

Part Three

Indices

Disorders Index

Disorders	Disorder No.
A	
Abetalipoproteinemia	28.9
Aceruloplasminemia	33.6
α -N-Acetylgalactosamine 4-sulfatase deficiency	18.9
Acetyl CoA: α -glucosaminide N-acetyltransferase deficiency	18.5
Acid maltase deficiency	15.9
Aconitase deficiency	27.8
Acrodermatitis enteropathica	33.4
Acute intermittent porphyria	31.2
α -N-Acetyl-galactosaminidase deficiency	19.7
Acyl-CoA oxidase deficiency	25.9
ADA deficiency	23.1
Addison disease and cerebral sclerosis	25.8
Adenine phosphoribosyltransferase deficiency	23.5
Adenosine deaminase deficiency	23.1
Adenylosuccinate lyase deficiency	23.6
Adrenal hyperplasia I	29.1
Adrenal hyperplasia III	29.4
Adrenal hyperplasia IV	29.5
Adrenal hyperplasia V	29.10
AGAT deficiency	24.2
ALA-dehydratase deficiency	31.1
Alanine:glyoxylate aminotransferase deficiency	26.1
Aldosterone synthase deficiency	29.6
Alipoprotein E deficiency	28.3
Alkaptonuria	4.5
Alkyldihydroxyacetonephosphate synthase deficiency	25.7
2-Aminoadipate aminotransferase deficiency	12.3
2-Aminoadipic semialdehyde synthetase deficiency	12.2
δ -Aminolevulinic acid dehydratase deficiency	31.1
2-Amino-/2-oxoadipic aciduria	12.3
Amylo-1,6-glucosidase deficiency	15.10
Amylopectinosis	15.11
Andersen disease	15.11
Androgen Insensitivity syndrome	29.14
Apolipoprotein A-I deficiency	28.7
Apolipoprotein B-100 deficiency	28.6
Apolipoprotein C-II deficiency	28.2

Disorders	Disorder No.
Apparent cortisone reductase deficiency	29.9
Apparent mineralocorticoid excess	29.8
Arginase 1 deficiency	11.5
Argininemia	11.5
Argininosuccinate lyase deficiency	11.4
Argininosuccinate synthase deficiency	11.3
Arginosuccinic aciduria	11.4
Aromatase deficiency	29.13
Aromatic L-amino acid decarboxylase deficiency	2.2
Arylsulfatase A deficiency	22.3
Aspartoacylase deficiency	8.6
Aspartylglucosaminuria	19.6
ATP synthase, ATPase defect	27.17
ATP/ADP translocator defect	27.19
B	
Barth syndrome	6.5
Batten disease	22.4.3
Biotinidase deficiency	7.1
Branched-chain α -dehydrogenase deficiency	6.1
Brancher enzyme deficiency	15.11
Bronze Schidler syndrome	25.8
C	
Canavan disease	8.6
Carbamylphosphate synthase deficiency	11.1
Carnitine palmityltransferase 1 deficiency	14.2
Carnitine palmityltransferase 2 deficiency	14.4
Carnitine uptake defect	14.1
Carnitine-acylcarnitine carrier defect	14.3
Carnitine-acylcarnitine translocase deficiency	14.3
Cathepsin A deficiency	19.5
CblA deficiency	7.8
CblB deficiency	7.8
CblC deficiency	7.9
CblD deficiency	7.10
CDG-Ia	20.1
CDG-Ib	20.2
CDG-Ic	20.3
CDG-Id	20.4
CDG-Ie	20.5
CDG-IIa	20.6
CDG-IIb	20.7
Ceramidase deficiency	22.2
Cerebrohepatorenal syndrome	25.1
Cerebrotendinous xanthomatosis	32.3
Ceruloplasmin (ferroxidase) deficiency	33.6
Cherry red spot and myoclonus syndrome	19.4
CHILD syndrome	30.2
Chondrodysplasia punctata	25.5

Disorders	Disorder No.
Citrin deficiency	11.11
Citrullinemia I	11.3
Citrullinemia type II	11.11
Classical phenylketonuria	1.1
CLN1	22.4.1
CLN2	22.4.2
CLN3	22.4.3
Coenzyme Q deficiency	27.14
Complex I deficiency	27.13
Complex II deficiency	27.11
Complex III deficiency	27.15
Complex IV deficiency	27.16
Complex V deficiency	27.17
Congenital adrenal hyperplasia	29.2–5
Congenital erythropoietic porphyria	31.3
Congenital glucose/galactose malabsorption	16.1
Conradi-Hünemann syndrome	30.3
Cori disease	15.10
Corticosterone methyl oxidase II deficiency	29.6
Costeff optic atrophy syndrome	6.6
Coupling state defect	27.18
Creatine deficiency	24
Creatine transporter deficiency	24.3
Cu-binding P-type ATPase deficiency	33.1
γ -Cystathionase deficiency	10.3
Cystathionine gamma-lyase deficiency	10.3
Cystathionine β -synthase deficiency	10.2
Cystathioninuria	10.3
Cysteinyl leukotrienes (LTC) ₄ -synthase deficiency	34.1
Cysteinyl-glycinase deficiency	34.3
Cystinosis	21.1
Cystinuria	13.1
Cytochrome bc ₁ deficiency	27.15
Cytochrome c oxidase deficiency	27.16
D	
D-2-Hydroxyglutaric aciduria	8.5
D-Bifunctional enzyme deficiency	25.10
Desmosterolosis	30.4
D-Glycerate kinase deficiency	15.7
D-Glyceric acidemia	15.7
Dibasicaminoaciduria	11.7
Dicarboxylic aminoaciduria	13.2
Dihydrolipoyl transacetylase deficiency	27.3
Dihydropteridine reductase deficiency	1.4
Dihydropyrimidinase deficiency	23.14
Dihydropyrimidine dehydrogenase deficiency	23.13
Dihydroxyacetonephosphate acyltransferase deficiency	25.6
Dihydroxyadenine urolithiasis	23.5
Dimethylglycinuria	35.2

Disorders	Disorder No.
Dolicholphosphate-mannose synthase-1 deficiency	20.5
Dopamine β -hydroxylase deficiency	2.3
Dopa-responsive dystonia	1.6
Doss porphyria	31.1
E	
Erythropoietic protoporphyria	31.7
Estrogen resistance	29.15
F	
Fabry disease	22.1
Familial dysbetalipoproteinemia	28.3
Familial hypercholesterolemia	28.5
Fanconi-Bickel syndrome	15.17/16.4
Farber disease	22.2
Fish odor syndrome	35.1
Fish-eye disease	28.2.2
Folic acid transport defect	10.9
Forbe disease	15.10
Formiminoglutamic aciduria	5.3
Formiminotransferase deficiency	5.3
Fructose-1,6-diphosphatase deficiency	15.5
Fructose-1-phosphat aldolase deficiency	15.4
Fucosidosis	19.3
Fumarase deficiency	8.2/27.12
Fumaric aciduria	8.2
Fumarylacetoacetase deficiency	4.1
G	
GABA transaminase deficiency	3.1
Galactokinase deficiency	15.1
Galactose-1-phosphatase uridyltransferase deficiency	15.2
Galactosemia	15.2
Galactosialidosis	19.5
β -Galactosidase deficiency	19.8
β -D-Galactosidase A deficiency	18.8
α -Galactosidase A deficiency	22.1
β -Galactocerebrosidase deficiency	19.16
GAMT deficiency	24.1
Gaucher disease	19.13
Gephyrin deficiency	10.5.3/23.3
Glucocorticoid suppressible hyperaldosteronism	29.7
β -Glucocerebrosidase deficiency	19.13
β -D-Glucuronidase deficiency	18.10
Glucose transporter 1 defect	16.3
Glucose transporter 2 defect	15.17/16.4
Glucose transporter protein syndrome	16.3
Glucose-6-phosphatase deficiency	15.8
Glucose-6-phosphate translocase deficiency	15.8a
Glucosidase I deficiency	20.7

Disorders	Disorder No.
Glucosyltransferase I deficiency	20.3
Glutamate dehydrogenase 1 abnormality	11.10
γ -Glutamylcysteine synthetase deficiency	9.1
γ -Glutamyl transpeptidase deficiency	34.2
γ -Glutamyl transpeptidase deficiency	9.3/34.2
Glutaric aciduria I	12.7
Glutaryl-CoA dehydrogenase deficiency	12.7
Glutathione synthetase deficiency	9.2
Glutathionuria	9.3/34.2
Glycerol kinase deficiency	17.1
Glycine cleavage system deficiency	3.2
Glycine receptor defect	2.5
Glycogen storage disease type 0	15.16
Glycogen storage disease type 1	15.8
Glycogen storage disease type 2	15.9
Glycogen storage disease type 3	15.10
Glycogen storage disease type 4	15.11
Glycogen storage disease type 5	15.12
Glycogen storage disease type 6	15.13
Glycogen storage disease type 7	15.14
Glycogen storage disease type 9	15.15
Glycogen synthetase deficiency	15.16
Glycogenosis VIIa	15.15
Glycosylasparaginase deficiency	19.6
Glyoxylate reductase deficiency	26.2
GM1 gangliosidosis	19.8
GM2 gangliosidosis	19.9
Goldberg syndrome	19.5
GTP cyclohydrolase I deficiency	1.2/1.6
Guanidinoacetate methyltransferase deficiency	24.1
Günther's disease	31.3
H	
Hartnup disorder	13.3
Hawkinsinuria	4.4
Hemochromatosis	33.5
Heparin sulfamidase deficiency	18.3
Hepatic lipase deficiency	28.4
Hepatoerythropoietic porphyria	31.8
Hepatoerythropoietic protoporphyria	31.7
Hereditary coproporphyria	31.5
Hereditary folate malabsorption	10.9
Hereditary fructose intolerance	15.4
Hereditary progressive dystonia	1.6
Hers disease	15.13
β -Hexosaminidase A deficiency	19.9
Histidine ammonia-lyase deficiency	5.1
Histidinemia	5.1
Holocarboxylase synthetase deficiency	7.2
Homocystinuria	10.2/10.6/10.7

Disorders	Disorder No.
Homogentisate dioxygenase deficiency	4.5
Homozygous PCT	31.8
Hunter syndrome	18.2
Hurler syndrome	18.1a
Hurler-Scheie syndrome	18.1c
Hydroxykynureninuria	12.5
3-Hydroxyisobutyric aciduria	7.6
3-Hydroxyisobutyryl deacylase deficiency	7.11
17 α -Hydroxylase deficiency	29.2
21-Hydroxylase deficiency	29.4
11 β -Hydroxylase I/II deficiency	29.7
11 β -Hydroxylase type I deficiency	29.5
3-Hydroxy-3-methylglutaric aciduria	6.8
3-Hydroxy-3-methylglutaryl-CoA lyase deficiency	6.8
3-Hydroxy-3-methylglutaryl-CoA synthase deficiency	14.12
Hydroxylysine kinase deficiency	12.6
Hydroxylysinuria	12.6
Hydroxymethylbilane synthase deficiency	31.2
4-Hydroxyphenylpyruvate dioxygenase deficiency	4.3
Hydroxyproline oxidase deficiency	3.9
3 β -Hydroxy- Δ^5 -C ₂₇ -steroid dehydrogenase deficiency	32.1
11b-Hydroxysteroid dehydrogenase type I deficiency	29.9
3 β -Hydroxysteroid dehydrogenase type II deficiency	29.3
11 β -Hydroxysteroid dehydrogenase type II deficiency	29.8
17 β -Hydroxysteroid dehydrogenase type III deficiency	29.11
3 β -Hydroxysteroid dehydrogenase deficiency	30.2
3 β -Hydroxysteroid- Δ^8 , Δ^7 -isomerase deficiency	30.3
3 β -Hydroxysteroid- Δ^7 -reductase deficiency	30.5
3 β -Hydroxysteroid- Δ^{24} -reductase deficiency	30.4
Hyper IgD syndrome	30.1
Hypercholesterolemia	28.5/6
Hyperkplexia	2.5
Hyperlysinemia I	12.2.1
Hyperlysinemia II	12.2.2
Hypermethioninemia	10.1
Hyperornithinemia and gyrate atrophy (HOGA)	12.1
Hyperornithinemia-hyperammonemia-homocitrullinuria syndrome	11.8
Hyperpipecolic acidemia	25.4
Hyperprolinemia type 1	3.6
Hyperprolinemia type 2	3.7
Hypertriglyceridemia	28.1/2
Hypoapolipoproteinemia	28.7/8
Hypobetalipoproteinemia	28.10
Hypolipoproteinemia	28.9/10
Hypophosphatasia	35.3
Hypoxanthine phosphoribosyltransferase deficiency	23.4
I	
I-cell disease	19.10
Iduronate 2-sulfatase deficiency	18.2

Disorders	Disorder No.
<i>α</i> -L-Iduronidase deficiency	18.1
Iminoglycinuria	3.11
Infantile neuronal ceroid lipofuscinosis	22.4.1
Infantile Refsum disease	25.3
Intermittent acute porphyria	31.2
Isovaleric acidemia	6.2
Isovaleryl-CoA dehydrogenase deficiency	6.2
J	
Jaeken syndrome	20.1
Janky-Bielschowsky disease	22.4.2
Juvenile neuronal ceroid lipofuscinosis	22.4.3
K	
Kanzaki disease	19.7.2
Kelly-Seegmiller syndrome	23.4
2-Ketoglutarate dehydrogenase complex deficiency	8.1
Krabbe disease	19.16
KSS	27
Kynureninase deficiency	12.5
L	
L-2-Hydroxyglutaric aciduria	8.4
L-Arginine:glycine amidinotransferase deficiency	24.2
Late infantile neuronal ceroid lipofuscinosis	22.4.2
LCAT deficiency	28.8
Lecithin:cholesterol acyltransferase deficiency	28.8
Leigh syndrome	27
Lesch-Nyhan syndrome	23.4
LHON	27
Lipoamide dehydrogenase deficiency	27.4
Lipoid adrenal hyperplasia	29.1
Lipoprotein lipase deficiency	28.1
Liver phosphorylase deficiency	15.13
Liver phosphorylase kinase deficiency	15.15
Long QT syndrome	27
Long-chain 3-hydroxy-acyl-CoA dehydrogenase- α/β deficiency	14.8
Long-chain fatty acid transport defect	35.4
Low density lipoprotein receptor defect	28.5
17,20-Lyase deficiency	29.10
Lysinuric protein intolerance	11.7
Lysosomal membrane cystine transporter defect	21.1
Lysosomal α -glucosidase deficiency	15.9
Lysyl oxidase deficiency	33.3
M	
Malate/aspartate shuttle defect	27.20
Malonic aciduria	8.3
Malonyl-CoA decarboxylase deficiency	8.3
α -Mannosidosis	19.1

Disorders	Disorder No.
β -Mannosidosis	19.2
Mannosyltransferase VI deficiency	20.4
Maple syrup disease	6.1
Maroteaux-Lamy syndrome	18.9
McArdle disease	15.12
Medium-chain acyl-CoA dehydrogenase deficiency	14.6
MELAS	27
Membrane-bound dipeptidase (cysteinyl-glycinase) deficiency	34.3
Menkes disease	33.2
MERRF	27
Metachromatic leukodystrophy	22.3
Methionine adenosyltransferase deficiency	10.1
Methionine synthase deficiency	10.7.1
Methionine synthase deficiency (incl. methylmalonyl CoA mutase def.)	7.9/10.8
Methionine synthase reductase deficiency	10.7.2
α -Methylacyl-CoA racemase deficiency	25.12
2-Methylbutyryl-CoA dehydrogenase deficiency	7.12
3-Methylcrotonyl-CoA carboxylase deficiency	6.3
Methylenetetrahydrofolate reductase deficiency	10.6
3-Methylglutaconic aciduria type I	6.4
3-Methylglutaconic aciduria type II	6.5
3-Methylglutaconic aciduria type III	6.6
3-Methylglutaconic aciduria type IV (idiopathic)	6.7
3-Methylglutaconyl-CoA hydratase deficiency	6.4
2-Methyl-3-hydroxybutyric aciduria	7.7
2-Methyl-3-hydroxybutyryl CoA dehydrogenase deficiency	7.7
Methylmalonic acidemia	7.8
Methylmalonic acidemia and homocystinuria	7.9/7.10
Methylmalonyl CoA mutase deficiency (incl. methionine synthase def.)	7.9/10.8
Methylmalonyl semialdehyde dehydrogenase deficiency	7.5
Methylmalonyl-CoA mutase deficiency	7.8
Mevalonate kinase deficiency	30.1
Mevalonic aciduria	30.1
Mitochondrial trifunctional protein deficiency	14.8
MNGIE	23.12/27
Molybdenum cofactor deficiency	10.5/23.3
Monoamine oxidase A deficiency	2.4
Morquio syndrome, type A	18.7
Morquio syndrome, type B	18.8
MPS I-H	18.1a
MPS I-H/S	18.1c
MPS II	18.2
MPS IIIA	18.3
MPS IIIB	18.4
MPS IIIC	18.5
MPS IIID	18.6
MPS I-S	18.1b
MPS IVA	18.7
MPS IVB	18.8
MPS VI	18.9

Disorders	Disorder No.
MPS VII	18.10
Mucolipidin deficiency	19.12
Mucopolidosis I	19.4
Mucopolidosis II	19.10
Mucopolidosis III	19.11
Mucopolidosis IV	19.12
Multiple acyl-CoA dehydrogenation defect	14.10
Multiple carboxylase deficiency	7.1/7.2
Multiple sulfatase deficiency	19.17
Muscle phospho-fructokinase deficiency	15.14
Myoadenylate deaminase deficiency	23.7
Myopathy with lactic acidosis	27
Myophosphorylase deficiency	15.12
N	
N-Acetyl- α -D-glucosaminidase deficiency	18.4
N-Acetylaspartic aciduria	8.6
N-Acetylgalactosamine 6-sulfatase deficiency	18.7
N-acetylglucosamine 1-phospho transferase deficiency	19.10
N-Acetylglucosamine 6-Sulfatase deficiency	18.6
N-acetylglucosaminyltransferase II deficiency	20.6
N-Acetyl-glutamate synthase deficiency	11.6
NADH dehydrogenase	27.13
α -NAGA deficiency	19.7
NARP	27
Neonatal adrenoleukodystrophy	25.2
α -Neuraminidase deficiency	19.4
Niemann-Pick disease A/B	19.14
Niemann-Pick disease C	19.15
Nonketotic hyperglycinemia	3.2
O	
Occipital Horn Syndrome	33.3
Ornithine transcarbamylase deficiency	11.2
Ornithine-5-aminotransferase deficiency	12.1
2-Oxoadipate dehydrogenase deficiency	12.3
2-Oxoglutarate dehydrogenase complex deficiency	8.1/27.10
5-Oxoprolinase deficiency	9.4
Oxoprolinuria	9.2/9.4
Δ^4 -3-Oxosteroid 5 β -reductase deficiency	32.2
Oxysterol 7 α -hydroxylase deficiency	32.4
3-Oxothiolase deficiency	7.4
P	
Palmitoyl protein thioesterase 1 deficiency	22.4.1
PBG-deaminase deficiency	31.2
Pearson syndrome	27
PEO	27
Peroxisomal 3-ketothiolase deficiency	25.11
Phenylalanine-4-hydroxylase deficiency	1.1

Disorders	Disorder No.
3-Phosphoglycerate dehydrogenase deficiency	3.3
Phosphomannomutase 2 deficiency	20.1
Phosphomannose isomerase deficiency	20.2
Phosphoribosylpyrophosphate synthetase deficiency	23.8
Phytanyl-CoA hydroxylase deficiency	25.13
PKU	1.1
Pompe disease	15.9
Porphyria cutanea tarda	31.4
Porphyria variegata	31.6
PPCA deficiency	19.5
Primapterinuria	1.5
Primary hyperoxaluria type 1	26.1
Primary hyperoxaluria type 2	26.2
Progesterone resistance	29.16
Prolidase deficiency	3.8
Proline oxidase deficiency	3.6
Propionic acidemia	7.3
Propionyl-CoA carboxylase deficiency	7.3
Protoporphyrria	31.7
Pseudo-corpus luteum deficiency	29.16
Pseudo-neonatal adrenoleukodystrophy	25.9
Pseudo-vaginal perineoscrotal hypospadias	29.12
Pseudo-Zellweger syndrome	25.11
Pterin carbinolamine-4a-dehydratase deficiency	1.5
Purine nucleoside phosphorylase deficiency	23.2
Pyroglutamic aciduria	9.2/9.4
Δ^1 -Pyrroline-5-carboxylate dehydrogenase deficiency	3.7
Δ^1 -Pyrroline-5-carboxylate synthase deficiency	3.5/119
Pyruvate carboxylase deficiency	15.6
Pyruvate dehydrogenase complex deficiency	27.6
Pyruvate dehydrogenase deficiency	27.1/2
Pyruvate dehydrogenase phosphatase deficiency	27.7
6-Pyruvoyl-tetrahydropterin synthase deficiency	1.3
R	
5 α -Reductase type II deficiency	29.12
5 β -Reductase deficiency	32.2
Refsum disease	25.13
Renal glucosuria	16.2
Rhizomelic chondrodysplasia punctata Type 1	25.5
Rhizomelic chondrodysplasia punctata Type 2	25.6
Rhizomelic chondrodysplasia punctata Type 3	25.7
Riboflavin-responsive multiple acyl-CoA dehydrogenation defect	14.11
S	
Saccharopinuria	12.2.2
Salla disease	22.5
Sandhoff disease	19.9.3
Sanfilippo syndrome, type A	18.3
Sanfilippo syndrome, type B	18.4

Disorders	Disorder No.
Sanfilippo syndrome, type C	18.5
Sanfilippo syndrome, type D	18.6
Santavuori disease	22.4.1
Saposin B deficiency	22.3.3
Sarcosine dehydrogenase deficiency	3.10
Scheie syndrome	18.1b
Schindler disease	19.7.1
Segawa disease	1.6
Sepiapterin reductase deficiency	1.7
Short-chain 3-hydroxy-acyl-CoA dehydrogenase deficiency	14.9
Short-chain acyl-CoA dehydrogenase deficiency	14.7
Sialic acid storage disorder	22.5
Sialidosis	19.4
Sialin deficiency	22.5
Sialolipidosis	19.12
Sialuria	22.6
Sly syndrome	18.10
Smith-Lemli-Opitz syndrome	30.5
Sodium dependent glucose transporter 1 defect	16.1
Sodium dependent glucose transporter 2 defect	16.2
South African porphyria	31.6
Sphingolipid activated protein deficiency	19.13.4
Sphingomyelinase deficiency	19.14
StAR deficiency	29.1
Sterol 27-hydroxylase deficiency	32.3
Succinate dehydrogenase deficiency	27.11
Succininc semialdehyde dehydrogenase deficiency	3.4
Succinyl-CoA:3-oxoacid-CoA transferase deficiency	14.13
Sulfite oxidase deficiency (combined)	10.5/23.3
Sulfite oxidase deficiency (isolated)	10.4
Swedish porphyria	31.2
Systemic carnitine deficiency	14.1
T	
Tauri disease	15.14
Tay-Sachs disease	19.9.1
Testicular feminization syndrome	29.14
Tetrahydrobiopterin deficiency	1.2–1.7
Thiopurine methyltransferase deficiency	23.9
Thymidine phosphorylase deficiency	23.12
Transcobalamin II deficiency	10.10
Tricho-hepato-enteric syndrome	33.5.4
Trimethylaminuria	35.1
Tripeptidyl peptidase I deficiency	22.4.2
Tryptophan-2,3-dioxygenase deficiency	12.4
Tryptophanuria	12.4
Tyrosine aminotransferase deficiency (cytosolic)	4.2
Tyrosine hydroxylase deficiency	2.1
Tyrosinemia type I	4.1
Tyrosinemia type II	4.2
Tyrosinemia type III	4.3

Disorders	Disorder No.
U	
UDPGal-4-epimerase deficiency	15.3
UDP-N-acetylglucosamine 2-epimerase deficiency	22.6
UMP hydrolase deficiency	23.11a
UMP hydrolase superactivity	23.11b
UMP synthase deficiency	23.10
Ureidopropionase deficiency	23.15
Urocanase deficiency	5.2
Urocanic aciduria	5.2
Uro-cosynthase deficiency	31.3
Uroporphyrinogen-III-synthase deficiency	31.3
V	
Van Bogaert Bertrand disease	8.6
Variegate porphyria	31.6
VDAC	27.22
Very long-chain acyl-CoA dehydrogenase deficiency	14.5
Vitamin B12-binding protein deficiency	10.10
W	
Wilson disease	33.1
X	
X component of pyruvate dehydrogenase complex deficiency	27.5
Xanthine dehydrogenase deficiency	10.5/23.3
Xanthine oxidase deficiency	10.5/23.3
X-linked adrenoleukodystrophy	25.8
Z	
Zellweger syndrome	25.1