

GETDISEASEBYCLINICALSIGN
Abnormal colour of the urine...
10768
56425
Cold agglutinin disease
38720
Abnormal colour of the urine...
652
Occasional

GETDISEASECLINICALSIGNSNOLANG
6
3297
6
Isolated...
33100
Respiratory distress...
652
Occasional

OrphaData

MEDDRA
C0000727
LT
10000647
Acute abdomen

MEDDRA_ALL_INDICATIONS
CID1000000085
C0015544
text.mention
Failure to Thrive
LLT
C0015544
Failure to Thrive

MEDDRA_ALL_SE
CID1000000085
CID000010917
C0000729
LLT
C0000729
Abdominal cramps

MEDDRA_FREQ
CID1000000085
CID000010917
C0000737
21%
0.21
0.21
LLT
C0000737
Abdominal pain

Sider

OMIM
INHERITANCE:
- Autosomal recessive
HEAD AND NECK:
- [Eyes];
- Impaired pupillary...
- [Mouth];
- Dry mouth

OMIM_ONTO
http://...
BBS4 Gene
BBS4
?
false
C1412749
http://...

OMIM

STITCH
CIDm00028871
CIDs00028871
ATC
M01AB10

STITCH

ATC
A01AA01
Sodium fluoride

ATC

PHENOTYPE_ANNOTATION
DECIPHER
1
Wolf-Hirschhorn Syndrome
None
HP:0000252
DECIPHER:1
IEA
None
None
None
O
WOLF-HIRSCHHORN SYNDROME
2013.05.29
HPO:skoehler

HPO
id:HP:0000020
name: Urinary incontinence
alt_id: HP:0006942
alt_id: HP:0008681
def: "Loss of the ability..." [HPO:sdoelken, pmid:12559262]
comment: ...
synonym: "Bladder incontinence" EXACT []
synonym: "Loss of bladder control" EXACT layperson [...]
xref: MSH:D014549
xref: SNOMEDCT_US:165232002
xref: UMLS:C0042024
is_a: HP:0000009 ! Functional abnormality of the bladder

HPO