

Insulin-like Growth Factor-I ELISA E20

Insulin-like growth factors (IGF) I and II play a pivotal role in regulating the proliferation, differentiation and specific functions of many cell types. IGF-I has a molecular weight of 7.649 kDa. Its major regulators are growth hormone (GH) and nutrition. In contrast to many other peptide hormones, IGFs are avidly bound to specific binding proteins (IGFBP).

Physiological function of IGF-I is manifold. It is involved in several growth and maturation processes. Stimulating cell division and cell differentiation in target tissues. In human serum IGF-I is bound to its binding proteins and only a very small amount exists as free and biological active form.

Clinical Relevance

For clinical practice, one of the most important regulatory factors of IGF-I is GH. In growth disorders IGF-I serum concentration allows secure diagnostics, especially in connection with IGFBP-3 concentration. Deviant IGF-I levels are found in states of malnutrition / malabsorption, hypothyroidism, liver disease, untreated diabetes mellitus, chronic inflammatory disease, malignant disease or polytrauma.

Low levels, i.e. close to or below the age-related 5th percentile, would indicate the necessity of further diagnostic efforts. For differentiation of healthy short children without GH deficiency and children with "classical" GH deficiency, the 0.1st percentile proved to be an appropriate cut-off point, especially after the age of eight. In contrast, acromegaly is characterized by pathologically elevated IGF-I levels which apparently reflect the severity of the disease better than GH-levels.

Mediagnost IGF-I ELISA

Our ELISA is suited for measuring IGF-I in human **serum, and other human biological fluids** (e.g. follicular fluid) or cell culture supernatants and is **calibrated against** the international standard of the WHO (**WHO NIBSC 02/254**). **No cross reactivity** to other components of the IGF system of relevance is known. Small sample requirement is ideal for use in paediatrics.

Reference Values

Age	Percentile				
	0.1	5	50	95	99
0-2 y.	13	28	66	156	220
2-4 y.	20	40	87	189	260
4-6 y.	26	50	108	233	320
6-7 y.	34	62	124	248	332
7-8 y.	45	78	148	281	364
8-9 y. boys	54	90	160	284	362
8-9 y. girls	55	99	193	376	496
9-10 y. boys	63	102	176	304	379
9-10 y. girls	68	114	205	369	469
10-11 y. boys	77	117	189	305	370
10-11 y. girls	81	134	239	426	539
11-12 y. boys	85	129	209	339	413
11-12 y. girls	91	160	305	581	758
12-13 y. boys	881	141	243	419	525
12-13 y. girls	16	201	377	707	914
13-14 y. boys	111	179	311	540	677
13-14 y. girls	163	256	428	716	884
14-15 y. boys	140	229	404	691	896
14-15 y. girls	193	284	443	713	832
15-16 y. boys	176	269	433	697	849
15-16 y. girls	187	279	442	700	845
16-17 y. boys	178	267	424	673	814
16-17 y. girls	183	270	422	660	792
17-18 y. boys	173	243	358	527	618
17-18 y. girls	176	246	362	533	624
18-19 y. boys	167	235	347	512	600
18-19 y. girls	167	233	341	499	797
19-20 y.	158	220	322	471	550
20-30 y.	72	115	198	340	425
30-40 y.	68	109	188	324	404
40-50 y.	64	103	178	310	385
50-60 y.	60	97	169	292	369
60-70 y.	55	91	161	282	362
70-80 y.	25	47	98	207	276
>80 y.	21	40	85	184	245

Serum levels are given as ng/ml.

Mediagnost IGF-ELISA has a **European Approval for Clinical Diagnostics: EC Directive 98/79EG (CE)**.

Assay Features E20

- ✓ **No extraction required because of IGF-II excess!**
- ✓ Results on your hands **in less than 2h**
- ✓ Analytical sensitivity of **0.09 ng/ml**
- ✓ 5 single standards included (2-50 ng/ml)
- ✓ Assay range **0.09 to 1050 ng/ml** (with recommended dilution)
- ✓ 2 human native control sera included
- ✓ Minimal cross-reactivity to IGF-II (0.055%)

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Reconstitution / Dilution of Reagents		
Standards A-E	Reconstitution in Sample Buffer PP	500µl
Control Serum KS1	Reconstitution in Sample Buffer PP	500 µl
Control Serum KS2	Reconstitution in Sample Buffer PP	500 µl
Wash Buffer WP	dilute in A. dest. (e.g. total volume of 50 ml in a graduated flask and fill up to 1000 ml)	1:20
Sample + Control Sera KS1 and KS2: dilute 1:21 in Sample Buffer PP , mix immediately, incubate max. 2h. Use 20 µl for each well in the assay.		
Before conducting the assay equilibrate all reagents to room temperature.		

Assay Procedure for Double Determinations:

Pipette	Reagent	Position
80 µl	Antibody Conjugate AK	in all wells used
20 µl	Sample Buffer PP (blank)	A1 and A2
20 µl	Standard A (2 ng/ml)	B1 and B2
20 µl	Standard B (5 ng/ml)	C1 and C2
20 µl	Standard C (15 ng/ml)	D1 and D2
20 µl	Standard D (30 ng/ml)	E1 and E2
20 µl	Standard E (50 ng/ml)	F1 and F2
20 µl	Control Serum KS1	G1 and G2
20 µl	Control Serum KS2	H1 and H2
20 µl	Samples	following wells
Cover the wells with the sealing tape.		

Incubation: 1 h **at** RT, 350 rpm

5x 300 µl	Aspirate the contents of the wells and wash 5x with 300 µl Wash Buffer WP	each well
100 µl	Enzyme Conjugate EK	each well

Incubation: 30 min **at** RT, 350 rpm

5x 300 µl	Aspirate the contents of the wells and wash 5x with 300 µl Wash Buffer WP	each well
100 µl	Substrate S	each well

Incubation: 15 min in the dark **RT**

100 µl	Stop Solution SL	each well
Measure the absorbance within 30 min at 450 nm with ≥ 590 nm as reference wavelength.		