

Review Article

Vol. 9, No. 1, January 2024, Webpage: http://rheumres.org Email: <u>rheumres@gmail.com</u> ISSN:2476-5856 doi: <u>10.32592/RR.2024.9.1.62</u> ©2024, Iranian Rheumatology Association

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Serial or compare bone mineral densitometry: how to do it step by step

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Bone mineral densitometry (BMD) is the most valuable method for assessing bone and calculating fracture risk. Serial or comparative bone densitometry is important in rheumatologists' work on osteoporosis management. The response or lack of response to osteoporosis treatment based on densitometry scans is crucial. This paper examines the timing of scan requests concerning the history of glucocorticoid use, renal or other solid organ transplantation, malignancy, and other situations discussed. We encountered four types of compared scans based on the centers where BMD was performed and the precision of the devices used for this survey: Same Center, Same Device (SSSD), Same Center Different Devices (SCDD), Different Centers Same Devices (DCSD), and Different Centers Different Devices (DCDD). We discussed the principles of comparison and the key indicators.

Keywords: Bone densitometry; Bone density; Serial BMD; Compared BMD; Center; Device

Principles and methods

Definitely the most important role of bone mineral densitometry (BMD) is to compare with multiple results from similar patients for changes in bone density over time. Frequency and correct time of repeat of BMD based on risk factors showed in two Figures (Figures 1, 2) [1-8]. There are four types of compared BMD: (1) Same Center & Same Device (SCSD), (2) Same Center & Different Devices (SCDD), (3) Different Centers & Same Device (DCSD), (4) Different Centers & Different Devices (DCDD). In this article we discussed the principles of SCSD.

Same Center & Same Device (SCSD)

It is the best type and recommended form of BMD. Serial or compared BMD indications are (1) to monitor response to therapy, (2) to assess nonresponse by bone densitometry as one source of responsiveness, (3) to follow up patients who are not on treatment and are at risk of bone loss (such as steroid users, hyperparathyroidism). Same as other BMD reading and interpretation, SCSD have 5 steps.

Step I: ID Characters control

In this step we control the name, surnames, age, sex, height, weight and ultimately reference population (ethnicity) shown in Figures 3, 4. It should be remembered that any problems in above data, can be solved if both scans be available. About reference population, the International Society of Clinical Densitometry (ISCD) recommendation is selection of specific population such as White, Black, Hispanic, Asian, if patient is not in any of above population better

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Received: 3 June 2023; Accepted: 25 September 2023



Figure 1. Algorithm for repeat BMD in some conditions



Figure 2. Algorithm for serial BMD based of BMD result









to use Caucasian.

Step II: Control of good scan criteria

This step has 2 phases:

I. Checking the similarity of scan image (straightness, top, bottom, both sides)

II. Checking the good scan criteria in each scan

It's very important that takes the new scan looks as similar as possible to the previous scan (except artifacts). The straightness of limbs (for example spine or hip) is very important at this step (Figures 5, 6). Since the first (old) scan is the basis for the second (last) scan, it is important to teach the technician the criteria for a good scan as shown in Figures 7 and 8 [9-13].

Spine good scan criteria (Figure 7) are:

- 1- Lack of artifact
- 2- Spine should be straight (S)
- 3- Upper border: at least 1/2 of T12 (A)
- 4- Lower border: at least $\frac{1}{2}$ of L5 (B)
- 5- Each side: at least 2 cm at each side of spine (C)
- Hip good scan criteria (Figure 8) are:
- 1. Lack of artifact
- 2. Hip should be straight (S)

3. Upper border: at least 2 cm above greater trochanter (A)

4. Lower border: at least 1.5 cm below ramus of pubis (B)

5. Inner border:

- a. no or small size of lesser trochanter (E)
- b. visibility of a part of obturator foramen (D)
- c. distance between ramus and neck 1cm(C)

6. Outer border: at least 1 cm soft tissue (OB)

When this step successfully passed (similarity of each scan in previous and recent results such as Figure 4 for spine scans and Figure 5 for hip region), we can go to step III. In should be emphasized that the technician must be seen the previous scan and take the new one with maximum similarity to earlier scan [14-26].

Step III: Unification of Region of Interest (ROI) insertion

During this step, the ROI or area of both scans should be the same. In spine region it's necessary to uniform labeling of vertebrate in scans. For this purpose, the vertebrates should be labeled or numbered. Technically two ways for labeling exist:



Figure 5. Special attention to spine image for similarity of straightness, above, below & sides of both scan

A

OB



Figure 6. Compare the both hip image for mentioned items (straightness, above, below & sides)



Figure 7. Spine good scan criteria



1. Shape of vertebrae: as a rule, vertebrae of L1, L2 and L3 are U shape and L4 is "X" or "H" shape. L5 has "wm" so the above border of L5 is same "w" and below borders as "m" (Figures 9, 10).

2. Landmark use: iliac crest bone is parallel of L5 and rib connects to T12 and the third landmark is long transverse process that see on L3 (Figures 11 and $\underline{12}$).



Figure 9. Schematic scan labeling [12]

Figure 10. Real scan [12]



Figure 11. labeling of vertebrae based of landmarks

In compared BMD, unifying of both spine scan is very important and need to use of similar labeling of vertebrae correct passing of this step is obligatory for going to next step.



Figure 12. Long transverse process at L3

In the pelvic area, it is important to have the neck box (rectangular box) in the same location in both scans (Figure 13).For this purpose, four characters should be assessed:

1. Neck box should not be entered to head or greater trochanter region (H/GT at Figure 14)

The inner line of the neck box should cross the ramus pubis if it continues (NBIB at <u>Figure 15</u>).
 The outer line of neck box should not enter the greater trochanter region if it continues (NBOL at Figure 16).



Figure 13. Neck box (green box)

4. The ward triangle (the square box) should be near or attached to neck box (WBNB at Figure 17) and should not be separated (Figure 18).

If all the characters in the two images are similar, you can go to the next step. It should be mentioned again, we or technician can correct this step without repeating of scan [27-29].



Figure 14. Neck box between head (H) and greater trochanter (GT)



Figure 15. Inner line of neck box should be near or cross the ramus of pubis or ILNB (black line)



Figure 16. outer line of neck box should not be enter the greater trochanter or OLNB (red line)



Figure 17. ward box (blue) should be near of neck box or WBNB (black circle)

If all the characters in the two images are similar, you can go to the next step. It should be mentioned again, we or technician can correct this step without repeating of scan [27-29].

Step IV: Control of area & BMD in both scan

This step has 3 phases for spine region and two steps for hip and forearm regions. In *the spine scan* 3 phases should be considered:

Phase 1: selection of best region

For selection of best region, area and BMD rules should be considered. According to the area rule, the lumbar spine area gradually increases from L1 to L4 (L1 < L2 < L3 < L4) (Figure 19). The border of the vertebral region is determined by the technician, so it requires great care to clarify the top, bottom and both sides. On the other hand, the area rules are the most important duty of the technician, and compliance with the rules of the area in each scan (spine, hip, forearm and whole) is the main duty of the technician to reduce reporting errors.

According to the BMD rule, the lumbar spine BMD gradually increases from L1 to L3 and the decrease to L4 (L1 < L2 < L3 > L4) (Figure 19). Thus, if both rules (area & BMD) are kept, L1-4 selected and based on the rules three or two consecutive vertebrae should be elected (Figure 20 area error & Figure 21 BMD error). The best region to scan the spine for a patient is shown in



Figure 18: ward box not near the neck box

Figure 22.

Phase 2: selection of the common best region for both scan

If two scans have the same best region based of area and bone density, the same region is selected as the common best region. But if in the first scan L1- 3 is suitable and in the second L2-4 is suitable, then L2-3 is the common best region (Figure 23).

Phase 3: Controlling the area difference between two scans

At this phase, it is necessary to compare the common best region between the two scans and this difference should not be more than 2 square centimeters (Figure 24). It should be noted that if there is no problem in performing & analysis the scans, the device itself compares the L1-4 region (Figure 25).

In the hip scan 2 phases should be considered: *Phase 1: Choosing the total area as the common best region*

For comparison in hip scan, we should be used total region. If the total region cannot be used in decreasing frequency greater trochanter, inter trochanter, and finally neck regions may be used.

Phase 2: Controlling the area difference between two scans

In hip scan, the difference area should not be above 2 cm^2 (Figure 26). If the difference in the total area of two scans is more than 2 square centimeters, then it is necessary to correct this

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	The second s		5	Scan Info	ormation:		
	Z			ican Date: Scan Type: Analysis: Operator: Aodel: Comment:	07 April 2021 f Lumbar Spine 07 April 2021 11:4 Spine NB Discovery W (S/N	ID: A04 15 Version 13.6. 83167)	007210K
	1						
XA Resul	Its Summary	y: BMC (g)	BMD (g/cm²)	T - score	PR (%)	Z - score	AM (%)
XA Resul tegion	Area (cm ²) 13.63	y: (g) 9,88	BVID (g/cm ²) 0.725	T- score -3.2	PR (%) 68	Z score -3.2	AM (%) 68
KA Resul	Area (cm ²) 13.63 14.41	y: BMC (g) 9.88 11.02	BNID (g/cm²) 0.725 0.765	T - score -3.2 -3.0	PR (%) 68 70	Z - score -3.2 -3.0	AM (%) 68 70
KA Resul	Area (cm ²) 13.63 14.41 16.03	9.88 11.02 12.98	BVID (g/cm ²) 0.725 0.765 0.810	T - score -3.2 -3.0 -2.7	PR (%) 68 70 73	Z, - score -3.2 -3.0 -2.7	AM (%) 68 70 73
KA Resul	Area (cm ²) 13.63 14.41 16.03 16.17	9.88 11.02 12.98 12.65 20.00	BXID (g/cm ²) 0.725 0.765 0.810 0.782	T- score -3.2 -3.0 -2.7 -2.8	PR (%) 68 70 73 72 71	Z - score -3.2 -3.0 -2.7 -2.8	AM (%) 68 70 73 72
KA Resul egion 1 2 3 1 1-L2 1 1 2	Area (cm ²) 13.63 14.41 16.03 16.17 28.04 28.04	9.88 11.02 12.98 12.65 20.90 23.96	BXID (g/cm ²) 0.725 0.765 0.810 0.782 0.746	T- score -3.2 -3.0 -2.7 -2.8 -2.8	PR (%) 68 70 73 72 71 73	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8	AM (%) 68 70 73 72 71 73
CA Resul egion 1 2 3 4 1 1-L2 1.L3 1.L4	Area (cm ²) 13.63 14.41 16.03 14.41 16.03 14.41 16.03 14.41 16.03 14.41 16.03 14.41 16.03 14.41 16.03 16.17 28.04 29.66	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.86 22.84	BMD (g/cm ²) 0.725 0.765 0.810 0.782 0.746 0.771 0.755	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.6 -3.0	PR (%) 68 70 73 72 71 73 70	Z score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.6 -2.6	AM (%) 68 70 73 72 71 73 70
CA Resul egion 1 2 3 1 1-L2 1.L3 1.L4 2 1.2	ts Summary (cm ²) 13.63 14.41 16.03 16.17 28.04 29.66 29.80 30.44	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00	BMD (g/cm ²) 0.725 0.765 0.810 0.782 0.746 0.771 0.756 0.779	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.8 -2.8 -2.8 -2.8	PR (%) 68 70 73 72 71 73 70 72	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.6 -3.0 -2.8	(%) 68 70 73 71 73 70 72
KA Resul egion 1 2 3 1 1-1-2 1.L3 1.L4 2-L3 2-L4	Lts Summary Area (cm ²) 13.63 14.41 16.03 16.17 28.04 29.80 30.44 30.58	BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00 23.67	BAD (g/cm ²) 0.725 0.765 0.810 0.746 0.771 0.756 0.774	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.6 -3.0 -2.8 -3.2	PR (%) 68 70 73 72 71 73 70 70 72 69	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.6 -3.0 -2.8 -3.2	(%) 68 70 73 72 71 73 70 72 60
XA Resul egion 1 2 3 1 1-L2 1.L3 1.L4 2-L3 2.L4 2-L3 2.L4 3.L4	Area (mr) 13,63, 14,41, 16,03, 14,41, 16,03, 14,41, 16,03, 14,41, 16,03, 14,41, 28,04, 29,66, 29,80, 30,44, 30,58, 32,20,	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00 23.67 23.63	BMD (g/cm ²) 0.725 0.765 0.810 0.782 0.746 0.771 0.736 0.789 0.774 0.776	T- score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -3.0 -2.8 -3.2 -3.0	PR (%) 68 70 73 72 71 73 70 72 69 71	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -3.0 -2.8 -3.0 -2.8 -3.2 -3.0	AM (%) 68 70 73 71 73 70 70 72 69 71
egion 1 2 4 1-1-2 1.1.3 1,1.4 2-1.3 2,1.4 3-1.4 3-1.4 1-1.3 2,1.4 1-1.3 1,1.4 1,1.3 1,1.4 1,1.4 1,1.3 1,1.5	Its Summary Area (cm ²) 13.63 14.44 16.03 14.44 16.03 16.17 28.04 29.66 29.80 30.44 30.58 32.20 44.07	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00 23.67 25.63 33.89	BMD (g/cm ²) 0.725 0.765 0.810 0.782 0.774 0.775 0.774 0.778 0.774 0.779 0.774	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.8 -3.0 -2.8 -3.2 -3.0 -2.2	PR (%) 68 70 73 72 71 73 70 72 69 71 72	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.8 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.7	(%) 68 70 73 71 73 70 72 69 71 72
CA Result egion 1 2 3 4 4 1-L2 1.1.3 1.1.4 2-L3 3.1.4 3-1.4 3-1.4 1-1.2 1.4	Its Summary (cm ²) 13.63 14.44 16.03 16.17 28.04 29.80 30.44 30.58 32.20 44.07 44.21	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00 23.67 25.63 33.89 33.56	BYD (g/cm ²) 0.725 0.765 0.810 0.746 0.774 0.756 0.774 0.756 0.774 0.796 0.779	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.6 -3.0 -2.8 -3.2 -2.8 -3.2 -3.0 -2.7 -3.0 -2.7 -3.0	PR (%) 68 70 73 72 71 73 70 72 69 71 72 70 71 72 70	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -2.6 -3.0 -2.8 -3.2 -3.0 -2.7 -3.0	AM (%) 68 70 73 72 71 73 70 72 69 71 72 70
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XA Result tegion 1 2 3 3 4 4 1-12 2-13 2-13 2-14 3-14 3-14 1-123 1-123 1-123 1-123 1-123 1-123 1-123 1-124 2-14	Its Summary I.3.63 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 14.44 16.03 13.65 29.66 29.80 30.44 30.58 32.20 44.07 44.21 45.83 46.61	y: BMC (g) 9.88 11.02 12.98 12.65 20.90 22.86 22.53 24.00 23.67 25.63 33.89 33.56 35.52 36.65	BND (g/cm ³) 0.725 0.765 0.810 0.782 0.746 0.771 0.756 0.778 0.774 0.779 0.779 0.775 0.786	T - score -3.2 -3.0 -2.7 -2.8 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.7 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.2 -3.0 -2.7 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.2 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.9 -3.0 -2.9 -3.0 -3.0 -2.9 -3.0 -3.0 -3.0 -2.9 -3.0 -3.0 -3.0 -2.9 -3.0	PR (%) 68 70 73 72 71 73 70 72 69 71 72 70 71 71	Z - score -3.2 -3.0 -2.7 -2.8 -2.8 -2.8 -3.0 -2.8 -3.0 -2.8 -3.0 -2.7 -3.0 -2.9 -3.0	AM (%) 68 70 73 72 71 73 70 72 69 71 72 70 71 71

Figure 19: keeping of rules in area and BMD in all vertebrae from L1 to L4 and election of L1-4 for report



Figure 20. Area error in spine region and choosing of kept area rules

Figure 21. Failure of BMD rule and selection of areas that have complied the rule

L1 L2 L3

L4

L1-L2 L1,L3

L2.L4

AM (%)



Figure 22. Selected best region for old & new spine scan

L2-L3	28.06	Scan Date:	01 October 2020
L2-L3	30.83	Scan Date:	28 August 2016

Figure 24. The difference between the two area in the selected common region, should not be more than 2 cm²

difference by increasing or decreasing the area. Figure 27 shows the compared scan in hip region with high area difference and how to correct this problem [30,31].

Step V: interpretation

This step has 3 phases:

Phase 1: reporteach scan separately

In this phase based on the results (best selected region in step IV for spine, lower result between total and neck in hip scan and 1/3 of radius for forearm) of bone density are reported (Figure 28). It should be mentioned here that the final conclusionis the lowest result

105 x 134							
XA Resu Region	Area	BMC	BMD (g/cm ²)	T - score	PR (%)	Z - score	AM (%)
LI	13.95	11.38	0.816	-1.6	82	-0.2	98
L2	14.93	12.04	0.806	-2.0	78	-0.5	94
L3	15.89	13.04	0.820	-2.4	76	-0.8	91
L4	21.22	17.98	0.847	-1.9	80	-0.3	97
L1-L2	28.88	23.42	0.811	-1.5	83	-0.1	99
LI.L3	29.84	24.42	0.818	-1.8	81	-0.2	97
L1.L4	35.17	29.36	0.835	-1.8	80	-0.3	96
L2-L3	30.83	25.07	0.813	-2.2	77	-0.6	92
L2.L4	36.15	30.01	0.830	-2.2	77	-0.6	92
L3-L4	37.12	31.01	0.836	-2.4	76	-0.8	91
L1-L3	44.77	36.46	0.814	-1.9	80	-0.3	96
L1-L2,L4	50.10	41.40	0.826	-1.9	80	-0.3	96
L1.L3-L4	51.07	42.40	0.830	-2.0	79	-0.4	94
L2-L4	52.05	43.05	0.827	-2.3	77	-0.7	92
L1-L4	66.00	54.43	0.825	-2.0	79	-0.5	94

Figure 23. Common best region selection

between spine, total and neck in hip, so that if, for example the spine is osteoporotic and the neck is normal, the final conclusion is osteoporosis (Figure 29).

Phase 2: determine percentage change between two scans:

In spine, after finding of best region or best vertebrae in previous step we can take percentage change between scans based this formula:

Cbmd= new the common selected region bmd previous same the common selected region bmd/ previous same the common selected region bmd ×

100. For example, for case of Figure 22 after use of formula we reached to these results: Cbmd (L2-3) = bmd (L2-3: 2020) - bmd (L2-3:



Figure 25. Usual device comparison



Operator: Model:

Comment

(green line). DXA Results Summary

Region Area (cm²)

Troch

Total

4.74 10.24

21.41 18 55 0 866

A Discovery W (S/N 83167)

 BMC
 BMD

 (g)
 (g/cm²)

 3.60
 0.759

 6.23
 0.608

0.60 0.517

36.39 28.37 0.780

T -score PR (%) Z -score AN (% 1.3 0.6

-0.8 -0.9

-1.5 -1.3

-1.9 70 1.0 12

89 87

79 83 0.0 0.4 101 107

The difference in the area of two scans is more than 2 $\rm cm^2,$ and the reason for that is the lower line of the ROI box(blue box), should be placed a little higher

AM (%)

112 113

115 118

126

123



Image not for di 112 x 107 NECK: 49 x 15

Figure 27. The area difference of two scans is more than the rule

2016)/ bmd (L2-3: 2016) × 100 Cbmd (L2-3) = $0.774 - 0.813/0.813 \times 100$ Cbmd $(L2-3) = -4,8\% \sim -5\%$



Figure 26. The difference in the appropriate area in the hip scan



Figure 28. Algorithm of BMD report based the results of best region of spine and hip(total or neck), forearm(1/3 or 33%) & whole body (subtotal or TBLH)

Indication for densitometry:

Age>65 yrs

History of fragility fracture(s):

None

Bone mineral density results:

	BMD	T- score	Z- score	Result
<i>Spine (L1-L4)</i> ¹	0.809	- 2.2	0.3	Low Bone Mass
L-Hip (Total) ¹	0.549	- 3.2	- 1.4	Osteoporosis

Diagnosis by WHO criteria²:

According T-Score criteria this study indicates Osteoporosis.

Figure 29. Conclusion of BMD report as lowest result

Another way to measure Cbmd is to use standardized bmd or sBMD calculator. To interpret densitometry with two devices, it is necessary to first convert BMD into sBMD (Figure 30) which shows how to convert in three different devices used (we will fully discuss the use of SBMD and its value later in DCDD, SCDD, DCSD topics). When the information is put in the standardized BMD calculator, the result of the Figure 31 is obtained. In hip region, the best region for comparison is total hip and we can take percentage change between scans based on this formula:

Calculation of compared hip scan (based formula) = total hip bmd (new) - total hip bmd (previous) / total hip bmd (previous) × 100

For example, the calculation of compared scan in the hip scan based on the standardized BMD calculator for case <u>Figure 32</u> is as follows:

= total hip (2020) – total hip (2018) / total hip (2018) × 100

 $= 0.887 - 0.864 / 0.864 \times 100$

 $= 2.7\% \sim 3\%$

The bone density change percentage in the hip scan is often presented as a graph (Figure 33). Phase 3: Review of response to treatment: To check the response to treatment, we need at least two information:

1. Precision of the device

2. The Least Significance change (LSC) of the

It should be noted that the precision of the device is constant, but the LSC of the center is often variable, because the technicians and their ability can change over time.

How to calculate the summary LSC (sLSC) is as follows:

 $sLSC = (LSC C1 + LSC C2) \times (precision C1 + precision C2)$

Based on this, LSC C1 is the LSC of the first center, LSC C2 is the LSC of the second center, and precision C1 & C2 is the precision of the first and second centers. For example, with the following information, the result of the sLSC for hip & spine region is as follows:

Spine: LSC C1=1.8%, LSC C2= 2%, precision C1= 1%, precision C2= 1.2%

 $sLSC = (LSC C1 + LSC C2) \times (precision C1 + precision C2)$

 $=(1.8+2) \times (1+1.2) = 8.36 \sim 8\%$

For SCSD, the precision of the device should be considered constant, and often LSC is often constant, so the formula changes as follows:

 $sLSC = LSC \times precision$ (if LSC of first & second

is constant)

sLSC= (LSC first/ old scan + LSC second/new scan) × precision (if LSC of first & second scans changes)

FEMORAL NECK		TOTAL HIP			
First Measurement Hologic Lunar Norland 0.000 g / cm2	Second Measurement Hologic Lunar Norland 0.000 g / cm2	First Measurement Hologic Lunar Norland 0.000 g / cm2	Second Measurement Hologic Lunar Norland 0.000 g / cm2		
click to convert		click to convert			
mg/cm2	mg/cm2	mg/cm2	mg/cm2		
% change		% change			
LUMBAR SPINE					
First Measurement Hologic Lunar Norland 0.000 g / cm2	Second Measurement Hologic Lunar Norland 0.000 g / cm2				
click to convert					
mg/cm2	mg/cm2				
% change					

LUMBAR SPINE Second Measurement First Measurement Hologic Holoaic O Lunar 🔾 Lunar O Norland O Norland g / cm2 0.774 g/cm2 0.813 click to convert 876 mg/cm2 835 mg/cm2 -4.7 % change

Figure 31. Use of standardized BMD for case of Figure 22

TOTAL HIP	
First Measurement Hologic Lunar Norland 0.864 g / cm2 	Second Measurement Hologic Lunar Norland 0.887 g / cm2
877 mg/cm2	900 mg/cm2
2.6 % change	500 mg/cm2

Figure 32. Hip scan compared BMD calculator

Rajaie

	Scan Date	Age	BMD	Т-	BMD	Change
			(g/cm ²)	score	vs Baseline	vs Previous
Θ	24.11.2020	58	0.931	-0.1	3.5%*	3.5%*
	13.05.2018	56	0.899	-0.3		
ik de in de Age	* Denotes signific	ance at 95% c	onfidence level, LS	C is 0.027 g/cr	n²	
Trochanter						
	Scan Date	Age	BMD (g/cm ²)	T - score	BMD vs Baseline	Change vs Previous
0	24.11.2020	58	0.607	-0.9	-3.4%*	-3,4%*
	13.05.5010	56	0.629	-0.7		
່ນັກເປັນ່ວນນັ້ນ Age Inter	* Denotes signific	ance at 95% c	onfidence level, LS	C is 0.018 g/cr	D ²	
່ ຕໍ ຈຸ ໃດ ສໍ ຄຸ ດິ ພໍ ຍ Age Inter	* Denotes signific	ance at 95% co	BMD (a/cm ²)	C is 0.018 g/cm	n ² BMD vs Recelline	Change ys Previous
కరజరీపలనటిల Ave Inter 	* Denotes signific	ance at 95% co Age 58	BMD (g/cm ²) 1.103	C is 0.018 g/cm C is 0.018 g/cm T - score 0.0	n ² BMD vs Baseline 5,9%*	Change vs Previous 5.9%%
sésésésés Ane Inter Ag. 0	* Denotes signific * Denotes signific * 24,11.2020 13.05.2018	ance at 95% c Age 58 56	BMD (g/cm ²) 1.103 1.042	C is 0.018 g/cm C is 0.018 g/cm C is 0.018 g/cm C is 0.018 g/cm	n° BMD vs Baseline 5.9%%	Change vs Previous 5.9%*
a d b b a d b a Ase Noter a a a Ase d b b a d b a b Ase	* Denotes signific * Denotes signific * Denotes signific * Denotes signific	Age 58 56 ance at 95% c	BMD (g/cm ²) 1.103 1.042 onfidence level, LS	C is 0.018 g/cm T	p ² <u>BMD</u> vs Baseline 5.9%%	Change vs Previous 5.9%*
Apr Noter C. C. C	* Denotes signific * Denotes signific * Denotes signific * Denotes signific	Age 58 56 ance at 95% c	BMD (g/cm ²) 1.103 1.042 onfidence level, LS	T - score 0.0 -0.4 C is 0.025 g/cd	n² vs Baseline 5.9%%	Change vs Previous 5.9%
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	* Denotes signific * Denotes signific * Denotes signific * Denotes signific * Denotes signific	Age 58 56 ance at 95% c	BMD (g/cm ²) 1.103 1.042 ontificance level, L5 BMD (g/cm ²)	C is 0.018 g/cr T - score 0.0 -0.4 C is 0.025 g/cr T - score	n ² vs Baseline 5.9%s ⁴ n ² BMD vs Baseline	Change vs Previous 5.9%
6 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	* Denotes signific * Denotes signific * Denotes signific * Denotes signific * Denotes signific * Denotes signific	Age 58 56 ance at 95% c	BMD (g/cm ²) 1.103 1.042 onfidence level, LS (g/cm ²) 0.634	C is 0.018 g/cr T- score 0.0 -0.4 C is 0.025 g/cr C is 0.025 g/cr T- score -0.9	n ² vs Baseline 5.9% ⁴⁰ n ² vs Baseline -3.3%	Change vs Previous 5.9%* Change vs Previous -3.3%

Figure 33. Usual device comparison in hip scan

Evaluation of treatment response is defined in three ways:

I. Complete response: when it increases more than the sLSC, but if there is no information, the value of 7% is the criterion, which indicates a greater increase.

II. Failure to response OR no response: when the reduction is more than sLSC or 7% in cases where we do not have the device & center information, it indicates a lack of response.

III. Partial response: which is when neither the increase nor the decrease is greater than sLSC, and in cases where we do not have device & center information, it is between +7% and -7%.

Of course, it should be known that the definition of lack of treatment response or refractory osteoporosis treatment is different, and based on this; the definition is one of the following three situations: - Occurrence of two fragile (low trauma) vertebral fractures after 1 year of correct treatment or one vertebra & one another regions (forearm, rib, humerus, tibia, pelvis)

- Occurrence of fragility hip fracture after 1 year of correct treatment.

- Occurrence of one fragile vertebra fracture and decrease of above 7% BMD (with this sequence: spine, then hip (total, troch, intertrochanter/body)

Treatment outcome	Description
Response	Stability or significant increase of BMD with appropriate change of bone turnover marker level
Success	BMD increase to T-score > -2.0 when treatment is started because of T-score ≤ -2.5 , or T-score increase of at least 1.0 units when treatment is started with T-score > -2.5 , and no recent fracture (e.g., within 3 years)
Failure	Two or more incident fractures; or one incident fracture with significant decrease in BMD and/or lack of appropriate change of bone turnover marker level; or significant decrease in BMD and lack of appropriate change of bone turnover marker level

Figure 34. Consequences of osteoporosis treatment [33]

& forearm total) on SCSD after 1 year of correct treatment.

There are other definitions for examining response to treatment, an example of which can be seen in Figure 34. Based on the mentioned rules, for the case, the spine changes -5% and hip +3% were calculated. The final response is "partial response", which means that is not necessary to change the treatment [31-36].

Acknowledgement

Not applicable.

Conflict of Interests

The author declares no conflict of interest.

Funding

The authors confirm that this work was not funded.

References:

- Lenchik L, Kiebzak GM, Blunt BA. What is the role of serial bone mineral density measurements in patient management? *J Clin Densitom* 2002; 5 Suppl:S29-38. doi: 10.1385/jcd:5:3s:s29.
- Kendler DL, Compston J, Carey JJ, Wu CH, Ibrahim A, Lewiecki EM. Repeating Measurement of Bone Mineral Density when Monitoring with Dual-energy Xray Absorptiometry: 2019 ISCD Official Position. J Clin Densitom 2019; 22(4):489-500. doi: 10.1016/ j.jocd.2019.07.010.
- Hillier TA, Stone KL, Bauer DC, Rizzo JH, Pedula KL, Cauley JA. *et al.* Evaluating the value of repeat bone mineral density measurement and prediction of fractures in older women: the study of osteoporotic fractures. *Arch Intern Med* 2007; 167(2):155-60. doi: 10.1001/archinte.167.2.155.
- Chotiyarnwong P, McCloskey EV. Pathogenesis of glucocorticoid-induced osteoporosis and options for treatment. *Nat Rev Endocrinol* 2020; 16(8):437-47. doi: 10.1038/s41574-020-0341-0.
- Unal A, Kocyigit I, Sipahioglu MH, Tokgoz B, Kavuncuoglu F, Oymak O. *et al.* Loss of bone mineral density in renal transplantation recipients. *Transplant Proc* 2010; 42(9):3550-3. doi: 10.1016/ j.transproceed. 2010.07.106.
- Batteux B, Gras-Champel V, Lando M, Brazier F, Mentaverri R, Desailly-Henry I. *et al.* Early steroid withdrawal has a positive effect on bone in kidney transplant recipients: a propensity score study with inverse probability-of-treatment weighting. *Ther Adv Musculoskelet Dis* 2020; 12:1759720x20953357. doi: 10.1177/1759720x20953357.
- Hadji P, Aapro MS, Body JJ, Gnant M, Brandi ML, Reginster JY. *et al.* Management of Aromatase Inhibitor-Associated Bone Loss (AIBL) in postmenopausal women with hormone sensitive breast cancer: Joint position statement of the IOF, CABS, ECTS, IEG, ESCEO IMS, and SIOG. *J Bone Oncol* 2017; 7:1-12. doi: 10.1016/j.jbo.2017.03.001.
- Pazianas M, Miller PD. Osteoporosis and Chronic Kidney Disease-Mineral and Bone Disorder (CKD-MBD): Back to Basics. *Am J Kidney Dis* 2021; 78(4):582-89. doi: 10.1053/j.ajkd.2020.12.024.
- Official website of the International Society for Clinical Densitometry (ISCD), http://www.iscd.org/officialpositions/official-positions, last reviewed on 1/31/2021.
- 10. Bone density. [https://courses.washington.edu/ bonep hys/opbmd.html]
- Abdellah El M. Interpreting a DXA Scan in Clinical Practice. In: *Dual Energy X-Ray Absorptiometry*. Edited by Abdellah El M. Intech Open, Rijeka; 2012:

Ch.1.

- S.L. Bonnick and L.A. Lewis, Performing a DXA PA Lumbar Spine, Proximal Femur, or Forearm DXA Study, In: Bone Densitometry for Technologists, 3rd edition, 2013, P: 97-118.
- Lekamwasam S, Lenora RS. Effect of leg rotation on hip bone mineral density measurements. *J Clin Densitom* 2003; 6(4):331-6. doi: 10.1385/jcd:6:4:331.
- Faulkner KG, Genant HK, McClung M. Bilateral comparison of femoral bone density and hip axis length from single and fan beam DXA scans. *Calcif Tissue Int* 1995; 56(1):26-31. doi: 10.1007/bf00298740.
- Wong JC, Ong B. Evaluation of femur angle abduction/adduction and bone mineral density values. J Clin Densitom 2005; 8(4):472-5.
- Licata AA, Williams SE. A DXA Primer for the Practicing Clinician: A Case-Based Manual for Understanding and Interpreting Bone Densitometry, 1 edn New York, NY; 2014. doi: 10.1007/978-1-4419-1375-3 2
- Morgan SL, Lopez-Ben R, Nunnally N, Burroughs L, Fineberg N, Tubbs RS. *et al.* The effect of common artifacts lateral to the spine on bone mineral density in the lumbar spine. *J Clin Densitom* 2008; 11(2):243-9. doi: 10.1016/j.jocd.2007.11.004.
- Wahner H. Technical aspects and clinical interpretation of bone mineral measurements. *Public Health Rep* 1989; 104 Suppl(Suppl):27-30.
- Lentle BC, Prior JC. Osteoporosis: What a clinician expects to learn from a patient's bone density examination. *Radiology* 2003; 228(3):620-8. doi: 10. 1148/radiol.2283020093.
- Jacobson JA, Jamadar DA, Hayes CW. Dual X-ray absorptiometry: recognizing image artifacts and pathology. *AJR Am J Roentgenol* 2000; 174(6):1699-705. doi: 10.2214/ajr.174.6.1741699.
- Antonacci MD, Hanson DS, Leblanc A, Heggeness MH. Regional variation in vertebral bone density and trabecular architecture are influenced by osteoarthritic change and osteoporosis. *Spine (Phila Pa 1976)* 1997; 22(20):2393-401; discussion 401-2. doi: 10.1097/0000 7632-199710150-00014.
- 22. Briggs AM, Wark JD, Greig AM, Fazzalari NL, Kantor S, Bennell KL. Subregional bone mineral density measurement in the lumbar spine using DXA: Potential for the application to osteoporosis and vertebral fractures. In: Mattingly BE, Pillare AC, (eds) Osteoporosis: Etiology, Diagnosis and Treatment. Nova Publishers, New York, 2009; 1-50.
- Cvijanovic O, Bobinac D, Zoricic S, Ostojic Z, Maric I, Crncevic-Orlic Z. *et al.* Age- and region-dependent changes in human lumbar vertebral bone: a histomorphometric study. *Spine (Phila Pa 1976)* 2004; 29(21):2370-5. doi: 10.1097/01.brs. 0000143620. 9526 7.39.
- Keller TS, Moeljanto E, Main JA, Spengler DM. Distribution and orientation of bone in the human lumbar vertebral centrum. *J Spinal Disord* 1992; 5(1):60-74. doi: 10.1097/00002517-199203000-00008.

- Nepper-Rasmussen J, Mosekilde L. Local differences in mineral content in vertebral trabecular bone measured by dual-energy computed tomography. *Acta Radiol* 1989; 30(4):369-71.
- Oda K, Shibayama Y, Abe M, Onomura T. Morphogenesis of vertebral deformities in involutional osteoporosis. Age-related, three-dimensional trabecular structure. *Spine (Phila Pa 1976)* 1998; 23(9):1050-5. doi: 10.1097/00007632-199805010-00016.
- Kim DG, Hunt CA, Zauel R, Fyhrie DP, Yeni YN. The effect of regional variations of the trabecular bone properties on the compressive strength of human vertebral bodies. *Ann Biomed Eng* 2007; 35(11):1907-13. doi: 10.1007/s10439-007-9363-1.
- Briggs AM, Wark JD, Kantor S, Teh R, Greig AM, Fazzalari NL. *et al.* In vivo intrarater and interrater precision of measuring apparent bone mineral density in vertebral subregions using supine lateral dual-energy x-ray absorptiometry. *J Clin Densitom* 2005; 8(3):314-9. doi: 10.1385/jcd:8:3:314.
- 29. Sran MM, Khan KM, Keiver K, Chew JB, McKay HA, Oxland TR. Accuracy of DXA scanning of the thoracic spine: cadaveric studies comparing BMC, areal BMD and geometric estimates of volumetric BMD against ash weight and CT measures of bone volume. *Eur Spine J* 2005; 14(10):971-6. doi: 10.1007/s00586-004-0836-8.
- Homminga J, Weinans H, Gowin W, Felsenberg D, Huiskes R. Osteoporosis changes the amount of vertebral trabecular bone at risk of fracture but not the

vertebral load distribution. *Spine (Phila Pa 1976)* 2001; 26(14):1555-61. doi: 10.1097/00007632-200 107150-00010.

- Meng X-l, Rosenthal R, Rubin DB. Comparing correlated correlation coefficients. *Psychological Bulletin* 1992; 111(1):172-75. doi: 10.1037/0033-2909. 111.1.172.
- 32. Sornay-Rendu E, Boutroy S, Munoz F, Delmas PD. Alterations of cortical and trabecular architecture are associated with fractures in postmenopausal women, partially independent of decreased BMD measured by DXA: the OFELY study. *J Bone Miner Res* 2007; 22(3):425-33. doi: 10.1359/jbmr.061206.
- 33. Porter JL, Varacallo M. Osteoporosis. In: Stat Pearls. Stat Pearls Publishing Copyright © 2023, Stat Pearls Publishing LLC., Treasure Island (FL) ineligible companies. Disclosure: Matthew Varacallo declares no relevant financial relationships with ineligible companies; 2023.
- Utilization of DXA Bone Mineral Densitometry in Ontario: An Evidence-Based Analysis. Ont Health Technol Assess Ser 2006; 6(20):1-180.
- Bartl R. Monitoring of Patients on Treatment. In: Osteoporosis in Clinical Practice. Edited by Bartl R. Springer International Publishing, Cham; 2023: 121-25.
- Lewiecki EM. Osteoporosis Treatment Success and Failure. In: *Osteoporosis: A Clinical Casebook.* Edited by Cusano NE. Springer International Publishing, Cham; 2021: 185-95.