

Applications

GRAS reagent crystallization screen for proteins, including monoclonal antibodies, where salt is the primary, and medium molecular weight Polyethylene glycol the secondary reagent, sampling pH 4-9.

Features

- Generally Recognized As Safe reagent formulation
- Samples pH 4.0 - 9.0 without an added buffer
- 24 unique salts - primary reagent
- Polyethylene glycol 1,000, MME 2000, 3,350, & 4,000 - secondary reagent
- Vapor diffusion, microbatch, free interface diffusion

Refer to the enclosed GRAS Screen 4 Reagent Formulation for more information.

General Description

GRAS Screen™ 4 was developed by Hampton Research for the crystallization of proteins, including monoclonal antibodies. Each of the chemicals in GRAS Screen 4 has been used under one or more of the following categories. As (1) a Generally Recognized As Safe (GRAS) substance, (2) a pharmaceutical excipient, (3) a normal physiological constituent, (4) a metabolic byproduct, and/or (5) a Everything Added to Food in the United States (EAFUS) substance. GRAS Screen 4 samples twenty four salts, encompassing pH 4-9 versus four medium molecular weight Polyethylene glycols (1,000, MME 2,000, 3,350 & 4,000). GRAS Screen 4 is supplied in a 96 Deep Well block format and is compatible with robotic and multi-channel pipet liquid handling systems. GRAS Screen 4 is compatible with vapor diffusion, free interface diffusion, and microbatch crystallization methods. For research use only.

Sample Preparation

The protein sample should be homogenous, as pure as is practically possible (>95%), and free of amorphous material. Remove amorphous material by centrifugation or microfiltration prior to use. The recommended sample concentration is 5 to 25 mg/ml in dilute (25 mM or less) buffer. For initial screens, the sample should be free of unnecessary additives in order to observe the effect of the GRAS Screen 4 reagents. However, agents that promote and preserve sample solubility, stability, and homogeneity can and should be included in the sample buffer. For additional sample preparation recommendations see Hampton Research Crystal Growth 101 - Preliminary Sample Preparation.

Preparing the Deep Well Block for Use

Allow the Deep Well Block and reagents to stabilize at room temperature, then centrifuge at 500 rpm for 5 minutes to remove stray drops from the film before removing the sealing film. The film can be removed by grasping a corner of the film and gently peeling the film from the plate. Alternatively, the film can be left intact, pierced to access reagents, and resealed using AlumaSeal II Sealing Film.

Performing the Screen

Automated Method - Sitting Drop Vapor Diffusion

The Deep Well block is compatible with the SBS standard 96 well microplate format and is compatible with numerous automated liquid handling systems that accept 8 x 12, 96 well assay blocks. Follow the automation manufacturer's recommendation for handling Deep Well blocks.

1. Using a 96 well sitting drop vapor diffusion plate, dispense the recommended volume (typically 50 to 100 microliters) of crystallization reagent from the Deep Well block into the reagent reservoirs of the crystallization plate.
2. Dispense the desired volume of crystallization reagent (typically 50 to 200 nanoliters) from the crystallization plate reservoir to the sitting drop well.
3. Transfer the equivalent volume of sample to the reagent drop in the sitting drop well.
4. Seal the crystallization plate using a clear sealing tape or film. View and score the experiment. See Hampton Research Crystal Growth 101 - Viewing Crystallization Experiments for more information.
5. Seal the remaining reagent in the Deep Well block using AlumaSeal II Sealing Film.

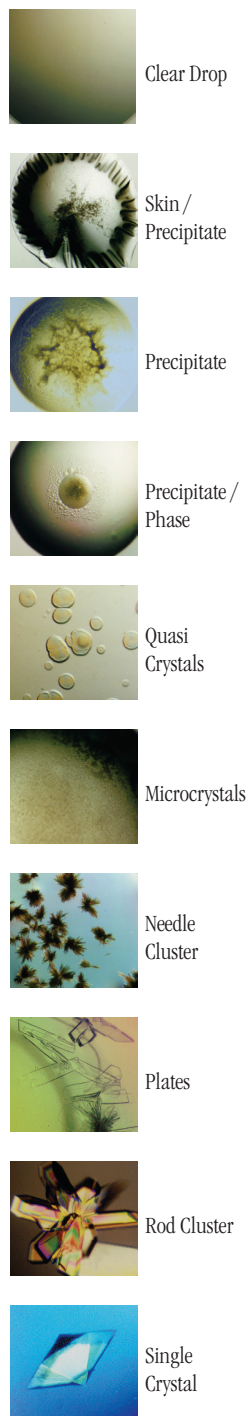
Manual Method - Sitting Drop Vapor Diffusion

1. Using a 96 well sitting drop vapor diffusion plate, pipet the recommended volume (typically 50 to 100 microliters) of crystallization reagent from the Deep Well block into the reagent reservoirs of the crystallization plate. The Deep Well block is compatible with 8, 12, and 96 channel automated and manual pipettors. Use clean pipet tips for each reagent set, transfer and change pipet tips when changing reagents. For an 8 channel pipet, transfer reagents A1-H1 to reservoirs A1-H1 of the crystallization plate. Repeat this procedure for reagent columns 2 through 12. Change pipet tips when moving between reagent columns. For a 12 channel pipet, transfer reagents A1-A12 to reservoirs A1-A12 of the crystallization plate. Repeat this procedure for reagent rows B through H.
2. Using clean pipet tips, pipet the desired volume of crystallization reagent (typically 0.05 to 2 microliters) from the crystallization plate reservoir to the sitting drop well. Some 96 well crystallization plates allow this procedure to be performed using a multichannel pipet where other plates require the use of a single channel pipet. Change the pipet tip between reagents.
3. Using a clean pipet tip, pipet the same volume (typically 0.05 to 2 microliters) of sample to the reagent drop in the sitting drop well. Work carefully but quickly to minimize evaporation from the crystallization plate.
4. Seal the crystallization plate using an optically clear sealing film or tape. Seal the remaining reagent in the Deep Well block using AlumaSeal II sealing film.

Examine the Drop

Carefully examine the drops under a stereo microscope (10 to 100x magnification) after setting the screen. Record all observations and be particularly careful to scan the focal plane for small crystals. Observe the drops once each day for the first week, then once a week thereafter for up to 60 days, or until the drop dries out. Records should indicate whether the drop is clear, contains precipitate, and/or crystals. It is helpful to describe the drop contents using descriptive terms. Adding magnitude is also helpful. Example: 4+ yellow/brown fine precipitate, 3+ needle shaped crystals in 1+ white precipitate. One may

Figure 1
Typical observations in a crystallization experiment



also employ a numerical scoring scheme (Clear = 0, Crystal = 1, Precipitate = 2). Figure 1 shows typical examples of what one might observe in a crystallization experiment.

Interpreting GRAS Screen 4

Clear drops indicate that either the relative supersaturation of the sample and reagent is too low or the drop has not yet completed equilibration. If the drop remains clear after 3 to 4 weeks consider repeating the screen condition and doubling the sample concentration. If more than 70 of the 96 drops are clear, then consider doubling the sample concentration and repeating the entire screen.

Drops containing precipitate indicate either the relative supersaturation of the sample and reagent is too high, the sample has denatured, or the sample is heterogeneous. To reduce the relative supersaturation, dilute the sample twofold with sample buffer and repeat the screen condition. If more than 70 of the 96 drops contain precipitate and no crystals are present, then consider diluting the sample concentration in half by adding an equal volume of sample buffer to the sample and repeating the entire screen. If sample denaturation is suspect, take measures to stabilize the sample (add reducing agent, ligands, additives, salt, or other stabilizing agents). If the sample is impure, aggregated, or heterogeneous take measures to pursue homogeneity. It is possible to obtain crystals from precipitate so do not discard nor ignore a drop containing precipitate. If possible, examine drops containing precipitate under polarizing or UV optics to differentiate precipitate from microcrystals.

If the drop contains a macromolecular crystal the relative supersaturation of the sample and reagent is appropriate for crystal nucleation and growth. The next step is to optimize the preliminary conditions by varying salt and/or PEG concentration, screen pH, vary temperature between 4 and 30°C, screen additives, and evaluate other crystallization variables including sample construct, purity, stability, and homogeneity in order to achieve the desired crystal size and quality.

When sample quantity permits, set GRAS Screen 4 in duplicate (4°C and 25°C) or triplicate (10°C and 20°C and 30°C) to evaluate the effect of temperature on crystallization. Compare the observations between the different temperatures to determine the effect of temperature on sample solubility. Different results in the same drops at different temperatures indicate that sample solubility is temperature dependent and that one should include temperature as a variable in subsequent screens and optimization experiments.

When sample quantity permits, set GRAS Screen 4 using multiple drops and drop ratios, such as 1:2, 1:1, and 2:1. See Hampton Research Crystal Growth 101: Drop Ratio for details.

GRAS Screen 4 Formulation

Crystallization reagents are formulated using the highest purity chemicals, ultrapure water (Formulated in Type 1+ ultrapure water: 18.2 megaohm-cm resistivity at 25°C, < 5 ppb Total Organic Carbon, bacteria free (<1 Bacteria (CFU/ml)), pyrogen free (<0.03 Endotoxin (EU/ml)), RNase-free (< 0.01 ng/mL) and DNase-free (< 4 pg/μL)) and are sterile filtered using 0.22 micron filters into sterile Deep Well blocks (no preservatives added). Store at -20°C. Best if used within 12 months of receipt.

Crystallization reagents can be reproduced using Hampton Research Optimize™ and StockOptions™ polyethylene glycols and salts.

Recommended Reading

1. Introduction to protein crystallization. Alexander McPherson and Jose A. Gavira. *Acta Crystallographica Section F* Volume 70, Issue 1, pages 2–20, January 2014.
2. Optimization of crystallization conditions for biological macromolecules. Alexander McPherson and Bob Cudney. *Acta Crystallographica Section F* Volume 70, Issue 11, pages 1445–1467, November 2014.
3. Crystallization of intact monoclonal antibodies. Harris LJ, Skaltsky E, McPherson A. *Proteins*. 1995 Oct;23(2):285-9.
4. Crystalline monoclonal antibodies for subcutaneous delivery. Yang MX1, Shenoy B, Distler M, Patel R, McGrath M, Pechenov S, Margolin AL. *Proc Natl Acad Sci U S A*. 2003 Jun 10;100(12):6934-9.
5. Fast and Scalable Purification of a Therapeutic Full-Length Antibody Based on Process Crystallization. Dariusch Hekmat et al, *Biotechnology and Bioengineering*, Vol. 110, No. 9, September, 2013.
6. Towards Protein Crystallization as a Process Step in Downstream Processing of Therapeutic Antibodies: Screening and Optimization at Microbatch Scale. Yuguo Zang et al, *PLoS One*. 2011; 6(9): e25282.
7. Crystallization and Liquid-Liquid Phase Separation of Monoclonal Antibodies and Fc-Fusion Proteins: Screening Results. Suresh Vunnum et al, *Biotechnol Prog*. 2011 Jul;27(4):1054-67.

Hampton Research
34 Journey

Aliso Viejo, CA 92656-3317 U.S.A.

Tel: (949) 425-1321 • Fax: (949) 425-1611

Technical Support e-mail: tech@hrmail.com

Website: www.hamptonresearch.com

Well #	Salt	Well #	PEG	Well #	pH ◇
1. (A1)	3.0 M Ammonium acetate	1. (A1)	2% w/v Polyethylene glycol 1,000	1. (A1)	7.3
2. (A2)	3.0 M Ammonium acetate	2. (A2)	2% w/v Polyethylene glycol monomethyl ether 2,000	2. (A2)	7.3
3. (A3)	3.0 M Ammonium acetate	3. (A3)	2% w/v Polyethylene glycol 3,350	3. (A3)	7.4
4. (A4)	3.0 M Ammonium acetate	4. (A4)	2% w/v Polyethylene glycol 4,000	4. (A4)	7.4
5. (A5)	3.0 M Ammonium chloride	5. (A5)	2% w/v Polyethylene glycol 1,000	5. (A5)	4.5
6. (A6)	3.0 M Ammonium chloride	6. (A6)	2% w/v Polyethylene glycol monomethyl ether 2,000	6. (A6)	4.8
7. (A7)	3.0 M Ammonium chloride	7. (A7)	2% w/v Polyethylene glycol 3,350	7. (A7)	4.8
8. (A8)	3.0 M Ammonium chloride	8. (A8)	2% w/v Polyethylene glycol 4,000	8. (A8)	4.5
9. (A9)	1.5 M Ammonium citrate dibasic	9. (A9)	2% w/v Polyethylene glycol 1,000	9. (A9)	4.8
10. (A10)	1.5 M Ammonium citrate dibasic	10. (A10)	2% w/v Polyethylene glycol monomethyl ether 2,000	10. (A10)	4.8
11. (A11)	1.5 M Ammonium citrate dibasic	11. (A11)	2% w/v Polyethylene glycol 3,350	11. (A11)	4.8
12. (A12)	1.5 M Ammonium citrate dibasic	12. (A12)	2% w/v Polyethylene glycol 4,000	12. (A12)	4.8
13. (B1)	0.8 M Ammonium citrate tribasic	13. (B1)	2% w/v Polyethylene glycol 1,000	13. (B1)	7.3
14. (B2)	0.8 M Ammonium citrate tribasic	14. (B2)	2% w/v Polyethylene glycol monomethyl ether 2,000	14. (B2)	7.3
15. (B3)	0.8 M Ammonium citrate tribasic	15. (B3)	2% w/v Polyethylene glycol 3,350	15. (B3)	7.3
16. (B4)	0.8 M Ammonium citrate tribasic	16. (B4)	2% w/v Polyethylene glycol 4,000	16. (B4)	7.2
17. (B5)	6.0 M Ammonium formate	17. (B5)	2% w/v Polyethylene glycol 1,000	17. (B5)	6.8
18. (B6)	6.0 M Ammonium formate	18. (B6)	2% w/v Polyethylene glycol monomethyl ether 2,000	18. (B6)	6.8
19. (B7)	6.0 M Ammonium formate	19. (B7)	2% w/v Polyethylene glycol 3,350	19. (B7)	6.8
20. (B8)	6.0 M Ammonium formate	20. (B8)	2% w/v Polyethylene glycol 4,000	20. (B8)	6.8
21. (B9)	0.9 M Ammonium phosphate dibasic	21. (B9)	2% w/v Polyethylene glycol 1,000	21. (B9)	7.9
22. (B10)	0.9 M Ammonium phosphate dibasic	22. (B10)	2% w/v Polyethylene glycol monomethyl ether 2,000	22. (B10)	7.9
23. (B11)	0.9 M Ammonium phosphate dibasic	23. (B11)	2% w/v Polyethylene glycol 3,350	23. (B11)	7.9
24. (B12)	0.9 M Ammonium phosphate dibasic	24. (B12)	2% w/v Polyethylene glycol 4,000	24. (B12)	7.9
25. (C1)	1.5 M Ammonium phosphate monobasic	25. (C1)	2% w/v Polyethylene glycol 1,000	25. (C1)	3.9
26. (C2)	1.5 M Ammonium phosphate monobasic	26. (C2)	2% w/v Polyethylene glycol monomethyl ether 2,000	26. (C2)	3.9
27. (C3)	1.5 M Ammonium phosphate monobasic	27. (C3)	2% w/v Polyethylene glycol 3,350	27. (C3)	3.9
28. (C4)	1.5 M Ammonium phosphate monobasic	28. (C4)	2% w/v Polyethylene glycol 4,000	28. (C4)	3.9
29. (C5)	1.1 M Ammonium sulfate	29. (C5)	2% w/v Polyethylene glycol 1,000	29. (C5)	5.2
30. (C6)	1.1 M Ammonium sulfate	30. (C6)	2% w/v Polyethylene glycol monomethyl ether 2,000	30. (C6)	5.3
31. (C7)	1.1 M Ammonium sulfate	31. (C7)	2% w/v Polyethylene glycol 3,350	31. (C7)	5.3
32. (C8)	1.1 M Ammonium sulfate	32. (C8)	2% w/v Polyethylene glycol 4,000	32. (C8)	5.2
33. (C9)	1.2 M Ammonium tartrate dibasic	33. (C9)	2% w/v Polyethylene glycol 1,000	33. (C9)	6.8
34. (C10)	1.2 M Ammonium tartrate dibasic	34. (C10)	2% w/v Polyethylene glycol monomethyl ether 2,000	34. (C10)	6.8
35. (C11)	1.2 M Ammonium tartrate dibasic	35. (C11)	2% w/v Polyethylene glycol 3,350	35. (C11)	6.7
36. (C12)	1.2 M Ammonium tartrate dibasic	36. (C12)	2% w/v Polyethylene glycol 4,000	36. (C12)	6.8
37. (D1)	1.0 M DL-Malic acid pH 7.0	37. (D1)	2% w/v Polyethylene glycol 1,000	37. (D1)	6.6
38. (D2)	1.0 M DL-Malic acid pH 7.0	38. (D2)	2% w/v Polyethylene glycol monomethyl ether 2,000	38. (D2)	6.9
39. (D3)	1.0 M DL-Malic acid pH 7.0	39. (D3)	2% w/v Polyethylene glycol 3,350	39. (D3)	6.8
40. (D4)	1.0 M DL-Malic acid pH 7.0	40. (D4)	2% w/v Polyethylene glycol 4,000	40. (D4)	6.8
41. (D5)	0.7 M Potassium phosphate dibasic	41. (D5)	2% w/v Polyethylene glycol 1,000	41. (D5)	9.2
42. (D6)	0.7 M Potassium phosphate dibasic	42. (D6)	2% w/v Polyethylene glycol monomethyl ether 2,000	42. (D6)	9.2
43. (D7)	0.7 M Potassium phosphate dibasic	43. (D7)	2% w/v Polyethylene glycol 3,350	43. (D7)	9.2
44. (D8)	0.7 M Potassium phosphate dibasic	44. (D8)	2% w/v Polyethylene glycol 4,000	44. (D8)	9.2
45. (D9)	0.9 M Potassium sodium tartrate tetrahydrate	45. (D9)	2% w/v Polyethylene glycol 1,000	45. (D9)	8.4
46. (D10)	0.9 M Potassium sodium tartrate tetrahydrate	46. (D10)	2% w/v Polyethylene glycol monomethyl ether 2,000	46. (D10)	8.5
47. (D11)	0.9 M Potassium sodium tartrate tetrahydrate	47. (D11)	2% w/v Polyethylene glycol 3,350	47. (D11)	8.2
48. (D12)	0.9 M Potassium sodium tartrate tetrahydrate	48. (D12)	2% w/v Polyethylene glycol 4,000	48. (D12)	8.4

Reagents formulated in Type 1+ ultrapure grade water

◇ Measured pH at 25°C, no pH adjustment made to reagent

Well #	Salt	Well #	PEG	Well #	pH ◇
49. (E1)	1.8 M Sodium acetate trihydrate	49. (E1)	2% w/v Polyethylene glycol 1,000	49. (E1)	9.1
50. (E2)	1.8 M Sodium acetate trihydrate	50. (E2)	2% w/v Polyethylene glycol monomethyl ether 2,000	50. (E2)	9.3
51. (E3)	1.8 M Sodium acetate trihydrate	51. (E3)	2% w/v Polyethylene glycol 3,350	51. (E3)	9.2
52. (E4)	1.8 M Sodium acetate trihydrate	52. (E4)	2% w/v Polyethylene glycol 4,000	52. (E4)	9.3
53. (E5)	3.0 M Sodium chloride	53. (E5)	2% w/v Polyethylene glycol 1,000	53. (E5)	5.9
54. (E6)	3.0 M Sodium chloride	54. (E6)	2% w/v Polyethylene glycol monomethyl ether 2,000	54. (E6)	5.9
55. (E7)	3.0 M Sodium chloride	55. (E7)	2% w/v Polyethylene glycol 3,350	55. (E7)	5.9
56. (E8)	3.0 M Sodium chloride	56. (E8)	2% w/v Polyethylene glycol 4,000	56. (E8)	6.2
57. (E9)	0.6 M Sodium citrate tribasic dihydrate	57. (E9)	2% w/v Polyethylene glycol 1,000	57. (E9)	8.3
58. (E10)	0.6 M Sodium citrate tribasic dihydrate	58. (E10)	2% w/v Polyethylene glycol monomethyl ether 2,000	58. (E10)	8.3
59. (E11)	0.6 M Sodium citrate tribasic dihydrate	59. (E11)	2% w/v Polyethylene glycol 3,350	59. (E11)	8.3
60. (E12)	0.6 M Sodium citrate tribasic dihydrate	60. (E12)	2% w/v Polyethylene glycol 4,000	60. (E12)	8.3
61. (F1)	3.0 M Sodium formate	61. (F1)	2% w/v Polyethylene glycol 1,000	61. (F1)	8.0
62. (F2)	3.0 M Sodium formate	62. (F2)	2% w/v Polyethylene glycol monomethyl ether 2,000	62. (F2)	8.1
63. (F3)	3.0 M Sodium formate	63. (F3)	2% w/v Polyethylene glycol 3,350	63. (F3)	8.0
64. (F4)	3.0 M Sodium formate	64. (F4)	2% w/v Polyethylene glycol 4,000	64. (F4)	8.0
65. (F5)	0.9 M Sodium malonate pH 7.0	65. (F5)	2% w/v Polyethylene glycol 1,000	65. (F5)	6.9
66. (F6)	0.9 M Sodium malonate pH 7.0	66. (F6)	2% w/v Polyethylene glycol monomethyl ether 2,000	66. (F6)	6.9
67. (F7)	0.9 M Sodium malonate pH 7.0	67. (F7)	2% w/v Polyethylene glycol 3,350	67. (F7)	6.9
68. (F8)	0.9 M Sodium malonate pH 7.0	68. (F8)	2% w/v Polyethylene glycol 4,000	68. (F8)	6.9
69. (F9)	0.6 M Sodium phosphate dibasic dihydrate	69. (F9)	2% w/v Polyethylene glycol 1,000	69. (F9)	8.9
70. (F10)	0.6 M Sodium phosphate dibasic dihydrate	70. (F10)	2% w/v Polyethylene glycol monomethyl ether 2,000	70. (F10)	8.9
71. (F11)	0.6 M Sodium phosphate dibasic dihydrate	71. (F11)	2% w/v Polyethylene glycol 3,350	71. (F11)	8.9
72. (F12)	0.6 M Sodium phosphate dibasic dihydrate	72. (F12)	2% w/v Polyethylene glycol 4,000	72. (F12)	8.9
73. (G1)	1.6 M Sodium phosphate monobasic monohydrate	73. (G1)	2% w/v Polyethylene glycol 1,000	73. (G1)	3.9
74. (G2)	1.6 M Sodium phosphate monobasic monohydrate	74. (G2)	2% w/v Polyethylene glycol monomethyl ether 2,000	74. (G2)	3.9
75. (G3)	1.6 M Sodium phosphate monobasic monohydrate	75. (G3)	2% w/v Polyethylene glycol 3,350	75. (G3)	3.9
76. (G4)	1.6 M Sodium phosphate monobasic monohydrate	76. (G4)	2% w/v Polyethylene glycol 4,000	76. (G4)	3.9
77. (G5)	0.6 M Sodium sulfate decahydrate	77. (G5)	2% w/v Polyethylene glycol 1,000	77. (G5)	5.8
78. (G6)	0.6 M Sodium sulfate decahydrate	78. (G6)	2% w/v Polyethylene glycol monomethyl ether 2,000	78. (G6)	5.9
79. (G7)	0.6 M Sodium sulfate decahydrate	79. (G7)	2% w/v Polyethylene glycol 3,350	79. (G7)	6.2
80. (G8)	0.6 M Sodium sulfate decahydrate	80. (G8)	2% w/v Polyethylene glycol 4,000	80. (G8)	5.9
81. (G9)	0.9 M Sodium tartrate dibasic dihydrate	81. (G9)	2% w/v Polyethylene glycol 1,000	81. (G9)	8.1
82. (G10)	0.9 M Sodium tartrate dibasic dihydrate	82. (G10)	2% w/v Polyethylene glycol monomethyl ether 2,000	82. (G10)	8.1
83. (G11)	0.9 M Sodium tartrate dibasic dihydrate	83. (G11)	2% w/v Polyethylene glycol 3,350	83. (G11)	8.1
84. (G12)	0.9 M Sodium tartrate dibasic dihydrate	84. (G12)	2% w/v Polyethylene glycol 4,000	84. (G12)	8.2
85. (H1)	4.8 M Sodium thiocyanate	85. (H1)	2% w/v Polyethylene glycol 1,000	85. (H1)	7.8
86. (H2)	4.8 M Sodium thiocyanate	86. (H2)	2% w/v Polyethylene glycol monomethyl ether 2,000	86. (H2)	7.6
87. (H3)	4.8 M Sodium thiocyanate	87. (H3)	2% w/v Polyethylene glycol 3,350	87. (H3)	7.2
88. (H4)	4.8 M Sodium thiocyanate	88. (H4)	2% w/v Polyethylene glycol 4,000	88. (H4)	8.0
89. (H5)	0.7 M Succinic acid pH 7.0	89. (H5)	2% w/v Polyethylene glycol 1,000	89. (H5)	7.0
90. (H6)	0.7 M Succinic acid pH 7.0	90. (H6)	2% w/v Polyethylene glycol monomethyl ether 2,000	90. (H6)	7.0
91. (H7)	0.7 M Succinic acid pH 7.0	91. (H7)	2% w/v Polyethylene glycol 3,350	91. (H7)	7.0
92. (H8)	0.7 M Succinic acid pH 7.0	92. (H8)	2% w/v Polyethylene glycol 4,000	92. (H8)	7.0
93. (H9)	30% v/v Tacsimate™ pH 7.0	93. (H9)	2% w/v Polyethylene glycol 1,000	93. (H9)	6.9
94. (H10)	30% v/v Tacsimate™ pH 7.0	94. (H10)	2% w/v Polyethylene glycol monomethyl ether 2,000	94. (H10)	6.9
95. (H11)	30% v/v Tacsimate™ pH 7.0	95. (H11)	2% w/v Polyethylene glycol 3,350	95. (H11)	6.9
96. (H12)	30% v/v Tacsimate™ pH 7.0	96. (H12)	2% w/v Polyethylene glycol 4,000	96. (H12)	6.9

Reagents formulated in Type 1+ ultrapure grade water

◇ Measured pH at 25°C, no pH adjustment made to reagent

Sample: _____ Sample Concentration: _____
 Sample Buffer: _____ Date: _____
 Reservoir Volume: _____ Temperature: _____
 Drop Volume: Total _____ μ l Sample _____ μ l Reservoir _____ μ l Additive _____ μ l

- 1 Clear Drop
- 2 Phase Separation
- 3 Regular Granular Precipitate
- 4 Birefringent Precipitate or Microcrystals
- 5 Posettes or Spherulites
- 6 Needles (1D Growth)
- 7 Plates (2D Growth)
- 8 Single Crystals (3D Growth < 0.2 mm)
- 9 Single Crystals (3D Growth > 0.2 mm)

GRAS Screen™ 4 - HR2-454 Scoring Sheet

Date: _____ Date: _____ Date: _____ Date: _____

1. (A1)	3.0 M Ammonium acetate, 2% w/v Polyethylene glycol 1,000				
2. (A2)	3.0 M Ammonium acetate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
3. (A3)	3.0 M Ammonium acetate, 2% w/v Polyethylene glycol 3,350				
4. (A4)	3.0 M Ammonium acetate, 2% w/v Polyethylene glycol 4,000				
5. (A5)	3.0 M Ammonium chloride, 2% w/v Polyethylene glycol 1,000				
6. (A6)	3.0 M Ammonium chloride, 2% w/v Polyethylene glycol monomethyl ether 2,000				
7. (A7)	3.0 M Ammonium chloride, 2% w/v Polyethylene glycol 3,350				
8. (A8)	3.0 M Ammonium chloride, 2% w/v Polyethylene glycol 4,000				
9. (A9)	1.5 M Ammonium citrate dibasic, 2% w/v Polyethylene glycol 1,000				
10. (A10)	1.5 M Ammonium citrate dibasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
11. (A11)	1.5 M Ammonium citrate dibasic, 2% w/v Polyethylene glycol 3,350				
12. (A12)	1.5 M Ammonium citrate dibasic, 2% w/v Polyethylene glycol 4,000				
13. (B1)	0.8 M Ammonium citrate tribasic, 2% w/v Polyethylene glycol 1,000				
14. (B2)	0.8 M Ammonium citrate tribasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
15. (B3)	0.8 M Ammonium citrate tribasic, 2% w/v Polyethylene glycol 3,350				
16. (B4)	0.8 M Ammonium citrate tribasic, 2% w/v Polyethylene glycol 4,000				
17. (B5)	6.0 M Ammonium formate, 2% w/v Polyethylene glycol 1,000				
18. (B6)	6.0 M Ammonium formate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
19. (B7)	6.0 M Ammonium formate, 2% w/v Polyethylene glycol 3,350				
20. (B8)	6.0 M Ammonium formate, 2% w/v Polyethylene glycol 4,000				
21. (B9)	0.9 M Ammonium phosphate dibasic, 2% w/v Polyethylene glycol 1,000				
22. (B10)	0.9 M Ammonium phosphate dibasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
23. (B11)	0.9 M Ammonium phosphate dibasic, 2% w/v Polyethylene glycol 3,350				
24. (B12)	0.9 M Ammonium phosphate dibasic, 2% w/v Polyethylene glycol 4,000				
25. (C1)	1.5 M Ammonium phosphate monobasic, 2% w/v Polyethylene glycol 1,000				
26. (C2)	1.5 M Ammonium phosphate monobasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
27. (C3)	1.5 M Ammonium phosphate monobasic, 2% w/v Polyethylene glycol 3,350				
28. (C4)	1.5 M Ammonium phosphate monobasic, 2% w/v Polyethylene glycol 4,000				
29. (C5)	1.1 M Ammonium sulfate, 2% w/v Polyethylene glycol 1,000				
30. (C6)	1.1 M Ammonium sulfate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
31. (C7)	1.1 M Ammonium sulfate, 2% w/v Polyethylene glycol 3,350				
32. (C8)	1.1 M Ammonium sulfate, 2% w/v Polyethylene glycol 4,000				
33. (C9)	1.2 M Ammonium tartrate dibasic, 2% w/v Polyethylene glycol 1,000				
34. (C10)	1.2 M Ammonium tartrate dibasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
35. (C11)	1.2 M Ammonium tartrate dibasic, 2% w/v Polyethylene glycol 3,350				
36. (C12)	1.2 M Ammonium tartrate dibasic, 2% w/v Polyethylene glycol 4,000				
37. (D1)	1.0 M DL-Malic acid pH 7.0, 2% w/v Polyethylene glycol 1,000				
38. (D2)	1.0 M DL-Malic acid pH 7.0, 2% w/v Polyethylene glycol monomethyl ether 2,000				
39. (D3)	1.0 M DL-Malic acid pH 7.0, 2% w/v Polyethylene glycol 3,350				
40. (D4)	1.0 M DL-Malic acid pH 7.0, 2% w/v Polyethylene glycol 4,000				
41. (D5)	0.7 M Potassium phosphate dibasic, 2% w/v Polyethylene glycol 1,000				
42. (D6)	0.7 M Potassium phosphate dibasic, 2% w/v Polyethylene glycol monomethyl ether 2,000				
43. (D7)	0.7 M Potassium phosphate dibasic, 2% w/v Polyethylene glycol 3,350				
44. (D8)	0.7 M Potassium phosphate dibasic, 2% w/v Polyethylene glycol 4,000				
45. (D9)	0.9 M Potassium sodium tartrate tetrahydrate, 2% w/v Polyethylene glycol 1,000				
46. (D10)	0.9 M Potassium sodium tartrate tetrahydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
47. (D11)	0.9 M Potassium sodium tartrate tetrahydrate, 2% w/v Polyethylene glycol 3,350				
48. (D12)	0.9 M Potassium sodium tartrate tetrahydrate, 2% w/v Polyethylene glycol 4,000				



Solutions for Crystal Growth

34 Journey
 Aliso Viejo, CA 92656-3317 U.S.A.
 Tel: (949) 425-1321 • Fax: (949) 425-1611
 e-mail: tech@hamptron.com
 Website: www.hamptronresearch.com

© 1991 - 2019 Hampton Research Corp. all rights reserved
 Printed in the United States of America. This guide or parts thereof may not
 be reproduced in any form without the written permission of the publishers.

Sample: _____ Sample Concentration: _____
 Sample Buffer: _____ Date: _____
 Reservoir Volume: _____ Temperature: _____
 Drop Volume: Total _____ μ l Sample _____ μ l Reservoir _____ μ l Additive _____ μ l

- 1 Clear Drop
- 2 Phase Separation
- 3 Regular Granular Precipitate
- 4 Birefringent Precipitate or Microcrystals

- 5 Posettes or Spherulites
- 6 Needles (1D Growth)
- 7 Plates (2D Growth)
- 8 Single Crystals (3D Growth < 0.2 mm)
- 9 Single Crystals (3D Growth > 0.2 mm)

GRAS Screen™ 4 - HR2-454 Scoring Sheet

Date: _____ Date: _____ Date: _____ Date: _____

49. (E1)	1.8 M Sodium acetate trihydrate, 2% w/v Polyethylene glycol 1,000				
50. (E2)	1.8 M Sodium acetate trihydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
51. (E3)	1.8 M Sodium acetate trihydrate, 2% w/v Polyethylene glycol 3,350				
52. (E4)	1.8 M Sodium acetate trihydrate, 2% w/v Polyethylene glycol 4,000				
53. (E5)	3.0 M Sodium chloride, 2% w/v Polyethylene glycol 1,000				
54. (E6)	3.0 M Sodium chloride, 2% w/v Polyethylene glycol monomethyl ether 2,000				
55. (E7)	3.0 M Sodium chloride, 2% w/v Polyethylene glycol 3,350				
56. (E8)	3.0 M Sodium chloride, 2% w/v Polyethylene glycol 4,000				
57. (E9)	0.6 M Sodium citrate tribasic dihydrate, 2% w/v Polyethylene glycol 1,000				
58. (E10)	0.6 M Sodium citrate tribasic dihydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
59. (E11)	0.6 M Sodium citrate tribasic dihydrate, 2% w/v Polyethylene glycol 3,350				
60. (E12)	0.6 M Sodium citrate tribasic dihydrate, 2% w/v Polyethylene glycol 4,000				
61. (F1)	3.0 M Sodium formate, 2% w/v Polyethylene glycol 1,000				
62. (F2)	3.0 M Sodium formate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
63. (F3)	3.0 M Sodium formate, 2% w/v Polyethylene glycol 3,350				
64. (F4)	3.0 M Sodium formate, 2% w/v Polyethylene glycol 4,000				
65. (F5)	0.9 M Sodium malonate pH 7.0, 2% w/v Polyethylene glycol 1,000				
66. (F6)	0.9 M Sodium malonate pH 7.0, 2% w/v Polyethylene glycol monomethyl ether 2,000				
67. (F7)	0.9 M Sodium malonate pH 7.0, 2% w/v Polyethylene glycol 3,350				
68. (F8)	0.9 M Sodium malonate pH 7.0, 2% w/v Polyethylene glycol 4,000				
69. (F9)	0.6 M Sodium phosphate dibasic dihydrate, 2% w/v Polyethylene glycol 1,000				
70. (F10)	0.6 M Sodium phosphate dibasic dihydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
71. (F11)	0.6 M Sodium phosphate dibasic dihydrate, 2% w/v Polyethylene glycol 3,350				
72. (F12)	0.6 M Sodium phosphate dibasic dihydrate, 2% w/v Polyethylene glycol 4,000				
73. (G1)	1.6 M Sodium phosphate monobasic monohydrate, 2% w/v Polyethylene glycol 1,000				
74. (G2)	1.6 M Sodium phosphate monobasic monohydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
75. (G3)	1.6 M Sodium phosphate monobasic monohydrate, 2% w/v Polyethylene glycol 3,350				
76. (G4)	1.6 M Sodium phosphate monobasic monohydrate, 2% w/v Polyethylene glycol 4,000				
77. (G5)	0.6 M Sodium sulfate decahydrate, 2% w/v Polyethylene glycol 1,000				
78. (G6)	0.6 M Sodium sulfate decahydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
79. (G7)	0.6 M Sodium sulfate decahydrate, 2% w/v Polyethylene glycol 3,350				
80. (G8)	0.6 M Sodium sulfate decahydrate, 2% w/v Polyethylene glycol 4,000				
81. (G9)	0.9 M Sodium tartrate dibasic dihydrate, 2% w/v Polyethylene glycol 1,000				
82. (G10)	0.9 M Sodium tartrate dibasic dihydrate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
83. (G11)	0.9 M Sodium tartrate dibasic dihydrate, 2% w/v Polyethylene glycol 3,350				
84. (G12)	0.9 M Sodium tartrate dibasic dihydrate, 2% w/v Polyethylene glycol 4,000				
85. (H1)	4.8 M Sodium thiocyanate, 2% w/v Polyethylene glycol 1,000				
86. (H2)	4.8 M Sodium thiocyanate, 2% w/v Polyethylene glycol monomethyl ether 2,000				
87. (H3)	4.8 M Sodium thiocyanate, 2% w/v Polyethylene glycol 3,350				
88. (H4)	4.8 M Sodium thiocyanate, 2% w/v Polyethylene glycol 4,000				
89. (H5)	0.7 M Succinic acid pH 7.0, 2% w/v Polyethylene glycol 1,000				
90. (H6)	0.7 M Succinic acid pH 7.0, 2% w/v Polyethylene glycol monomethyl ether 2,000				
91. (H7)	0.7 M Succinic acid pH 7.0, 2% w/v Polyethylene glycol 3,350				
92. (H8)	0.7 M Succinic acid pH 7.0, 2% w/v Polyethylene glycol 4,000				
93. (H9)	30% v/v Tacsimate™ pH 7.0, 2% w/v Polyethylene glycol 1,000				
94. (H10)	30% v/v Tacsimate™ pH 7.0, 2% w/v Polyethylene glycol monomethyl ether 2,000				
95. (H11)	30% v/v Tacsimate™ pH 7.0, 2% w/v Polyethylene glycol 3,350				
96. (H12)	30% v/v Tacsimate™ pH 7.0, 2% w/v Polyethylene glycol 4,000				