WHAT IS CLAIMED IS:

1. A conjugate having a formula

$$B\text{-}L\text{-}(D)_n$$

or a pharmaceutically acceptable salt thereof;

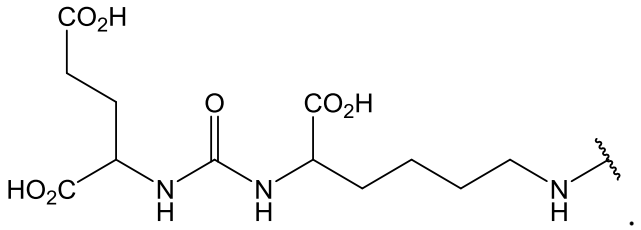
wherein B comprises a urea of lysine and glutamic acid;

wherein L is a non-releasable, polyvalent linker comprising a cycloalkylene, an alkylenecycloalkyl, or a cycloalkylenecarbonyl;

wherein D comprises a radioactive isotope of a metal coordinated to a chelating group; and

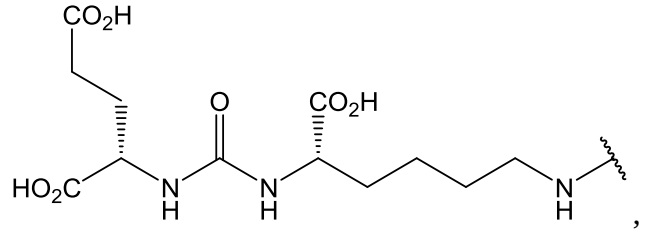
wherein n is 1.

2. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein B comprises a structure

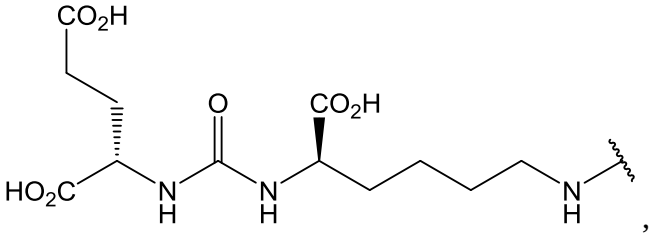


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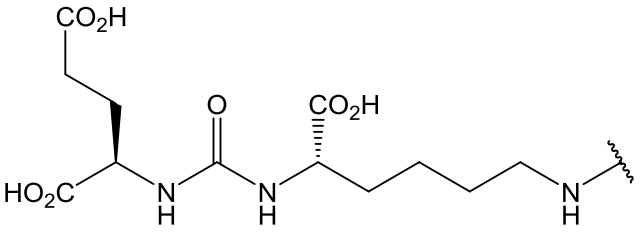
3. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of



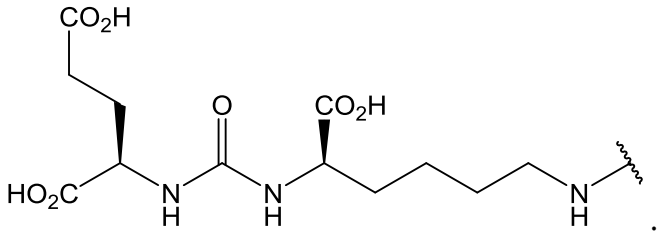
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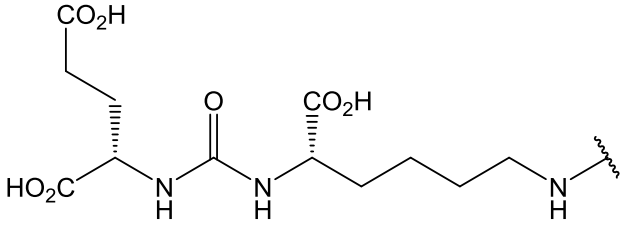


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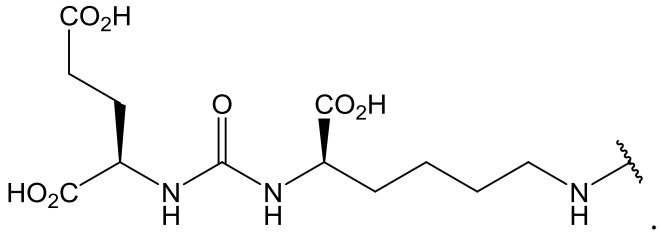


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4. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of

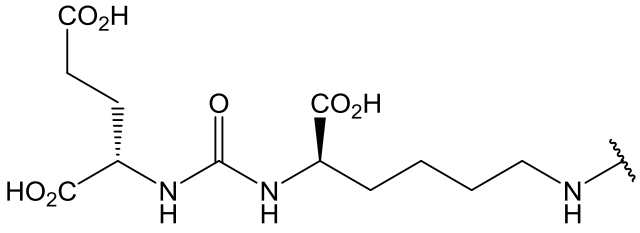


and

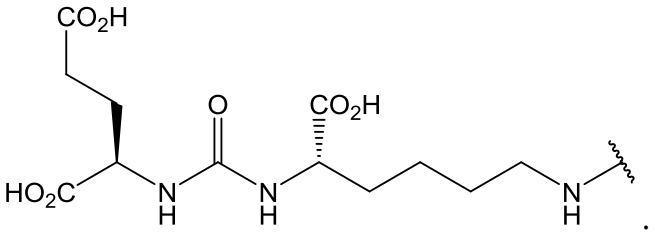


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5. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of



and



6. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein L forms an amide with the lysine.

7. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein L comprises a chain of at least about 7 atoms.

8. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein L comprises a chain of at least about 8 atoms.

9. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein L comprises a chain of at least about 9 atoms.

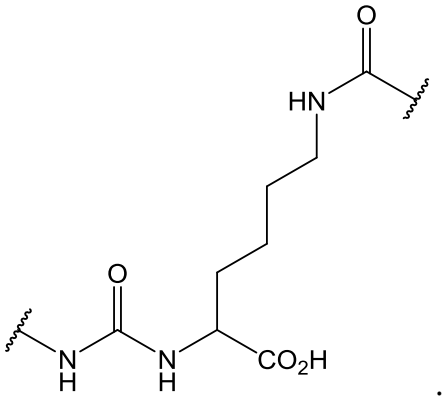
10. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein L comprises a chain of at least 10 atoms.

11. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein the linker comprises an alkylene, or an alkylenecarbonyl.

12. The conjugate of claim 11 or the pharmaceutically acceptable salt thereof, wherein the alkylene, or alkylenecarbonyl are each independently optionally substituted with one or more substituents X^1 selected from the group consisting of aryl, substituted aryl, arylalkyl, and substituted arylaklyl.

13. The conjugate of claim 12 or the pharmaceutically acceptable salt thereof, wherein X^1 comprises naphthyl.

14. The conjugate of claim 1 or the pharmaceutically acceptable salt thereof, wherein B-L comprises a diradical of the formula

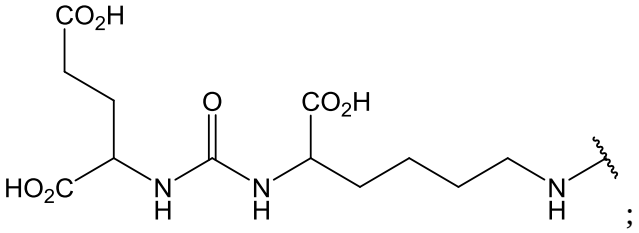


15. A conjugate having a formula

B-L-D

or a pharmaceutically acceptable salt thereof;

wherein B comprises a urea of lysine and glutamic acid, and wherein B comprises a structure



wherein L is a non-releasable, polyvalent linker, wherein L forms an amide with the lysine, and wherein L comprises a cycloalkylene, an alkylenecycloalkyl, or a cycloalkylenecarbonyl; and

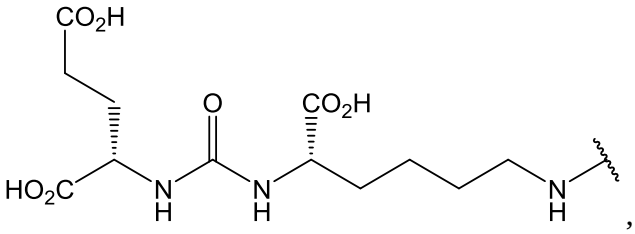
wherein D comprises a radioactive isotope of a metal coordinated to a chelating group.

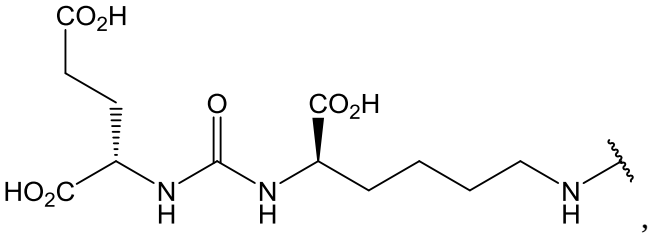
16. The conjugate of claim 15 or the pharmaceutically acceptable salt thereof, wherein the the cycloalkylene, the alkylenecycloalkyl, and the cycloalkylenecarbonyl are each independently optionally substituted with one or more substituents X^1 selected from the group consisting of aryl, substituted aryl, arylalkyl, and substituted arylaklyl.

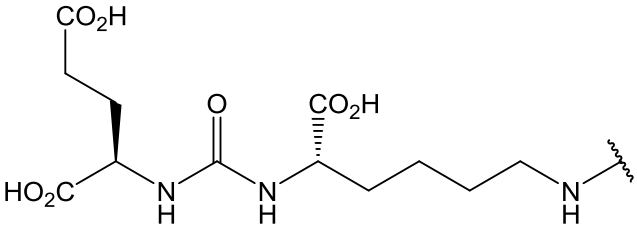
17. The conjugate of claim 16 or the pharmaceutically acceptable salt thereof, wherein X^1

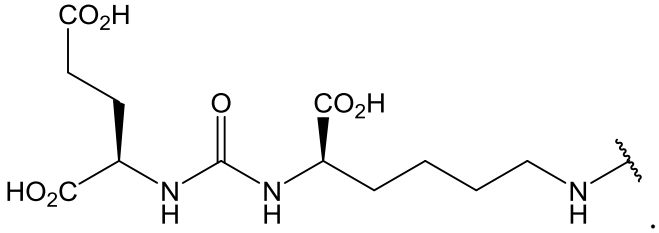
comprises naphthyl.

18. The conjugate of claim 15 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of

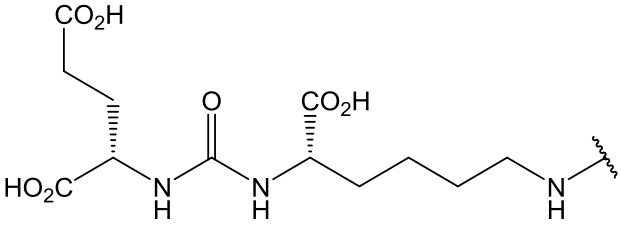

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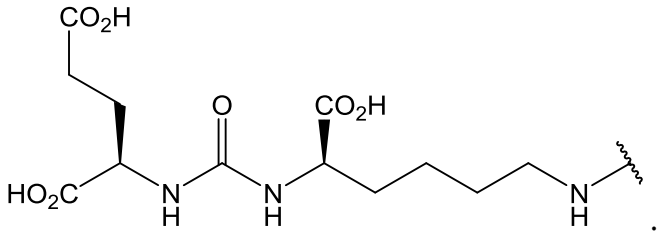

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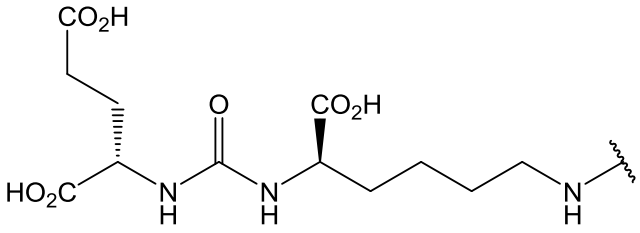

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19. The conjugate of claim 15 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of


and



20. The conjugate of claim 15 or the pharmaceutically acceptable salt thereof, wherein B is selected from the group consisting of



and

