81z |
| [x]Mild Learning Disability | Eu816 | XaREt |
| [x]Moderate Learning Disability | Eu814 | XaQZ3 |
| [x]Severe Learning Disability | Eu815 | XaQZ4 |
| [x]Profound Learning Disability | Eu817 | XaREu |
| Learning Disabilities | Eu8 % | XaQX4, XaREu, XaREt, XaQZ3 |
| Specific learning disability | Eu818 | E3 % |

Every effort has been made to ensure that this information is correct at the time of the review. Please note that mentioned syndromes will affect ones cognitive ability in different ways and to different extent.

WILL syndrome (WILL indicate a learning disability) – includes a list of syndromes that would indicate high probability of person having a learning disability.

MAY syndrome (MAY indicate a learning disability) – includes a list of syndromes that would indicate that a proportion of people with such syndrome may have a learning disability.

| WILL indicate a learning disability. | Version 2 | Version 3 | Comments |
|--|-----------------|-----------|---|
| Angelman's syndrome | PKyz7/ Pkyz7 | PKyz5 | |
| Coffin lowry syndrome | PKy5F | Xa0Zc | |
| Cornelia de Lange syndrome | PKy60 | PKy60 | |
| Cohen syndrome | PKy5K | XaZWI | Very rare, approx. 1000 people diagnosed |
| Cri du chat syndrome | PJ31. | PJ31. | |
| Down's syndrome | PJ0 % | X78EI | People with Down's Syndrome will typically have a learning disability but there may be rare exceptions. |
| <ul style="list-style-type: none"> • Trisomy 21- meiotic nondisjunction | PJ00. | PJ00. | |
| <ul style="list-style-type: none"> • Trisomy 21- mitotic nondisjunction mosaicism | PJ01. | PJ01. | 2 in 100 people with Down's Syndrome will have mosaic Down's Syndrome. |
| <ul style="list-style-type: none"> • Down's syndrome NOS | PJ0z. | X78Ek | |
| <ul style="list-style-type: none"> • Down's syndrome - trisomy 21 | | | |
| <ul style="list-style-type: none"> • Trisomy 21, translocation | PJ02. | | |
| <ul style="list-style-type: none"> • Mongolism | PJ0.. | / | |
| Dubowitz syndrome | PKy66 | PKy66 | |
| Edwards syndrome | PJ2.. | PJ2.. | |
| <ul style="list-style-type: none"> • Edward's syndrome NOS | PJ2z. | X78Em | |
| Hurler's syndrome | C3751 | X40Vy | |
| Laurence-Moon syndrome | PKy1. | X00e1 | |
| Mowat-Wilson syndrome | PJ9.. | XaNWy | |
| Patau's syndrome | PJ1.. | PJ1.. | |
| <ul style="list-style-type: none"> • Pataus Syndrome NOS | PJ1z. | X78Eo | |
| <ul style="list-style-type: none"> • Partial trisomy 13 in Patau's | | X78Ep | |

| | | | |
|----------------------------|-------|-------|--|
| syndrome | | | |
| Rett syndrome | Eu842 | X005S | |
| Smith-Magenis syndrome | PJ333 | XalwZ | affects an estimated 1 in 25,000 individuals |
| Wolff - Hirschorn syndrome | PJ32. | | |

| MAY indicate a learning disability. | Version 2 | Version 3 | Comments |
|---|------------------|------------------|---|
| Acrodysostosis | / | X78Ak | |
| Aircardi syndrome | Xa0OH | P2283 | |
| Aicardi Goutieres syndrome | F1306 | X004C | |
| Alpha thalassaemia-mental retardation syndrome | | Xa0Yy | People can be a carrier. |
| Anterior chamber cleavage syndrome | | XaEvv | |
| <ul style="list-style-type: none"> Peters-plus syndrome | P3423 | | |
| Apert syndrome | / | XE1Lu | |
| Autism | E140% | E1401 % | Some people may have Autism and a learning disability; some may only have Autism. |
| <ul style="list-style-type: none"> Kanners syndrome | | XE2v2 | |
| <ul style="list-style-type: none"> [X]Kanner's syndrome | Eu840 | | |
| <ul style="list-style-type: none"> Childhood autism | Eu840 | XE2v2 | |
| <ul style="list-style-type: none"> Atypical autism | Eu841 | X00TN | |
| <ul style="list-style-type: none"> Autistic spectrum disorder | Eu84z | X00TM | |
| <ul style="list-style-type: none"> Autistic spectrum disorder with isolated skills | / | Ub1Ts | |
| Bannayan-Riley-Ruvalcaba syndrome | PKy00 | X207k | |
| Cerebral palsy | | XE2Q8 | Learning Disability will be |
| <ul style="list-style-type: none"> Double athetosis | | XE15M | |

| | | | |
|---|-------|-------|---|
| | | | present in approximately 50% of people with Cerebral Palsy. |
| • Spastic cerebral palsy | | X00En | |
| • Dyskinetic cerebral palsy | | X00Eu | |
| • Ataxic cerebral palsy | | Xa0IM | |
| • Hypotonic cerebral palsy | | F23y1 | |
| • Other congenital cerebral palsy | | F23y. | |
| • Congenital cerebral palsy NOS | F23z. | F23z. | |
| • Congenital cerebral palsy | F23 | XM1Pu | |
| • Cerebral palsy, not congenital or infantile, acute | G669. | XaBE2 | |
| • Athetoid cerebral palsy | F1370 | | |
| • Congenital diplegia | F230. | | |
| • Congenital paraplegia | F2300 | | |
| • Cerebral palsy with spastic diplegia | F2301 | | |
| • Congenital diplegia NOS | F230z | | |
| • Congenital hemiplegia | F231. | | |
| • Congenital quadriplegia | F232. | | |
| Coffin-Siris syndrome | | X50HX | |
| Congenital megalocornea | P3221 | XE1Jt | |
| Foetal alcohol syndrome | PK80. | PK80. | |
| Fragile X syndrome | PJyy4 | X78FB | |
| • Family history of fragile X syndrome in first degree relative | XaX8x | 12J80 | |
| Galactosaemia | C311. | C311. | |
| Jacobsen syndrome | PJ334 | XSDcM | |
| Johanson blizzard syndrome | PKyM. | X50Kj | |

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|---|-------|-------|--|
| Klinefelter syndrome | PJ7.. | PJ7.. | |
| <ul style="list-style-type: none"> • Klinefelter's syndrome NOS | | PJ7z. | |
| <ul style="list-style-type: none"> • XXY Klinefelter's syndrome | | PJ70. | |
| <ul style="list-style-type: none"> • Klinefelter's syndrome XXY | | XM1MJ | |
| <ul style="list-style-type: none"> • Klinefelter's syndrome XXXY | | XM1MK | |
| <ul style="list-style-type: none"> • Klinefelter's syndrome, XYY | PJ73. | PJ73. | |
| <ul style="list-style-type: none"> • Klinefelter's syndrome NOS | PJ7z. | PJ7z. | |
| Menkes syndrome | | PKy92 | |
| <ul style="list-style-type: none"> • Kinky hair syndrome | PKy92 | | |
| Noonan Syndrome | PKy80 | PKy80 | |
| Phenylketonuria | C301. | C301. | |
| <ul style="list-style-type: none"> • Classical phenylketonuria | | Xa0IA | |
| Prader-Willi syndrome | PKy93 | PKy93 | |
| Trisomy 6 | | PJ500 | |
| Tuberous sclerosis | PK5.. | PK5.. | |
| Williams syndrome | PKy4. | PKy4. | |